# 2019 Stock Assessment and Fishery Evaluation Report for the Tanner Crab Fisheries of the Bering Sea and Aleutian Islands Regions 

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## Executive Summary

## 1. Stock: species/area.

Southern Tanner crab (Chionoecetes bairdi) in the eastern Bering Sea (EBS).

## 2. Catches: trends and current levels.

Legal-sized male Tanner crab are caught and retained in the directed (male-only) Tanner crab fishery in the EBS. The NPFMC annually determines the overfishing limit (OFL) and acceptable biological catch (ABC) levels for Tanner crab in the EBS, while the Alaska Department of Fish and Game (ADFG) determines the total allowable catch (TAC) separately for areas east and west of $166^{\circ} \mathrm{W}$ longitude in the Eastern Subdistrict of the Bering Sea District Tanner crab Registration Area J. Following rationalization of the Bering Sea and Aleutian Islands (BSAI) crab fisheries in 2005/06, the directed fishery for Tanner crab was open through 2009/10, after which time it was determined that the stock was overfished in the EBS and directed fishing was closed. Prior to the closure, the retained catch averaged 770 t per year between 2005/06-2009/10. The directed fishery was re-opened in 2013/14 following determinations by NMFS in 2012 that the stock was rebuilt and no longer overfished and by ADFG that the stock met state harvest guidelines for opening the fishery. ADFG set the TAC at 1,645,000 lbs ( 746 t ) for the area west of $166^{\circ} \mathrm{W}$ and at $1,463,000 \mathrm{lbs}(664 \mathrm{t})$ for the area east of $166^{\circ} \mathrm{W}$. On closing, $79.6 \%(594 \mathrm{t})$ of the TAC was taken in the western area while $98.6 \%$ ( 654 t ) was taken in the eastern area.

TACs were steadily increased for the next two years, with concomitant increasing harvests. In 2014/15, TAC was set at $6,625,000 \mathrm{lbs}(2,329 \mathrm{t})$ for the area west of $166^{\circ} \mathrm{W}$ and at $8,480,000 \mathrm{lbs}(3,829 \mathrm{t})$ for the area east of $166^{\circ} \mathrm{W}$. On closing, $77.5 \%(2,329 \mathrm{t})$ of the TAC was taken in the western area while $99.6 \%$ $(3,829 \mathrm{t})$ were taken in the eastern area. In 2015/16, TAC was set at $8,396,000 \mathrm{lbs}(3,808 \mathrm{t})$ for the western area and $11,272,000 \mathrm{lbs}(5,113 \mathrm{t}$ ) for the eastern area. On closing, essentially $100 \%$ of the TAC was taken in both areas $(8,373,493 \mathrm{lbs}[3,798 \mathrm{t}]$ in the western area, $11,268,885 \mathrm{lbs}[5,111 \mathrm{t}]$ in the eastern area based on the 5/20/2016 in-season catch report).

Although the NPFMC determined an OFL of almost 60,000,000 lbs ( $\sim 25,000 \mathrm{t}$ ) based on the 2016 assessment (Stockhausen, 2016), mature female Tanner crab biomass fell below the threshold set in the State of Alaska's harvest strategy for opening the fishery; consequently, the fishery was closed and the TAC was set to 0 . Thus, no directed harvest occurred in 2016/17. In 2017/18, ADFG determined that a directed fishery could occur in the area west of $166^{\circ} \mathrm{W}$ longitude. The TAC was set at $2,500,200 \mathrm{lbs}$ ( $1,130 \mathrm{t}$ ), of which $100 \%$ was taken. A similar situation occurred in 2018/19, with only the area west of $166^{\circ} \mathrm{W}$ open to directed fishing. The TAC for 2018/19 was $2,439,000 \mathrm{lbs}(1,106 \mathrm{t})$, with slightly more actually harvested ( $2,441,201 \mathrm{lbs}[1,107 \mathrm{t}]$ ).

In addition to legal-sized males, females and sub-legal males are taken in the directed fishery as bycatch and must be discarded. Discarding of legal-sized males also occurs, primarily because the minimum size
preferred by processors is larger than the minimum legal size but also because "old shell" crab can be less desirable than "new shell" males. Tanner crab are also taken as bycatch in the snow crab and Bristol Bay red king crab fisheries, in the groundfish fisheries and, to a very minor extent, in the scallop fishery. Over the last five years, the snow crab fishery has been the major source of Tanner crab bycatch among these fisheries, averaging $\sim 3,000 t$ for the 5 -year period 2013/14-2017/18. Bycatch in the snow crab fishery in 2018/19 was 888 t . The groundfish fisheries have been the next major source of Tanner crab bycatch over the same five year time period, averaging 325 t . Bycatch in the groundfish fisheries in 2018/19 was 191 t . Excluding the scallop fishery, the Bristol Bay red king crab fishery has typically been the smallest source of Tanner crab bycatch among these fisheries, averaging 202 t over the 5 -year time period. In 2018/19, this fishery accounted for only 74 t of Tanner crab bycatch.

In order to account for mortality of discarded crab, handling mortality rates are assumed to be $32.1 \%$ for Tanner crab discarded in the crab fisheries, $50 \%$ for Tanner crab in the groundfish fisheries using fixed gear, and $80 \%$ for Tanner crab discarded in the groundfish fisheries using trawl gear to account for differences in gear and handling procedures used in the various fisheries.
3. Stock biomass: trends and current levels relative to virgin or historic levels

For EBS Tanner crab, spawning stock biomass is expressed as mature male biomass (MMB) at the time of mating (mid-February). From the author's preferred model (M19F03), estimated MMB for 2018/19 was 79.5 thousand t (Table 47; Figure 61). MMB has been on a declining trend since 2014/15 when it peaked at 135.8 thousand $t$, and it is approaching the very low levels seen in the mid-1990s to early 2000s (1993 to 2003 average: 55.1 thousand t). However, it is considerably below model-estimated historical levels in the late 1970s (1975-1980 average: 215.9 thousand t) before it declined through 1985.

## 4. Recruitment: trends and current levels relative to virgin or historic levels.

From the author's preferred model (M19F03), the estimated total recruitment for 2018/19 (the number of crab entering the population on July 1) is $1,234.9$ million crab (Table 50; Figure 59). Although this value is highly uncertain, it follows two years of similarly high estimates for 2016/17 and 2017/18 (647 and 677 million crab, respectively). The average 5 -year recruitment prior to 2016/17 was only 108 million crab while the longterm (1982+) mean is 394 million crab.

## 5. Management performance

Historical status and catch specifications for eastern Bering Sea Tanner crab.
(a) in 1000 's t.

| Year | MSST | Biomass <br> (MMB) | TAC <br> (East + West) | Retained <br> Catch | Total Catch <br> Mortality | OFL | ABC |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2015 / 16$ | 12.82 | 73.93 | 8.92 | 8.91 | 11.38 | 27.19 | 21.75 |
| $2016 / 17$ | 14.58 | 77.96 | 0.00 | 0.00 | 1.14 | 25.61 | 20.49 |
| $2017 / 18$ | 15.15 | 64.09 | 1.13 | 1.13 | 2.37 | 25.42 | 20.33 |
| $2018 / 19$ | 20.54 | 82.61 | 1.11 | 1.11 | 1.90 | 20.87 | 16.70 |
| $2019 / 20$ |  | 39.55 |  |  |  | 28.86 | 23.09 |

(b) in millions lbs.

| Year | MSST | Biomass <br> (MMB) | TAC <br> (East + West) | Retained <br> Catch | Total Catch <br> Mortality | OFL | ABC |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2015 / 16$ | 28.27 | 162.99 | 19.67 | 19.64 | 25.09 | 59.94 | 47.95 |
| $2016 / 17$ | 32.15 | 171.87 | 0.00 | 0.00 | 2.52 | 56.46 | 45.17 |
| $2017 / 18$ | 33.40 | 95.49 | 2.50 | 2.50 | 5.22 | 56.03 | 44.83 |
| $2018 / 19$ | 45.27 | 182.09 |  |  |  | 46.01 | 36.82 |
| $2019 / 20$ |  | 87.18 |  |  |  | 63.62 | 50.89 |

Shaded values are new estimates or projections based on the current assessment. Other table entries are based on historical assessments and are not updated except for retained catch and total catch mortality.

## 6. Basis for the OFL

a) in 1000's t.

| Year | Tier ${ }^{\text {A }}$ | $\mathrm{B}_{\mathrm{MSY}}{ }^{\text {a }}$ | $\begin{gathered} \text { Current } \\ \text { MMB }^{\mathbf{A}} \\ \hline \end{gathered}$ | B/B MSY $^{\text {a }}$ | $\begin{gathered} \mathbf{F}_{\mathbf{O F L}^{\mathbf{A}}}^{\left(\mathrm{yr}^{-1}\right)} \\ \hline \end{gathered}$ | $\begin{gathered} \hline \text { Years to } \\ \text { define } \\ \mathbf{B}_{\mathrm{MSY}^{4}}{ }^{4} \\ \hline \end{gathered}$ | Natural Mortality $\left(\mathbf{y r}^{-1}\right)^{, B}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2015/16 | 3a | 26.79 | 53.70 | 2.00 | 0.58 | 1982-2015 | 0.23 |
| 2016/17 | 3 a | 25.65 | 45.34 | 1.77 | 0.79 | 1982-2016 | 0.23 |
| 2017/18 | 3 a | 29.17 | 47.04 | 1.49 | 0.75 | 1982-2017 | 0.23 |
| 2018/19 | 3 a | 21.87 | 23.53 | 1.08 | 0.93 | 1982-2018 | 0.23 |
| 2019/20 | 3 b | 41.07 | 39.55 | 0.96 | 1.08 | 1982-2019 | 0.23 |

b) in millions lbs.

| Year | Tier $^{\mathbf{A}}$ | $\mathbf{B}_{\mathbf{M S Y}^{\mathbf{A}}}$ | Current <br> $\mathbf{M M B}^{\mathbf{A}}$ | $\mathbf{B}^{\mathbf{B} / \mathbf{B M S Y}^{\mathbf{A}}}$ | $\mathbf{F}_{\mathbf{\text { OFL }}}{ }^{\mathbf{A}}$ <br> $\left.\mathbf{y r}^{-1}\right)$ | Years to <br> define <br> $\mathbf{B M S Y}^{\mathbf{A}}$ | Natural <br> $\mathbf{M o r t a l i t y ~}^{\mathbf{A}, \mathbf{B}}$ <br> $\left(\mathbf{y r}^{-1}\right)$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2015 / 16$ | 3 a | 59.06 | 118.38 | 2.00 | 0.58 | $1982-2015$ | 0.23 |
| $2016 / 17$ | 3 a | 56.54 | 99.95 | 1.77 | 0.79 | $1982-2016$ | 0.23 |
| $2017 / 18$ | 3 a | 64.30 | 103.70 | 1.49 | 0.75 | $1982-2017$ | 0.23 |
| $2018 / 19$ | 3 a | 48.21 | 51.87 | 1.08 | 0.93 | $1982-2018$ | 0.23 |
| $2019 / 20$ | 3 b | 90.53 | 87.18 | 0.96 | 1.08 | $1982-2019$ | 0.23 |

A-Calculated from the assessment reviewed by the Crab Plan Team in 20XX of 20XX/(XX+1) or based on the author's preferred model for 2019/20.
B-Nominal rate of natural mortality. Actual rates used in the assessment are estimated and may be different.
Current male spawning stock biomass (MMB), as projected for 2019/20, is estimated at 39.55 thousand t . $B_{\text {MSY }}$ for this stock is calculated to be 41.07 thousand $t$, so MSST is 20.54 thousand $t$. Because current MMB $>$ MSST, the stock is not overfished. Total catch mortality (retained + discard mortality in all fisheries, using a discard mortality rate of 0.321 for pot gear and 0.8 for trawl gear) in 2018/19 was 1.90
thousand $t$, which was less than the OFL for 2017/18 (20.97 thousand $t$ ); consequently overfishing did not occur. The OFL for 2019/20 based on the author's preferred model (M19F03) is 28.86 thousand $t$. The $\mathrm{ABC}_{\text {max }}$ for 2019/20, based on the p ${ }^{*} \mathrm{ABC}$, is 28.79 thousand t . In 2014, the SSC adopted a $20 \%$ buffer to calculate ABC for Tanner crab to incorporate concerns regarding model uncertainty for this stock. Based on this buffer, the ABC would be 23.09 thousand t .

## 7. Rebuilding analyses summary.

The EBS Tanner crab stock was found to be above MSST (and $\mathrm{B}_{\mathrm{MSY}}$ ) in the 2012 assessment (Rugolo and Turnock, 2012b) and was subsequently declared rebuilt. The stock remains not overfished. Consequently no rebuilding analyses were conducted.

## A. Summary of Major Changes

1. Changes (if any) to the management of the fishery.

At the March, 2015 SOA Board of Fish (BOF) meeting, the Board adopted a revised harvest strategy for Tanner crab in the Bering Sea District ${ }^{1}$, wherein the TAC for the area east of $166^{\circ} \mathrm{W}$ longitude would be based on a minimum preferred harvest size of 127 mm CW ( 5.0 inches), including the lateral spines. Formerly, this calculation was based on a minimum preferred size of 140 mm CW ( 5.5 inches). The TAC in the area west of $166^{\circ} \mathrm{W}$ longitude continues to be based on a minimum preferred harvest size of 127 mm CW (including lateral spines).

The directed Tanner crab fishery east of $166^{\circ} \mathrm{W}$ longitude has been closed since 2016/17 because mature female Tanner crab biomass in the area has failed to meet the criteria defined in the SOA's harvest strategy to open the fishery. The directed fishery west of $166^{\circ} \mathrm{W}$ longitude was also closed in 2016/17, but has since been prosecuted in 2017/18 and 2018/19.

## 2. Changes to the input data

The following table summarizes data sources that have been updated for this assessment:

[^0]Updated data sources.

| Description | Data types | Time frame | Notes | Source |
| :---: | :---: | :---: | :---: | :---: |
| NMFS EBS Bottom Trawl Survey | area-swept abundance, biomass size compositions male maturity data | $\begin{gathered} 1975-2019 \\ 1975-2019 \\ 2006+ \end{gathered}$ | recalculated, new recalculated, new new | NMFS |
| NMFS/BSFRF | molt-increment data | 2015-17, 2019 | same as 2017 | NMFS, BSFRF |
| BSFRF SBS Bottom Trawl Survey | area-swept abundance, biomass size compositions | $\begin{aligned} & 2013-17 \\ & 2013-17 \end{aligned}$ | $\begin{aligned} & \text { new } \\ & \text { new } \end{aligned}$ | BSFRF |
| Directed fishery | historical retained catch (numbers, biomass) historical retained catch size compositions retained catch (numbers, biomass) retained catch size compositions total catch (abundance, biomass) total catch size compositions | $\begin{aligned} & \hline 1965 / 66-1996 / 97 \\ & 1980 / 81-2009 / 10 \\ & 2005 / 06-2018 / 19 \\ & 2013 / 14-2018 / 19 \\ & 1991 / 92-2017 / 18 \\ & 1991 / 92-2017 / 18 \\ & \hline \end{aligned}$ | not updated not updated updated, new updated, new revised, new revised, new | 2018 assessment 2018 assessment ADFG ADFG ADFG ADFG |
| Snow Crab Fishery | historical effort <br> effort <br> total bycatch (abundance, biomass) <br> total bycatch size compositions | $\begin{aligned} & 1978 / 79 / 1989 / 90 \\ & 1990 / 91-2018 / 19 \\ & 1990 / 91-2018 / 19 \\ & 1990 / 91-2018 / 19 \\ & \hline \end{aligned}$ | not updated revised, new revised, new revised, new | 2018 assessment <br> ADFG <br> ADFG <br> ADFG |
| Bristol Bay Red King Crab Fishery | historical effort effort total bycatch (abundance, biomass) total bycatch size compositions | $\begin{aligned} & \hline 1953 / 54-1989 / 90 \\ & 1990 / 91-2018 / 19 \\ & 1990 / 91-2018 / 19 \\ & 1990 / 91-2018 / 19 \\ & \hline \end{aligned}$ | not updated revised, new revised, new revised, new | 2018 assessment <br> ADFG <br> ADFG <br> ADFG |
| Groundfish Fisheries <br> (all gear types) | historical total bycatch (abundance, biomass) hostorical total bycatch size compositions total bycatch (abundance, biomass) total bycatch size compositions | $\begin{aligned} & \hline 1973 / 74-1990 / 91 \\ & 1973 / 74-1990 / 91 \\ & 1991 / 92-2017 / 18 \\ & 1991 / 92-2017 / 18 \end{aligned}$ | not updated not updated revised, new updated, new | 2018 assessment NMFS/AKFIN |

Changes of note include the incorporation of BSFRF bottom trawl survey data from the "side-by-side" (SBS) catchability studies jointly conducted with the NMFS EBS bottom trawl survey in 2013-2017, the addition of new molt increment (growth) data, and the use of revised estimates by ADFG of total catch/bycatch data from at-sea observer sampling in the crab fisheries. Otherwise, the changes consist of finalized catch data for 2017/18 and new catch data for 2018/19.

## 3. Changes to the assessment methodology.

Following a considerable development effort and substantial review by the CPT at the January 2017 Modeling Workshop and the May 2017 CPT Meeting, with additional review by the SSC at its February and June 2017 meetings, a new modeling "framework", TCSAM02, was recommended by the CPT at its May 2017 meeting (and approved by the SSC at its June 2017 meeting) for use in the 2017/18 assessment. This framework was used again in 2018/19 and is the basis for this assessment. TCSAM02, while based on the previous assessment model (TCSAM2013), constitutes a completely rewritten code library for the Tanner crab assessment model. Results presented at the May 2017 CPT meeting demonstrated that TCSAM02 could be configured to exactly match results from the TCSAM2013 code, thus providing continuity with the old model code.

The 2017 assessment model ("B2b" in that assessment), built on the 2016 model by: 1) fitting EBS model-increment data inside the model to inform growth parameters, b) estimating separate retention functions for three time periods (pre-1997/98, 2005/06-2009/10, and 2013/14-2015/16), and c) estimating the asymptotic value for the fraction of male crab retained in the directed fishery (in the same three time periods as (b)), rather than assuming it was 1 (i.e., $100 \%$ retention at large sizes). This was also the model (with updated data for 2017/18, referred to in that assessment as "18AM17") selected by the CPT and SSC for the 2018 assessment. This model is referred to here as "M19F00" as the base model scenario for this assessment.

The author-recommended model scenario proposed here, "M19F03", differs rather substantially from the 2017 and 2018 assessment models by: 1) adding a likelihood component to fit annual male maturity ogives determined from chela height-to-carapace width ratios in the NMFS survey; 2) eliminating fits to survey biomass and size composition data for male crab classified as mature/immature based on a maturity ogive determined outside the model; and 3) instead fitting to time series of undifferentiated male survey biomass, abundance, and size compositions. In addition, this scenario fits the revised time series data for retained and total catch biomass since 1990/91 provided by ADFG for the directed Tanner crab, snow crab and Bristol Bay red king crab fisheries.

## 4. Changes to the assessment results

Revisions to the input crab fishery data used in the assessment model have had a large effect (almost 2x) on the estimated scale of the population, although the trends are very similar. Average recruitment (1982present) was estimated at 224 million in last year's model, whereas it is estimated at 394 million in the author's preferred model this year. $\mathrm{F}_{\text {MSY }}$ is larger this year ( $1.18 \mathrm{yr}^{-1}$ this year vs. $0.74 \mathrm{yr}^{-1}$ last year), as is $B_{\text {MSY }}(40.75$ thousand $t$ vs. 30.29 thousand $t$ ). The stock remains in Tier 3, but it is now classified as " 3 b " rather than " 3 a " (its classification last year) because the ratio of projected MMB to $\mathrm{B}_{\text {MSY }}$ is 0.95 , i.e. less than 1. Last year the ratio was 1.19.

## B. Responses to SSC and CPT Comments

1. Responses to the most recent two sets of SSC and CPT comments on assessments in general.

## June 2019 SSC Meeting

SSC Comment: The SSC reminded authors to use the model numbering protocols that allow the SSC to understand the year in which a particular version of the model was first introduced.
Response: The Tanner crab assessment has not fully implemented this suggestion. The 2018 assessment model was labeled 18AM17, which does not follow the guidelines. Here, that model is referred to as M19F00 (" 00 " designating the base model from which other scenarios proceed in the 2019 assessment, " F " denoting the "final" scenarios proposed in May). This also does not reflect the requested model numbering. However, the model numbering adopted herein should allow subsequent model numbering to follow the guidelines (so that the author's preferred model M19F03 would become 19.03 in the future).

## May 2019 Crab Plan Team Meeting

## No general comments.

## October 2018 SSC Meeting

SSC Comment: The SSC reminded authors to use the model numbering protocols that allow the SSC to understand the year in which a particular version of the model was first introduced.
Response: Model numbering was consistent with this guideline for the model scenarios presented by the author to the CPT in September 2018. However, the CPT recommended a model based on the 2017 assessment which was labeled 18AM17 to designate the 2017 assessment model updated with 2018 data, which did not follow the guidelines.

SSC Comment: The SSC encourages authors (using VAST estimates of survey biomass) to consider whether or not the apparent reduction in uncertainty in survey biomass is appropriately accounted for with their models/
Response: The Tanner crab assessment does not yet use VAST-based estimates of survey biomass.

## September 2018 Crab Plan Team Meeting

No general comments.
2. Responses to the most recent two sets of SSC and CPT comments specific to the assessment. [Note: for continuity with the previous assessment, the following includes comments prior to the most recent two sets of comments.]

## June 2019 SSC Meeting

The SSC endorsed the CPT suggestions from its May meeting.
Response: none.
The SSC requested an evaluation of all parameters estimated to be at or very near bounds, or substantially limited by priors (unless those priors can be logically defended).
Response: Two tables of parameters estimated at or near their bounds are provided (Tables 18 and 19). These parameters are estimated at their bounds in all (or nearly all) of the scenarios examined here. The parameters include one related to peak retention in the directed fishery prior to 1997 (at its upper bound on the logit scale, implying full retention of large legal males) and two related to the probability of undergoing terminal molt (effectively 1 for males in the largest model size bin and 0 for females in the smallest model size bin). These could be fixed in future models (the latter two are in several scenarios here). Survey catchability parameters for the 1975-1981 time period were also estimated at their lower bound (0.5). This might not be unreasonable given the reduced areal coverage of these surveys relative to later surveys and the spatial limits of the Tanner crab stock. However, it would be worthwhile to explore the effect of reducing these bounds. The remaining parameters are related to selectivity functions describing the size-specific capture efficiency of the fisheries and surveys. Two at their lower bounds are probably inconsequential ( $\mathrm{pS} 2[10]$ and $\mathrm{pS} 4[1]$ ) and are related to the ascending and descending slopes of the dome-shaped selectivity describing male bycatch in the snow crab fishery prior to 1997. A doublenormal is used to describe the dome shape, but an alternative function (e.g., a single normal) might have better estimation properties. The size at $50 \%$ selected was estimated at its upper bound ( 90 mm CW ) for NMFS survey selectivity in the 1975-1981 time period $\mathrm{pS} 1[1]$ ). This results in an almost linear function, rather than asymptotic, across the size range. This result may reflect the changing interaction between the areas surveyed (availability) and the gear selectivity in this time period as the survey gradually extended from the southeast shelf and Bristol Bay where adult males were prevalent to the north and west where more immature males would be encountered, effectively "seeing" relatively more large males than small males. Two other survey-related selectivity parameters, describing the size difference between crab at $50 \%$ and $95 \%$ selected) were estimated at their upper bounds for the both males and females in the NMFS EBS trawl survey in the 1982-present time period ( pS 2 [2] and $\mathrm{pS} 2[4]$ ). The selectivity functions are assumed to be logistic, with the other estimated parameter being the size at $95 \%$ selected. The practical consequence of this is that small crab (females in particular) are described as fairly well-selected ( $>50 \%$ for females) relative to fully-selected (sex-specific) large crab. This result may reflect conflicts from between the model assumption of equal sex ratios for recruitment in the $25-40 \mathrm{~mm}$ CW range, apparent equal abundances and spatial patterns for males and females at small sizes in the NMFS EBS survey, and assumed logistic selectivity. The selectivity parameter describing the size at $50 \%$ selected for males in the groundfish fisheries during 1987-1996 was estimated in all scenarios at its lower bound ( 40 mm CW ), probably a consequence of fairly substantial catches of small crab in some years (e.g., 1993, Figure 12). Finally, three parameters at their upper bounds $(\mathrm{pS} 1[23], \mathrm{pS1}[24]$, and $\mathrm{pS1}[27])$ are related to the size at $95 \%$ selected in the BBRKC fishery in the 1997-2004 (males) and 2005+ (males and females) time periods. The upper bounds ( 180 for males, 140 for females) were selected to reflect the largest possible sizes reasonably expected in the model, so the resulting selectivity functions are essentially positivelysloped linear functions with values fixed at 0.95 at the parameter bound because the other estimated logistic parameter estimates a large size at $50 \%$ selected (see selectivity curves in Figure 46).

## May2019 Crab Plan Team Meeting

The CPT accepted the author's recommended models for presentation in September 2019.
Response: The model runs with the recommended scenarios were run for this assessment, and the results are presented herein. The CPT (and assessment author) referred to these "final scenarios" as 19F.0, 19F.0a, 19F.1, 19F.2, 19F.3, 19F.4, and 19F.5. Here, they are referred to as M19F00, M19F00a, M19F01, M19F02, M19F03, M19F04, and M19F05 (which allowed for additional scenarios while maintaining folder/scenario order on computer disk).

CPT comment: compare the estimated selectivity to the ratio of NMFS to BSFRF numbers at length. Is estimated and empirical catchability/availability/selectivity the same? Does the empirical selectivity look logistic?"
Response: The model-estimated availability of Tanner crab to the survey gears in the side-by-side (SBS) study areas was compared to "empirical" estimates of availability using the ratio of numbers-at-size in the NMFS SBS datasets to those form the full NMFS EBS survey. The results are shown in Figure 53. While there are some similarities between the two sets, there are also substantial differences when conceptually they should be the same. Results for the empirical size-specific relative catchability (the ratio of NMFS to BSFRF estimated abundance at size) are shown in Figure 65, but are not compared directly to the estimated selectivity. The mean curves appear reasonably logistic, with approximate asymptotes of $\sim 0.6$ for males and $\sim 0.4$ for females. If the BSFRF surveys are regarded as providing estimates of absolute abundance (catchability $=1$ for all sizes), this would suggest fully-selected NMFS survey " q "'s are $\sim 0.6$ for males and $\sim 0.4$ for females--which are about $50 \%$ higher than the estimates ( 0.43 and 0.24 , respectively) from the assessment model, but within the $95 \%$ confidence intervals for males (0.37-0.49) (but not females: 0.19-0.29).

CPT comment: show the fits to the BSFRF length composition data by year as well as in aggregate. Response: These fits are shown in Appendix B.

CPT comment: check the bounds of parameters when estimating the BSFRF data.
Response: Fitting the BSFRF data results in no better, or worse, performance in terms of parameters hitting their bounds.

## CPT comment: indicate whether or not Hessians were produced.

Response Hessians were produced for the "best" model runs for all scenarios and .std files were obtained.
CPT comment: Suggest rationale for chosen weighting for the second difference smoothing on the availability curve.
Response: The rationale for the selected weighting is that it reflects a preference toward a smoothlyvarying function, reflecting an assumption that crab of similar sizes would tend to be found together with no abrupt dichotomies (which would justify a smaller smoothing weight) in spatial distribution with size. However, this assumption has not been examined in detail.

CPT comment: Compare trends in largest crab to fishing pressure and area occupied by stock. Response: This is a good suggestion that, time permitting, will be addressed before the January 2020 CPT meeting.

CPT comment: Compare the maximum sizes seen in the fishery to the survey.
Response: Another good suggestion that, time permitting, will be addressed before the January 2020 CPT meeting.

CPT comment: Consider blocking for estimation of growth and probability of maturing.
Response: This has been on the "to do" list for a while now, but with relatively low priority. The problem is that the principal data which the model relies on for estimating both processes is, except for size
compositions, only available (from a practical standpoint) since 2006 for male maturity ogives and since 2015 for (both sexes) molt increment data. The ability of the model to reliably estimate changes in these processes is thus somewhat doubtful.

CPT comment: Make incorporating chela height data in the assessment a priority because this might address changes in the probability of maturing over time
Response: Chela height data, in the form of male maturity ogives based on collections of chela heights since 2006, is incorporated in several model scenarios examined here, including the author-preferred scenario.

CPT comment: Provide retrospective analysis and calculate Mohn's rho for MMB
Response: Retrospective analyses for Tanner crab are complicated given the recent fishery closures and short time frames for molt increment and maturity ogive data. Time did not permit making retrospective analyses for the model scenarios considered herein. However, a retrospective analysis for the CPTselected assessment model could be presented at the January 2020 CPT meeting.

## October 2018 SSC Meeting

Comment: The SSC supports "the author's plans to investigate the sensitivity of the model to just a few early years of catch data".
Response: As described in Section 3.2, the apparent sensitivity of the model to changes in the early 1990s crab observer data was instead due to using erroneous input sample sizes for several years of fishery size composition data. After correcting these errors, the results using the revised crab fishery data are more reasonable, with less inflation of estimated population sizes. However, these sizes are still substantially larger than those obtained using the out-of-date fishery catch data. The author recommended adopting the revised crab fishery data, which was based on a painstaking reclassification of directed vs. incidental effort in the early Tanner and snow crab fisheries that more closely reflects current ADFG practices. Both the CPT and SSC concurred in May/June 2019 with this recommendation.

Comment: "The SSC continues to recommend that the authors try to resolve the parameters on the bounds issue by either simplifying the model or experimenting with removing the bounds".
Response: A number of formerly-estimated parameters related to the sex- and size-specific probability of undergoing the terminal molt to maturity have been eliminated because they were, unsurprisingly, estimated at their bounds (implying a probability of 0 for a terminal molt of very small immature crab or 1 for very large immature crab). This had no discernable effect on the MLE solution.

Comment: "The author should justify fitting both abundance and biomass indices in the model or fit only one index".
Response: The author sees no justification for fitting both abundance and biomass indices in the current model configuration and so will only include fits to one index (biomass) in the model optimization. Fits to the other index may provide a diagnostic capability.

Comment: "The team looks forward to seeing the BSFRF work included in the future If the catchability study is to be used to inform selectivity and catchability estimates in the model, it could be as a prior instead of as fixed inputs".
Response: After preliminary examination of this for the May 2019 CPT meeting, two model scenarios incorporating the BSFRF side-by-side (SBS) tow studies are considered in this assessment, using an approach similar to that used in the snow crab model. The use of the catchability study as a prior is an intriguing idea but would require substantial additional model development and remains to be explored. An alternative approach to the one applied here, which assumes that selectivity in the BSFRF studies is 1 and estimates availability curves that are applied to both the BSFRF and NMFS SBS simultaneously, is to
use the NMFS SBS data to estimate the availability curves outside the model using size-specific ratios between the NMFS SBS and full NMFS estimates of abundance-at-size. These could then be applied inside the model and would eliminate $\sim 50$ additional parameters per year of SBS data. However, issues associated with unobserved size ranges would need to be addressed.

September 2018 CPT Meeting Comment: None

## C. Introduction

## 1. Scientific name.

Chionocoetes bairdi.Tanner crab is one of five species in the genus Chionoecetes (Rathbun, 1924). The common name "Tanner crab" for C. bairdi (Williams et al. 1989) was recently modified to "southern Tanner crab" (McLaughlin et al. 2005). Prior to this change, the term "Tanner crab" had also been used to refer to other members of the genus, or the genus as a whole. Hereafter, the common name "Tanner crab" will be used in reference to "southern Tanner crab".

## 2. Description of general distribution

Tanner crabs are found in continental shelf waters of the north Pacific. In the east, their range extends as far south as Oregon (Hosie and Gaumer 1974) and in the west as far south as Hokkaido, Japan (Kon 1996). The northern extent of their range is in the Bering Sea (Somerton 1981a), where they are found along the Kamchatka peninsula (Slizkin 1990) to the west and in Bristol Bay to the east.

In the eastern Bering Sea (EBS), the Tanner crab distribution may be limited by water temperature (Somerton 1981a). The unit stock is that defined across the geographic range of the EBS continental shelf, and managed as a single unit (Fig. 1). C. bairdi is common in the southern half of Bristol Bay, around the Pribilof Islands, and along the shelf break, although males less than the industry-preferred size ( $>125 \mathrm{~mm}$ CW) and ovigerous and immature females of all sizes are distributed broadly from southern Bristol Bay northwest to St. Matthew Island (Rugolo and Turnock, 2011a). The southern range of the cold water congener the snow crab, C. opilio, in the EBS is near the Pribilof Islands (Turnock and Rugolo, 2011). The distributions of snow and Tanner crab overlap on the shelf from approximately $56^{\circ}$ to $60^{\circ} \mathrm{N}$, and in this area, the two species hybridize (Karinen and Hoopes 1971).

## 3. Evidence of stock structure

Tanner crabs in the EBS are considered to be a separate stock distinct from Tanner crabs in the eastern and western Aleutian Islands (NPFMC 1998). Somerton (1981b) suggests that clinal differences in some biological characteristics may exist across the range of the unit stock. These conclusions may be limited since terminal molt at maturity in this species was not recognized at the time of that analysis, nor was stock movement with ontogeny considered. Biological characteristics estimated based on comparisons of length frequency distributions across the range of the stock, or on modal length analysis over time may be confounded as a result.

Although the State of Alaska's (SOA) harvest strategy and management controls for this stock are different east and west of $166^{\circ} \mathrm{W}$, the unit stock of Tanner crab in the EBS appears to encompass both regions and comprises crab throughout the geographic range of the NMFS bottom trawl survey. Strong evidence is lacking that the EBS shelf is home to two distinct, non-intermixing, non-interbreeding stocks that should be assessed and managed separately (G. Johnson, presentation at the May 2019 CPT meeting).

## 4. Life history characteristics

## a. Molting and Shell Condition

Tanner crabs, like all crustaceans, normally exhibit a hard exoskeleton of chitin and calcium carbonate. This hard exoskeleton requires individuals to grow through a process referred to as molting, in which the individual sheds its current hard shell, revealing a new, larger exoskeleton that is initially soft but which rapidly hardens over several days. Newly-molted crab in this "soft shell" phase can be vulnerable to predators because they are generally torpid and have few defenses if discovered. Subsequent to hardening, an individual's shell provides a settlement substrate for a variety of epifaunal "fouling" organisms such as barnacles and bryozoans. The degree of hard-shell fouling was once thought to correspond closely to post-molt age and led to a classification of Tanner crab by shell condition (SC) in survey and fishery data similar to that described in the following table (NMFS/AFSC/RACE, unpublished):

| Shell Condition <br> Class | $\quad$ Description |
| :---: | :--- |
| 0 | pre-molt and molting crab |
| 1 | carapace soft and pliable |
| 2 | carapace firm to hard, clean |
| 3 | carapace hard; topside usually yellowish brown; thoracic sternum and underside of legs yellow <br> with numerous scratches; pterygostomial and bronchial spines worn and polished; dactyli on <br> meri and metabranchial region rounded; epifauna (barnacles and leech cases) usually present <br> but not always. |
| 5 | carapace hard, topside yellowish-brown to dark brown; thoracic sternum and undersides of legs <br> data yellow with many scratches and dark stains; pterygostomial and branchial spines rounded <br> with tips sometimes worn off; dactyli very worn, sometimes flattened on tips; spines on meri <br> and metabranchial region worn smooth, sometimes completely gone; epifauna most always <br> present (large barnacles and bryozoans). |
| 5 | conditions described in Shell Condition 4 above much advanced; large epifauna almost <br> completely covers crab; carapace is worn through in metabranchial regions, pterygostomial <br> branchial spines, or on meri; dactyli flattened, sometimes worn through, mouth parts and eyes <br> sometimes nearly immobilized by barnacles. |

Although these shell classifications continue to be applied to crab in the field, it has been shown that there is little real correspondence between post-molt age and shell classifications SC 3 through 5, other than that they indicate that the individual has probably not molted within the previous year (Nevisi et al, 1996). In this assessment, crab classified into SCs 3-5 have been aggregated as "old-shell" crab, indicating that these are crab likely to have not molted within the previous year. In a similar fashion, crab classified in SCs $0-2$ have been combined as "new shell" crab, indicating that these are crab have certainly (SCs 0 and 1 ), or are likely to have (SC 2), molted within the previous year.

## b. Growth

Work by Somerton (1981a) estimated growth for EBS Tanner crab based on modal size frequency analysis of Tanner crab in survey data assuming no terminal molt at maturity. Somerton's approach did not directly measure molt increments and his findings are constrained by not considering that the progression of modal lengths between years was biased because crab ceased growing after their terminal molt to maturity.

Growth in immature Tanner crab larger than approximately 25 mm CW proceeds by a series of annual molts, up to a final (terminal) molt to maturity (Tamone et al., 2007). Rugolo and Turnock (2012a) derived growth relationships for male and female Tanner crab used as priors for estimated growth parameters in this (and previous) assessments from data on observed growth in males to approximately 140 mm carapace width (CW) and in females to approximately 115 mm CW that were collected near Kodiak Island in the Gulf of Alaska (Munk, unpublished.; Donaldson et al. 1981). Rugolo and Turnock (2010) compared the resulting growth per molt (gpm) relationships with those of Stone et al. (2003) for Tanner crab in southeast Alaska in terms of the overall pattern of gpm over the size range of crab and found that the pattern of gpm for both males and females was characterized by a higher rate of growth to an intermediate size ( $90-100 \mathrm{~mm} \mathrm{CW}$ ) followed by a decrease in growth rate from that size thereafter. Similarly-shaped growth curves were found by Somerton (1981a) and Donaldson et al. (1981), as well.

Molt increment data was collected for Tanner crab in the EBS during 2015, 2016, 2017 and 2019 in cooperative research between NMFS and the Bering Sea Research Foundation (R. Foy and E. Fedewa, NMFS, pers. comm.s). Previous analysis of the data suggests it is not substantially different from that obtained near Kodiak Island (Stockhausen, 2017). The EBS molt increment data is incorporated in the assessment model to inform inferred growth trajectories in all of the alternative models evaluated in this assessment.

## c. Weight at Size

Weight-at-size relationships used in this assessment were revised in 2014 based on a comprehensive reevaluation of data from the NMFS EBS Bottom Trawl Survey (Daly et al., 2014). Weight-at-size is described by a power-law model of the form $w=a \cdot z^{b}$, where $w$ is weight in kg and $z$ is size in mm CW (Daly et al., 2016; table below). Parameter values are presented in the following table:

| sex | maturity | $a$ | $b$ |
| :---: | :---: | :---: | :---: |
| males |  | 0.000270 | 3.022134 |
| females | immature <br> (non-ovigerous) <br> mature <br> (ovigerous) | 0.000562 | 2.816928 |

## d. Maturity and Reproduction

It is now generally accepted that both Tanner crab males (Tamone et al. 2007) and females (Donaldson and Adams 1989) undergo a terminal molt to maturity, as in most majid crabs. Maturity in females can be determined visually rather unambiguously from the relative size of the abdomen. Females usually undergo their terminal molt from their last juvenile, or pubescent, instar while being grasped by a male (Donaldson and Adams 1989). Subsequent mating takes place annually in a hard shell state (Hilsinger 1976) and after extruding the female's clutch of eggs. While mating involving old-shell adult females has been documented (Donaldson and Hicks 1977), fertile egg clutches can be produced in the absence of males by using sperm stored in the spermathacae (Adams and Paul 1983, Paul and Paul 1992). Two or more consecutive egg fertilization events can follow a single copulation using stored sperm to selffertilize the new clutch (Paul 1982, Adams and Paul 1983), although egg viability decreases with time and age of the stored sperm (Paul 1984).

Maturity in males can be classified either physiologically or morphometrically, but is not as easily determined as with females. Physiological maturity refers to the presence or absence of spermataphores in the gonads whereas morphometric maturity refers to the presence or absence of a large claw (Brown and Powell 1972). During the molt to morphometric maturity, there is a disproportionate increase in the size of the chelae in relation to the carapace (Somerton 1981a). The ratio of chela height (CH) to carapace width (CW) has been used to classify male Tanner crab as to morphometric maturity. While many earlier studies on Tanner crabs assumed that morphometrically mature male crabs continued to molt and grow, there is now substantial evidence supporting a terminal molt for males (Otto 1998, Tamone et al. 2007). A consequence of the terminal molt in male Tanner crab is that a substantial portion of the population may never achieve legal size (NPFMC 2007). In this assessment, several model scenarios are considered in which size-specific annual proportions of mature, new shell male crab to all new shell male crab in the NMFS EBS bottom trawl survey, based on classification using CH:CW ratios, are fit to inform sizespecific probabilities of terminal molt.

Although observations are lacking in the EBS, seasonal differences have been observed between mating periods for pubescent and multiparous females in the Gulf of Alaska and Prince William Sound. There, pubescent molting and mating takes place over a protracted period from winter through early summer, whereas multiparous mating occurs over a relatively short period during mid April to early June (Hilsinger 1976, Munk et al. 1996, and Stevens 2000). In the EBS, egg condition for multiparous Tanner crabs assessed between April and July 1976 also suggested that hatching and extrusion of new clutches for this maturity state began in April and ended sometime in mid-June (Somerton 1981a).

## e. Fecundity

A variety of factors affect female fecundity, including somatic size, maturity status (primiparous vs. multiparous), age post terminal molt, and egg loss (NMFS 2004). Of these factors, somatic size is the most important, with estimates of 89 to 424 thousand eggs for females 75 to 124 mm CW , respectively
(Haynes et al. 1976). Maturity status is another important factor affecting fecundity, with primiparous females being only $\sim 70 \%$ as fecund as equal size multiparous females (Somerton and Meyers 1983). The number of years post maturity molt, and whether or not, a female has had to use stored sperm from that first mating can also affect egg counts (Paul 1984, Paul and Paul 1992). Additionally, older senescent females often carry small clutches or no eggs (i.e., are barren) suggesting that female crab reproductive output is a concave function of age (NMFS 2004).

## f. Size at Maturity

Rugolo and Turnock (2012b) estimated size at $50 \%$ mature for females (all shell classes combined) from data collected in the NMFS bottom trawl survey at 68.8 mm CW, and 74.6 mm CW for new shell females. For males, Rugolo and Turnock (2012a) estimated classification lines using mixture-of-tworegressions analysis to define morphometric maturity for the unit Tanner crab stock, and for the sub-stock components east and west of $166^{\circ} \mathrm{W}$, based on chela height and carapace width data collected during the 2008 NMFS bottom trawl survey. These rules were then applied to historical survey data from 1990-2007 to apportion male crab as immature or mature based on size (Rugolo and Turnock, 2012b). Rugolo and Turnock (2012a) found no significant differences between the classification lines of the sub-stock components (i.e., east and west of $166^{\circ} \mathrm{W}$ ), or between the sub-stock components and that of the unit stock classification line. Size at $50 \%$ mature for males (all shell condition classes combined) was estimated at 91.9 mm CW, and at 104.4 mm CW for new shell males. By comparison, Zheng and Kruse (1999) used knife-edge maturity at $>79 \mathrm{~mm}$ CW for females and $>112 \mathrm{~mm} \mathrm{CW}$ for males in development of the current SOA harvest strategy.

The Rugolo-Turnock classification approach is referred to herein as the "Rugolo-Turnock male maturity ogive". In this and previous assessments, the Rugolo-Turnock maturity ogive has been used to fix the proportions of immature and mature, new shell male crab in size composition data from the entire NMFS EBS bottom trawl survey dataset and to subsequently provide survey biomass estimates of abundance and biomass aggregated over all size classes. The NMFS survey datasets that use this approach to characterize male maturity outside the assessment model are identified here as "NMFS 0". The assessment model has used the resulting annual estimates of immature and new shell mature male crab abundance, biomass and size compositions as "data" to inform the model's estimates of population size and processes, including the probability of immature male crab within a given model size bin undergoing the terminal molt to maturity. This is somewhat circular in nature, and several model scenarios in this assessment fit directly to annual observed (i.e., classifying crab based on $\mathrm{CH}: \mathrm{CW}$ ratios) proportions of new shell mature males to all new shell males by size bin without classifying new shell males as immature or mature outside the model.

## g. Mortality

Due to the lack of age information for crab, Somerton (1981a) estimated mortality separately for individual EBS cohorts of immature and adult Tanner crab. Somerton postulated that age five crab (mean $\mathrm{CW}=95 \mathrm{~mm}$ ) were the first cohort to be fully recruited to the NMFS trawl survey sampling gear and estimated an instantaneous natural mortality rate of 0.35 for this size class using catch curve analysis. Using this analysis with two different data sets, Somerton estimated natural mortality rates of adult male crab from the fished stock to range from 0.20 to 0.28 . When using CPUE data from the Japanese fishery, estimates of M ranged from 0.13 to 0.18 . Somerton concluded that estimates of M from 0.22 to 0.28 obtained from models that used both the survey and fishery data were the most representative.

Rugolo and Turnock (2011a) examined empirical evidence for reliable estimates of oldest observed age for male Tanner crab. Unlike its congener the snow crab, information on longevity of the Tanner crab is lacking. They reasoned that longevity in a virgin population of Tanner crab would be analogous to that of the snow crab, where longevity would be at least 20 years, given the close analogues in population dynamic and life-history characteristics (Turnock and Rugolo 2011a). Employing 20 years as a proxy for
longevity and assuming that this age represented the upper 98.5th percentile of the distribution of ages in an unexploited population, M was estimated to be 0.23 based on Hoenig's (1983) method. If 20 years was assumed to represent the $95 \%$ percentile of the distribution of ages in the unexploited stock, the estimate for M was 0.15 . Rugolo and Turnock (2011a) adopted $\mathrm{M}=0.23$ for both male and female Tanner because the value corresponded with the range estimated by Somerton (1981a), as well as the value used in the analysis to estimate new overfishing definitions underlying Amendment 24 to the Crab Fishery Management Plan (NPFMC 2007).

## 5. Brief summary of management history.

A complete summary of the management history is provided in the ADFG Area Management Report appended to the annual SAFE. Fisheries have historically taken place for Tanner crab throughout their range in Alaska, but currently only the fishery in the EBS is managed under a federal Fishery Management Plan (FMP; NPFMC 2011). The plan defers certain management controls for Tanner crab to the State of Alaska (SOA), with federal oversight (Bowers et al. 2008). The SOA manages Tanner crab based on registration areas divided into districts. Under the FMP, the state can adjust districts as needed to avoid overharvest in a particular area, change size limits from other stocks in the registration area, change fishing seasons, or encourage exploration (NPFMC 2011).

The Bering Sea District of Tanner crab Registration Area J (Figure 1) includes all waters of the Bering Sea north of Cape Sarichef at $54^{\circ} 36^{\prime} \mathrm{N}$ and east of the U.S.-Russia Maritime Boundary Line of 1991. This district is divided into the Eastern and Western Subdistricts at $173^{\circ} \mathrm{W}$. The Eastern Subdistrict is further divided at the Norton Sound Section north of the latitude of Cape Romanzof and east of $168^{\circ} \mathrm{W}$ and the General Section to the south and west of the Norton Sound Section (Bowers et al. 2008). In this report, the terms "east region" and "west region" are used in shorthand fashion to refer to the regions demarcated by $166^{\circ} \mathrm{W}$ longitude.

In March 2011, the Alaska Board of Fisheries (BOF) approved a new minimum size limit harvest strategy for Tanner crab effective for the 2011/12 fishery. Prior to this change, the minimum legal size limit was 5.5 " ( 138 mm CW) throughout the Bering Sea District. The new regulations established different minimum size limits east and west of $166^{\circ} \mathrm{W}$. The minimum size limit for the fishery to the east of $166^{\circ} \mathrm{W}$ is now $4.8^{\prime \prime}(122 \mathrm{~mm} \mathrm{CW})$ and that to the west is $4.4^{\prime \prime}(112 \mathrm{~mm} \mathrm{CW})$, where the size measurement includes the lateral spines. For economic reasons, fishers may adopt larger minimum sizes for retention of crab in both areas, and the SOA's harvest strategy and total allowable catch (TAC) calculations are based on assumed minimum preferred sizes that are larger than the legal minimums. In 2011, these minimum preferred sizes were set at 5.5 " ( 140 mm CW ) in the east and 5 " ( 127 mm CW ) in the west, including the lateral spines. In 2015, following a petition by the crab industry, the BOF revised the minimum preferred size for TAC calculations in the area east of $166^{\circ} \mathrm{W}$ longitude to 5 " ( 127 mm CW ), the same as that in the western area. These new "preferred" sizes were used to set the TAC for the 2015/16 fishery season.

In assessments prior to 2016, the term "legal males" was used to refer to male crab $\geq 138 \mathrm{~mm} \mathrm{CW}$ (not including the lateral spines), although this was not strictly correct as it referred to the industry's "preferred" crab size in the east region, as well as to the minimum size in the east used in the SOA's harvest strategy for TAC setting. In subsequent assessments (and this one), the term "legal males" is used to refer to male crab 125 mm CW (the current minimum "preferred" size for both eastern and western areas used in the SOA's harvest strategy) and larger.

Landings of Tanner crab in the Japanese pot and tangle net fisheries were reported in the period 19651978, peaking at 19.95 thousand t in 1969. The Russian tangle net fishery was prosecuted during 19651971 with peak landings in 1969 at 7.08 thousand t . Both the Japanese and Russian Tanner crab fisheries were displaced by the domestic fishery by the late-1970s (Table 1; Figure 2). Foreign fishing for Tanner crab ended in 1980.

The domestic Tanner crab pot fishery developed rapidly in the mid-1970s (Tables 1 and 2; Figure 3). Domestic US landings were first reported for Tanner crab in 1968 at 0.46 thousand $t$ taken incidentally to the EBS red king crab fishery. Tanner crab was targeted thereafter by the domestic fleet and landings rose sharply in the early 1970s, reaching a high of 30.21 thousand $t$ in 1977/78. Landings fell sharply after the peak in 1977/78 through the early 1980s, and domestic fishing was closed in 1985/86 and 1986/87 due to depressed stock status. In 1987/88, the fishery reopened and landings rose again in the late-1980s to a second peak in 1990/91 at 16.61 thousand t , and then fell sharply through the mid-1990s. The domestic Tanner crab fishery was closed between 1997/98 and 2004/05 as a result of conservation concerns regarding the depressed status of the stock. It re-opened in 2005/06 and averaged 0.77 thousand $t$ retained catch between 2005/06-2009/10 (Tables 1 and 2). The SOA closed directed commercial fishing for Tanner crab during the 2010/11-2012/13 seasons because estimated female stock metrics fell below thresholds adopted in the state harvest strategy. However, these thresholds were met in fall 2013 and the directed fishery was opened in 2013/14. TAC was set at $1,645,000 \mathrm{lbs}\left(746 \mathrm{t}\right.$ ) for the area west of $166^{\circ} \mathrm{W}$ and at $1,463,000 \mathrm{lbs}(664 \mathrm{t})$ for the area east of $166^{\circ} \mathrm{W}$ in the Eastern Subdistrict of Tanner crab Registration Area J. The fisheries opened on October 15 and closed on March 31. On closing, 79.6\% (594 t) of the TAC had been taken in the western area while $98.6 \%$ (654 t) had been taken in the eastern area. Prior to the closures, the retained catch averaged 770 t per year between 2005/06-2009/10. In 2014, TAC was set at $6,625,000 \mathrm{lbs}(3,005 \mathrm{t})$ for the area west of $166^{\circ} \mathrm{W}$ and at $8,480,000 \mathrm{lbs}(3,846 \mathrm{t})$ for the area east of $166^{\circ} \mathrm{W}$. On closing, $77.5 \%(2,329 \mathrm{t})$ of the TAC was taken in the western area while $99.6 \%(3,829$ t) were taken in the eastern area. In 2015, TAC was set at $8,396,000 \mathrm{lbs}(3,808 \mathrm{t})$ in the western area and $11,272,000 \mathrm{lbs}(5,113 \mathrm{t})$ in the eastern area. On closing, essentially $100 \%$ of the TAC was taken in each area ( $3,798 \mathrm{t}$ in the west, $5,111 \mathrm{t}$ in the east). The total retained catch in 2015/16 ( $8,910 \mathrm{t}$ ) was the largest taken in the fishery since 1992/93 (Tables 1, 2; Figure 2). The directed fisheries in both areas were closed in 2016/17 because mature female biomass in the NMFS EBS Bottom Trawl Survey did not exceed the threshold set in the SOA's harvest strategy to allow them to open. Total retained catch was thus 0 in 2016/17. In 2017/18, the SOA allowed a limited directed fishery west of $166^{\circ} \mathrm{W}$ longitude but closed the fishery east of $166^{\circ}$ W. Essentially, the entire TAC ( 1,130 t) was taken in 2017/18. The 2018/19 season followed a similar pattern, with the directed fishery closed in the eastern area and open in the western area (with a TAC of 1.106 thousand t ). The entire TAC was again harvested in 2018/19.

Bycatch and discard losses of Tanner crab originate from the directed pot fishery, non-directed snow crab and Bristol Bay red king crab pot fisheries, and the groundfish fisheries (Table 3; Figure 3). Within the assessment model, bycatch estimates are converted to discard mortality using assumed handling mortality rates of $32.1 \%$ for bycatch in the crab fisheries and $80 \%$ for bycatch in the groundfish fisheries. Bycatch was persistently high during the early-1970s; a subsequent peak mode of discard losses occurred in the early-1990s. In the early-1970s, the groundfish fisheries contributed significantly to total bycatch losses (although bycatch in the crab fisheries was undocumented at the time). From 1992/93 (when reliable crab fishery bycatch estimates are considered to be first available) to 2004/05, the groundfish fisheries accounted for the largest proportion of discard mortality. Since 2005/06, however, the crab fisheries have accounted for the largest proportion.

## D. Data

For several years now, NMFS has annually provided a standardized version of the EBS bottom trawl survey for Tanner and other crab stocks for surveys from 1975 to the present. Similarly, estimates from the NMFS Regional Office for crab bycatch in the groundfish fisheries since 1990 have been provided by AKFIN. Standard procedure in this assessment has been to update all the data used in the assessment model based on these sources each year, so that the data used in the assessment remains consistent with the survey and groundfish bycatch data provided by NMFS and AKFIN (see below).

However, this was not done with the retained catch and bycatch data provided annually by ADFG due mainly to inconsistency between years in the formats in which the data were provided. More recently
(starting in 2017), ADFG has provided datasets in more consistent formats, allowing development of stable R code to extract the data required for the assessment in a repeatable fashion, rather than doing it by hand or in "one-off" code for a particular assessment. Thus, prior to 2018 the corresponding data in the assessment tended to be added for the current and only updated for the previous year (if necessary). Following the 2017 assessment in the course of developing R code to extract the data to a format compatible with the assessment, it was noted that discrepancies had accrued primarily between the total catch biomass data used in the assessment and those provided by ADFG for fisheries conducted in the 1990s, although there were also some (much smaller) discrepancies later in the time series and in the retained catch data as well (Tables 4-7). The discrepancies in the total catch estimates in the 1990s were traced back to a substantial reclassification of directed fishing effort and at-sea observer sampling by Doug Pengilly in 2015 that primarily affected the expansion of sampled catch by at-sea observers to total catch estimates in the early 1990s; these had not been updated in the assessment (pending a review). The smaller discrepancies later in the time series may have been due to a change in the size-weight relationships used to calculate average catch weight when CPUE was expanded to total catch biomass. The main discrepancies in retained catch occurred in 2013/14 and 2014/15 and were the result of incidental retained catch of Tanner crab in the snow and BBRKC fisheries inadvertently not being aggregated into the values for the directed fishery provided to the assessment author (Table 8). For the 2018 assessment, the "current" crab fisheries data differed from "historical" data (i.e., used in the 2017 assessment) as summarized in the following table:

| data type | years not updated | years updated |
| :--- | :---: | :---: |
| effort in the BBRKC fishery | $1953 / 54-1989 / 90$ | $1990 / 91$ to present |
| effort in the snow crab fishery | $1978 / 79-1989 / 90$ | $1990 / 91$ to present |
| retained catch abundance, biomass | $1965 / 66-1996 / 97$ | $2005 / 06$ to present |
| retained catch size compositions | $1980 / 81-1989 / 90$ | $1990 / 91$ to present |
| total catch abundance, biomass (all fisheries) | -- | $1990 / 91$ to present |
| total catch size compositions (all fisheries) | -- | $1990 / 91$ to present |

Unfortunately, the CPT and SSC did not have the opportunity to approve the use of the "current" version of data from the crab fisheries prior to the 2018 assessment; thus, the 2018 assessment was based on the "historical" version, with the addition of 2017/18 data. However, the "current" version was reviewed by the CPT in May 2019 and approved for use in this assessment (to which the SSC concurred at the June 2019 Council meeting).

## 1. Summary of new information

ADFG provided revised values for retained catch abundance and biomass from fish ticket data for 2005/06-2017/18, with new values for 2018/19. This included a breakout of incidental retained Tanner crab catch in the snow crab and BBRKC fisheries; prior to the 2018/19 assessment only total retained catch (assumed taken in the directed fishery) had been provided. In general, incidental retained catch of Tanner crab in the snow crab and BBRKC fisheries has been very small compared with that from the directed fishery and continues to be "lumped" with that for the directed fishery. Revised retained catch size composition data from "dockside" observer sampling in the directed fishery were provided by ADFG last year for 1989/90-2017/18 and updated by ADFG this year for 2013/14-2017/18, with new data for 2018/19.

Revised estimates of total Tanner crab catch and bycatch in the directed, snow crab, and BBRKC fisheries provided by ADFG for 1990/91-2017/18 were incorporated into the assessment. ADFG provided updated values for total catch in the crab fisheries for 2017/18 and new values for 2018/19.

Tanner crab bycatch data in the groundfish fisheries (abundance, biomass, size compositions) were extracted for 1991/92-2018/19 from the groundfish observer and AKRO databases on AKFIN. Although
the bycatch data in the groundfish fisheries is available by gear type, all model scenarios examined here fit the data aggregated over gear types.

Swept-area abundance, biomass and size composition data from the 2019 NMFS EBS Bottom Trawl Survey were added to the assessment. Survey results for the assessment were calculated directly from the survey "crab haul" data files and station strata file to incorporate assessment criteria (e.g., excluding crab $<25 \mathrm{~mm} \mathrm{CW}$, aggregating crab $>185 \mathrm{~mm}$ CW into the upper-most size bin in size compositions) and facilitate comparisons across multiple areas and population categories.

Molt increment data from growth studies conducted in the EBS as cooperative research by NMFS and BSFRF are fit in the model scenarios included in this assessment, with new data from studies in 2017 and 2019 included in this assessment.

Annual male maturity ogives based on classification of male crab in the NMFS EBS bottom trawl survey using $\mathrm{CH}: \mathrm{CW}$ ratios are fit in a number of the model scenarios considered in this assessment. Existing and new (2019) chela height data sets were analyzed to provide estimates of the fraction of new shell mature males to all new shell male crab by 10 mm size bin (J. Richar, NMFS, pers. comm.). Data from collections since 2006, when chela heights were first measured to 0.1 mm , are included in the assessment.

Finally, data for Tanner crab from the joint BSFRF-NMFS comparative catchability ("side-by-side") studies in 2013-2017 are included in the assessment for the first time.

The following table summarizes data sources that have been updated for this assessment:

| Description | Data types | Time frame | Notes | Source |
| :---: | :---: | :---: | :---: | :---: |
| NMFS EBS Bottom Trawl Survey | area-swept abundance, biomass size compositions male maturity data | $\begin{gathered} 1975-2019 \\ 1975-2019 \\ 2006+ \end{gathered}$ | recalculated, new recalculated, new new | NMFS |
| NMFS/BSFRF | molt-increment data | 2015-17, 2019 | same as 2017 | NMFS, BSFRF |
| BSFRF SBS Bottom Trawl Survey | area-swept abundance, biomass size compositions | $\begin{aligned} & 2013-17 \\ & 2013-17 \end{aligned}$ | $\begin{aligned} & \text { new } \\ & \text { new } \end{aligned}$ | BSFRF |
| Directed fishery | historical retained catch (numbers, biomass) historical retained catch size compositions retained catch (numbers, biomass) retained catch size compositions total catch (abundance, biomass) total catch size compositions | $\begin{aligned} & 1965 / 66-1996 / 97 \\ & 1980 / 81-2009 / 10 \\ & 2005 / 06-2018 / 19 \\ & 2013 / 14-2018 / 19 \\ & 1991 / 92-2017 / 18 \\ & 1991 / 92-2017 / 18 \\ & \hline \end{aligned}$ | not updated not updated updated, new updated, new revised, new revised, new | 2018 assessment 2018 assessment ADFG ADFG ADFG ADFG |
| Snow Crab Fishery | historical effort effort total bycatch (abundance, biomass) total bycatch size compositions | $\begin{aligned} & 1978 / 79 / 1989 / 90 \\ & 1990 / 91-2018 / 19 \\ & 1990 / 91-2018 / 19 \\ & 1990 / 91-2018 / 19 \end{aligned}$ | not updated revised, new revised, new revised, new | 2018 assessment <br> ADFG <br> ADFG <br> ADFG |
| Bristol Bay Red King Crab Fishery | historical effort effort total bycatch (abundance, biomass) total bycatch size compositions | $\begin{aligned} & \hline 1953 / 54-1989 / 90 \\ & 1990 / 91-2018 / 19 \\ & 1990 / 91-2018 / 19 \\ & 1990 / 91-2018 / 19 \end{aligned}$ | not updated revised, new revised, new revised, new | 2018 assessment <br> ADFG <br> ADFG <br> ADFG |
| Groundfish Fisheries <br> (all gear types) | historical total bycatch (abundance, biomass) hostorical total bycatch size compositions total bycatch (abundance, biomass) total bycatch size compositions | $\begin{aligned} & 1973 / 74-1990 / 91 \\ & 1973 / 74-1990 / 91 \\ & 1991 / 92-2017 / 18 \\ & 1991 / 92-2017 / 18 \end{aligned}$ | not updated not updated revised, new updated, new | 2018 assessment NMFS/AKFIN |

The following table summarizes the data coverage in the assessment model (color shading highlights different model time periods and data components):


## 2. Data presented as time series

For the data presented in this document, the convention is that 'year' refers to the year in which the NMFS bottom trawl survey was conducted (nominally July 1, yyyy), and fishery data are those subsequent to the survey (July 1, yyyy to June 30, yyyy+1)--e.g., 2015/16 indicates the 2015 bottom trawl survey and the winter 2015/16 fishery.

## a. Retained catch

Retained catch in the directed fisheries for Tanner crab conducted by the foreign fisheries (Japan and Russia) and the domestic fleet, starting in 1965/66, is presented in Table 1 and Figures 2 and 3 by fishery year. More detailed information on retained catch in the directed domestic pot fishery is provided in Table 2, which lists total annual catches in numbers of crab and biomass (in lbs), as well as the SOA's Guideline Harvest Level (GHL) or Total Allowable Catch (TAC), number of vessels participating in the directed fishery, and the fishery season. Information from the Community Development Quota (CDQ) is included in the totals starting in 2005/06.

Directed fisheries for Tanner crab in the EBS began in 1965. Retained catch has followed a "boom-andbust" cycle over the years, with the fishery experiencing periods of rapidly increasing catches followed by rapidly declining ones, after which it is closed for a time during which the stock partially recovers. Retained catch increased rapidly from 1965 to 1975 , reaching $\sim 25,000 \mathrm{t}$ in 1970. It declined to $\sim 13,000 \mathrm{t}$ in 1973/74 coinciding with the termination of Russian fishing and the beginning of the domestic pot fishery. It increased again, this time to its highest level, in 1977/78 ( $\sim 35,000 \mathrm{t}$ ) as the domestic fishery developed rapidly, but it subsequently declined again and the fishery was closed in 1985/86 and 1986/87. In the late 1980s and early 1990s, the fishery experienced another, somewhat smaller, "boom" followed by a "bust" and closure of the fishery from 1997/98 to 2004/05. From 2005/06 to 2009/10, the fishery experienced its smallest boom-and-bust cycle, peaking at only $\sim 1,000 \mathrm{t}$ retained catch, and was closed again from 2010/11 to 2012/13. The fishery was re-opened in 2013/14, and retained catch increased each subsequent year until 2016/17 as TACs increased (Figures 2 and 3). The retained catch for 2015/16 (8,910 t) was the largest since 1992/1993 (15,920 t; Table 1). However, ADFG closed the directed fishery in both areas for the 2016/17 fishing season because mature female biomass in the 2016 NMFS EBS bottom trawl survey did not meet the SOA's criteria for opening the fisheries. In 2017/18, ADFG allowed the fishery to commence in the western area (TAC was set at $1,130 \mathrm{t}$ ) but was closed in the eastern area. The directed fishery essentially caught the entire TAC. The 2018/19 fishery was similar to that in 2017/18 in that the eastern area was closed and the entire TAC $(1,100 \mathrm{t})$ was taken west of $166^{\circ} \mathrm{W}$ longitude.

## b. Information on bycatch and discards

Total catch estimates for Tanner crab in the directed Tanner crab, the snow crab, and the BBRKC fisheries are provided in Table 3 and Figure 4 based on ADFG "at-sea" crab observer sampling starting in 1990/91. Annual bycatch in the groundfish fisheries, based on NMFS groundfish observer programs, is also available starting in 1973/74, but sex is undifferentiated. A value of 0.321 is used in the assessment model for "handling mortality" in the crab fisheries to convert observed bycatch to (unobserved) mortality (Stockhausen, 2014). For the groundfish fisheries, a value of 0.8 is used for handling mortality aggregated across gear types to reflect differences in groundfish gear effects and on-deck operations compared with the crab fleets. In previous assessments, estimates of "discards" were provided rather than estimates for "total catch", which allowed mortality associated with the handling process to be estimated outside the assessment model. While this generally remains true for bycatch in the groundfish and non-directed crab fisheries (most or all Tanner crab bycatch is discarded), "discard mortality" cannot be estimated outside the assessment model for males in the directed fishery.

Estimated bycatch mortality in the groundfish fisheries (without distinguishing gear type) was highest ( $\sim 15,000 \mathrm{t}$ ) in the early 1970 s , but was substantially reduced by 1977 to $\sim 2,000 \mathrm{t}$ with the curtailment of foreign fishing fleets (Stockhausen, 2017). It declined further in the 1980s (to $\sim 500 \mathrm{t}$ ) but increased
somewhat in the late 1980s to a peak of $\sim 2,000 t$ in the early 1990s before undergoing a gradual decline until 2008, after which it has fluctuated annually below $\sim 300 t$ to the present ( 150 t in 2018/19).

In the crab fisheries, the largest component of bycatch occurs on males. In the early 1990s, female bycatch ranged between 6 and $40 \%$ of the bycatch in the directed and snow crab fisheries. Since the directed fishery re-opened in 2013/14, the fraction of bycatch that is female has ranged between $2 \%$ and $6 \%$ in the directed fishery, between 0.3 and $3 \%$ in the BBRKC fishery, and has been below $1 \%$ in the snow crab fishery. Estimates of total groundfish bycatch are not currently available by sex.
c. Catch-at-size for fisheries, bycatch, and discards

Retained (male) catch-at-size in the directed Tanner crab fishery from ADFG dockside observer sampling is shown in Figure 5 by fishery region and shell condition since the fishery re-opened in 2013/14. These appear to indicate a shift to retaining somewhat smaller minimum sizes since 2013/14, compared with 2005/06-2009/10 (Stockhausen, 2017). In fact, the BOF in 2014/15, in response to a petition by industry, changed its harvest strategy for calculating TACs to reflect a smaller minimum industry-preferred size of 125 mm CW east of $166^{\circ} \mathrm{W}$ longitude. In addition, the proportion of old shell crab retained appears to have increased over the past few years and substantially exceeded that of new shell crab across the retained size range.

Normalized total catch (retained + discards) size compositions from at-sea crab fishery observer sampling are presented by sex and fishery in Figures 6-11. The snow crab fishery, conducted primarily in the northern and western parts of the EBS shelf, catches predominantly small males while the BBRKC fishery, conducted to the south and east in Bristol Bay, predominantly catches large males. The size compositions in the snow crab fishery clearly reflect some sort of "dome-shaped" selectivity pattern (as assumed in the assessment model), with selectivity small for small and large males and highest for intermediate-sized males. In contrast, selectivity in the BBRKC fishery appears more consistent with asymptotic selection. The directed fishery, which extends across the shelf from west of the Pribilof Islands into Bristol Bay in the east catches primarily intermediate-sized males, with about half the new shell males caught larger than the industry-preferred size of 125 mm CW . Similar patterns are apparent for females, as well.

Sex-specific size compositions from observer sampling for bycatch in the groundfish fisheries, expanded to total bycatch, are shown in Figures 112-13 for 1991/92 to 2018/19. These fisheries, targeting a variety of groundfish stocks and using a variety of gear types, take a much larger size range of Tanner crab as bycatch than does the pot gear used in the crab fisheries-perhaps even providing support for recruitment events (see, e.g., the peaks in relative abundance at small sizes in the size compositions for 2003/04 and 2004/05 in Figure 12).

Raw and input sample sizes (number of individuals measured) for the various fisheries are presented in Tables 9-13.

## d. Survey biomass estimates

Time series trends from the NMFS EBS bottom trawl survey suggest the Tanner crab stock in the EBS has undergone decadal-scale fluctuations (Tables 14-15, Figures14-15). Estimated biomass of male crab in the survey time series started at its maximum ( $295,000 \mathrm{t}$ ) in 1975, decreased rapidly to a low $(15,000 \mathrm{t}$ ) in 1985, and rebounded quickly to a smaller peak (146,000 t) in 1991 (Table 14). After 1991, male survey biomass decreased again, reaching a minimum of $14,600 \mathrm{t}$ in 1997. Recovery following this decline was slow and male survey biomass did not peak again until 2007 (104,000 t), after which it has fluctuated more rapidly-decreasing within two years by over $50 \%$ to a minimum in $2009(47,000 \mathrm{t})$, followed by a doubling to a peak in 2014 ( $109,000 \mathrm{t}$ ). Since 2014 the trend has been a steady decline, with male biomass currently at its lowest point ( $28,000 \mathrm{t}$ ) since 2000 (Table 14). Trends in the male and female components of survey biomass have primarily been in synchrony with one another, as have changes in the eastern and
western management regions (east and west of $166^{\circ} \mathrm{W}$ longitude), although the magnitudes differ (Figure 14). Preferred-size male survey biomass has been declining east of $166^{\circ} \mathrm{W}$ (and in the EBS as a whole) since 2014, but was increasing up to 2016 in the west. In the west, it declined in 2017, remained essentially unchanged in 2018, and dropped by over $50 \%$ from 2018 to 2019 (Table 15, Figure 15). The ratio of new shell to old shell preferred-size males crab across the EBS has dropped dramatically since 2015, when the ratio was almost 1:1. In 2019, the ratio was almost 1:20 new shell to old shell crab biomass.

Data from the BSFRF-NMFS cooperative side-by-side (SBS) catchability studies are incorporated into several model scenarios in this assessment for the first time. During the SBS catchability studies, NMFS performed standard survey tows (e.g., 83-122 trawl gear, 30 minute tow duration) as part of its annual EBS bottom trawl survey while BSFRF performed parallel tows within 0.5 nm using a nephrops trawl and 5 minute tow duration. Because the nephrops trawl has better bottom-tending performance than the 83112 gear, the BSFRF tows are hypothesized to catch all crab within the net path (i.e., to have selectivity equal to 1 at all crab sizes) and thus provide a measure of absolute abundance/biomass. The spatial footprints of the SBS studies for 2013-2017 are illustrated in Figure 16, while estimates of area-swept biomass for the study areas are compared in Figure 17 for the BSFRF and NMFS tows. Although the BSFRF gear is assumed to provide estimates of absolute abundance with the area surveyed, the relationship between these estimates and Tanner crab stock biomass is confounded by changes in the availability of Tanner crab to the BSFRF gear because the studies did not sample across the entire spatial extent of the population (in contrast to the full NMFS EBS bottom trawl survey).

## e. Survey catch-at-length

Bubble plots of NMFS EBS bottom survey size compositions for Tanner crab by sex and fishery region are shown in Figure 18. Distinct recruitment events (late 1970s, early 1990s, mid-2000s, early 2010s and possibly late 2010s) and subsequent cohort progression are evident in the plots, particularly in the western area. The absence of small male crab in the 2010-2016 period is notable, although there is evidence for new recruitment in the western area in 2016-2109, with perhaps some spillover to the eastern area lagged by a year at slightly larger sizes .

Based on the total abundance size compositions from the BSFRF-NMFS SBS studies (Figure 19), the BSFRF nephrops gear is in general (as expected) more selective for Tanner crab, particularly at smaller sizes ( $<60 \mathrm{~mm}$ CW) , than is the NMFS 83-112 gear. However, the size-specific catch ratio of the BSFRF survey to the NMFS survey appears to vary substantially across years, which one would not expect if gear-specific selectivity were, in general, constant. It is worth noting that the nephrops gear appear to give a much better indication of recruitment than the 83-112 gear does (e.g., Figure 19, survey year 2017).

Observed sample sizes for the NMFS survey size compositions, aggregated to the EBS regional level used in the assessment, are presented in Table 16. Given the large number of individuals sampled, a sample size of 200 is used to fit survey size compositions in the assessment model to prevent convergence issues associated with using the actual sample sizes.

## f. Other time series data.

Spatial patterns of abundance in the 2014-2019 NMFS bottom trawl surveys are shown in Figures 20-22 for immature males, mature males, immature females, mature females and legal males. There has been some suggestions that an extensive cold pool in the middle region of the EBS shelf may act to diminish relative crab densities in this region, particularly for mature males. The cold pool on the EBS shelf was extensive during the 2017 survey and absent during the 2018 and 2019 surveys, but the distribution of mature males did not change remarkably.

Annual maturity ogives for new shell males, based on chela height collections from the NMFS EBS bottom trawl survey, are shown in Figure 23 for years in which chela heights were measured to 0.1 mm
precision (i.e., since 2006). For each year, chela height:carapace width ratios for individual new shell crab were binned into 10 mm size bins, with the data split based on which management area (east or west of 1660 W longitude) it was collected in. The resulting histograms were analyzed to determine threshold sizes to discriminate mature from immature crab, and the fraction of mature crab was taken as the value of the resulting maturity ogive in the associated size bin (J. Richar, NMFS, pers. comm.). The areaspecific ogives were combined to obtain one for the entire EBS by weighting each by the estimated abundance of new shell males in each area by size bin.

Annual effort in the snow crab and BBRKC fisheries is used in the model to "project" bycatch fishing mortality rates backward in time from the period when data on bycatch in these fisheries exists (1992present). A table of annual effort (number of potlifts) is provided for the snow crab and BBRKC fisheries (Table 17).

## 3. Data which may be aggregated over time:

a. Growth-per-molt

Molt increment data collected for Tanner crab in the EBS in 2015-2017 and 2019 (Figure 24) is included in the parameter optimization for every model scenario considered in this assessment and is assumed to reflect growth rates over the entire model period.
b. Weight-at size

Weight-at-size relationships used in the assessment model for males, immature females, and mature females is depicted in Figure 25.
c. Size distribution at recruitment

The assumed size distribution for recruits to the population in the assessment model is presented in Figure 26.
4. Information on any data sources that were available, but were excluded from the assessment. The 1974 NMFS trawl survey was dropped entirely from the standardized survey dataset in 2015 due to inconsistencies in spatial coverage with the standardized dataset. Molt increment data from the Kodiak area in the Gulf of Alaska were not included in the assessment given the current use of molt increment data from the EBS to inform growth estimates. BSFRF survey data focused on Tanner crab recruitment (size compositions) have not yet been incorporated into the assessment.

## E. Analytic Approach

## 1. History of modeling approaches for this stock

Prior to the 2012 stock assessment, Tanner crab was managed as a Tier-4 stock using a survey-based assessment approach (Rugolo and Turnock 2011b). The Tier 3 Tanner Crab Stock Assessment Model (TCSAM) was developed by Rugolo and Turnock and presented for review in February 2011 to the Crab Modeling Workshop (Martel and Stram 2011), to the SSC in March 2011, to the CPT in May 2011, and to the CPT and SSC in September 2011. The model was revised after May 2011 and the report to the CPT in September 2011 (Rugolo and Turnock 2011a) described the developments in the model per recommendations of the CPT, SSC and Crab Modeling Workshop through September 2011. In January 2012, the TCSAM was reviewed at a second Crab Modeling Workshop. Model revisions were made during the Workshop based on consensus recommendations. The model resulting from the Workshop was presented to the SSC in January 2012. Recommendations from the January 2012 Workshop and the SSC, as well as the authors' research plans, guided changes to the model. A model incorporating all revisions recommended by the CPT, the SSC and both Crab Modeling Workshops was presented to the SSC in March 2012.

In May 2012 and June 2012, respectively, the TCSAM was presented to the CPT and SSC to determine its suitability for stock assessment and the rebuilding analysis (Rugolo and Turnock 2012b). The CPT agreed that the model could be accepted for management of the stock in the 2011/12 cycle, and that the stock should be promoted to Tier-3 status. The CPT also agreed that the TCSAM could be used as the basis for rebuilding analyses to underlie a rebuilding plan developed in 2012. In June 2012, the SSC reviewed the model and accepted the recommendations of the CPT. The Council subsequently approved the SSC recommendations in June 2012. For 2011/12, the Tanner crab was assessed as a Tier-3 stock and the model was used for the first time to estimate status determination criteria and overfishing levels.

Modifications have been made to the TCSAM computer code to improve code readability, computational speed, model output, and user friendliness without altering its underlying dynamics and overall framework. A detailed description of the 2013 model (TCSAM2013) is presented in Appendix 3 of the 2014 SAFE chapter (Stockhausen, 2014). Following the 2014 assessment, the model code was put under version control using "git" software and is publicly available for download from the GitHub website ${ }^{2}$.

A new model "framework", TCSAM02, was reviewed by the CPT and SSC in May/June 2017 and adopted for use in subsequent assessments as a transition to Gmacs. The new framework is a completelyrewritten basis for the Tanner crab model: substantially different model scenarios can be created and run by editing model configuration files rather than modifying the underlying code itself. Most importantly, no time blocks are "hard-wired" into the code-any time blocks are defined in the configuration files. In addition, the new frame work incorporates new data types (e.g., molt increment data, male maturity ogives), new survey data (e.g., the BSFRF surveys), and new fishery data (e.g., bycatch in the groundfish fisheries by gear type). The new model framework also incorporates status determination and OFL calculations directly within a model run, so a follow-on, stand-alone projection model does not need to be run (as was the case with TCSAM2013). This approach has the added benefit of allowing a more complete characterization of model uncertainty in the OFL calculation, because the OFL calculations are now included in Markov Chain Monte Carlo (MCMC) evaluation of a model's posterior probability distribution.

Most recently, the model code has been modified to allow fitting to molt increment observations, chela height data, and male maturity ogives. It has also been restructured to function in a management strategy evaluation (MSE) mode. The code for the TCSAM02 model framework is publicly available on GitHub ${ }^{3}$.

## 2. Model Description

## a. Overall modeling approach

TCSAM02 is a stage/size-based population dynamics model that incorporates sex (male, female), shell condition (new shell, old shell), and maturity (immature, mature) as different categories into which the overall stock is divided on a size-specific basis. For details of the model, the reader is referred to Appendix A.

In brief, crab enter the modeled population as recruits following the size distribution in Figure 26. An equal (50:50) sex ratio is generally assumed at recruitment (although can be set otherwise or estimated), and all recruits begin as immature, new shell crab. Within a model year, new shell, immature recruits are added to the population numbers-at-sex/shell condition/maturity state/size remaining on July 1 from the previous year. These are then projected forward to Feb. $15(\delta t=0.625 \mathrm{yr})$ and reduced for the interim effects of natural mortality. Subsequently, the various fisheries that either target Tanner crab or catch them as bycatch are prosecuted as pulse fisheries (i.e., instantaneously). Catch by sex/shell condition/maturity state/size in the directed Tanner crab, snow crab, BBRKC, and groundfish fisheries is

[^1]calculated based on fishery-specific stage/size-based selectivity curves and fully-selected fishing mortalities and removed from the population. The numbers of surviving immature, new shell crab that will molt to maturity are then calculated based on sex/size-specific probabilities of maturing, and growth (via molt) is calculated for all surviving new shell crab. Crab that were new shell, mature crab become old shell, mature crab (i.e., they don't molt) and old shell crab remain old shell. Population numbers are then adjusted for the effects of maturation, growth, and change in shell condition. Finally, population numbers are reduced for the effects of natural mortality operating from Feb. 15 to July 1 ( $\delta t=0.375 \mathrm{yr}$ ) to calculate the population numbers (prior to recruitment) on July 1.

Model parameters are estimated using a maximum likelihood approach, with Bayesian-like priors on some parameters and penalties for smoothness and regularity on others. Data components in the base model entering the likelihood include fits to mature survey biomass, survey size compositions, retained catch, retained catch size compositions, bycatch mortality in the bycatch fisheries, and bycatch size compositions in the bycatch fisheries.

## b. Changes since the previous assessment.

The model code has been revised to function in a management strategy evaluation mode, with specific computational routes defined for use as an operational model and as an estimation model. Fits to annual male maturity ogives were incorporated into the model last year, but with the assumption that these data would be aggregated to the same size bins as used in the model and other data. Now, this requirement has been loosened and the model can now fit ogives given using any size bin width. Finally, the model now allows specification and estimation of "availability" functions, similar to selectivity functions, that reflect the size-specific fraction of a section of the population (defined by sex, maturity state and shell condition) that can be encountered within a specific survey collection. This was necessary to incorporate the BSFRF SBS data into the assessment framework because these collections, in contrast to the complete NMFS EBS bottom trawl survey, do not encompass the entire Tanner crab stock.

## i. Methods used to validate the code used to implement the model

The TCSAM02 model framework was demonstrated to produce results that were exactly equivalent to those from the 2016 assessment model incorporating the changes listed in the previous table. TCSAM02 also underwent a review in July 2017 conducted by the Center for Independent Experts and has been further reviewed by the CPT in May 2017 and September 2017. Changes to model code are validated against results from the previous assessment model to ensure that modifications do not change the results of the previous assessment.

## 3. Model Selection and Evaluation

a. Description of alternative model configurations

The model selected for the 2018 assessment (Model 18AM17 from Stockhausen, 2018) provides the baseline model configuration for subsequent alternative model scenarios evaluated in this assessment. Here, the 2018 assessment model is designated "M19F00". The following tables provide a summary of the baseline model configuration, M19F00, for this assessment.

Model M19F00: Description of model population processes and survey characteristics.

| process | time blocks | description |
| :---: | :---: | :---: |
| Population rates and quantities |  |  |
| Population built from annual recruitment |  |  |
| Recruitment | 1949-1974 | In-scale mean + annual devs constrained as AR1 process |
|  | 1975+ | In-scale mean + annual devs |
| Growth | 1949+ | sex-specific |
|  |  | mean post-molt size: power function of pre-molt size |
|  |  | post-molt size: gamma distribution conditioned on pre-molt size |
| Maturity | 1949+ | sex-specific |
|  |  | size-specific probability of terminal molt |
|  |  | logit-scale parameterization |
| Natural mortalty | 1949-1979, | estimated sex/maturity state-specific multipliers on base rate |
|  | 1985+ | priors on multipliers based on uncertainty in max age |
|  | 1980-1984 | estimated "enhanced mortality" period multipliers |
| Surveys |  |  |
| NMFS EBS trawl survey |  |  |
| male survey q | 1975-1981 | In-scale |
|  | 1982+ | In-scale w/ prior based on Somerton's underbag experiment |
| female survey q | 1975-1981 | In-scale |
|  | 1982+ | In-scale w/ prior based on Somerton's underbag experiment |
| male selectivity | 1975-1981 | ascending logistic |
|  | 1982+ | ascending logistic |
| female selectivity | 1975-1981 | ascending logistic |
|  | 1982+ | ascending logistic |

Model M19F00: Description of model fishery characteristics.


Model M19F00: Description of model likelihood components.

| Component | Type | included in optimization | Distribution | Likelihood |
| :---: | :---: | :---: | :---: | :---: |
| TCF: retained catch | abundance | no | lognormal | males only |
|  | biomass | yes | norm2 | males only |
|  | size comp.s | yes | multinomial | males only |
| TCF: total catch | abundance | no | lognormal | by sex |
|  | biomass | yes | norm2 | by sex |
|  | size comp.s | yes | multinomial | by sex |
| SCF: total catch | abundance | no | lognormal | by sex |
|  | biomass | yes | norm2 | by sex |
|  | size comp.s | yes | multinomial | by sex |
| RKF: total catch | abundance | no | lognormal | by sex |
|  | biomass | yes | norm2 | by sex |
|  | size comp.s | yes | multinomial | by sex |
| GTF: total catch | abundance | no | lognormal | by sex |
|  | biomass | yes | norm2 | by sex |
|  | size comp.s | yes | multinomial | by sex |
| NMFS "0" survey | abundance | no | lognormal | by sex |
|  | biomass | yes | lognormal | by sex, for mature crab only |
|  | size comp.s | yes | multinomial | by sex/maturity |
|  | chela height data | no | -- | -- |
| NMFS "M" survey (males only, no maturity) | abundance <br> biomass <br> size comp.s | $\begin{aligned} & \text { no } \\ & \text { no } \\ & \text { no } \\ & \hline \end{aligned}$ | lognormal lognormal multinomial | all males <br> all males <br> all males |
| NMFS "F" survey (females only, w/ maturity) | abundance <br> biomass <br> size comp.s | $\begin{aligned} & \text { no } \\ & \text { no } \end{aligned}$ no | lognormal lognormal multinomial | by maturity classification by maturity classification by maturity classification |
| growth data | EBS only | yes | gamma | by sex |

The NMFS " 0 " survey refers to the "flavor" of the NMFS EBS bottom trawl survey data which has been fit in previous assessment models: maturity state (immature/mature) is determined outside the model for females based on morphological identification and for males on a size-dependent proportional basis using the Rugolo-Turnock maturity ogive. The NMFS "M" survey refers to a new, male-only "flavor" of the NMFS survey data in which maturity is not determined outside the model (males in the M survey have "undetermined" maturity). The NMFS " $F$ " survey is simply the female portion of the NMFS " 0 " survey data configured as a separate data file to accompany the NMFS "M" survey data file.

As per CPT recommendation, the following model scenarios were evaluated as part of this assessment:

| model <br> scenario | number of <br> parameters | objective <br> function value | max <br> gradient | Jitter <br> runs | \# runs <br> converged <br> to MLE | scenario description |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| M19F00 | 357 | $2,962.17$ | 0.0004 | -- | -- | 2018 assessment model (18AM17) |
| M19F00a | 357 | $3,025.43$ | 0.0003 | -- | -- | M19F00 with revised ADFG data for 1990+ crab fisheries |
| M19F01 | 363 | $3,368.11$ | 0.0002 | 3,000 | 94 | M19F00a updated for 2018/19 (base model for 2019) |
| M19F02 | 363 | $3,521.89$ | 0.0004 | -- | -- | M19F01 + 2006+ observed male maturity data |
| M19F03 | 343 | $3,467.75$ | 0.0013 | 3,000 | 72 | M19F02 - male maturity characterized by Rugolo/Turnock maturity ogive |
| M19F04 | 628 | $3,578.47$ | 0.0004 | 3,000 | 7 | M19F01 + 2013-2017 BSFRF/NMFS side-by-side data |
| M19F05 | 608 | $3,674.61$ | 0.0004 | 3,000 | 5 | M19F03 + 2013-2017 BSFRF/NMFS side-by-side data |

As noted previously, M19F00 is the 2018 assessment model and data ("18AM17" in the 2018 assessment). For M19F00a, the "historical" crab fishery catch data is replaced with the "current" data provided by ADFG through 2017/18. This represents a bridging scenario to the 2019 assessment and allows a characterization of the effects of the changes in fishery data on model outcomes without the confounding effects of new data for 2018/19. M19F01 is M19F00a updated with 2018/19 data. It represents "business as usual" in terms of the development of the assessment model. M19F02 includes fits to the male maturity ogive data developed from 2006-onward chela height data collections during the NMFS EBS bottom trawl survey. It also, however, fits the male data in the NMFS " 0 " dataset with male maturity determined outside the assessment model using the Rugolo-Turnock maturity ogive. This is a bridging scenario that provides a transition to M19F03, which drops the fits to the male data in the NMFS " 0 " dataset and relies strictly on the male maturity ogive data (and the size composition data) to inform the model estimates of the size-specific probability of terminal molt for males. M19F04 constitutes a different development "fork" based on M19F01, and includes fits to the biomass and size composition data from 2013-2017 BSFRF and NMFS side-by-side studies. In this scenario, the BSFRF survey is assumed to be fully-selected across the size ranges in the model ( $>25 \mathrm{~mm} \mathrm{CW}$ ) and thus provides estimates of absolute size-specific abundance within a given study area. Sex-specific "availability" functions are estimated in the model to relate the size-specific study-area abundance estimates to population abundance. The final scenario, M19F05, reflects a merging of the M19F01-M19F02-M19F03 fork with the M19F01-M19F04 fork.

The number of estimated parameters, the final value of the objective function for each converged scenario and the maximum gradient of the objective function at the converged solution are listed as well in the table above. The total objective function values, however, cannot be directly compared between scenarios because each scenario fits different datasets. Convergence for the four scenarios under consideration for status determination and OFL-setting ( M19F01, M19F03, M19F04, and M19F05) was evaluated using parameter jittering, with a total of 3,000 runs initiated for each scenario. Of these runs, generally a large number failed to converge at all and a smaller number converged to local minima smaller than the maximum likelihood (ML) solution. About $3 \%$ of the runs found the (presumed) ML solution in M19F01, about $2.4 \%$ for M19F03, and only about $0.2 \%$ in M19F04 and M19F05. In the interest of time and computing resources, the bridging scenarios were not jittered but instead were initialized using the final parameter estimates from the base scenario in the bridge.

M19F03 is the author's preferred model, as explained below.
b. Progression of results from the previous assessment to the preferred base model

The following table summarizes basic model results from the 2018 assessment model (M19F00) and the 6 scenarios considered here (results from the author's preferred model are highlighted):

| Model <br> Scenario | average <br> recruitment <br> millions | Final MMB | BO | Bmsy | Fmsy | MSY | Fofl | OFL | projected <br> MMB | projected MMB <br> $/ B m s y ~$ |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| M19F00 | 223.63 | 66.64 | 86.55 | 30.29 | 0.74 | 12.75 | 0.74 | 20.87 | 35.95 | 1.19 |
| M19F00a | 284.28 | 82.05 | 94.24 | 32.99 | 0.89 | 14.58 | 0.89 | 27.90 | 41.52 | 1.26 |
| M19F01 | 316.79 | 68.79 | 100.85 | 35.30 | 0.81 | 15.58 | 0.81 | 22.54 | 35.66 | 1.01 |
| M19F02 | 367.48 | 71.54 | 105.59 | 36.96 | 1.11 | 17.89 | 1.03 | 24.75 | 34.63 | 0.94 |
| M19F03 | 393.84 | 82.61 | 118.96 | 41.64 | 1.18 | 19.49 | 1.12 | 29.48 | 39.68 | 0.95 |
| M19F04 | 377.28 | 74.03 | 106.76 | 37.37 | 0.87 | 16.87 | 0.87 | 24.87 | 37.50 | 1.00 |
| M19F05 | 418.73 | 80.33 | 116.44 | 40.75 | 1.21 | 19.40 | 1.14 | 28.58 | 38.42 | 0.94 |

## c. Evidence of search for balance between realistic (but possibly over-parameterized) and simpler

 (but not realistic) models.It was noted at the May 2018 CPT meeting that it was not biologically realistic that male Tanner crab less than 60 mm CW had undergone their terminal molt, although there were a few males collected in the NMFS EBS bottom trawl surveys below 60 mm CW that were classified as mature using the chela height data. It was similarly recognized that it was probably biologically unrealistic for female crab less than 40 mm CW to have undergone terminal molt. This actually resulted in simpler, but more realistic models, in scenarios where these constraints were implemented (scenarios M19F03 and M19F05).

A future avenue for exploration in this regard is to estimate the "availability" functions outside the model that are required to allow the SBS data to inform NMFS survey catchability, rather than estimating these functions inside the model. Because the availability functions are estimated in the model using a nonparametric approach to allow for an arbitrary, but smoothly-varying, shape, this adds 48 additional parameters per included SBS study ( 32 for the male availability function, 16 for the female one). Instead, the availability functions can be estimated outside the model using the size-specific ratios of the size composition data from each NMFS SBS dataset to the corresponding data from the full NMFS dataset, perhaps with a smoothing penalty applied to the resulting curve. In this respect, there would be no need to fit the NMFS SBS data within the model (as is done now) at all.

## d. Convergence status and convergence criteria

As noted above, convergence in all non-bridging models was assessed by running each model 3,000 times with randomly-selected ("jittered") initial parameter values for each run. For each model, a number of these jitter runs failed, primarily because the initial values for the growth parameters resulted in the mean post-molt size being smaller than the pre-molt size. Of those that converged, the run with the smallest objective function value and smallest maximum gradient was selected as the "converged" model, if it was also possible to invert the associated hessian and obtain standard deviation estimates for parameter values. Theoretically, all gradients at a minimum of the objective function would be zero. However, because numerical methods have finite precision, the numerical search for the minimum is terminated after either achieving a minimum threshold for the maximum gradient or exceeding the maximum number of iterations. As noted previously, many more runs converged to the final (presumably) ML solution for scenarios M19F01 and M19F03 than for M19F04 and M19F05, but this is not too surprising given the much larger number of estimated parameters for the latter two scenarios.

## e. Sample sizes assumed for the compositional data

Input sample sizes used for compositional data are listed in Tables 9-13 for fishery-related size compositions. Input sample sizes for all survey size compositions were set to 200 , which was also the
maximum allowed for the fishery-related sample sizes. Otherwise, input sample sizes were scaled as described in Stockhausen (2014, Appendix 5):

$$
S S_{y}^{i n p}=\min \left(200, \frac{S S_{y}}{(\overline{S S} / 200)}\right)
$$

where $\overline{S S}$ was the mean sample size for all males from dockside sampling in the directed fishery.

## f. Parameter sensibility

Limits were placed on all estimated parameters in all model scenarios primarily to provide ranges for jittering initial parameter values. Although these limits, for the most part, did not constrain parameter estimates in the converged models, some parameters were found to be at, or very close, to one of the bounds placed on them. These parameters are listed for the scenarios in Tables 18 and 19. The CPT and SSC have both expressed concerns regarding parameters estimated at their bounds, as such results frequently violate assumptions regarding model convergence, parameter uncertainty estimates, and suggest that model suitability may be improved by widening the bounds or re-parameterizing the model. The logit-scale parameter describing the retention of male crab at large (asymptotic) sizes prior to 1997 was estimated at its upper bound (15) in all model scenarios. Because retention can only go as high as 1 on the arithmetic scale, and a logit-scale value of 15 corresponds to an arithmetic scale value of 0.9999997 , this parameter should be fixed in future models. In a similar fashion, the logit-scale parameter describing the probability of terminal molt for males in the largest size bin $(180+\mathrm{mm}$ CW $)$ reached its bound of 15 in scenarios M19F00a, M19F01, M19F02, and M19F04 while that describing the probability of terminal molt for females in the smallest size bin ( $25-30 \mathrm{~mm} \mathrm{CW}$ ) reached its lower bound ( -15 ). These were fixed in M19F03 and M19F05, based on assumptions of minimum and maximum sizes for immature crab at terminal molt, such that the corresponding probabilities of terminal molt in these size bins were 1 or 0 .

Survey catchability parameters for the 1975-1981 time period (pQ[1] and pQ[3]) were also estimated at their lower bound (0.5). This might not be unreasonable given the reduced areal coverage of these surveys relative to later surveys and the spatial limits of the Tanner crab stock. However, it would be worthwhile to explore the effect of reducing these bounds. The remaining parameters are related to selectivity functions describing the size-specific capture efficiency of the fisheries and surveys. Two at their lower bounds are probably inconsequential ( $\mathrm{pS} 2[10]$ and $\mathrm{pS} 4[1]$ ) and are related to the ascending and descending slopes of the dome-shaped selectivity describing male bycatch in the snow crab fishery prior to 1997. A double-normal is used to describe the dome shape, but an alternative function (e.g., a single normal) might have better estimation properties. The size at $50 \%$ selected was estimated at its upper bound ( 90 mm CW ) for NMFS survey selectivity in the 1975-1981 time period $\mathrm{pS1}$ [1]). This results in an almost linear function, rather than asymptotic, across the size range. This result may reflect the changing interaction between the areas surveyed (availability) and the gear selectivity in this time period as the survey gradually extended from the southeast shelf and Bristol Bay where adult males were prevalent to the north and west where more immature males would be encountered, effectively "seeing" relatively more large males than small males. Two other survey-related selectivity parameters, describing the size difference between crab at $50 \%$ and $95 \%$ selected) were estimated at their upper bounds for the both males and females in the NMFS EBS trawl survey in the 1982-present time period (pS2[2] and pS2[4]). The selectivity functions are assumed to be logistic, with the other estimated parameter being the size at $95 \%$ selected. The practical consequence of this is that small crab (females in particular) are described as fairly well-selected ( $>50 \%$ for females) relative to fully-selected (sex-specific) large crab. This result may reflect conflicts from between the model assumption of equal sex ratios for recruitment in the 25-40 mm CW range, apparent equal abundances and spatial patterns for males and females at small sizes in the NMFS EBS survey, and assumed logistic selectivity. The selectivity parameter describing the size at $50 \%$ selected for males in the groundfish fisheries during 1987-1996 was estimated in all scenarios at its lower
bound ( 40 mm CW ), probably a consequence of fairly substantial catches of small crab in some years (e.g., 1993, Figure 12). Finally, three parameters at their upper bounds ( pS 1 [23], $\mathrm{pS1}[24]$, and $\mathrm{pS1}$ [27]) are related to the size at $95 \%$ selected in the BBRKC fishery in the 1997-2004 (males) and 2005+ (males and females) time periods. The upper bounds ( 180 for males, 140 for females) were selected to reflect the largest possible sizes reasonably expected in the model, so the resulting selectivity functions are essentially positively-sloped linear functions with values fixed at 0.95 at the parameter bound because the other estimated logistic parameter estimates a large size at $50 \%$ selected (see selectivity curves in Figure 46).

Estimates of parameter uncertainty, approximations calculated by inverting the model hessian and using the "delta" method, were obtained from each converged model's ADMB "std" file (Tables 20-33). Extremely large uncertainties were obtained for parameters related to the NMFS trawl survey selectivity for females after 1981 for all scenarios that estimated these parameters, unless the estimates hit one of the bounds (Table19). A number of other selectivity-related parameters, while not at one of their bounds, have large uncertainties associated with the estimates (e.g., the $95 \%$-selected size for female bycatch in the BBRKC fishery, Table 31). These may reflect indeterminacy between the estimated capture rates for fully-selected crab and these parameters in determining the effective capture rates on large crab.

Unweighted negative log-likelihoods (NLLs) and their associated (weighted) components in the model objective function are compared for fits to data for the scenarios with 2018/19 data in Tables 34-36. Comparison of the unweighted versions gives some insight into the tradeoffs between fitting to different datasets in the model scenarios. For example, M19F03 doesn't actually fit the NMFS "0" dataset mature male biomass (i.e., the likelihood is not included in the objective function that is optimized) whereas M19F00 does, while the latter doesn't fit the NMFS "M" dataset biomass and the former does. The NLL for M19F00 from the NMFS " 0 " biomass is $\sim 17$ likelihood units better than that for M19F03 but the NLL for M19F03 is $\sim 50$ likelihood units better than that for M19F00. Another way of assessing model fit is to examine the average root mean square errors (RMSE) associated with differences between observed and predicted values (Table 38). In this regard, M19F03 fits NMFS "0" male size compositions (rmse=490.64) slightly worse than M19F01 (rmse=487.07) but fits the NMFS "M" size compositions better (185.98 vs. 195.51).

## g. Criteria used to evaluate the model or to choose among alternative models

None of the model scenarios evaluated in this assessment were directly comparable using likelihood criteria because different combinations of datasets were fit in each scenario. Consequently, the criteria used to evaluate the alternative models were based primarily on: 1) goodness of fit (assessed using the unweighted NLLs and RMSEs for different datasets, even when the datasets were not included in the likelihood), 2) parameter sensibility, and 3) biological realism.

## h. Residual analysis

Standardized residuals to model fits were plotted and examined for all data components, including datasets that were not included (weighted 0 ) in the model objective function.

## i. Evaluation of the model(s)

All scenarios fit their respective catch biomass data quite well (noting that different crab fishery data is fit in M19F00 and the other scenarios; Figures 27-30), although scenarios M19F01-M19F05 slightly underestimate total bycatch biomass in the groundfish fisheries from 1991-2013. The model fits to fishery size compositions are similar in quality to what has been obtained in previous assessments: not terrible, but not really great either. The fits to retained catch size compositions are the best overall and are essentially identical for all the scenarios excluding M19F00 (Appendix B, Figures 42-45). Some less good fits seem to be associated with a closure of one of the management areas (e.g., 2005, 2009, 2018). Fits to total catch size compositions in the directed fishery (Appendix B, Figures 46-51) are reasonably
good (except for 1996, when sample sizes were very small) but continue to somewhat overestimate the catch of large males since 2013. Again, however, the fits are almost identical among the scenarios. The fits to the total catch size compositions from the snow crab fishery are somewhat worse, particularly in the early 1990s, than those for the directed fishery-to be expected given the differences in the numbers of crab sampled. Some slight differences can be seen among the scenarios in the fits to the total catch size compositions form the groundfish fisheries (Appendix B, Figures 58-67), but the fits themselves are not particularly good. Selectivity functions for the groundfish fisheries are estimated for three different time periods between 1973 and the present, but underlying changing in areas targeted and gear composition may occur on shorter time scales that contribute to the lack of fit. The fits to total catch in the BBRKC fishery (Appendix B, Figures 68-73) are the poorest, consistent with the low observer sample sizes for Tanner crab, particularly females, in this fishery. However, the disagreement between predicted and observed male compositions in the early 1990s is rather puzzling.

The fits to survey biomass (Figures 31-34) are somewhat poorer than those to fishery catch biomass, a not unexpected result because fitting to the fishery catch biomass data was weighted heavily (20x) in the model objective functions. The most notable differences among the fits are that scenarios M19F03 and M19F05 (which fit the male maturity ogive data) both follow the low female biomasses in the 1980's better than the other scenarios do. While all the scenarios estimate declining trends in mature survey biomass starting in 2014, all are biased somewhat high relative to the data.

In general, the predicted survey size compositions are remarkably similar among the scenarios (Appendix B, Figures 1-41), but scenarios that fit the male maturity ogive data (M19F02, M19F03, and M19F05) tend to estimate slightly higher proportions of "mature" (as categorized by the Rugolo-Turnock maturity ogive) males at smaller sizes, lower proportions of immature females at small sizes, and lower proportions of mature females at larger sizes, than occurs in the other scenarios. Somewhat surprisingly, the proportions for "immature" males are almost identical among the scenarios. The two scenarios that fit the SBS datasets also estimate almost identical size compositions which fit the observed ones fairly well for both the NMFS and BSFRF data. In particular, the models capture the recruitment event in 2017 well in both datasets, although it occurs much more strongly in the BSFRF dataset.

All the scenarios fit female growth equally well, but over-predict male molt increments at larger sizes, with M19F03 and M19F05 being the most biased (Figure 35). In contrast, M19F03 and M19F05 fit the male maturity ogive data better than the other scenarios, but all scenarios tend to underestimate the fraction mature in any size bin, although this is not true in all cases (Figure 36).

Estimated capture rates in the directed fishery (Figure 37) follow the same temporal patterns in all scenarios, with the largest peak in 1979 and a lesser peak in 1991 or 1992. However, the relative levels vary among the scenarios, reflecting differences in recruitment (see below) rather than differences in estimated size-specific capture functions (Figures 38-41) or retention functions (Figure 42), which are essentially identical (the differences between M19F00 and M19F00a in 1990 and 1991 are primarily due to changes in the underlying snow crab data).

Estimated capture rates in the snow crab (Figure 43) also exhibited similar temporal patterns. Scenarios M19F00 and M19F00a differ substantially in level due to changes in the underlying crab data, which changes the selectivity function estimated for the early 1990s, as well as differences in recruitment. The capture rates estimated in the other scenarios are much more similar to one another, and primarily reflect smaller differences in estimated recruitment. Estimated selectivity functions for these scenarios were almost identical for the three time periods in which they were estimated, with the only substantial difference being that the curves estimated in M19F04 and M19F05 for the pre-1997 time period were right-shifted to larger sizes by one or two size bins (Figure 44).

Estimated capture rates in the BBRKC fishery (Figure 45) exhibited similar temporal patterns among the scenarios, as well. Scenarios M19F00 and M19F00a were much more similar in level than was the case for the snow crab fishery because the underlying data was not substantially changed. The levels of the capture rates for females in scenarios M19F00a-M19F05 appear fairly variable, but the absolute scale is very small (on the order of 0.04 relative to 0.5 for the directed fishery) and the variability is primarily due to the small scale of the associated catches. The estimated selectivity functions (Figure 46) were also slightly different among the scenarios for females, while those for males were basically identical.

As with the other fisheries, estimated capture rates in the groundfish fisheries (Figure 47) exhibited similar temporal patterns in all scenarios but differed somewhat in absolute level. In addition, M19F00a exhibited substantially higher levels in the 1991-2012 time period than did scenarios M19F01-M19F05, which were all quite similar in level. Estimated male selectivity curves exhibited a fair amount of variation among scenarios during the 1997-2004 time period, while the selectivity curves for both males and females in the 2005+ period exhibited somewhat less variability and those in the pre-1997 period were essentially the same (Figure 48).

The change in the crab fishery data had fairly large effects on estimates of survey catchability and selectivity functions (Figures 49-51). Although estimated catchability was the same for scenarios M19F00 and M19F00a in the pre-1982 time period, the estimated size-at-50\% selected for the male and female selectivity functions shifted substantially to larger sizes (more so for males than females) such that many more small and intermediate size crab were "invisible" to the survey during this time period. In the 1982+ time period, catchability was estimated to be smaller in M19F00a while the selectivity functions remained similar (the male function was slightly shifted toward larger sizes), with the result that crab of all sizes were effectively invisible to the survey in M19F00a. Survey catchability in the pre-1982 time period did not change in the M19F01-M19F05 scenarios, no did the male selectivity function, but the female selectivity function shifted to somewhat larger sizes in scenarios M19F03 and M19F05. Survey catchabilities did change in all of these scenarios in the $1982+$ time period for both males and females, with the largest values estimated in M19F00a while the smallest value for females was estimated in M19F02 and the smallest for males in M19F03. In general, including the male maturity ogive data in the model fit decreased the catchability for both sexes. Selectivity for males in the pre-1982 time period was essentially unchanged among M19F01-M19F05 scenarios, while including the male maturity ogive data shifted female selectivity $\sim 5 \mathrm{~mm}$ to larger sizes. The selectivity functions for both sexes differed among these scenarios for the $1982+$ time period, shifting the $50 \%$-selected size substantially to larger size for females in scenarios M19F03, M19F04, and M19F05 but only slightly to larger size for males.

Survey availability, estimated in scenarios M19F04 and M19F05 for the SBS datasets, were similar to one another (Figure 52). Curves for females were fairly similar for 2013-15, increasing with size, but different from those for 2016 and 2017, which decreased with size. For males, larger males in the 100-150 mm CW range tended to be most available to the survey. In 2013-15, small males were mostly unavailable while in 2016-17 the smallest were much more available while intermediate-sized males were relatively less available. It is possible to estimate empirical versions for the availability functions using the ratio of crab abundance in the NMFS SBS dataset to that in the NMFS " 0 " dataset by size bin. These empirical availability functions provide a check on the estimated versions. However, they do not particularly resemble the estimated versions (Figure 53), except for females in 2013.

Another effect of the revised crab fishery data is to slightly increase the estimated rate of M on mature mature females and to slightly decrease them on males, outside the 1980-84 "enhanced mortality" period when the effect is to increase the rates for both sexes (M19F00a compared with M19F00; Figure 54). Fitting the male maturity ogive data rather than mature male survey biomass based on the RugoloTurnock maturity ogive (M19F03, M19F05) results in a much reduced estimate of M on mature males in the enhanced mortality period while it is elevated for mature females.

The estimated probability of terminal molt by size is almost the same for all scenarios, but is shifted to smaller sizes by $\sim 5 \mathrm{~mm}$ CW for the scenarios that fit the male maturity ogive data (scenarios M19F02, M19F03, and M19F05; Figure 55). Mean growth, as well, is similar across all scenarios for females while the scenarios that fit the male maturity ogive data yield slightly higher estimates of growth for males at large pre-molt sizes (Figure 56).

Estimated recruitment time series exhibit similar temporal patterns in all scenarios, but differ in overall scale, with the largest difference occurring between M19F00 (the 2018 assessment model) and M19F00a, the 2018 assessment model with the revised crab fishery data (Figures 57-58). The good news for the stock a few years in the future is that all the scenarios estimate recruitment during 2016-18 was much larger than during 2011-2015. The bad news is that all the scenarios estimated a declining trend in mature male and female biomass (MMB and MFB, evaluated on Feb. 15 for each year) over the past 4-5 years since a recent high in 2014 (or 2015, depending on scenario; Figures 59-60). Across the time series, the estimated trajectories for mature biomass also follow similar temporal trends but differ in scale. Unsurprisingly, similar trends were also estimated for the mature components of population biomass (evaluated on July 1 for each year; Figure 61). However, trends in immature biomass reflect the estimated recent recruitment trends and have been increasing in all scenarios for the past two years following a low point not seen since the early 1990s.

The author's preferred model, M19F03, fits all of the datasets reasonably well and includes fits to "observed" new shell male maturity ogives derived for years after 2005 when chela height data was collected in the NMFS EBS bottom trawl survey. It also drops the fits to immature/mature male categories created by applying the Rugolo-Turnock maturity ogive to male abundance and biomass by size outside the model. It does not fit the BSFRF SBS datasets, but doing so (i.e. M19F04, M19F05) does not seem to substantially change the estimates of catchability for the NMFS EBS bottom trawl survey or population quantities such as recruitment and mature biomass in the manner one would expect (higher estimates of catchability, lower estimates for population quantities). In addition, the manner in which "availability" is handled in the scenarios that fit the SBS data is somewhat problematic in terms of potential confounding between the ability to estimate availabilities for the BSFRF surveys and the ability to estimate catchabilities for the NMFS surveys. Finally, the estimated availability functions are somewhat inconsistent with empirical versions derived from the full NMFS survey and the NMFS SBS studies.

## 4. Results (best model(s))

Scenario M19F03 was selected as the author's preferred model for the 2019 assessment.

## a. List of effective sample sizes, the weighting factors applied when fitting the indices, and the

 weighting factors applied to any penalties.Input and effective sample sizes for size composition data fit in the model are listed in Tables 40-45 from the 2018 assessment model and scenario M19F03. A weighting factor of 20 (corresponding to a standard deviation of 0.158 ) was applied to all fishery catch biomass likelihood components to achieve close fits to catch biomass time series.

## b. Tables of estimates:

i. All parameters

Parameter estimates and associated standard errors, based on inversion of the converged model's Hessian, are listed in Tables 20-34.

## ii. Abundance and biomass time series, including spawning biomass and MMB.

Estimates for mature survey biomass, by sex, are listed in Table 46and for mature biomass at mating, by sex, in Table 47 for the 2018 assessment model and the author's preferred model, M19F03. Due to the
size of the tables, the numbers at size for females and males by year in 5 mm CW size bins for scenario M19F03 are available online as zipped csv files (see Tables 48 and 49, respectively).

## iii. Recruitment time series

The estimated recruitment time series from the 2018 assessment and M19F03 are listed in Table 50.
iv. Time series of catch divided by biomass.

A comparison of catch divided by biomass (i.e., exploitation rate) from the 2017 assessment and 18C2a is listed in Table 51.
c. Graphs of estimates

Graphs of estimates from the preferred scenario, M19F03 have been discussed above in the "Model Selection" section.

## i. Fishery and survey selectivities, molting probabilities, and other schedules depending on parameter estimates.

Graphs of estimated selectivity curves for the directed fishery are shown in Figures 39-42, for the snow crab fishery in Figure 45, for the BBRKC fishery in Figure 47, and for the groundfish fisheries in Figure 49. Estimated retention curves are shown in Figure 43. Graphs of selectivity curves for the NMFS survey are shown Figure 51; graphs of estimated availability curves from the NMFS SBS studies are shown in Figure 53. Natural mortality estimates are shown in Figure 55, terminal molt probabilities are shown in Figure 56, and mean growth rates (molt increments) are shown in Figure 57.
iii. Estimated full selection F over time

Graphs of time series of estimated fully-selected F (total catch capture rates, not mortality) on males in the directed fishery and bycatch in the snow crab, BBRKC and groundfish fisheries are shown in Figures $38,44,46$, and 48.
ii. Estimated male, female, mature male, total and effective mature biomass time series Estimates of the time trends in population biomass for mature and immature components of the stock are shown by sex in Figure 62. Mature male and female biomass trends (MMB and MFB) are shown in Figures 60 and 61.
iv. Estimated fishing mortality versus estimated spawning stock biomass

See Figure 65.

## v. Fit of a stock-recruitment relationship, if feasible.

Not available.

## e. Evaluation of the fit to the data:

i. Graphs of the fits to observed and model-predicted catches

Graphs of fits to observed catches are provided in Figure 26 for retained and total catch in the directed fishery and in Figures 27-29 for total catch in the snow crab, BBRKC, and groundfish fisheries. Fits to NMFS survey biomass are shown for mature crab and all males and females by maturity state in Figures 30 and 31, respectively. Fits to survey biomass in the SBS studies are shown in Figures 32 and 33.

## ii. Graphs of model fits to survey numbers

Not available.
iii. Graphs of model fits to catch proportions by size class

See Appendix B for model fits to annual catch proportions by size class.
iv. Graphs of model fits to survey proportions by size class

See Appendix B for model fits to annual survey proportions by size class.
v. Marginal distributions for the fits to the compositional data.

See Appendices C and D for marginal distributions of fits to the compositional data.
vi. Plots of implied versus input effective sample sizes and time-series of implied effective sample sizes.
See Appendices C and D for plots of implied and input sample sizes. For the most part, the implied effective sample sizes tend to be substantially larger than the input values.
vii. Tables of the RMSEs for the indices (and a comparison with the assumed values for the coefficients of variation assumed for the indices).
RMSEs for fits to various datasets are provided in Tables 37-39.
viii. Quantile-quantile ( $q-q$ ) plots and histograms of residuals (to the indices and compositional data) to justify the choices of sampling distributions for the data.
Due to time constraints, quantile-quantile ( $\mathrm{q}-\mathrm{q}$ ) plots and histograms of residuals were not completed for the assessment.
f. Retrospective and historic analyses (retrospective analyses involve taking the "best" model and truncating the time-series of data on which the assessment is based; a historic analysis involves plotting the results from previous assessments).
i. Retrospective analysis (retrospective bias in base model or models).

Due to time constraints, retrospective analyses were not completed for the assessment.
ii. Historical analysis (plot of actual estimates from current and previous assessments).

Due to time constraints, an historical analysis was not completed for the assessment.
g. Uncertainty and sensitivity analyses

MCMC runs were completed for scenario M19F03 to explore model uncertainty. The model was run for four chains, which 10 million iterations each, with a burn-in period of $1,000,000$ iterations and keeping results from every $1,000^{\text {th }}$ iteration to reduce serial autocorrelation, which yielded 4000 samples per chain. Mixing appeared to be sufficient, but this can be difficult to evaluate. This run provides empirical posterior distributions for model parameters and selected derived quantities, including OFL-related quantities.

Time constraints (the MCMC run took several days to complete) did not allow a full exploration of the MCMC results. Summary results for the objective function and OFL-related quantities (Figure 62) indicates that they are reasonably well-behaved and normally-distributed, and do not exhibit unexpected correlation structures (e.g., Fofl $_{\text {OF }}$ and $\mathrm{F}_{\text {MSY }}$ are expected to be highly correlated, so this is not cause for concern).

## F. Calculation of the OFL and ABC

## 1. Status determination and OFL calculation

EBS Tanner crab was elevated to Tier 3 status following acceptance of the TCSAM by the CPT and SSC in 2012. Based upon results from the model, the stock was subsequently declared rebuilt and not overfished. Consequently, EBS Tanner crab is assessed as a Tier 3 stock for status determination and OFL setting.

The (total catch) OFL for 2018/19 was 20.87 thousand t while the total catch mortality was 1.90 thousand $t$, based on applying mortality rates of 1.000 for retained catch, 0.321 to bycatch in the crab fisheries, and 0.800 to bycatch in the groundfish fisheries to the model-estimated catch by fleet for 2018/19. Therefore overfishing did not occur.

Amendment 24 to the NPFMC fishery management plan (NPFMC 2007) revised the definitions for overfishing for EBS crab stocks. The information provided in this assessment is sufficient to estimate overfishing limits for Tanner crab under Tier 3. The OFL control rule for Tier 3 is (Figure 63):

$$
\begin{array}{llcc}
\hline B, F_{35 \%}, B_{35 \%} \cdot{ }^{3} & \text { a. } \frac{B}{B_{35 \% * *}}>1 & F_{\text {OFL }}=F_{35 \%} * & \\
& \text { b. } \beta<\frac{B}{B_{35 \%} *} \leq 1 & F_{\text {OFL }}=F^{*} * \frac{B}{3_{35} \%} \frac{B}{B_{35 \%}}-\alpha & \text { ABC } \leq(1-\text {-by }) * \text { OFL } \\
& \text { c. } \frac{B}{B_{35 \%} *} \leq \beta & \begin{array}{c}
\text { Directed fishery } F=0 \\
F_{\text {ofL }} \leq F_{\text {MSY }}{ }^{\dagger}
\end{array} & \\
\hline
\end{array}
$$

and is based on an estimate of "current" spawning biomass at mating ( $B$ above, taken as the projected MMB at mating in the assessment year) and spawning biomass per recruit (SBPR)-based proxies for $\mathrm{F}_{\text {MSY }}$ and $B_{\text {MSY }}$. In the above equations, $\alpha=0.1$ and $\beta=0.25$. For Tanner crab, the proxy for $\mathrm{F}_{\text {MSY }}$ is $\mathrm{F}_{35 \%}$, the fishing mortality that reduces the SBPR to $35 \%$ of its value for an unfished stock. Thus, if $\phi(F)$ is the SBPR at fishing mortality $F$, then $\mathrm{F}_{35 \%}$ is the value of fishing mortality that yields $\phi(F)=0.35 \cdot \phi(0)$. The Tier 3 proxy for $\mathrm{B}_{\mathrm{MSY}}$ is $\mathrm{B}_{35 \%}$, the equilibrium biomass achieved when fishing at $\mathrm{F}_{35 \%}$, where $\mathrm{B}_{35 \%}$ is simply $35 \%$ of the unfished stock biomass. Given an estimate of average recruitment, $\bar{R}$, then $B_{35 \%}=$ $0.35 \cdot \bar{R} \cdot \phi(0)$.

Thus Tier 3 status determination and OFL setting for 2019/20 require estimates of $B=\mathrm{MMB}_{2019 / 20}$ (the projected MMB at mating time for the coming year), $\mathrm{F}_{35 \%}$, spawning biomass per recruit in an unfished stock $(\phi(0))$, and $\bar{R}$. Current stock status is determined by the ratio $B / \mathrm{B}_{35 \%}$ for Tier 3 stocks. If the ratio is greater than 1 , then the stock falls into Tier 3 a and $\mathrm{F}_{\text {OFL }}=\mathrm{F}_{\mathrm{MSY}}=\mathrm{F}_{35 \%}$. If the ratio is less than one but greater than $\beta$, then the stock falls into Tier 3 b and $\mathrm{F}_{\text {OFL }}$ is reduced from $\mathrm{F}_{35 \%}$ following the descending limb of the control rule (Figure 19). If the ratio is less than $\beta$, then the stock falls into Tier 3c and directed fishing must cease. In addition, if $B$ is less than $1 / 2 \mathrm{~B}_{35 \%}$ (the minimum stock size threshold, MSST), the stock must be declared overfished and a rebuilding plan subsequently developed.

In 2015, the SOA's Board of Fish, under petition from the commercial Tanner crab fishing industry, changed the minimum preferred size for crab in the area east of $166^{\circ} \mathrm{W}$ longitude in calculations used for setting TACs from 138 mm CW (not including lateral spines) to 125 mm CW. The minimum preferred size in the area west of $166^{\circ} \mathrm{W}$ remained the same ( 125 mm CW). In assessments before 2017, an attempt was made to account for retention of slightly ( 10 mm CW) smaller crab in the directed fishery in the western area. Because the preferred size is now the same in both areas, the OFL is calculated assuming both selectivity (as previously) and retention (new) curves are the same in both areas.

In assessments before 2017, a separate "projection model" was used to determine OFL based on results from the assessment model. The estimated coefficient of variation for the estimate of final MMB was used to characterize model uncertainty and provided a calculational basis for determining an empirical probability density function (pdf) for OFL based on sampling final MMB from its assumed pdf. Since the transition to TCSAM02 in 2017, the OFL is calculated within the assessment model based on equilibrium calculations for Fofl and projecting the state of the population at the end of the modeled time period one
year forward assuming fishing mortality at $\mathrm{F}_{\text {ofl }}$. Using MCMC, one can thus estimate the pdf of OFL (and related quantities of interest) incorporating full model uncertainty.

To calculate the $\mathrm{F}_{\text {OFL }}$, the fishery capture rate for males in the directed fishery is adjusted until the longterm (equilibrium) MMB-at-mating is $35 \%$ of its unfished value. This calculation also depends on the assumed bycatch F's on Tanner crab in the snow crab, BBRKC and groundfish fisheries. As with last year, the average F over the last 5 years for each of the bycatch fisheries is used in these calculations (in previous years, a different approach was used to determine the F to use for the snow crab fishery - see e.g., Stockhausen, 2016).

Selectivity curves in the bycatch fisheries were set using the average curves over the last 5 years for each fishery, the same approach as in previous assessments (Stockhausen 2017).

The determination of $\mathrm{B}_{\mathrm{MSY}}=\mathrm{B}_{35 \%}$ for Tanner crab depends on the selection of an appropriate time period over which to calculate average recruitment $(\bar{R})$. Following discussion in 2012 and 2013, the SSC endorsed an averaging period of 1982+. This issue was revisited at the May 2018 CPT meeting with regard to the final year to be included in the calculation, but no definitive were made. Starting the average recruitment period in 1982 is consistent with a 5-6 year recruitment lag from 1976/77, when a wellknown climate regime shift occurred in the EBS (Rodionov and Overland, 2005) that may have affected stock productivity. The value of $\bar{R}$ for this period from MCMC runs of the author's preferred model is 373.96 million. The estimates of average recruitment from the author's preferred model (M19F03), as well as all the other models based on the "current" ADFG fishery data, are substantially higher than from the 2018 assessment model ( 224 million; see Table 52). The value of $\mathrm{B}_{\mathrm{MSY}}=\mathrm{B}_{35 \%}$ for $\bar{R}$ is 41.07 thousand t , which is larger than that from the 2018 assessment ( 21.87 thousand t ).

Once $\mathrm{F}_{\text {OFL }}$ is determined using the control rule (Figure 63), the (total catch) OFL can be calculated based on projecting the population forward one year assuming that $F=\mathrm{F}_{\text {OFL }}$. In the absence of uncertainty, the OFL would then be the predicted total catch taken when fishing at $F=$ Fofl. When uncertainty (e.g. assessment uncertainty, variability in future recruitment) is taken into account, the OFL is taken as the median total catch when fishing at $F=\mathrm{F}_{\text {ofL }}$.

The total catch (biomass), including all bycatch of both sexes from all fisheries, was estimated using

$$
C=\sum_{f} \sum_{x} \sum_{z} \frac{F_{f, x, z}}{F_{,, x, z}} \cdot\left(1-e^{-F_{,, x, z}}\right) \cdot w_{x, z} \cdot\left[e^{-M_{x} \cdot \delta t} \cdot N_{x, Z}\right]
$$

where $C$ is total catch (biomass), $F_{f, x, z}$ is the fishing mortality in fishery $f$ on crab in size bin $z$ by sex $(x)$, $F_{, x, z}=\sum_{f} F_{f, x, z}$ is the total fishing mortality by sex on crab in size bin $z, w_{x, z}$ is the mean weight of crab in size bin $z$ by sex, $M_{x}$ is the sex-specific rate of natural mortality, $\delta t$ is the time from July 1 to the time of the fishery ( 0.625 yr ), and $N_{x, z}$ is the numbers by sex in size bin $z$ on July 1,2019 as estimated by the assessment model.

Assessment model uncertainty was included in the calculation of OFL using MCMC. Conceptually, a random draw from the assessment model's joint posterior distribution for the estimated parameters was taken, and the $\bar{R}, \mathrm{~B}_{0}, \mathrm{~F}_{\text {MSY }}, \mathrm{B}_{\text {MSY }}, \mathrm{F}_{\text {OFL }}$, OFL, and "current" MMB for 2019/20 were calculated based on resulting model parameter values. This would be repeated a large number of times to approximate the distribution of OFL given the full model uncertainty. For this assessment, four chains of 10 million MCMC steps were generated, with the OFL and associated quantities calculated at each step. The chain was initialized from the converged model state using a "burn in" of 2,000,000 steps and subsequently thinned by a factor of 1,000 to reduce serial autocorrelation in the MCMC sampling. This resulted in about $20,000 \mathrm{MCMC}$ samples with which to characterize the distribution of the OFL. The median value
of this distribution was taken as the OFL for 2019/20. The OFL for 2019/20 from the author's preferred model (Model M19F03) is 28.86 thousand $\mathbf{t}$ (Figure 64).

The $\mathrm{B}_{\text {MSY }}$ proxy, $\mathrm{B}_{35 \%}$, from the author's preferred model is 41.07 thousand t , so MSST $=0.5 \mathrm{~B}_{\mathrm{MSY}}=$ 20.54 thousand t . Because current projected $B=39.55$ thousand $\mathrm{t}>$ MSST, the stock is not overfished. However, because current projected $B<\mathrm{B}_{\mathrm{MSY}}$, the stock falls into Tier 3b. The population state (directed F vs. MMB) is plotted for each year from 1965/66-2018/19 in Figure 65 against the Tier 3 harvest control rule.

## 2. ABC calculation

Amendments 38 and 39 to the Fishery Management Plan (NPFMC 2010) established methods for the Council to set Annual Catch Limits (ACLs). The Magnuson-Stevens Act requires that ACLs be established based upon an acceptable biological catch (ABC) control rule that accounts for scientific uncertainty in the OFL such that $\mathrm{ACL}=\mathrm{ABC}$ and the total allowable catch (TAC) and guideline harvest levels (GHLs) be set below the ABC so as not to exceed the ACL. ABCs must be recommended annually by the Council's SSC.

Two methods for establishing the $A B C$ control rule are: 1) a constant buffer where the $A B C$ is set by applying a multiplier to the OFL to meet a specified buffer below the OFL; and 2) a variable buffer where the ABC is set based on a specified percentile ( $\mathrm{P}^{*}$ ) of the distribution of the OFL that accounts for uncertainty in the OFL. $\mathrm{P}^{*}$ is the probability that ABC would exceed the OFL and overfishing occur. In 2010, the NPFMC prescribed that ABCs for BSAI crab stocks be established at $\mathrm{P}^{*}=0.49$ (following Method 2). Thus, annual $\mathrm{ACL}=\mathrm{ABC}$ levels should be established such that the risk of ovefishing, $\mathrm{P}[\mathrm{ABC}>\mathrm{OFL}]$, is $49 \%$. In 2014, however, the SSC adopted a buffer of $20 \%$ on OFL for the Tanner crab stock for calculating ABC. Here, ABCs are provided based on both methods.

For the author's preferred scenario, M19F03, the $\mathrm{P}^{*} \mathrm{ABC}\left(\mathrm{ABC}_{\max }\right)$ is 28.79 thousand t while the $20 \%$ Buffer ABC is 23.09 thousand $t$. The author remains concerned that the OFL calculation, based on $\mathrm{F}_{35 \%}$ as a proxy for $\mathrm{F}_{\mathrm{MSY}}$, is overly optimistic regarding the actual productivity of the stock. Fishery-related mortality similar to the $P^{*}$ ABC level has occurred only in the latter half of the 1970s and in 1992/93, coincident with collapses in stock biomass to low levels. This suggests that $\mathrm{F}_{35 \%}$ may not be a realistic proxy for $\mathrm{F}_{\text {MSY }}$ and/or that MMB may not be a good proxy for reproductive success, as are currently assumed for this stock. In addition, the estimates of survey catchability for this stock remain problematic and contribute to this year's inflated OFL recommendation (relative to last year's) despite a continued decline in survey biomass across the last few years. Given this uncertainty concerning the stock, the author recommends using the $\mathbf{2 0 \%}$ buffer previously adopted by the SSC for this stock to calculate ABC. Consequently, the author's recommended ABC is $\mathbf{2 3 . 0 9}$ thousand $\mathbf{t}$.

## G. Rebuilding Analyses

Tanner crab is not currently under a rebuilding plan. Consequently no rebuilding analyses were conducted.

## H. Data Gaps and Research Priorities

Information on growth-per-molt has been collected in the EBS on Tanner crab and incorporated into the assessment. It would be helpful to have more information on growth associated with the terminal molt, because it seems likely this is has different characteristics than previous molts. Additionally, more data regarding temperature-dependent effects on molting frequency would be helpful to assess potential impacts of the EBS cold pool on the stock and potentially improve recruitment estimates. Information on temperature-dependent changes in crab movement and survey catchability would also be of value. In addition, it would be worthwhile to develop a "better" index of reproductive potential than MMB that can be calculated in the assessment model, as well as to revisit the issue of MSY proxies for this stock.

The characterization of fisheries in the assessment model needs to be carefully reconsidered. How, and whether or not, the differences in the directed fishery in areas east and west $166^{\circ} \mathrm{W}$ longitude should be explicitly represented in the assessment model should be addressed. The question of whether or not bycatch in the groundfish fisheries should be split into pot- and trawl-related components should be revisited. Also, the appropriate weight for male maturity ogives based on NMFS survey data in the model likelihood needs to be further explored.

Incorporating the BSFRF side-by-side (SBS) surveys into the assessment in the best way possible is also a matter for further exploration. There appears to be conflicting information from the NMFS and BSFRF SBS surveys regarding "availability" relative to the full NMFS survey, so estimating availability in the assessment model by fitting SBS data from both surveys (as was done here in Scenarios M19F04 and M19F05) may not be the best approach to incorporating the BSFRF surveys, which are assumed to provide absolute estimates of crab abundance within the area(s) in which the SBS surveys are conducted.

Development of a Gmacs version of the Tanner crab model is also a priority, but will await development of a Gmacs snow crab model.

## I. Ecosystem Considerations

Mature male biomass is currently used as the "currency" of Tanner crab spawning biomass for assessment purposes. However, its relationship to stock-level rates of egg production, a better measure of stock-level reproductive capacity, is unclear. Thus, use of MMB to reflect Tanner crab reproductive potential may be misleading as to stock health. Nor is it likely that mature female biomass has a clear relationship to annual egg production. For Tanner crab, the fraction of barren mature females by shell condition appears to vary at decadal time scales (Rugolo and Turnock, 2012), suggesting a climatic driver.

## 1. Ecosystem Effects on Stock

Time series trends in prey availability or abundance are generally unknown for Tanner crab because typical survey gear is not quantitative for Tanner crab prey. On the other hand, Pacific cod (Gadus macrocephalus) is thought to account for a substantial fraction of annual mortality on Tanner crab (Aydin et al., 2007). Total P. cod biomass is estimated to have been slowly declining from 1990 to 2008, during the time frame of a collapse in the Tanner crab stock, but has been increasing rather rapidly since 2008 (Thompson and Lauth, 2012). This suggests that the rates of "natural mortality" used in the stock assessment for the period post-1980 may be underestimates (and increasingly biased low if the trend in P. cod abundance continues). This trend is definitely one of potential concern.

## 2. Effects of Tanner crab fishery on ecosystem

Potential effects of the Tanner crab fishery on the ecosystem are considered in the following table:

| Effects of Tanner crab fishery on ecosystem |  |  |  |
| :---: | :---: | :---: | :---: |
| Indicator | Observation | Interpretation | Evaluation |
| Fishery contribution to bycatch |  |  |  |
| Prohibited species | salmon are unlikely to be trapped inside a pot when it is pulled, although halibut can be | unlikely to have substantial effects at the stock level | minimal to none |
| Forage (including herring, Atka mackerel, cod and pollock) | Forage fish are unlikely to be trapped inside a pot when it is pulled crab pots have a very | unlikely to have substantial effects unlikely to be having | minimal to none |
| HAPC biota | small footprint on the bottom | substantial effects postrationalization | minimal to none |


| Marine mammals and birds | crab pots are unlikely to attract birds given the depths at which they are fished | unlikely to have substantial effects | minimal to none |
| :---: | :---: | :---: | :---: |
| Sensitive non-target species | Non-targets are unlikely to be trapped in crab pot gear in substantial numbers | unlikely to have substantial effects | minimal to none |
| Fishery concentration in space and time | substantially reduced in time following rationalization of the fishery | unlikely to be having substantial effects | probably of little concern |
| Fishery effects on amount of large size target fish | Fishery selectively removes large males | May impact stock reproductive potential as large males can mate with a wider range of females | possible concern |
| Fishery contribution to discards and offal production | discarded crab suffer some mortality | May impact female spawning biomass and numbers recruiting to the fishery | possible concern |
| Fishery effects on age-atmaturity and fecundity | none | unknown | possible concern |

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## Table captions

Table 1. Retained catch (males) in directed Tanner crab fisheries (1965/66-2000/01). Catch units are metric tons. Asterisks denote a closure of the directed domestic fishery. ..... 53
Table 2. Retained catch (males) in the US domestic pot fishery. Information from the CommunityDevelopment Quota (CDQ) fisheries is included in the table for fishery years 2005/06 to the present.Number of crabs caught and harvest includes deadloss. The "Fishery Year" YYYY/YY+1 runs from July1, YYYY to June 30, YYYY+1. The ADFG year (in parentheses, if different from the "Fishery Year")indicates the year ADFG assigned to the fishery season in compiled reports.55
Table 3. Total catch (retained + discarded) of Tanner crab in various fisheries, as estimated from observerdata. Units are 1000's t. TCF: directed Tanner crab fishery; SCF: snow crab fishery; RKF: Bristol Bay redking crab fishery; GTF: groundfish fisheries56
Table 4. Comparison of retained catch abundance and biomass used in the previous assessment ("historical") with "current" catch abundance and biomass. Only values since 2005 (highlighted in grey) have been changed. ..... 58
Table 5. Comparison of total catch biomass in the directed Tanner crab fisheries used in the previous assessment ("historical") with "current" catch biomass dataset. See text for details ..... 59
Table 6. Comparison of Tanner crab bycatch biomass in the snow crab fisheries used in the previous assessment ("historical") with the "current" catch biomass dataset. See text for details ..... 60
Table 7. Comparison of Tanner crab bycatch biomass in the BBRKC fishery used in the previous assessment ("historical") with the "current" catch biomass dataset. See text for details ..... 61
Table 8. Retained catch biomass in the directed Tanner crab (TCF), snow crab (SCF), and BBRKC (RKF)fisheries since 2005. The directed fishery was completely closed from 2010/11 to 2012/13, and in2016/17. Legal-sized Tanner crab can be incidentally-retained in the snow crab and BBRKC fisheries upto a cap of $5 \%$ the target catch.62
Table 9. Sample sizes for retained catch-at-size in the directed fishery. $\mathrm{N}=$ number of individuals. $\mathrm{N}^{`}=$ scaled sample size used in assessment. The directed fishery was closed in 2016/17. ..... 62
Table 10. Sample sizes for total catch-at-size in the directed fishery from crab observer sampling. $\mathrm{N}=$ number of individuals. $\mathrm{N}^{`}=$ scaled sample size used in assessment. ..... 63
Table 11. Sample sizes for total bycatch-at-size in the snow crab fishery, from crab observer sampling. N $=$ number of individuals. $\mathrm{N}^{`}=$ scaled sample size used in assessment ..... 64
Table 12. Sample sizes for total bycatch-at-size in the BBRKC fishery, from crab observer sampling. $\mathrm{N}=$number of individuals. $\mathrm{N}^{`}=$ scaled sample size used in assessment.65
Table 13. Sample sizes for total catch-at-size in the groundfish fisheries, from groundfish observer sampling. $\mathrm{N}=$ number of individuals. $\mathrm{N}^{`}=$ scaled sample size used in the assessment. ..... 66
Table 14. Trends in Tanner crab biomass (metric tons) in the NMFS EBS summer bottom trawl survey, by sex and area. ..... 67
Table 15. Trends in biomass for preferred-size ( $>125 \mathrm{~mm} \mathrm{CW}$ ) male Tanner crab in the NMFS EBS summer bottom trawl survey (in metric tons). ..... 69
Table 16. Sample sizes for NMFS survey size composition data. In the assessment model, an input samplesize of 200 is used for all survey-related compositional data. ..................................................................... 71Table 17. Effort data (potlifts) in the crab fisheries, by area. TCF: directed Tanner crab fishery; SCF:snow crab fishery; RKF: Bristol Bay red king crab fishery.73
Table 18.Non-selectivity parameters from all model scenarios that were estimated within $1 \%$ of bounds.75
Table 19. Selectivity-related parameters from all model scenarios estimated within $1 \%$ of bounds. ..... 76
Table 20. Estimated growth, natural mortality, and non-vector recruitment parameters for all model scenarios ..... 78
Table 21. Historical recruitment devs estimates (1948-1974) for all model scenarios. ..... 79
Table 22. Current recruitment devs estimates (1975-2019) for all model scenarios. ..... 80
Table 23. Logit-scale parameters for the probability of terminal molt for males for all model scenarios. The (arithmetic) probability of terminal molt was fixed at 0 for males less than 60 mm CW and at 1 for males greater than 145 mm CW in Scenarios M19F03 and M19F05.81
Table 24. Logit-scale parameters for the probability of terminal molt for females for all model scenarios. The (arithmetic) probability of terminal molt was fixed at 0 for females less than 50 mm CW in Scenarios M19F03 and M19F05 and at 1 for females greater than 105 mm CW for all scenarios. ..... 82
Table 25. Log-scale NMFS survey catchability and selectivity parameters for all model scenarios. ..... 83
Table 26. BSFRF SBS (side-by-side) male availability parameters for all model scenarios in which they were estimated. ..... 84
Table 27. BSFRF SBS (side-by-side) female availability parameters for all model scenarios. in which they were estimated. ..... 85
Table 28. Mean capture rate, selectivity and retention parameter estimates for the directed fishery (TCF) for all model scenarios. ..... 86
Table 29. Log-scale male capture rate dev parameter estimates for the directed fishery (TCF) for all model scenarios ..... 87
Table 30. Comparison of mean capture rate, In-scale capture rate devs, and selectivity parameter estimates for the snow crab fishery (SCF) for all model scenarios. ..... 88
Table 31. Comparison of mean capture rate, ln -scale capture rate devs, and selectivity parameters estimates for the BBRKC fishery (RKF) for all model scenarios. ..... 89
Table 32. Comparison of mean capture rate and selectivity parameters estimates for the groundfish fisheries (GTF). ..... 90
Table 33. Log-scale capture rate dev parameter estimates for the groundfish fisheries (GTF) for all model scenarios. ..... 91
Table 34. (Unweighted) negative log-likelihoods and (weighted) objective function values for fishery- related data components from the model scenarios. TCF: directed Tanner crab fishery; SCF: snow crab fishery; RKF: BBRKC fishery; GTF: groundfish fisheries. ..... 92Table 35. (Unweighted) negative log-likelihoods and (weighted) objective function values for survey-related data components from the model scenarios. Rows consisting of all zero values indicate a datacomponent which was not included in any of the models. Blank cells indicate a data component (row) thatwas not included in the associated scenario (column)93
Table 36. (Unweighted) negative log-likelihoods and (weighted) objective function values for fits to growth (molt increment) and male maturity ogive data components from the model scenarios. ..... 93
Table 37. Root mean square errors (RMSE) for fishery-related data components from the modelscenarios. TCF: directed Tanner crab fishery; SCF: snow crab fishery; RKF: BBRKC fishery; GTF:groundfish fisheries. Rows consisting of all zero values indicate a data component which was notincluded in any of the models.94
Table 38. Average root mean square errors (RMSE) for survey-related data components from the modelscenarios. Rows consisting of all zero values indicate a data component which was not included in any ofthe models. Blank cells indicate a data component (row) that was not included in the likelihood in theassociated scenario (column).95
Table 39. Root mean square errors (RMSE) for fits to growth (molt increment) and male maturity ogive data components from the model scenarios ..... 95Table 40. Effective sample sizes used for NMFS 0 EBS trawl survey size composition data for the 2018assessment model (M19F00) and the author's preferred model (M19F03). Effective sample sizes wereestimated using the McAllister-Ianelli approach. Note that, while effective N's were calculated for thisdataset in MF 1903, it was not included in the model objective function (the weight in the likelihood wasset to 0). Input sample sizes were set at 200 .96Table 41. Effective sample sizes used for retained catch size composition data from the directed fisheryfor the 2018 assessment model (M19F00) and the author's preferred model (M19F03). Effective samplesizes were estimated using the McAllister-Ianelli approach.97

Table 42. Effective sample sizes used for total catch size composition data from the directed fishery for the 2018 assessment model (M19F00) and the author's preferred model (M19F03). Effective sample sizes were estimated using the McAllister-Ianelli approach.98 Table 43. Effective sample sizes used for bycatch size composition data from the snow crab fishery for
the 2018 assessment model (M19F00) and the author's preferred model (M19F03). Effective sample sizes
were estimated using the McAllister-Ianelli approach....................................................................... 99 Table 44. Effective sample sizes used for bycatch size composition data from the BBRKC fishery for the 2018 assessment model (M19F00) and the author's preferred model (M19F03. Effective sample sizes were estimated using the McAllister-Ianelli approach.
Table 45. Effective sample sizes used for bycatch size composition data from the groundfish fisheries for the 2018 assessment model (M19F00) and the author's preferred model (M19F03). Effective sample sizes were estimated using the McAllister-Ianelli approach. 101
Table 46. Comparison of fits to mature survey biomass by sex (in 1000's $t$ ) from the 2018 assessment
model (M19F00) and the author's preferred model (M19F03)....................................................... 102
Table 47. Comparison of estimates of mature biomass-at-mating by sex (in 1000's t) from the 2018 assessment model (M19F00) and the author's preferred model (M19F03).
Table 48. Estimated population size (millions) for females on July 1 of year. from the author's preferred model, Model M19F03 ..... 103
Table 49. Estimated population size (millions) for males on July 1 of year. from the author's preferred mode, Model M19F03. ..... 103
Table 50. Comparison of estimates of recruitment (in millions) from the 2018 assessment model (M19F00) and the author's preferred model (M19F03), ..... 104
Table 51. Comparison of exploitation rates (i.e., catch divided by biomass) from the 2018 assessment model (M19F00) and the author's preferred model (M19F03) ..... 105Table 52. Values required to determine Tier level and OFL for the models considered here. These valuesare presented only to illustrate the effect of incremental changes in the model scenarios. Results from theauthor's preferred model (M19F03) are highlighted in green.106

## Figure captions

Figure 1. Eastern Bering Sea District of Tanner crab Registration Area J including sub-districts and sections (from Bowers et al. 2008). 107
Figure 2. Upper: retained catch (males, 1000's t) in the directed fisheries (US pot fishery [green bars], Russian tangle net fishery [red bars], and Japanese tangle net fisheries [blue bars]) for Tanner crab since 1965/66. Lower: Retained catch (males, 1000's $t$ ) in directed fishery since 2001/02. The directed fishery was closed from 1996/97 to 2004/05, from 2010/11 to 2012/13, and in 2016/17. 108
Figure 3. Time series of retained catch biomass ( 1000 's $t$ ) in the directed Tanner crab (TCF: red; eastern area: triangles; western area: circles; all EBS: squares), snow crab (SCF: green), and BBRKC (RFF: blue) fisheries since 2005. The directed fishery was closed from 2010/11 to 2012/13, and in 2016/17. Legalsized Tanner crab can be incidentally-retained in the snow crab and BBRKC fisheries up to a cap of $5 \%$ the target catch.
Figure 4. Upper: total catch (retained + discards) of Tanner crab (males and females, 1000's $t$ ) in the directed Tanner crab, snow crab, Bristol Bay red king crab, and groundfish fisheries. Bycatch reporting began in 1973 for the groundfish fisheries and in 1992 for the crab fisheries. Lower: detail since 2005.110 Figure 5. Retained catch size compositions in the directed Tanner crab fisheries since the fishery reopened in 2013/14 (red: western area, green: eastern area; blue: all EBS)
Figure 6. Total catch (retained + discards) size compositions for males, normalized by fleet, during
1990/91-1999/2000 in the directed Tanner crab (aggregated across areas, TCF: blue), snow crab (SCF: green), and BBRKC (RKF: red) fisheries (solid line: new shell crab; dotted line: old shell crab).
Figure 7. Total catch (retained + discards) size compositions for males, normalized by fleet, during 2000/01-2009/10 in the directed Tanner crab (aggregated across areas, TCF: blue), snow crab (SCF: green), and BBRKC (RKF: red) fisheries (solid line: new shell crab; dotted line: old shell crab). The directed fishery was closed in 2000/01-2004/05 and was open only in the western area in 2005/06 and in the eastern area in 2009/10.
Figure 8. Total catch (retained + discards) size compositions for males, normalized by fleet, during 2010/11-2018/19 in the directed Tanner crab (aggregated across areas, TCF: blue), snow crab (SCF: green), and BBRKC (RKF: red) fisheries (solid line: new shell crab; dotted line: old shell crab). The directed fishery was closed in 2010/11-2012/13 and 206/17, and was open only in the western area in 2017/18 and 2018/19.
Figure 9. Bycatch size compositions for females, normalized by fleet, during 1990/91-1999/2000 in the directed Tanner crab (aggregated across areas, TCF: blue), snow crab (SCF: green), and BBRKC (RKF: red) fisheries (solid line: new shell crab; dotted line: old shell crab). 115
Figure 10. Bycatch size compositions for females, normalized by fleet, during 2000/01-2009/10 in the directed Tanner crab (aggregated across areas, TCF: blue), snow crab (SCF: green), and BBRKC (RKF: red) fisheries (solid line: new shell crab; dotted line: old shell crab). 116
Figure 11. Bycatch size compositions for females, normalized by fleet, during 2010/11-2018/19 in the directed Tanner crab (aggregated across areas, TCF: blue), snow crab (SCF: green), and BBRKC (RKF: red) fisheries (solid line: new shell crab; dotted line: old shell crab). 117
Figure 12. Annual bycatch size compositions in the groundfish fisheries by sex, expanded to total
bycatch, during 1991/92-2006/07. Red lines: females; green lines: males...................................... 118
Figure 13. Annual bycatch size compositions in the groundfish fisheries by sex, expanded to total bycatch, during 2007/08-2018/19. Red lines: females; green lines: males. 119
Figure 14. Annual estimates of area-swept biomass from the NMFS EBS bottom trawl survey, by sex, maturity state, and management area. Red lines: total biomass; green lines: biomass in the eastern area; blue: biomass in the western area.
Figure 15. Annual estimates of area-swept biomass from the NMFS EBS bottom trawl survey for preferred-size ( $>125 \mathrm{~mm}$ CW) legal males. Red lines: total biomass; green lines: biomass in the eastern area; blue: biomass in the western area.

Figure 16. Spatial footprints (stations occupied in green) during the BSFRF-NMFS cooperative side-byside (SBS) catchability studies in 2013-2017. Squares and circles represent stations in the standard NMFS EBS bottom trawl survey (which extends beyond the area shown in the maps).
Figure 17. Annual estimates of area-swept biomass from the BSFRF-NMFS cooperative side-by-side
(SBS) catchability studies in 2013-2017. The SBS studies had different spatial footprints each year, so
annual changes in biomass do not necessarily reflect underlying population trends. Red lines: BSFRF;
green lines: NMFS. ............................................................................................................................. 123
Figure 18. Size compositions from the NMFS EBS bottom trawl survey for 1975-2019. 124
Figure 19. Annual size compositions of area-swept abundance by sex from the BSFRF-NMFS cooperative side-by-side (SBS) catchability studies in 2013-2015. Red lines: BSFRF; green lines: NMFS.

Figure 20. Annual estimates of area-swept abundance (blue circles) from the NMFS EBS bottom trawl
survey, by sex and maturity state for 2014 and 2015. Local abundance scales with symbol area. The
background "heatmap" represents bottom water temperatures at the time of the survey....................... 127127
Figure 21. Annual estimates of area-swept abundance (blue circles) from the NMFS EBS bottom trawl survey, by sex and maturity state for 2016 and 2017. Local abundance scales with symbol area. The background "heatmap" represents bottom water temperatures at the time of the survey. ..... 128
Figure 22. Annual estimates of area-swept abundance (blue circles) from the NMFS EBS bottom trawl survey, by sex and maturity state for 2018 and 2019. Local abundance scales with symbol area. The background "heatmap" represents bottom water temperatures at the time of the survey. ..... 129
Figure 23. Male maturity ogives (the fraction of new shell mature males, relative to all new shell males)as determined from chela height:carapace width ratios from the NMFS EBS bottom trawl survey for yearswhen chela heights were collected with 0.1 mm precision.130
Figure 24. Molt increment data collected collaboratively by NMFS, BSFRF, and ADFG ..... 130
Figure 25. Size-weight relationships developed from NMFS EBS summer trawl survey data. ..... 131
Figure 26. Assumed size distribution for recruits entering the population. ..... 131
Figure 27. Fits to retained and total catch biomass in the directed fishery from all model scenarios. ..... 132
Figure 28. Fits to total catch biomass in the snow crab fishery from all scenarios ..... 133
Figure 29. Fits to total catch biomass in the BBRKC fishery from all scenarios. ..... 133
Figure 30. Fits to total catch biomass in the groundfish fisheries for all scenarios. ..... 134
Figure 31. Fits to mature biomass from the NMFS " 0 " EBS bottom trawl survey data for all. Note that scenarios M19F03 and M19F05 do not include the mature male component in the likelihood (they fit totalmale biomass) and fit both mature and immature biomass for females.135
Figure 32. Fits to mature biomass from the NMFS "M" and NMFS "F" EBS bottom trawl survey data forscenarios M19F00a, M19F01, M19F02, M19F02, M19F03, M19F04, and M19F05. Note that onlyscenarios M19F03 and M19F05 include these data components in the model objective function.136
Figure 33. Fits to survey biomass from the NMFS SBS bottom trawl survey data for scenarios M19F04 and M19F05 ..... 137
Figure 34. Fits to survey biomass from the BSFRF SBS bottom trawl survey data for scenarios M19F04 and M19F05 ..... 138
Figure 35. Fits to molt increment data for scenarios M19F00a, M19F01, M19F02, M19F02, M19F03, M19F04, and M19F05 ..... 139
Figure 36. Fits to male maturity ogive data for scenarios M19F00a, M19F01, M19F02, M19F02, M19F03, M19F04, and M19F05. Note that only scenarios M1902, M19F03, and M19F05 include the datain the likelihood.140
Figure 37. Directed fishery catchability (capture rates) from all model scenarios. ..... 141
Figure 38. Directed fishery selectivity curves from all scenarios for the pre-1991 time period and 1991- 1994. The $50 \%$-selected parameter varies annually for $1991+$ ..... 142
Figure 39. Directed fishery selectivity curves from all scenarios for 1995-1996 and 2005-2007. The 50\%-selected parameter varies annually for $1991+$.143
Figure 40. Directed fishery selectivity curves from all scenarios for 2008-2009 and 2013-2015. The 50\%selected parameter varies annually for 1991+144
Figure 41. Directed fishery selectivity curves from all scenarios for 2008-2009 and 2013-2015. The 50\%- selected parameter varies annually for 1991+ ..... 145
Figure 42. Directed fishery retention curves from all scenarios for the pre-1991, 1991-1996, and post- 2004 time periods ..... 146
Figure 43. Snow crab fishery catchability (capture rates) from all scenarios. ..... 147
Figure 44. Snow crab fishery selectivity curves from all scenarios for 3 time periods: pre-1997, 1997- 2004, 2005+. ..... 148
Figure 45. BBRKC fishery catchability (capture rates) from all scenarios. ..... 149
Figure 46. BBRKC fishery selectivity curves from all scenarios for 3 time periods: pre-1997, 1997-2004, 2005+. ..... 150
Figure 47. Catchability (capture rates) in the groundfish fisheries from all scenarios. ..... 151
Figure 48. Groundfish fisheries selectivity curves from all scenarios estimated for 3 time periods: pre- 1987, 1987-1996, 1997+ ..... 152
Figure 49. NMFS " 0 " survey catchabilities for all scenarios for the 1975-1981 and 1982+ time periods.153
Figure 50. NMFS " 0 " survey selectivity functions for all scenarios for the 1975-1981 and 1982+ time periods ..... 154
Figure 51. NMFS " 0 " survey capture probabilities (i.e., catchability $x$ selectivity) for all scenarios for the 1975-1981 and 1982+ time periods ..... 155
Figure 52. Survey availabilities from scenarios M19F04 and M19F05 for the 2013-2017 SBS studies.. 156Figure 53. Comparison of empirical "observed" and predicted availability in the 2013-2017 SBS studiesfrom scenario M19F04. The "observed" availability is the ratio of abundance in the NMFS SBS survey tothat in the full NMFS survey by size bin. Observed: red points, lines. Red fills are from loess smoothingof the observed availability. Predicted: green points, lines.157
Figure 54. Estimates of natural mortality from all scenarios. ..... 158
Figure 55 . Estimates of the probability of terminal molt from all scenarios. ..... 159
Figure 56. Estimates of mean growth from all scenarios. Dashed line is $1: 1$ ..... 160
Figure 57. Estimated recruitment time series from all scenarios ..... 161
Figure 58. Estimated recent recruitment time series from all scenarios ..... 162
Figure 59. Estimated (Feb. 15) mature biomass time series from all scenarios. ..... 163
Figure 60. Estimated recent (Feb. 15) mature biomass time series from all scenarios. ..... 164
Figure 61. Estimated (July 1) biomass time series by population category for all scenarios. ..... 165
Figure 62. MCMC results from scenario M19F03, the author's preferred model, for OFL-related quantities. ..... 166
Figure 63. The $\mathrm{F}_{\text {ofl }}$ harvest control rule. ..... 167
Figure 64. The OFL and ABC from the author's preferred model, scenario M19F03. ..... 167
Figure 65. Quad plot for the author's preferred model, scenario M19F03. ..... 168
Figure 65 . The ratio of estimated abundance by size from the NMFS and BSFRF side-by-side catchabilitystudies. The heavy green line is the size-specific mean over the 5 years. These represent simple empiricalestimates of the size-specific catchability of the NMFS survey gear relative to the BSFRF gear. If theBSFRF survey gear is assumed to capture all crab within the area swept, these curves represent empiricalestimates of the size-specific NMFS survey gear catchability (i.e., fully selected catchability [q] $x$selectivity).169

Tables
Table 1. Retained catch (males) in directed Tanner crab fisheries (1965/66-2000/01). Catch units are metric tons. Asterisks denote a closure of the directed domestic fishery.

| year | US | Japan | Russia | Total |
| :---: | :---: | :---: | :---: | :---: |
| 1965 | 0 | 1,170 | 750 | 1,920 |
| 1966 | 0 | 1,690 | 750 | 2,440 |
| 1967 | 0 | 9,750 | 3,840 | 13,590 |
| 1968 | 460 | 13,590 | 3, 960 | 18,010 |
| 1969 | 460 | 19,950 | 7,080 | 27,490 |
| 1970 | 80 | 18,930 | 6,490 | 25,500 |
| 1971 | 50 | 15, 900 | 4,770 | 20,720 |
| 1972 | 100 | 16,800 | 0 | 16,900 |
| 1973 | 2, 290 | 10, 740 | 0 | 13,030 |
| 1974 | 3, 300 | 12,060 | 0 | 15, 360 |
| 1975 | 10, 120 | 7,540 | 0 | 17, 660 |
| 1976 | 23, 360 | 6,660 | 0 | 30,020 |
| 1977 | 30,210 | 5,320 | 0 | 35,530 |
| 1978 | 19, 280 | 1,810 | 0 | 21,090 |
| 1979 | 16, 600 | 2,400 | 0 | 19,000 |
| 1980 | 13, 426 | 0 | 0 | 13, 426 |
| 1981 | 4,990 | 0 | 0 | 4,990 |
| 1982 | 2,390 | 0 | 0 | 2,390 |
| 1983 | 549 | 0 | 0 | 549 |
| 1984 | 1,429 | 0 | 0 | 1,429 |
| 1985* | 0 | 0 | 0 | 0 |
| 1986* | 0 | 0 | 0 | 0 |
| 1987 | 998 | 0 | 0 | 998 |
| 1988 | 3,180 | 0 | 0 | 3,180 |
| 1989 | 11,113 | 0 | 0 | 11, 113 |
| 1990 | 18,189 | 0 | 0 | 18,189 |
| 1991 | 14, 424 | 0 | 0 | 14, 424 |
| 1992 | 15, 921 | 0 | 0 | 15, 921 |
| 1993 | 7,666 | 0 | 0 | 7,666 |
| 1994 | 3,538 | 0 | 0 | 3,538 |
| 1995 | 1,919 | 0 | 0 | 1,919 |
| 1996 | 821 | 0 | 0 | 821 |
| 1997* | 0 | 0 | 0 | 0 |
| 1998* | 0 | 0 | 0 | 0 |
| 1999* | 0 | 0 | 0 | 0 |
| 2000* | 0 | 0 | 0 | 0 |

Table 1 (cont.). Retained catch (males) in directed Tanner crab fisheries (2001/02-2018/19). Catch units are metric tons. Asterisks denote a closure of the directed domestic fishery; retained catch in these years represent incidentally retained Tanner crab in the snow crab and Bristol Bay red king crab fisheries.

| year | US | Japan | Russia | Total |
| :--- | ---: | ---: | ---: | ---: |
| $2001^{*}$ | 0 | 0 | 0 | 0 |
| $2002^{*}$ | 0 | 0 | 0 | 0 |
| $2003^{*}$ | 0 | 0 | 0 | 0 |
| $2004^{*}$ | 0 | 0 | 0 | 0 |
| 2005 | 432 | 0 | 0 | 432 |
| 2006 | 963 | 0 | 0 | 963 |
| 2007 | 956 | 0 | 0 | 956 |
| 2008 | 880 | 0 | 0 | 880 |
| 2009 | 603 | 0 | 0 | 603 |
| $2010^{*}$ | 1 | 0 | 0 | 1 |
| $2011^{*}$ | 2 | 0 | 0 | 2 |
| $2012^{*}$ | 1 | 0 | 0 | 1 |
| 2013 | 1,264 | 0 | 0 | 1,264 |
| 2014 | 6,216 | 0 | 0 | 6,216 |
| 2015 | 8,910 | 0 | 0 | 8,910 |
| $2016^{*}$ | 1 | 0 | 0 | 1 |
| 2017 | 1,133 | 0 | 0 | 1,133 |
| 2018 | 1,107 | 0 | 0 | 1,107 |

Table 2. Retained catch (males) in the US domestic pot fishery. Information from the Community Development Quota (CDQ) fisheries is included in the table for fishery years 2005/06 to the present. Number of crabs caught and harvest includes deadloss. The "Fishery Year" YYYY/YY+1 runs from July 1, YYYY to June 30, YYYY+1. The ADFG year (in parentheses, if different from the "Fishery Year") indicates the year ADFG assigned to the fishery season in compiled reports.

| year <br> (ADFG year) | Total Crab (no.) | Total Harvest <br> (lbs) | $\begin{array}{r} \text { GHL/TAC } \\ \text { (millions lbs) } \end{array}$ | Vessels (no.) | Season |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1968/69 (1969) | 353,300 | 1,008,900 |  |  |  |
| 1969/70 (1970) | 482,300 | 1,014,700 |  |  |  |
| 1970/71 (1971) | 61,300 | 166,100 |  |  |  |
| 1971/72 (1972) | 42,061 | 107,761 |  |  |  |
| 1972/73 (1973) | 93,595 | 231,668 |  |  |  |
| 1973/74 (1974) | 2,531,825 | 5,044,197 |  |  |  |
| 1974/75 | 2,773,770 | 7,028,378 |  | 28 |  |
| 1975/76 | 8,956,036 | 22,358,107 |  | 66 |  |
| 1976/77 | 20,251,508 | 51,455,221 |  | 83 |  |
| 1977/78 | 26,350,688 | 66,648,954 |  | 120 |  |
| 1978/79 | 16,726,518 | 42,547,174 |  | 144 |  |
| 1979/80 | 14,685,611 | 36,614,315 | 28-36 | 152 | 11/01-05/11 |
| 1980/81 (1981) | 11,845,958 | 29,630,492 | 28-36 | 165 | 01/15-04/15 |
| 1981/82 (1982) | 4,830,980 | 11,008,779 | 12-16 | 125 | 02/15-06/15 |
| 1982/83 (1983) | 2,286,756 | 5,273,881 | 5.6 | 108 | 02/15-06/15 |
| 1983/84 (1984) | 516,877 | 1,208,223 | 7.1 | 41 | 02/15-06/15 |
| 1984/85 (1985) | 1,272,501 | 3,036,935 | 3 | 44 | 01/15-06/15 |
| 1985/86 (1986) |  |  | -clo |  |  |
| 1986/87 (1987) | ------ |  | ---clo |  |  |
| 1987/88 (1988) | 957,318 | 2,294,997 | 5.6 | 98 | 01/15-04/20 |
| 1988/89 (1989) | 2,894,480 | 6,982,865 | 13.5 | 109 | 01/15-05/07 |
| 1989/90 (1990) | 9,800,763 | 22,417,047 | 29.5 | 179 | 01/15-04/24 |
| 1990/91 | 16,608,625 | 40,081,555 | 42.8 | 255 | 11/20-03/25 |
| 1991/92 | 12,924,102 | 31,794,382 | 32.8 | 285 | 11/15-03/31 |
| 1992/93 | 15,265,865 | 35,130,831 | 39.2 | 294 | 11/15-03/31 |
| 1993/94 | 7,235,898 | 16,892,320 | 9.1 | 296 | 11/01-11/10, 11/20-01/01 |
| 1994/95 (1994) | 3,351,639 | 7,766,886 | 7.5 | 183 | 11/01-11/21 |
| 1995/96 (1995) | 1,877,303 | 4,233,061 | 5.5 | 196 | 11/01-11/16 |
| 1996/97 (1996) | 734,296 | 1,806,077 | 6.2 | 196 | 11/01-11/05, 11/15-11/27 |
| 1997/98-2004/05 |  |  | --clo |  |  |
| 2005/06 | 443,978 | 952,887 | 1.7 | 49 | 10/15-03/31 |
| 2006/07 | 927,086 | 2,122,589 | 3.0 | 64 | 10/15-03/31 |
| 2007/08 | 927,164 | 2,106,655 | 5.7 | 50 | 10/15-03/31 |
| 2008/09 | 830,363 | 1,939,571 | 4.3 | 53 | 10/15-03/31 |
| 2009/10 | 485,676 | 1,327,952 | 1.3 | 45 | 10/15-03/31 |
| 2010/11 |  |  | ---clo |  |  |
| 2011/12 |  |  | -----clo |  |  |
| 2012/13 |  |  | -----clo | ---- |  |
| 2013/14 | 1,426,670 | 2,751,124 | 3.108 | 32 | 10/15-03/31 |
| 2014/15 | 7,442,931 | 13,576,105 | 15.105 | 100 | 10/15-03/31 |
| 2015/16 | 10,856,418 | 19,642,462 | 19.668 | 112 | 10/15-03/31 |
| 2016/17 | ------------- | -------------- | ---------clo | -------- | --- |
| 2017/18 | 1,340,394 | 2,497,033 | 2.500 | 34 | 10/15-03/31 |
| 2018/19 | 1,381,008 | 2,441,201 | 2.439 | 36 | 10/15-03/31 |

Table 3. Total catch (retained + discarded) of Tanner crab in various fisheries, as estimated from observer data. Units are 1000's t. TCF: directed Tanner crab fishery; SCF: snow crab fishery; RKF: Bristol Bay red king crab fishery; GTF: groundfish fisheries.

| year | TCF |  |  |  | $\begin{aligned} & \text { SCF } \\ & \text { all EBS } \end{aligned}$ |  | $\begin{aligned} & \text { RKF } \\ & \text { all EBS } \end{aligned}$ |  | $\begin{aligned} & \text { GTF } \\ & \text { all EBS } \end{aligned}$ | Total all EBS |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | male | female | male | female | male | female | male | female | all | all |
| 1973 | - | - | - | - | - | - | - | - | 17.7355 | 17.7355 |
| 1974 | - | - | - | - | - | - | - | - | 24.4486 | 24.4486 |
| 1975 | - | - | - | - | - | - | - | - | 9.4075 | 9.4075 |
| 1976 | - | - | - | - | - | - | - | - | 4.6992 | 4.6992 |
| 1977 | - | - | - | - | - | - | - | - | 2.7760 | 2.7760 |
| 1978 | - | - | - | - | - | - | - | - | 1.8688 | 1.8688 |
| 1979 | - | - | - | - | - | - | - | - | 3.3974 | 3.3974 |
| 1980 | - | - | - | - | - | - | - | - | 2.1137 | 2.1137 |
| 1981 | - | - | - | - | - | - | - | - | 1.4742 | 1.4742 |
| 1982 | - | - | - | - | - | - | - | - | 0.4491 | 0.4491 |
| 1983 | - | - | - | - | - | - | - | - | 0.6713 | 0.6713 |
| 1984 | - | - | - | - | - | - | - | - | 0.6441 | 0.6441 |
| 1985* | - | - | - | - | - | - | - | - | 0.3992 | 0.3992 |
| 1986* | - | - | - | - | - | - | - | - | 0.6486 | 0.6486 |
| 1987 | - | - | - | - | - | - | - | - | 0.6396 | 0.6396 |
| 1988 | - | - | - | - | - | - | - | - | 0.4627 | 0.4627 |
| 1989 | - | - | - | - | - | - | - | - | 0.6713 | 0.6713 |
| 1990 | - | - | - | - | 7.0812 | 0.1057 | 3.7224 | 0.0356 | 0.9435 | 11.8885 |
| 1991 | 6.2206 | 0.4408 | 19.5967 | 1.4452 | 8.3602 | 0.1440 | 1.9703 | 0.0272 | 2.5432 | 40.7482 |
| 1992 | 7.3470 | 0.5996 | 29.6604 | 1.1040 | 2.4872 | 0.1625 | 1.3167 | 0.0190 | 2.7596 | 45.4561 |
| 1993 | 1.6439 | 0.1361 | 10.2100 | 0.8601 | 2.8744 | 0.4004 | 3.1308 | 0.1493 | 1.7580 | 21.1630 |
| 1994 | 0.3573 | 0.1124 | 6.9581 | 0.7293 | 1.3451 | 0.1942 | - | - | 2.0960 | 11.7924 |
| 1995 | 0.6503 | 0.1407 | 4.4152 | 0.9242 | 1.0210 | 0.1209 | - | - | 1.5249 | 8.7973 |
| 1996 | 0.0718 | - | 0.2286 | 0.0567 | 1.9607 | 0.1196 | 0.2700 | 0.0024 | 1.5945 | 4.3044 |
| 1997* | - | - | - | - | 1.9637 | 0.0927 | 0.1601 | 0.0017 | 1.1800 | 3.3981 |
| 1998* | - | - | - | - | 0.6559 | 0.0804 | 0.1152 | 0.0017 | 0.9350 | 1.7882 |
| 1999* | - | - | - | - | 0.1318 | 0.0112 | 0.0751 | 0.0022 | 0.6306 | 0.8509 |
| 2000* | - | - | - | - | 0.3128 | 0.0061 | 0.0664 | 0.0014 | 0.7415 | 1.1282 |

Table 3 (cont.). Total catch (retained + discarded) of Tanner crab in various fisheries, as estimated from observer data. Units are 1000's t. TCF: directed Tanner crab fishery; SCF: snow crab fishery; RKF: Bristol Bay red king crab fishery; GTF: groundfish fisheries.

| year | TCF |  |  |  | $\begin{gathered} \text { SCF } \\ \text { all EBS } \end{gathered}$ |  | $\begin{aligned} & \text { RKF } \\ & \text { all EBS } \end{aligned}$ |  | $\begin{aligned} & \text { GTF } \\ & \text { all EBS } \end{aligned}$ | Total all EBS |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | male | female | male | female | male | female | male | female | all | all |
| 2001* | - | - | - | - | 0.545308 | 0.020530 | 0.042200 | 0.000963 | 1.185191 | 1.794192 |
| 2002* | - | - | - | - | 0.167178 | 0.013815 | 0.061253 | 0.001580 | 0.719068 | 0.962894 |
| 2003* | - | - | - | - | 0.064743 | 0.007011 | 0.054937 | 0.001847 | 0.423801 | 0.552339 |
| 2004* | - | - | - | - | 0.134619 | 0.039899 | 0.049761 | 0.001650 | 0.675058 | 0.900987 |
| 2005 | 0.684588 | 0.023750 | - | - | 1.162843 | 0.016258 | 0.041416 | 0.000991 | 0.621172 | 2.551018 |
| 2006 | 0.579229 | 0.072287 | 1.132145 | 0.048832 | 1.527248 | 0.085518 | 0.029515 | 0.001481 | 0.717134 | 4.193389 |
| 2007 | 0.679879 | 0.014809 | 1.779104 | 0.029297 | 1.861591 | 0.052063 | 0.060557 | 0.001422 | 0.694930 | 5.173652 |
| 2008 | 0.119145 | 0.001495 | 1.177782 | 0.006659 | 1.100270 | 0.024925 | 0.279901 | 0.002541 | 0.532864 | 3.245582 |
| 2009 | - | - | 0.664586 | 0.002270 | 1.559556 | 0.015674 | 0.186506 | 0.001139 | 0.374187 | 2.803918 |
| 2010* | - | - | - | - | 1.453261 | 0.009179 | 0.031920 | 0.000553 | 0.231367 | 1.726280 |
| 2011* | - | - | - | - | 2.141349 | 0.013272 | 0.017470 | 0.000072 | 0.203984 | 2.376147 |
| 2012* | - | - |  | - | 1.564344 | 0.010297 | 0.042113 | 0.001314 | 0.153263 | 1.771331 |
| 2013 | 0.933101 | 0.011362 | 0.746213 | 0.012106 | 1.841754 | 0.015630 | 0.128942 | 0.001265 | 0.348367 | 4.038740 |
| 2014 | 3.057006 | 0.030467 | 5.306589 | 0.008767 | 5.330041 | 0.050675 | 0.305409 | 0.000997 | 0.435732 | 14.525683 |
| 2015 | 5.467550 | 0.029386 | 6.761436 | 0.028221 | 3.919177 | 0.016818 | 0.204958 | 0.005581 | 0.361220 | 16.794347 |
| $2016{ }^{*}$ | - | - | - | - | 2.575704 | 0.016695 | 0.175692 | 0.004222 | 0.310121 | 3.082434 |
| 2017 | 1.362519 | 0.038489 | - | - | 1.081659 | 0.006841 | 0.183555 | 0.001433 | 0.167927 | 2.842423 |
| 2018 | 1.598424 | 0.034668 | - | - | 0.879726 | 0.008857 | 0.074017 | 0.000131 | 0.190972 | 2.786795 |

Table 4. Comparison of retained catch abundance and biomass used in the previous assessment ("historical") with "current" catch abundance and biomass. Only values since 2005 (highlighted in grey) have been changed.

| year | abundance (num. crab) |  | biomass (millions lbs) |  |
| :---: | :---: | :---: | :---: | :---: |
|  | historical | current | historical | current |
| 1965 | 1,558,362 | 1,558,362 | 4.24 | 4.24 |
| 1966 | 1,981,280 | 1,981,280 | 5.39 | 5.39 |
| 1967 | 11,032,652 | 11,032,652 | 29.98 | 29.98 |
| 1968 | 14,576,228 | 14,576,228 | 39.69 | 39.69 |
| 1969 | 22,394,986 | 22,394,986 | 60.60 | 60.60 |
| 1970 | 22,004,597 | 22,004,597 | 56.20 | 56.20 |
| 1971 | 17,820,914 | 17,820,914 | 45.66 | 45.66 |
| 1972 | 14,906,645 | 14,906,645 | 37.27 | 37.27 |
| 1973 | 12,000,825 | 12,000,825 | 28.72 | 28.72 |
| 1974 | 13,404,770 | 13,404,770 | 33.60 | 33.60 |
| 1975 | 15,603,036 | 15,603,036 | 38.92 | 38.92 |
| 1976 | 26,120,508 | 26,120,508 | 66.17 | 66.17 |
| 1977 | 26,821,995 | 26,821,995 | 78.32 | 78.32 |
| 1978 | 18,780,962 | 18,780,962 | 46.50 | 46.50 |
| 1979 | 16,805,611 | 16,805,611 | 41.90 | 41.90 |
| 1980 | 12,928,112 | 12,928,112 | 29.60 | 29.60 |
| 1981 | 4,830,980 | 4,830,980 | 11.00 | 11.00 |
| 1982 | 2,286,756 | 2,286,756 | 5.27 | 5.27 |
| 1983 | 516,877 | 516,877 | 1.21 | 1.21 |
| 1984 | 1,272,501 | 1,272,501 | 3.15 | 3.15 |
| 1987 | 957,318 | 957,318 | 2.20 | 2.20 |
| 1988 | 2,894,480 | 2,894,480 | 7.01 | 7.01 |
| 1989 | 10,672,607 | 10,672,607 | 24.50 | 24.50 |
| 1990 | 16,609,286 | 16,609,286 | 40.10 | 40.10 |
| 1991 | 12,924,102 | 12,924,102 | 31.80 | 31.80 |
| 1992 | 15,265,865 | 15,265,865 | 35.10 | 35.10 |
| 1993 | 7,236,054 | 7,236,054 | 16.90 | 16.90 |
| 1994 | 3,351,639 | 3,351,639 | 7.80 | 7.80 |
| 1995 | 1,881,525 | 1,881,525 | 4.23 | 4.23 |
| 1996 | 734,303 | 734,303 | 1.81 | 1.81 |
| 2005 | 443,865 | 443,977 | 0.95 | 0.95 |
| 2006 | 926,101 | 926,103 | 2.12 | 2.12 |
| 2007 | 927,164 | 927,164 | 2.11 | 2.11 |
| 2008 | 830,363 | 830,369 | 1.94 | 1.94 |
| 2009 | 485,963 | 485,963 | 1.33 | 1.33 |
| 2013 | 1,426,670 | 1,445,768 | 2.75 | 2.79 |
| 2014 | 7,442,931 | 7,522,844 | 13.58 | 13.70 |
| 2015 | 10,856,418 | 10,856,418 | 19.64 | 19.64 |
| 2017 | 1,340,394 | 1,340,394 | 2.50 | 2.50 |
| 2018 | -- | 1,381,008 | -- | 2.44 |

Table 5. Comparison of total catch biomass in the directed Tanner crab fisheries used in the previous assessment ("historical") with "current" catch biomass dataset. See text for details.

| biomass (millions lbs) <br> males |  |  |  | females <br> year |  |  | historical | current | historical | current |
| ---: | ---: | ---: | ---: | ---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1991 | -- | 56.92 | -- | 4.16 |  |  |  |  |  |  |
| 1992 | 48.71 | 81.59 | 2.21 | 3.76 |  |  |  |  |  |  |
| 1993 | 25.43 | 26.13 | 2.27 | 2.20 |  |  |  |  |  |  |
| 1994 | 14.70 | 16.13 | 2.80 | 1.86 |  |  |  |  |  |  |
| 1995 | 10.32 | 11.17 | 3.88 | 2.35 |  |  |  |  |  |  |
| 1996 | 2.07 | 0.66 | 0.10 | 0.12 |  |  |  |  |  |  |
| 2005 | 1.97 | 1.51 | 0.10 | 0.05 |  |  |  |  |  |  |
| 2006 | 5.14 | 3.77 | 0.78 | 0.27 |  |  |  |  |  |  |
| 2007 | 6.61 | 5.42 | 0.21 | 0.10 |  |  |  |  |  |  |
| 2008 | 2.89 | 2.86 | 0.03 | 0.02 |  |  |  |  |  |  |
| 2009 | 1.49 | 1.47 | 0.01 | 0.01 |  |  |  |  |  |  |
| 2013 | 3.60 | 3.70 | 0.05 | 0.05 |  |  |  |  |  |  |
| 2014 | 19.12 | 18.44 | 0.09 | 0.09 |  |  |  |  |  |  |
| 2015 | 26.35 | 26.96 | 0.13 | 0.13 |  |  |  |  |  |  |
| 2017 | 4.66 | 3.00 | 0.13 | 0.08 |  |  |  |  |  |  |
| 2018 | -- | 3.52 | -- | 0.08 |  |  |  |  |  |  |

Table 6. Comparison of Tanner crab bycatch biomass in the snow crab fisheries used in the previous assessment ("historical") with the "current" catch biomass dataset. See text for details.

| year | biomass (millions lbs) |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
|  | males |  | females |  |
|  | historical | current | historical | current |
| 1990 | -- | 15.61 | -- | 0.23 |
| 1991 | -- | 18.43 | -- | 0.32 |
| 1992 | 56.79 | 5.48 | 3.94 | 0.36 |
| 1993 | 32.03 | 6.34 | 4.00 | 0.88 |
| 1994 | 15.71 | 2.97 | 2.80 | 0.43 |
| 1995 | 10.58 | 2.25 | 3.88 | 0.27 |
| 1996 | 1.84 | 4.32 | 0.51 | 0.26 |
| 1997 | 3.86 | 4.33 | 0.50 | 0.20 |
| 1998 | 4.38 | 1.45 | 0.39 | 0.18 |
| 1999 | 1.53 | 0.29 | 0.32 | 0.02 |
| 2000 | 0.32 | 0.69 | 0.05 | 0.01 |
| 2001 | 0.71 | 1.20 | 0.02 | 0.05 |
| 2002 | 1.23 | 0.37 | 0.08 | 0.03 |
| 2003 | 0.43 | 0.14 | 0.06 | 0.02 |
| 2004 | 0.17 | 0.30 | 0.03 | 0.09 |
| 2005 | 2.13 | 2.56 | 0.09 | 0.04 |
| 2006 | 3.22 | 3.37 | 0.37 | 0.19 |
| 2007 | 4.13 | 4.10 | 0.22 | 0.11 |
| 2008 | 2.47 | 2.43 | 0.11 | 0.05 |
| 2009 | 2.92 | 3.44 | 0.03 | 0.03 |
| 2010 | 2.96 | 3.20 | 0.03 | 0.02 |
| 2011 | 4.67 | 4.72 | 0.03 | 0.03 |
| 2012 | 2.62 | 3.45 | 0.02 | 0.02 |
| 2013 | 4.04 | 4.06 | 0.03 | 0.03 |
| 2014 | 11.87 | 11.75 | 0.11 | 0.11 |
| 2015 | 8.64 | 8.64 | 0.04 | 0.04 |
| 2016 | 5.68 | 5.68 | 0.04 | 0.04 |
| 2017 | 2.45 | 2.38 | 0.02 | 0.02 |
| 2018 | -- | 1.94 | -- | 0.02 |

Table 7. Comparison of Tanner crab bycatch biomass in the BBRKC fishery used in the previous assessment ("historical") with the "current" catch biomass dataset. See text for details.

| biomass (millions lbs) <br> males |  |  |  | females <br> year |  |  | historical | current | historical | current |
| :---: | :---: | :---: | :---: | ---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1990 | -- | 8.21 | -- | 0.08 |  |  |  |  |  |  |
| 1991 | -- | 4.34 | -- | 0.06 |  |  |  |  |  |  |
| 1992 | 2.62 | 2.90 | 0.06 | 0.04 |  |  |  |  |  |  |
| 1993 | 6.54 | 6.90 | 0.44 | 0.33 |  |  |  |  |  |  |
| 1996 | 0.06 | 0.60 | 0.01 | 0.01 |  |  |  |  |  |  |
| 1997 | 0.36 | 0.35 | 0.01 | 0.00 |  |  |  |  |  |  |
| 1998 | 0.26 | 0.25 | 0.01 | 0.00 |  |  |  |  |  |  |
| 1999 | 0.17 | 0.17 | 0.01 | 0.00 |  |  |  |  |  |  |
| 2000 | 0.15 | 0.15 | 0.01 | 0.00 |  |  |  |  |  |  |
| 2001 | 0.09 | 0.09 | 0.00 | 0.00 |  |  |  |  |  |  |
| 2002 | 0.14 | 0.14 | 0.01 | 0.00 |  |  |  |  |  |  |
| 2003 | 0.12 | 0.12 | 0.01 | 0.00 |  |  |  |  |  |  |
| 2004 | 0.11 | 0.11 | 0.01 | 0.00 |  |  |  |  |  |  |
| 2005 | 0.09 | 0.09 | 0.00 | 0.00 |  |  |  |  |  |  |
| 2006 | 0.06 | 0.07 | 0.01 | 0.00 |  |  |  |  |  |  |
| 2007 | 0.12 | 0.13 | 0.02 | 0.00 |  |  |  |  |  |  |
| 2008 | 0.59 | 0.62 | 0.01 | 0.01 |  |  |  |  |  |  |
| 2009 | 0.33 | 0.41 | 0.00 | 0.00 |  |  |  |  |  |  |
| 2010 | 0.07 | 0.07 | 0.00 | 0.00 |  |  |  |  |  |  |
| 2011 | 0.04 | 0.04 | 0.00 | 0.00 |  |  |  |  |  |  |
| 2012 | 0.09 | 0.09 | 0.00 | 0.00 |  |  |  |  |  |  |
| 2013 | 0.25 | 0.28 | 0.00 | 0.00 |  |  |  |  |  |  |
| 2014 | 0.65 | 0.67 | 0.00 | 0.00 |  |  |  |  |  |  |
| 2015 | 0.45 | 0.45 | 0.01 | 0.01 |  |  |  |  |  |  |
| 2016 | 0.39 | 0.39 | 0.01 | 0.01 |  |  |  |  |  |  |
| 2017 | 0.40 | 0.40 | 0.00 | 0.00 |  |  |  |  |  |  |
| 2018 | -- | 0.16 | -- | 0.00 |  |  |  |  |  |  |

Table 8. Retained catch biomass in the directed Tanner crab (TCF), snow crab (SCF), and BBRKC (RKF) fisheries since 2005. The directed fishery was completely closed from 2010/11 to 2012/13, and in 2016/17. Legal-sized Tanner crab can be incidentally-retained in the snow crab and BBRKC fisheries up to a cap of $5 \%$ the target catch.

| year | TCF |  |  |  |  |  | $\begin{gathered} \text { SCF } \\ \text { all EBS } \end{gathered}$ |  | $\begin{aligned} & \text { RKF } \\ & \text { all EBS } \end{aligned}$ |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | West 166 W |  | East 166 W |  | all EBS |  |  |  |  |  |
|  | Abundance | Bionuss (kg) | Abundance | Biomass (kg) | Abundance | Biommss (kg) | Abundance | Biommss (kg) | Abundance | Biommss (kg) |
| 2005 | 376.080 | 365.110 | 0 | 0 | 376.080 | 365.110 | 67.897 | 67.112 | 0 | 0 |
| 2006 | 393,508 | 320. 187 | 583, 650 | 633,937 | 917.158 | 954.124 | 7,115 | 6.784 | 1,830 | 1,883 |
| 2007 | 232.345 | 228,829 | 679.137 | 711,640 | 911.482 | 940,469 | 9.328 | 8,761 | 6. 354 | 6,334 |
| 2008 | 48,171 | 47,157 | 760, 166 | 809, 022 | 808, 337 | 856.179 | 3.300 | 2,535 | 18.732 | 21,068 |
| 2009 | 0 | 0 | 476,668 | 592,417 | 476,668 | 592,417 | 2,544 | 1,714 | 6,751 | 8,402 |
| 2010 | 0 | 0 | 0 | 0 | 0 | 0 | 1,689 | 1,154 | 6 | 3 |
| 2011 | 0 | 0 | 0 | 0 | 0 | 0 | 3.095 | 2,092 | 0 | 0 |
| 2012 | 0 | 0 | 0 | 1 | 0 | 0 | 1,643 | 1,111 | 4 | 3 |
| 2013 | 722,469 | 593,617 | 704, 201 | 654.271 | 1,426,670 | 1,247,888 | 13,256 | 9,882 | 5,842 | 6,322 |
| 2014 | 3, 121,442 | 2,368,693 | 4.378, 199 | 3.829.288 | $7.499,641$ | 6.197 .981 | 19,512 | 14,458 | 3.691 | 3.792 |
| 2015 | 4,817,145 | 3,770.319 | 5,998,876 | 5, 107,722 | 10,816,021 | 8,878,041 | 39.011 | 30, 252 | 1,386 | 1,350 |
| 2016 | 0 | 0 | 0 | 0 | 0 | -0 | 1.733 | 1.177 | 33 | 21 |
| 2017 | 1,322,542 | 1,117, 483 | 139 | 119 | 1,322,681 | 1,117,602 | 17,688 | 15,018 | 25 | 17 |
| 2018 | 1,376,977 | 1,103,903 | 0 | 0 | 1,376,977 | 1,103,903 | 4,013 | 3, 409 | 18 | 12 |

Table 9. Sample sizes for retained catch-at-size in the directed fishery. $\mathrm{N}=$ number of individuals. $\mathrm{N}^{`}=$ scaled sample size used in assessment. The directed fishery was closed in 2016/17.

| year | new +old shell |  |
| :---: | ---: | ---: |
|  | N | $\mathrm{N}^{\prime}$ |
| $1980 / 81$ | 13,310 | 104.6 |
| $1981 / 82$ | 11,311 | 88.9 |
| $1982 / 83$ | 13,519 | 106.2 |
| $1983 / 84$ | 1,675 | 13.2 |
| $1984 / 85$ | 2,542 | 20.0 |
| $1988 / 89$ | 12,380 | 97.3 |
| $1989 / 90$ | 4,123 | 32.4 |
| $1990 / 91$ | 120,676 | 200.0 |
| $1991 / 92$ | 126,299 | 200.0 |
| $1992 / 93$ | 125,193 | 200.0 |
| $1993 / 94$ | 71,622 | 200.0 |
| $1994 / 95$ | 27,658 | 200.0 |
| $1995 / 96$ | 1,525 | 12.0 |
| $1996 / 97$ | 4,430 | 34.8 |
| $2005 / 06$ | 705 | 5.5 |
| $2006 / 07$ | 2,940 | 23.1 |
| $2007 / 08$ | 6,935 | 45.2 |
| $2008 / 09$ | 3,490 | 27.4 |
| $2009 / 10$ | 2,417 | 19.0 |
| $2013 / 14$ | 4,760 | 35.8 |
| $2014 / 15$ | 14,055 | 113.7 |
| $2015 / 16$ | 24,420 | 190.3 |
| $2016 / 17$ | -- | -- |
| $2017 / 18$ | 3,470 | 27.3 |
| $2018 / 19$ | 3,306 | 26.0 |

Table 10. Sample sizes for total catch-at-size in the directed fishery from crab observer sampling. $\mathrm{N}=$ number of individuals. $\mathrm{N}^{`}=$ scaled sample size used in assessment.

|  | $N$ |  | $N^{\prime}$ |  |
| :---: | ---: | ---: | ---: | ---: |
| year | males | females | males | females |
| $1991 / 92$ | 31,252 | 5,605 | 200.0 | 44.0 |
| $1992 / 93$ | 54,836 | 8,755 | 200.0 | 68.8 |
| $1993 / 94$ | 40,388 | 10,471 | 200.0 | 82.3 |
| $1994 / 95$ | 5,792 | 2,132 | 45.5 | 16.7 |
| $1995 / 96$ | 5,589 | 3,119 | 43.9 | 24.5 |
| $1996 / 97$ | 352 | 168 | 2.8 | 1.3 |
| $2005 / 06$ | 19,715 | 1,107 | 154.9 | 8.7 |
| $2006 / 07$ | 24,226 | 4,432 | 190.3 | 34.8 |
| $2007 / 08$ | 61,546 | 3,318 | 200.0 | 26.1 |
| $2008 / 09$ | 29,166 | 646 | 200.0 | 5.1 |
| $2009 / 10$ | 17,289 | 147 | 135.8 | 1.2 |
| $2013 / 14$ | 17,291 | 710 | 135.8 | 5.6 |
| $2014 / 15$ | 85,116 | 1,191 | 200.0 | 9.4 |
| $2015 / 16$ | 119,843 | 1,622 | 200.0 | 12.8 |
| $2016 / 17$ | -- |  | -- | -- |
| $2017 / 18$ | 18,785 | 1,721 | 147.6 | 13.5 |
| $2018 / 19$ | 28,338 | 2,036 | 200.0 | 16.0 |

Table 11. Sample sizes for total bycatch-at-size in the snow crab fishery, from crab observer sampling. N $=$ number of individuals. $\mathrm{N}^{\wedge}=$ scaled sample size used in assessment.

| year | N |  | $\mathrm{N}^{\prime}$ |  |
| :---: | :---: | :---: | :---: | :---: |
|  | males | females | males | females |
| 1990/91 | 14,032 | 478 | 110.2 | 3.8 |
| 1991/92 | 11,708 | 686 | 92.0 | 5.4 |
| 1992/93 | 6,280 | 859 | 49.3 | 6.7 |
| 1993/94 | 6,969 | 1,542 | 54.7 | 12.1 |
| 1994/95 | 2,982 | 1,523 | 23.4 | 12.0 |
| 1995/96 | 1,898 | 428 | 14.9 | 3.4 |
| 1996/97 | 3,265 | 662 | 25.6 | 5.2 |
| 1997/98 | 3,970 | 657 | 31.2 | 5.2 |
| 1998/99 | 1,911 | 324 | 15.0 | 2.5 |
| 1999/00 | 976 | 82 | 7.7 | 0.6 |
| 2000/01 | 1,237 | 74 | 9.7 | 0.6 |
| 2001/02 | 3,113 | 160 | 24.5 | 1.3 |
| 2002/03 | 982 | 118 | 7.7 | 0.9 |
| 2003/04 | 688 | 152 | 5.4 | 1.2 |
| 2004/05 | 833 | 707 | 6.5 | 5.6 |
| 2005/06 | 9,807 | 368 | 77.0 | 2.9 |
| 2006/07 | 10,391 | 1,256 | 81.6 | 9.9 |
| 2007/08 | 13,797 | 728 | 108.4 | 5.7 |
| 2008/09 | 8,455 | 722 | 66.4 | 5.7 |
| 2009/10 | 11,057 | 474 | 86.9 | 3.7 |
| 2010/11 | 12,073 | 250 | 94.8 | 2.0 |
| 2011/12 | 9,453 | 189 | 74.3 | 1.5 |
| 2012/13 | 11,004 | 270 | 86.4 | 2.1 |
| 2013/14 | 12,935 | 356 | 101.6 | 2.8 |
| 2014/15 | 24,878 | 804 | 195.4 | 6.3 |
| 2015/16 | 19,839 | 230 | 155.9 | 1.8 |
| 2016/17 | 16,369 | 262 | 128.6 | 2.1 |
| 2017/18 | 5,598 | 109 | 44.0 | 0.9 |
| 2018/19 | 6,145 | 233 | 48.3 | 1.8 |

Table 12. Sample sizes for total bycatch-at-size in the BBRKC fishery, from crab observer sampling. $\mathrm{N}=$ number of individuals. $\mathrm{N}^{`}=$ scaled sample size used in assessment.

| year | N |  | $\mathrm{N}^{\prime}$ |  |
| :---: | :---: | :---: | :---: | :---: |
|  | males | females | males | females |
| 1990/91 | 1,580 | 43 | 12.4 | 0.3 |
| 1991/92 | 2,273 | 89 | 17.9 | 0.7 |
| 1992/93 | 2,056 | 105 | 16.2 | 0.8 |
| 1993/94 | 7,359 | 1,196 | 57.8 | 9.4 |
| 1997/98 | 1,030 | 41 | 8.1 | 0.3 |
| 1998/99 | 457 | 20 | 3.6 | 0.2 |
| 1999/00 | 207 | 14 | 1.6 | 0.1 |
| 2000/01 | 845 | 44 | 6.6 | 0.3 |
| 2001/02 | 456 | 39 | 3.6 | 0.3 |
| 2002/03 | 750 | 50 | 5.9 | 0.4 |
| 2003/04 | 555 | 46 | 4.4 | 0.4 |
| 2004/05 | 487 | 44 | 3.8 | 0.3 |
| 2005/06 | 983 | 70 | 7.7 | 0.5 |
| 2006/07 | 746 | 68 | 5.9 | 0.5 |
| 2007/08 | 1,360 | 89 | 10.7 | 0.7 |
| 2008/09 | 3,797 | 121 | 29.8 | 1.0 |
| 2009/10 | 2,871 | 70 | 22.6 | 0.5 |
| 2010/11 | 582 | 28 | 4.6 | 0.2 |
| 2011/12 | 323 | 4 | 2.5 | 0.0 |
| 2012/13 | 618 | 48 | 4.9 | 0.4 |
| 2013/14 | 2,110 | 60 | 16.6 | 0.5 |
| 2014/15 | 3,110 | 32 | 24.4 | 0.3 |
| 2015/16 | 2,175 | 186 | 17.1 | 1.5 |
| 2016/17 | 3,220 | 246 | 25.3 | 1.9 |
| 2017/18 | 3,782 | 86 | 29.7 | 0.7 |
| 2018/19 | 1,283 | 6 | 10.1 | 0.0 |

Table 13. Sample sizes for total catch-at-size in the groundfish fisheries, from groundfish observer sampling. $\mathrm{N}=$ number of individuals. $\mathrm{N}^{`}=$ scaled sample size used in the assessment.

| year | N |  | $\mathrm{N}^{\prime}$ |  |
| :---: | :---: | :---: | :---: | :---: |
|  | males | females | males | females |
| 1973/74 | 3,155 | 2,277 | 24.8 | 17.9 |
| 1974/75 | 2,492 | 1,600 | 19.6 | 12.6 |
| 1975/76 | 1,251 | 839 | 9.8 | 6.6 |
| 1976/77 | 6,950 | 6,683 | 54.6 | 52.5 |
| 1977/78 | 10,685 | 8,386 | 83.9 | 65.9 |
| 1978/79 | 18,596 | 13,665 | 146.1 | 107.4 |
| 1979/80 | 19,060 | 11,349 | 149.7 | 89.2 |
| 1980/81 | 12,806 | 5,917 | 100.6 | 46.5 |
| 1981/82 | 6,098 | 4,065 | 47.9 | 31.9 |
| 1982/83 | 13,439 | 8,006 | 105.6 | 62.9 |
| 1983/84 | 18,363 | 8,305 | 144.3 | 65.2 |
| 1984/85 | 27,403 | 13,771 | 200.0 | 108.2 |
| 1985/86 | 23,128 | 12,728 | 181.7 | 100.0 |
| 1986/87 | 14,860 | 7,626 | 116.7 | 59.9 |
| 1987/88 | 23,508 | 15,857 | 184.7 | 124.6 |
| 1988/89 | 10,586 | 7,126 | 83.2 | 56.0 |
| 1989/90 | 59,943 | 41,234 | 200.0 | 200.0 |
| 1990/91 | 23,545 | 11,212 | 185.0 | 88.1 |
| 1991/92 | 6,817 | 3,479 | 53.6 | 27.3 |
| 1992/93 | 3,128 | 1,175 | 24.6 | 9.2 |
| 1993/94 | 1,217 | 358 | 9.6 | 2.8 |
| 1994/95 | 3,628 | 1,820 | 28.5 | 14.3 |
| 1995/96 | 3,904 | 2,669 | 30.7 | 21.0 |
| 1996/97 | 8,306 | 3,400 | 65.3 | 26.7 |
| 1997/98 | 9,949 | 3,900 | 78.2 | 30.6 |
| 1998/99 | 12,105 | 4,440 | 95.1 | 34.9 |
| 1999/00 | 11,053 | 4,522 | 86.8 | 35.5 |
| 2000/01 | 12,895 | 3,087 | 101.3 | 24.3 |
| 2001/02 | 15,788 | 3,083 | 124.0 | 24.2 |
| 2002/03 | 15,401 | 3,249 | 121.0 | 25.5 |
| 2003/04 | 9,572 | 2,733 | 75.2 | 21.5 |
| 2004/05 | 13,844 | 4,460 | 108.8 | 35.0 |
| 2005/06 | 17,785 | 3,709 | 139.7 | 29.1 |
| 2006/07 | 15,903 | 3,047 | 124.9 | 23.9 |
| 2007/08 | 16,148 | 3,819 | 126.9 | 30.0 |
| 2008/09 | 26,171 | 4,235 | 200.0 | 33.3 |
| 2009/10 | 19,075 | 2,704 | 149.9 | 21.2 |
| 2010/11 | 15,131 | 2,275 | 118.9 | 17.9 |
| 2011/12 | 16,119 | 4,244 | 126.6 | 33.3 |
| 2012/13 | 12,987 | 3,083 | 102.0 | 24.2 |
| 2013/14 | 28,782 | 6,064 | 200.0 | 47.6 |
| 2014/15 | 39,119 | 4,212 | 200.0 | 33.1 |
| 2015/16 | 27,428 | 5,735 | 200.0 | 45.1 |
| 2016/17 | 18,313 | 4,299 | 143.9 | 33.8 |
| 2017/18 | 12,541 | 1,229 | 98.5 | 9.7 |
| 2018/19 | 7,004 | 1,227 | 55.0 | 9.6 |

Table 14. Trends in Tanner crab biomass (metric tons) in the NMFS EBS summer bottom trawl survey, by sex and area.

|  | male |  |  |  | female |  |  |  |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: | :---: | :---: |
| year | W166 | E166 | all EBS | W166 | E166 | all EBS |  |  |
| 1975 | 80,689 | 214,202 | 294,891 | 13,374 | 27,594 | 40,968 |  |  |
| 1976 | 55,092 | 101,958 | 157,050 | 12,140 | 25,420 | 37,560 |  |  |
| 1977 | 51,038 | 87,463 | 138,501 | 21,613 | 31,435 | 53,048 |  |  |
| 1978 | 25,394 | 72,913 | 98,308 | 14,167 | 18,406 | 32,574 |  |  |
| 1979 | 32,058 | 17,978 | 50,036 | 19,701 | 3,448 | 23,149 |  |  |
| 1980 | 103,505 | 48,979 | 152,484 | 64,420 | 12,883 | 77,303 |  |  |
| 1981 | 56,540 | 23,390 | 79,930 | 35,525 | 8,577 | 44,102 |  |  |
| 1982 | 49,255 | 16,602 | 65,856 | 57,757 | 8,107 | 65,864 |  |  |
| 1983 | 24,708 | 13,337 | 38,045 | 17,418 | 5,350 | 22,769 |  |  |
| 1984 | 18,490 | 12,020 | 30,510 | 12,358 | 4,800 | 17,158 |  |  |
| 1985 | 6,676 | 8,231 | 14,907 | 3,393 | 3,160 | 6,554 |  |  |
| 1986 | 11,986 | 9,625 | 21,612 | 2,570 | 3,504 | 6,074 |  |  |
| 1987 | 16,648 | 28,863 | 45,511 | 5,137 | 15,009 | 20,146 |  |  |
| 1988 | 41,093 | 58,130 | 99,223 | 12,668 | 22,885 | 35,553 |  |  |
| 1989 | 45,106 | 87,718 | 132,824 | 12,254 | 18,975 | 31,230 |  |  |
| 1990 | 55,539 | 76,879 | 132,418 | 22,532 | 25,022 | 47,554 |  |  |
| 1991 | 55,986 | 89,825 | 145,811 | 20,445 | 31,341 | 51,787 |  |  |
| 1992 | 37,674 | 89,918 | 127,592 | 16,857 | 11,358 | 28,215 |  |  |
| 1993 | 19,877 | 53,394 | 73,271 | 7,382 | 5,325 | 12,707 |  |  |
| 1994 | 16,032 | 32,303 | 48,335 | 5,716 | 5,332 | 11,048 |  |  |
| 1995 | 15,310 | 19,672 | 34,982 | 7,474 | 5,982 | 13,456 |  |  |
| 1996 | 10,790 | 19,979 | 30,770 | 4,470 | 6,548 | 11,019 |  |  |
| 1997 | 5,561 | 9,088 | 14,649 | 1,893 | 2,914 | 4,806 |  |  |
| 1998 | 6,604 | 8,404 | 15,008 | 2,489 | 1,752 | 4,241 |  |  |
| 1999 | 6,719 | 14,835 | 21,554 | 3,347 | 3,360 | 6,708 |  |  |
| 2000 | 6,903 | 16,429 | 23,332 | 2,999 | 3,613 | 6,613 |  |  |

Table 14 (cont). Trends in Tanner crab biomass (metric tons) in the NMFS EBS summer bottom trawl survey, by sex and area.

|  |  | male |  |  |  | female |  |  |
| :--- | :--- | :--- | ---: | ---: | ---: | ---: | :---: | :---: |
| year | W166 | E166 | all EBS | W166 | E166 | all EBS |  |  |
| 2001 | 13,089 | 16,231 | 29,320 | 6,989 | 3,931 | 10,920 |  |  |
| 2002 | 13,010 | 14,402 | 27,411 | 6,499 | 3,469 | 9,968 |  |  |
| 2003 | 20,661 | 17,164 | 37,825 | 10,297 | 2,795 | 13,092 |  |  |
| 2004 | 26,468 | 12,455 | 38,923 | 7,731 | 1,131 | 8,862 |  |  |
| 2005 | 46,313 | 17,443 | 63,756 | 17,469 | 4,493 | 21,962 |  |  |
| 2006 | 72,907 | 28,636 | 101,543 | 21,723 | 6,476 | 28,198 |  |  |
| 2007 | 76,285 | 27,938 | 104,223 | 12,465 | 6,612 | 19,076 |  |  |
| 2008 | 47,736 | 37,177 | 84,913 | 9,444 | 5,079 | 14,523 |  |  |
| 2009 | 32,653 | 14,786 | 47,439 | 6,495 | 4,553 | 11,048 |  |  |
| 2010 | 34,601 | 14,426 | 49,027 | 6,366 | 2,910 | 9,276 |  |  |
| 2011 | 39,321 | 23,390 | 62,712 | 9,190 | 6,615 | 15,805 |  |  |
| 2012 | 34,764 | 45,367 | 80,131 | 9,787 | 14,245 | 24,032 |  |  |
| 2013 | 38,839 | 64,580 | 103,420 | 10,866 | 13,398 | 24,264 |  |  |
| 2014 | 50,739 | 58,196 | 108,936 | 8,728 | 8,648 | 17,377 |  |  |
| 2015 | 39,158 | 35,093 | 74,251 | 7,574 | 5,304 | 12,878 |  |  |
| 2016 | 43,315 | 25,520 | 68,835 | 7,133 | 1,479 | 8,612 |  |  |
| 2017 | 29,685 | 23,952 | 53,637 | 6,274 | 2,144 | 8,418 |  |  |
| 2018 | 32,734 | 13,769 | 46,503 | 8,213 | 1,588 | 9,801 |  |  |
| 2019 | 17,503 | 10,790 | 28,293 | 7,452 | 2,133 | 9,585 |  |  |

Table 15. Trends in biomass for preferred-size ( $>125 \mathrm{~mm} \mathrm{CW}$ ) male Tanner crab in the NMFS EBS summer bottom trawl survey (in metric tons).

| year | W166 |  |  | E166 |  |  | all EBS |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | new shell | old shell | all | new shell | old shell | all | new shell | old shell | all |
| 1975 | 56,181 | 2,509 | 58,691 | 152,683 | 6,522 | 159,205 | 208,864 | 9,032 | 217,896 |
| 1976 | 38,107 | 1,534 | 39,640 | 57,034 | 9,674 | 66,709 | 95,141 | 11,208 | 106,349 |
| 1977 | 26,511 | 6,808 | 33,319 | 50,855 | 7,543 | 58,399 | 77,366 | 14,351 | 91,717 |
| 1978 | 3,221 | 6,626 | 9,847 | 40,633 | 9,780 | 50,413 | 43,853 | 16,406 | 60,259 |
| 1979 | 4,115 | 3,745 | 7,860 | 9,767 | 3,426 | 13,192 | 13,882 | 7,171 | 21,052 |
| 1980 | 11,210 | 1,677 | 12,887 | 23,184 | 10,857 | 34,041 | 34,394 | 12,534 | 46,927 |
| 1981 | 5,884 | 2,167 | 8,050 | 3,445 | 11,286 | 14,731 | 9,329 | 13,452 | 22,781 |
| 1982 | 5,763 | 5,859 | 11,622 | 3,009 | 4,851 | 7,860 | 8,772 | 10,710 | 19,481 |
| 1983 | 2,416 | 3,240 | 5,655 | 5,151 | 2,082 | 7,233 | 7,566 | 5,322 | 12,889 |
| 1984 | 571 | 3,159 | 3,730 | 4,348 | 3,077 | 7,424 | 4,919 | 6,236 | 11,154 |
| 1985 | 588 | 870 | 1,458 | 4,055 | 1,046 | 5,101 | 4,642 | 1,917 | 6,559 |
| 1986 | 142 | 674 | 816 | 734 | 2,546 | 3,280 | 876 | 3,219 | 4,096 |
| 1987 | 3,505 | 658 | 4,163 | 4,911 | 3,473 | 8,385 | 8,416 | 4,132 | 12,548 |
| 1988 | 9,690 | 929 | 10,618 | 15,698 | 2,715 | 18,413 | 25,387 | 3,644 | 29,031 |
| 1989 | 13,758 | 2,741 | 16,499 | 37,364 | 3,740 | 41, 104 | 51,122 | 6,481 | 57,603 |
| 1990 | 21,082 | 3,274 | 24,356 | 35,903 | 7,084 | 42,987 | 56,985 | 10,358 | 67,343 |
| 1991 | 13,386 | 8,430 | 21,816 | 32,973 | 14,476 | 47,449 | 46,359 | 22,906 | 69,265 |
| 1992 | 9,851 | 6,461 | 16,311 | 41,423 | 16,242 | 57,665 | 51,274 | 22,703 | 73,977 |
| 1993 | 3,716 | 2,596 | 6,312 | 22,942 | 11,990 | 34,932 | 26,658 | 14,586 | 41,244 |
| 1994 | 1,248 | 4,143 | 5,391 | 10,000 | 13,912 | 23,912 | 11,248 | 18,054 | 29,303 |
| 1995 | 370 | 5,392 | 5,761 | 1,241 | 13,516 | 14,757 | 1,611 | 18,907 | 20,518 |
| 1996 | 100 | 3,580 | 3,680 | 330 | 13,912 | 14,242 | 430 | 17,492 | 17,922 |
| 1997 | 163 | 958 | 1,121 | 316 | 4,245 | 4,561 | 478 | 5,203 | 5,681 |
| 1998 | 441 | 644 | 1,085 | 1,001 | 2,604 | 3,605 | 1,442 | 3,247 | 4,689 |
| 1999 | 256 | 356 | 612 | 1,645 | 1,838 | 3,483 | 1,902 | 2,194 | 4,095 |
| 2000 | 250 | 377 | 627 | 4,484 | 3,045 | 7,529 | 4,734 | 3,422 | 8,156 |

Table 15 (cont.). Trends in biomass for preferred-size (> 125 mm CW) male Tanner crab in the NMFS EBS summer bottom trawl survey (in metric tons).

| year | W166 |  |  | E166 |  |  | all EBS |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | new shell | old shell | all | new shell | old shell | all | new shell | old shell | all |
| 2001 | 418 | 1,361 | 1,780 | 4,473 | 3,600 | 8,073 | 4,892 | 4,961 | 9,853 |
| 2002 | 384 | 838 | 1,222 | 944 | 7,102 | 8,046 | 1,328 | 7,940 | 9,268 |
| 2003 | 434 | 2,227 | 2, 661 | 1,558 | 6,433 | 7,991 | 1,992 | 8,660 | 10,652 |
| 2004 | 980 | 1,825 | 2,805 | 1,597 | 4,916 | 6,513 | 2,577 | 6,741 | 9,318 |
| 2005 | 8,776 | 5,062 | 13,839 | 2,368 | 5,822 | 8,190 | 11,145 | 10,884 | 22,029 |
| 2006 | 3,755 | 15,328 | 19,083 | 2,134 | 6,794 | 8,927 | 5,889 | 22,122 | 28,011 |
| 2007 | 8,523 | 7,757 | 16,281 | 4,143 | 5,314 | 9,457 | 12,666 | 13,071 | 25,737 |
| 2008 | 8,688 | 4,457 | 13,145 | 15,476 | 3,288 | 18,764 | 24, 163 | 7,745 | 31,909 |
| 2009 | 6,657 | 4,156 | 10,812 | 2,644 | 5,139 | 7,783 | 9,300 | 9,295 | 18,595 |
| 2010 | 9,593 | 4,867 | 14,460 | 3,006 | 4,576 | 7,582 | 12,599 | 9,443 | 22,042 |
| 2011 | 9, 023 | 6,637 | 15,660 | 1,513 | 6,987 | 8,500 | 10,536 | 13,624 | 24,160 |
| 2012 | 2,368 | 3,997 | 6,365 | 3,352 | 5,026 | 8,378 | 5,720 | 9,023 | 14,743 |
| 2013 | 5,383 | 2,837 | 8,220 | 10,871 | 3,527 | 14,397 | 16,254 | 6,364 | 22,618 |
| 2014 | 7,163 | 4,604 | 11,766 | 14,899 | 9,310 | 24,210 | 22,062 | 13,914 | 35, 976 |
| 2015 | 8,380 | 5,925 | 14,306 | 9,084 | 10,217 | 19,301 | 17,464 | 16,143 | 33,607 |
| 2016 | 5,799 | 12,527 | 18,326 | 2,640 | 8,055 | 10,695 | 8,439 | 20,582 | 29, 021 |
| 2017 | 894 | 11,659 | 12,553 | 1,629 | 10,841 | 12,470 | 2, 523 | 22,500 | 25,024 |
| 2018 | 996 | 11,875 | 12,871 | 102 | 7,253 | 7,355 | 1,097 | 19,128 | 20,225 |
| 2019 | 202 | 4,799 | 5,001 | 315 | 4,455 | 4,769 | 517 | 9,254 | 9,771 |

Table 16. Sample sizes for NMFS survey size composition data. In the assessment model, an input sample size of 200 is used for all survey-related compositional data.

| year | number of hauls | females |  |  |  |  |  | males |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | immature |  | mature |  |  |  | immature |  | mature |  |  |  |
|  |  | $\begin{array}{cc}\begin{array}{c}\text { number of } \\ \text { nonzero hauls }\end{array} & \begin{array}{c}\text { number of } \\ \text { crab }\end{array}\end{array}$ |  | number of nonzero hauls | number of | number of | number of crab | number of number of <br> nonzero hauls crab |  | number of | number of | number of | number of |
| 1975 | 136 | 73 | 1,047 | 91 | 1,861 | 39 | 706 | 127 | 2,895 | 127 | 3,993 | 80 | 399 |
| 1976 | 214 | 88 | 1,097 | 91 | 1,304 | 39 | 311 | 130 | 2,023 | 130 | 2,469 | 47 | 242 |
| 1977 | 155 | 69 | 776 | 76 | 1,183 | 60 | 738 | 114 | 1,778 | 114 | 1,971 | 79 | 485 |
| 1978 | 230 | 88 | 1,949 | 82 | 638 | 65 | 1,307 | 147 | 2,957 | 147 | 1,570 | 104 | 700 |
| 1979 | 307 | 74 | 733 | 62 | 735 | 42 | 341 | 138 | 1,805 | 138 | 808 | 68 | 306 |
| 1980 | 320 | 103 | 1,491 | 95 | 1,471 | 49 | 570 | 164 | 4,602 | 164 | 2,359 | 71 | 569 |
| 1981 | 305 | 71 | 579 | 79 | 1,319 | 94 | 1,206 | 158 | 3,809 | 158 | 2,293 | 116 | 886 |
| 1982 | 342 | 87 | 823 | 72 | 457 | 103 | 2,384 | 181 | 1,751 | 181 | 1,371 | 147 | 2,082 |
| 1983 | 353 | 102 | 2,113 | 56 | 201 | 102 | 2,154 | 166 | 2,484 | 166 | 983 | 132 | 1,181 |
| 1984 | 355 | 135 | 1,879 | 53 | 284 | 94 | 1,531 | 171 | 1,965 | 171 | 490 | 126 | 1,399 |
| 1985 | 353 | 141 | 847 | 52 | 228 | 65 | 601 | 179 | 1,060 | 179 | 381 | 86 | 459 |
| 1986 | 353 | 162 | 1,588 | 64 | 191 | 68 | 331 | 213 | 2,141 | 213 | 528 | 115 | 468 |
| 1987 | 355 | 189 | 4,230 | 105 | 445 | 73 | 392 | 226 | 4,659 | 226 | 1,306 | 103 | 498 |
| 1988 | 370 | 206 | 3,735 | 149 | 1,753 | 100 | 530 | 252 | 5,627 | 252 | 2,210 | 101 | 475 |
| 1989 | 373 | 204 | 3,271 | 144 | 1,241 | 108 | 882 | 237 | 4,977 | 237 | 3,201 | 135 | 1,067 |
| 1990 | 370 | 198 | 3,114 | 155 | 1,502 | 126 | 1,511 | 247 | 5,107 | 247 | 3,149 | 151 | 1,342 |
| 1991 | 371 | 163 | 2,259 | 138 | 1,283 | 141 | 2,568 | 227 | 4,361 | 227 | 2,692 | 181 | 2,893 |
| 1992 | 355 | 107 | 1,494 | 119 | 820 | 123 | 2,205 | 215 | 2,958 | 215 | 2,047 | 177 | 1,924 |
| 1993 | 374 | 99 | 869 | 96 | 545 | 122 | 1,337 | 207 | 2,051 | 207 | 1,677 | 180 | 1,865 |
| 1994 | 374 | 97 | 921 | 52 | 148 | 104 | 1,293 | 175 | 1,281 | 175 | 724 | 174 | 1,827 |
| 1995 | 375 | 115 | 834 | 35 | 140 | 107 | 1,057 | 153 | 958 | 153 | 220 | 137 | 1,611 |
| 1996 | 374 | 115 | 883 | 57 | 109 | 98 | 963 | 148 | 1,069 | 148 | 222 | 134 | 1,414 |
| 1997 | 375 | 116 | 1,329 | 62 | 168 | 83 | 504 | 161 | 1,336 | 161 | 289 | 125 | 582 |
| 1998 | 374 | 146 | 1,710 | 53 | 160 | 73 | 344 | 176 | 2,032 | 176 | 396 | 128 | 624 |
| 1999 | 372 | 138 | 2,628 | 52 | 255 | 85 | 510 | 170 | 2,816 | 170 | 550 | 124 | 567 |
| 2000 | 371 | 142 | 2,249 | 61 | 242 | 55 | 345 | 188 | 2,836 | 188 | 628 | 133 | 653 |

Table16 (cont.). Sample sizes for NMFS survey size composition data. In the assessment model, an input sample size of 200 is used for all surveyrelated compositional data.

| year | number of hauls | females |  |  |  |  |  | males |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | immature |  | mature |  |  |  | immature |  | new | shell | old s |  |
|  |  | new shell  <br> number of number of <br> nonzero hauls crab  |  | number of nonzero hauls | number of crab | number of nonzero hauls | number of crab | number of nonzero hauls | number of crab | number of nonzero hauls | number of crab | number of nonzero hauls | number of crab |
| 2001 | 374 | 164 | 3,678 | 83 | 364 | 72 | 644 | 211 | 4,036 | 211 | 629 | 145 | 817 |
| 2002 | 374 | 155 | 3,585 | 81 | 350 | 70 | 500 | 186 | 3,912 | 186 | 458 | 154 | 1,089 |
| 2003 | 375 | 153 | 2,834 | 111 | 923 | 83 | 752 | 203 | 4,754 | 203 | 900 | 153 | 1,349 |
| 2004 | 374 | 175 | 3,922 | 90 | 427 | 80 | 656 | 236 | 4,568 | 236 | 1,027 | 179 | 1,873 |
| 2005 | 372 | 201 | 3,352 | 103 | 634 | 74 | 928 | 254 | 4,496 | 254 | 1,280 | 185 | 1,753 |
| 2006 | 375 | 211 | 4,364 | 143 | 1,332 | 125 | 1,327 | 254 | 6,224 | 254 | 1,757 | 211 | 4,054 |
| 2007 | 375 | 186 | 2,430 | 138 | 1,311 | 136 | 1,396 | 261 | 4,697 | 261 | 1,982 | 201 | 2,907 |
| 2008 | 374 | 153 | 1,747 | 104 | 580 | 120 | 1,783 | 240 | 3,127 | 240 | 2,116 | 196 | 2,146 |
| 2009 | 375 | 171 | 2,408 | 75 | 363 | 115 | 1,317 | 216 | 2,879 | 216 | 1,144 | 187 | 1,954 |
| 2010 | 375 | 186 | 3,180 | 67 | 245 | 104 | 941 | 223 | 3,654 | 223 | 1,268 | 166 | 1,702 |
| 2011 | 375 | 193 | 5,044 | 90 | 471 | 102 | 705 | 210 | 6,095 | 210 | 1,115 | 167 | 1,941 |
| 2012 | 375 | 195 | 3,611 | 100 | 942 | 97 | 720 | 215 | 5,526 | 215 | 1,564 | 139 | 1,296 |
| 2013 | 375 | 163 | 2,917 | 116 | 1,417 | 101 | 1,002 | 207 | 5,592 | 207 | 2,675 | 137 | 1,344 |
| 2014 | 375 | 165 | 2,211 | 98 | 482 | 121 | 1,584 | 222 | 4,746 | 222 | 3,286 | 167 | 2,829 |
| 2015 | 375 | 118 | 1,455 | 60 | 445 | 94 | 1,363 | 225 | 2,737 | 225 | 1,859 | 200 | 2,817 |
| 2016 | 375 | 110 | 1,373 | 56 | 370 | 82 | 1,248 | 222 | 2,235 | 222 | 1,170 | 218 | 3,668 |
| 2017 | 375 | 131 | 2,033 | 50 | 213 | 99 | 1,125 | 186 | 2,241 | 186 | 424 | 205 | 3,541 |
| 2018 | 375 | 196 | 4,666 | 68 | 525 | 93 | 703 | 222 | 4,990 | 222 | 513 | 190 | 2,748 |
| 2019 | 375 | 181 | 3,810 | 85 | 649 | 55 | 541 | 208 | 4,216 | 208 | 522 | 169 | 1,175 |

Table 17. Effort data (potlifts) in the crab fisheries, by area. TCF: directed Tanner crab fishery; SCF: snow crab fishery; RKF: Bristol Bay red king crab fishery.

|  | SCF <br> year | RKF <br> all EBS |
| :---: | :---: | ---: |
| 1953 | - | 30,083 |
| 1954 | - | 17,122 |
| 1955 | - | 28,045 |
| 1956 | - | 41,629 |
| 1957 | - | 23,659 |
| 1958 | - | 27,932 |
| 1959 | - | 22,187 |
| 1960 | - | 26,347 |
| 1961 | - | 72,646 |
| 1962 | - | 123,643 |
| 1963 | - | 181,799 |
| 1964 | - | 180,809 |
| 1965 | - | 127,973 |
| 1966 | - | 129,306 |
| 1967 | - | 135,283 |
| 1968 | - | 184,666 |
| 1969 | - | 175,374 |
| 1970 | - | 168,059 |
| 1971 | - | 126,305 |
| 1972 | - | 208,469 |
| 1973 | - | 194,095 |
| 1974 | - | 212,915 |
| 1975 | - | 205,096 |
| 1976 | - | 321,010 |
| 1977 | - | 451,273 |
| 1978 | 190,746 | 406,165 |
| 1979 | 255,102 | 315,226 |
| 1980 | 435,742 | 567,292 |
| 1981 | 469,091 | 536,646 |
| 1982 | 287,127 | 140,492 |
| 1983 | 173,591 | - |
| 1984 | 370,082 | 107,406 |
| 1985 | 542,346 | 84,443 |
| 1986 | 616,113 | 175,753 |
| 1987 | 747,395 | 220,971 |
| 1988 | 665,242 | 146,179 |
| 1989 | 912,718 | 205,528 |
|  |  |  |
|  | - |  |

Table 17 (cont.). Effort data (potlifts) in the crab fisheries, by area. TCF: directed Tanner crab fishery; SCF: snow crab fishery; RKF: Bristol Bay red king crab fishery.

| year | West 166W | TCF <br> East 166W | all EBS | all EBS | RKF EBS |
| ---: | ---: | ---: | ---: | ---: | ---: |
| 1990 | 479 | 493,820 | 494,299 | $1,382,908$ | 262,761 |
| 1991 | 140,050 | 360,864 | 500,914 | $1,278,502$ | 227,555 |
| 1992 | 166,670 | 508,922 | 675,592 | 969,209 | 206,815 |
| 1993 | 40,100 | 286,620 | 326,720 | 716,524 | 254,389 |
| 1994 | 21,282 | 228,254 | 249,536 | 507,603 | 697 |
| 1995 | 46,454 | 201,988 | 248,442 | 520,685 | 547 |
| 1996 | 8,533 | 64,989 | 73,522 | 754,140 | 77,081 |
| 1997 | - | - | - | 930,794 | 91,085 |
| 1998 | - | - | - | 945,533 | 145,689 |
| 1999 | - | - | - | 182,634 | 151,212 |
| 2000 | - | - | - | 191,200 | 104,056 |
| 2001 | - | - | - | 326,977 | 66,947 |
| 2002 | - | - | - | 153,862 | 72,514 |
| 2003 | - | - | - | 123,709 | 134,515 |
| 2004 | - | - | - | 75,095 | 97,621 |
| 2005 | 6,346 | - | 6,346 | 117,375 | 116,320 |
| 2006 | 4,517 | 15,273 | 19,790 | 86,328 | 72,404 |
| 2007 | 7,268 | 26,441 | 33,709 | 140,857 | 113,948 |
| 2008 | 2,336 | 19,401 | 21,737 | 163,537 | 139,937 |
| 2009 | - | 6,635 | 6,635 | 137,292 | 119,261 |
| 2010 | - | - | - | 147,478 | 132,183 |
| 2011 | - | - | - | 270,602 | 45,784 |
| 2012 | - | - | - | 225,627 | 38,842 |
| 2013 | 23,062 | 16,613 | 39,675 | 225,245 | 46,589 |
| 2014 | 68,695 | 72,768 | 141,463 | 279,183 | 57,725 |
| 2015 | 84,933 | 130,302 | 215,235 | 202,526 | 48,763 |
| 2016 | - | - | - | 118,548 | 33,608 |
| 2017 | 19,284 |  | 11 | 19,295 | 114,673 |
| 2018 | 29,833 | - | 29,833 | 119,484 | 39,169 |
|  |  |  |  |  | 31,975 |

Table 18 . Non-selectivity parameters from all model scenarios that were estimated within $1 \%$ of bounds.

| category | name | index | scenario | which? | bound | description |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| fisheries | pLgtRet[1] | 1 | M19F00 | at upper bound | 15 | TCF: logit-scale max retention (pre-1997) |
|  |  |  | M19F00a | at upper bound | 15 | TCF: logit-scale max retention (pre-1997) |
|  |  |  | M19F01 | at upper bound | 15 | TCF: logit-scale max retention (pre-1997) |
|  |  |  | M19F02 | at upper bound | 15 | TCF: logit-scale max retention (pre-1997) |
|  |  |  | M19F03 | at upper bound | 15 | TCF: logit-scale max retention (pre-1997) |
|  |  |  | M19F04 | at upper bound | 15 | TCF: logit-scale max retention (pre-1997) |
|  |  |  | M19F05 | at upper bound | 15 | TCF: logit-scale max retention (pre-1997) |
| population processes | $\mathrm{pvLgtPrM} 2 \mathrm{M}[1]$ | 32 | M19F00 | at upper bound | 15 | males (entire model period) |
|  |  |  | M19F00a | at upper bound | 15 | males (entire model period) |
|  |  |  | M19F01 | at upper bound | 15 | males (entire model period) |
|  |  |  | M19F02 | at upper bound | 15 | males (entire model period) |
|  |  |  | M19F04 | at upper bound | 15 | males (entire model period) |
|  | pvLgtPrM2M[2] | 1 | M19F00 | at lower bound | -15 | females (entire model period) |
|  |  |  | M19F00a | at lower bound | -15 | females (entire model period) |
|  |  |  | M19F01 | at lower bound | -15 | females (entire model period) |
|  |  |  | M19F02 | at lower bound | -15 | females (entire model period) |
|  |  |  | M19F04 | at lower bound | -15 | females (entire model period) |
| surveys | $\mathrm{pQ}[1]$ | 1 | M19F00 | at lower bound | 0.5 | NMFS trawl survey: males, 1975-1981 |
|  |  |  | M19F00a | at lower bound | 0.5 | NMFS trawl survey: males, 1975-1981 |
|  |  |  | M19F01 | at lower bound | 0.5 | NMFS trawl survey: males, 1975-1981 |
|  |  |  | M19F02 | at lower bound | 0.5 | NMFS trawl survey: males, 1975-1981 |
|  |  |  | M19F03 | at lower bound | 0.5 | NMFS trawl survey: males, 1975-1981 |
|  |  |  | M19F04 | at lower bound | 0.5 | NMFS trawl survey: males, 1975-1981 |
|  |  |  | M19F05 | at lower bound | 0.5 | NMFS trawl survey: males, 1975-1981 |
|  | $\mathrm{pQ}[3]$ | 1 | M19F00 | at lower bound | 0.5 | NMFS trawl survey: females, 1975-1981 |
|  |  |  | M19F00a | at lower bound | 0.5 | NMFS trawl survey: females, 1975-1981 |
|  |  |  | M19F01 | at lower bound | 0.5 | NMFS trawl survey: females, 1975-1981 |
|  |  |  | M19F02 | at lower bound | 0.5 | NMFS trawl survey: females, 1975-1981 |
|  |  |  | M19F03 | at lower bound | 0.5 | NMFS trawl survey: females, 1975-1981 |
|  |  |  | M19F04 | at lower bound | 0.5 | NMFS trawl survey: females, 1975-1981 |
|  |  |  | M19F05 | at lower bound | 0.5 | NMFS trawl survey: females, 1975-1981 |

Table 19. Selectivity-related parameters from all model scenarios estimated within $1 \%$ of bounds.

|  | name |  | scenario | which? | bound | description |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| selectivity | pS1[1] | 1 | M19F00a | at upper bound | 90 | z50 for NMFS survey selectivity (males, pre-1982) |
|  |  |  | M19F01 | at upper bound | 90 | z50 for NMFS survey selectivity (males, pre-1982) |
|  |  |  | M19F02 | at upper bound | 90 | z50 for NMFS survey selectivity (males, pre-1982) |
|  |  |  | M19F03 | at upper bound | 90 | z50 for NMFS survey selectivity (males, pre-1982) |
|  |  |  | M19F04 | at upper bound | 90 | z50 for NMFS survey selectivity (males, pre-1982) |
|  |  |  | M19F05 | at upper bound | 90 | z50 for NMFS survey selectivity (males, pre-1982) |
|  | pS1[20] | 1 | M19F00 | at lower bound | 40 | z50 for GF.AllGear selectivity (males, 1987-1996) |
|  |  |  | M19F00a | at lower bound | 40 | z50 for GF.AllGear selectivity (males, 1987-1996) |
|  |  |  | M19F01 | at lower bound | 40 | z50 for GF.AllGear selectivity (males, 1987-1996) |
|  |  |  | M19F02 | at lower bound | 40 | z50 for GF.AllGear selectivity (males, 1987-1996) |
|  |  |  | M19F03 | at lower bound | 40 | z50 for GF.AllGear selectivity (males, 1987-1996) |
|  |  |  | M19F04 | at lower bound | 40 | z50 for GF.AllGear selectivity (males, 1987-1996) |
|  |  |  | M19F05 | at lower bound | 40 | z50 for GF.AllGear selectivity (males, 1987-1996) |
|  | $\mathrm{pS1}[23]$ | 1 | M19F00 | at upper bound | 180 | z95 for RKF selectivity (males, 1997-2004) |
|  |  |  | M19F00a | at upper bound | 180 | z95 for RKF selectivity (males, 1997-2004) |
|  |  |  | M19F01 | at upper bound | 180 | z95 for RKF selectivity (males, 1997-2004) |
|  |  |  | M19F02 | at upper bound | 180 | z95 for RKF selectivity (males, 1997-2004) |
|  |  |  | M19F03 | at upper bound | 180 | z95 for RKF selectivity (males, 1997-2004) |
|  |  |  | M19F04 | at upper bound | 180 | z95 for RKF selectivity (males, 1997-2004) |
|  |  |  | M19F05 | at upper bound | 180 | z95 for RKF selectivity (males, 1997-2004) |
|  | pS1[24] | 1 | M19F00 | at upper bound | 180 | z95 for RKF selectivity (males, 2005+) |
|  |  |  | M19F00a | at upper bound | 180 | z95 for RKF selectivity (males, 2005+) |
|  |  |  | M19F01 | at upper bound | 180 | z95 for RKF selectivity (males, 2005+) |
|  |  |  | M19F02 | at upper bound | 180 | z95 for RKF selectivity (males, 2005+) |
|  |  |  | M19F03 | at upper bound | 180 | z95 for RKF selectivity (males, 2005+) |
|  |  |  | M19F04 | at upper bound | 180 | z95 for RKF selectivity (males, 2005+) |
|  |  |  | M19F05 | at upper bound | 180 | z95 for RKF selectivity (males, 2005+) |
|  | pS1[27] | 1 | M19F00 | at upper bound | 140 | z95 for RKF selectivity (females, 2005+) |
|  |  |  | M19F00a | at upper bound | 140 | z95 for RKF selectivity (females, 2005+) |
|  |  |  | M19F01 | at upper bound | 140 | z95 for RKF selectivity (females, 2005+) |
|  |  |  | M19F02 | at upper bound | 140 | z95 for RKF selectivity (females, 2005+) |
|  |  |  | M19F03 | at upper bound | 140 | z95 for RKF selectivity (females, 2005+) |
|  |  |  | M19F04 | at upper bound | 140 | z95 for RKF selectivity (females, 2005+) |
|  |  |  | M19F05 | at upper bound | 140 | z95 for RKF selectivity (females, 2005+) |
|  | pS2[10] | 1 | M19F00a | at lower bound | 0.1 | ascending slope for SCF selectivity (males, pre-1997) |
|  |  |  | M19F01 | at lower bound | 0.1 | ascending slope for SCF selectivity (males, pre-1997) |

Table 19 (cont.).Selectivity-related parameters from all model scenarios estimated within $1 \%$ of bounds.

| name |  | 3ddobeiod | sthiowver bound | bdund | dssenipitigrslope for SCF selectivity (males, pre-1997) |
| :---: | :---: | :---: | :---: | :---: | :---: |
| $\mathrm{pS} 2[2]$ | 1 | M19F03 | at lower bound | 0.1 | ascending slope for SCF selectivity (males, pre-1997) |
|  |  | M19F04 | at lower bound | 0.1 | ascending slope for SCF selectivity (males, pre-1997) |
|  |  | M19F05 | at lower bound | 0.1 | ascending slope for SCF selectivity (males, pre-1997) |
|  |  | M19F01 | at upper bound | 100 | z95-z50 for NMFS survey selectivity (males, 1982+) |
|  |  | M19F02 | at upper bound | 100 | z95-z50 for NMFS survey selectivity (males, 1982+) |
|  |  | M19F03 | at upper bound | 100 | z95-z50 for NMFS survey selectivity (males, 1982+) |
|  |  | M19F04 | at upper bound | 100 | z95-z50 for NMFS survey selectivity (males, 1982+) |
|  |  | M19F05 | at upper bound | 100 | z95-z50 for NMFS survey selectivity (males, 1982+) |
| pS2[4] | 1 | M19F00 | at upper bound | 100 | z95-z50 for NMFS survey selectivity (females, 1982+) |
|  |  | M19F00a | at upper bound | 100 | z95-z50 for NMFS survey selectivity (females, $1982+$ ) |
|  |  | M19F01 | at upper bound | 100 | z95-z50 for NMFS survey selectivity (females, 1982+) |
|  |  | M19F02 | at upper bound | 100 | z95-z50 for NMFS survey selectivity (females, 1982+) |
|  |  | M19F03 | at upper bound | 100 | z95-z50 for NMFS survey selectivity (females, 1982+) |
|  |  | M19F04 | at upper bound | 100 | z95-z50 for NMFS survey selectivity (females, 1982+) |
|  |  | M19F05 | at upper bound | 100 | z95-z50 for NMFS survey selectivity (females, 1982+) |
| $\mathrm{pS} 4[1]$ | 1 | M19F00 | at upper bound | 0.5 | descending slope for SCF selectivity (males, pre-1997) |
|  |  | M19F00a | at lower bound | 0.1 | descending slope for SCF selectivity (males, pre-1997) |
|  |  | M19F01 | at lower bound | 0.1 | descending slope for SCF selectivity (males, pre-1997) |
|  |  | M19F02 | at lower bound | 0.1 | descending slope for SCF selectivity (males, pre-1997) |
|  |  | M19F03 | at lower bound | 0.1 | descending slope for SCF selectivity (males, pre-1997) |
|  |  | M19F04 | at lower bound | 0.1 | descending slope for SCF selectivity (males, pre-1997) |
|  |  | M19F05 | at lower bound | 0.1 | descending slope for SCF selectivity (males, pre-1997) |

Table 20. Estimated growth, natural mortality, and non-vector recruitment parameters for all model scenarios.

| process | description | parameter | phase |  | scale | M19F00 |  | M19FOOs |  | M19F01 |  | M19FO2 |  | M19F03 |  | M19FO4 |  | M19F05 |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  | value | std dev | value | std dev | value | std dev | value | std dev | value | std dev | value | std dev | value | std dev |
| growth | both sexes | pGrBeta[ $]$ | 1 | , |  | ARITHMETIC | $8.116 e-01$ | 0.12597 | $5.441 \mathrm{e}-01$ | 0.07594 | $5.822 e-01$ | 0.06559 | 7.936 e - 01 | 0.09799 | $9.035 \mathrm{e}-01$ | 0.11392 | 5.651 - 01 | 0.06168 | 9.061 - 01 | 0.11105 |
|  | females | pGral 2 ] | 1 | 4 | arithmetic | $3.446 e+01$ | 0.42304 | $3.359 e+01$ | ${ }^{0.35337}$ | $3.326 e+01$ | 0.25176 | $3.367 e+01$ | 0.29993 | $3.399 x+01$ | 0.33574 | $3.320 e+01$ | 0.24548 | $3.396 e+01$ | 0.33400 |
|  |  | pGrB[ 2 ] | 1 | 4 | arithmetic | $1.151 \mathrm{c}+02$ | 0.84195 | $1.151 e+02$ | 0.77050 | $1.152 e+02$ | 0.59572 | $1.151 e+02$ | 0.655101 | $1.143 \mathrm{c}+02$ | 0.64756 | $1.152 e+02$ | 0.55714 | $1.149 \mathrm{e}+02$ | 0.61193 |
|  | males | pGrail | 1 | 4 | arithmetic | $3.309 c+01$ | 0,35045 | $3.268 \mathrm{c}+01$ | 0.26962 | $3.258 \mathrm{c}+01$ | 0.23545 | $3.235 \mathrm{c}+01$ | 0.26033 | $3.274 c+01$ | 0.29222 | $3.245 \mathrm{e}+01$ | 0.23171 | $3.261 e+01$ | 0.28179 |
|  |  | $\mathrm{pGrB}[1]$ | 1 | 4 | arithmetic | $1.670 c+02$ | 1.07890 | $1.634 e+02$ | 1.10110 | $1.612 e+02$ | 1.02490 | $1.657 \mathrm{c}+02$ | 0.86298 | $1.666 e+02$ | 0.92085 | $1.612 e+02$ | 0.98847 | $1.670 \mathrm{c}+02$ | 0.94701 |
| natural mortality | 1980-1984 multiplier for mature females | pDM2[2] | 1 | 4 | arithmetic | $1.307 e+\infty$ | 0.09981 | $1.440 e+00$ | 0.0786 | $1.380 \mathrm{e}+00$ | 0.10693 | $1.353 \mathrm{e}+00$ | 0.10385 | $1.873 \mathrm{c}+00$ | 0.15471 | $1.356 \mathrm{e}+00$ | 0.1045 | $1.893 c+00$ | 0.15211 |
|  | 1980-1984 multiplier for mature males | pDM2[1] |  | 4 | arithmetic | $2.587{ }^{\text {e }}+00$ | 0.24183 | $2.8833^{+}+00$ | 0.25788 | $2.798 \mathrm{c}+00$ | 0.25750 | $2.620 \mathrm{c}+00$ | 0.22434 | $2.231 \mathrm{c}+00$ | 0.21496 | $2.720 e+00$ | 0.24396 | $2.286{ }^{2}+00$ | 0.21422 |
|  | base $\ln$-scale M | $\mathrm{pM}(1)$ | 1 | -1 | LOG | $-1.470 \mathrm{e}+00$ | NA | $-1.470 c+00$ | NA | $-1.470 e+00$ | NA | $-1.470 e^{+} 00$ | $N A$ | $-1.470 \mathrm{c}+00$ | NA | $-1.470 e^{+}+00$ | NA | $-1.470 \mathrm{c}+00$ | NA |
|  | multiplier for immature crab | pDM11] | 1 | 4 | ARITHMETIC | $1.002 c+00$ | 0.0504 | $9.997 e-01$ | 0.05066 | 9.901 c - 01 | 0.05042 | $9.756 \mathrm{c}-01$ | 0.05022 | 9.833 c - 01 | 0.05149 | $1.064 c+00$ | 0.04820 | $1.048 \mathrm{c}+00$ | 0.04979 |
|  | multiplier for mature femakes | pDM13] |  | 4 | arithmetic | $1.3866+00$ | 0.03557 | $1.341 e+00$ | 0.03710 | $1.328 \mathrm{c}+00$ | 0.03635 | $1.348 c+00$ | 0.03688 | $1.316 \mathrm{c}+00$ | 0.03885 | $1.327 e+00$ | 0.03567 | $1.3276+00$ | 0.03811 |
|  | multiplice for mature males | pDMi[ 2 ] | 1 | 4 | arithmetic | $1.152 e+00$ | 0.03952 | $1.221 \mathrm{c}+00$ | 0.00087 | $1.230 \mathrm{e}+00$ | 0.03938 | $1.352 c+00$ | 0.03742 | $1.292 \mathrm{e}+00$ | 0.04001 | $1.262 e+00$ | 0.03864 | $1.3180+00$ | 0.03873 |
| recruitment | current recruitment period | pLaR[2] | 1 | 1 | ARITHMETIC | $5.135 e+00$ | 0.07180 | $5.414 \mathrm{e}+00$ | 0.07889 | 5.484 + +00 | 0.08083 | $5.615 \mathrm{e}+00$ | 0.08148 | $5.691 \mathrm{le}+00$ | 0.08257 | $5.630 \mathrm{e}+00$ | 0.06936 | $5.740 \mathrm{e}+00$ | 0.06904 |
|  | fixed value | pRa[ [1] | 1 | -1 | LOG | $2.442 e+00$ | NA | $2.442 e+00$ | NA | $2.442 c+00$ | NA | $2.442 e+00$ | NA | $2.442 \epsilon+00$ | NA | $2.442 e+00$ | NA | $2.442 c+00$ | NA |
|  |  | pRb ${ }^{\text {[ }}$ ] | 1 | -1 | LOG | $1.3866+00$ | NA | $1.386 \mathrm{c}+00$ | NA | $1.386 e+00$ | NA | $1.3866+00$ | NA | $1.386 \mathrm{c}+00$ | NA | $1.386 \mathrm{e}+00$ | NA | $1.3866+00$ | NA |
|  | full model period | pRCV[1] | 1 | ${ }^{-1}$ | LOG | $-6.931 e-01$ | NA | $-6.931-01$ | NA | -6.931e-01 | NA | $-6.931 e-01$ | NA | $-6.931 e-01$ | NA | $-6.931 \mathrm{c}-01$ | NA | $-6.931 e-01$ | NA |
|  |  | pRX[1] | 1 | $\stackrel{-1}{1}$ | LOGIT | $-1.110 e-16$ | NA | $-1.110 e-16$ | NA | $-1.110 e-16$ | NA | $-1.110 e-16$ | NA | $-1.110 e-16$ | NA | $-1.110 c-16$ | NA | $-1.110 e-16$ | NA |
|  | historical recruitment period | pLar [1] | 1 | 1 | ARITHMETIC | $5.662 e+00$ | 0.40017 | $6.039 e+00$ | 0.38683 | $6.185 \mathrm{c}+00$ | 0.37589 | $6.251 e+00$ | 0.39915 | $6.281 e+00$ | 0.41654 | $6.315 \mathrm{e}+00$ | 0.37471 | $6.341 e+00$ | 0.41554 |

Table 21. Historical recruitment devs estimates (1948-1974) for all model scenarios.

| index |  |  | M19F00 |  | M19F00a |  | M19F01 |  | M19F02 |  | M19F03 |  | M19F04 |  | M19F05 |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | value | std dev | value | std dev | value | std dev | value | std dev | value | std dev | value | std dev | value | std dev |
| 1 | 1 | ARITHMETIC | -1.4120 | 1.4764 | -1.2802 | 1.4628 | -1.21227 | 1,4492 | -1.21416 | 1.4817 | $-1.39338$ | 1.5043 | -1.19404 | 1.4508 | -1.381738 | 1.5055 |
| 2 | 1 | ARITHMETIC | $-1.4120$ | 1.3270 | $-1.2826$ | 1.3122 | $-1.21620$ | 1.2978 | -1.21685 | 1.3320 | $-1.38912$ | 1.3562 | -1.19841 | 1.2993 | -1.378240 | 1.3574 |
| 3 | 1 | ARITHMETIC | $-1.4105$ | 1.1914 | -1.2858 | 1.1758 | -1.22223 | 1.1611 | -1.22085 | 1.1953 | -1.37960 | 1.2206 | $-1.20533$ | 1.1624 | $-1.370257$ | 1.2216 |
| 4 | 1 | ARITHMETIC | $-1.4057$ | 1.0721 | $-1.2877$ | 1.0561 | $-1.22784$ | 1.0417 | $-1.22425$ | 1.0745 | -1.36341 | 1.0997 | -1.21230 | 1.0428 | -1.356370 | 1.1006 |
| 5 | 1 | ARITHMETIC | $-1.3949$ | 0.9712 | $-1.2851$ | 0.9552 | $-1.22960$ | 0.9417 | $-1.22430$ | 0.9714 | $-1.33850$ | 0.9956 | $-1.21587$ | 0.9423 | -1.334528 | 0.9963 |
| 6 | 1 | ARITHMETIC | $-1.3746$ | 0.8894 | $-1.2738$ | 0.8738 | $-1.22273$ | 0.8615 | $-1.21710$ | 0.8872 | -1.30189 | 0.9094 | $-1.21124$ | 0.8616 | $-1.301728$ | 0.9098 |
| 7 | 1 | ARITHMETIC | $-1.3393$ | 0.8260 | $-1.2476$ | 0.8107 | -1.20041 | 0.7997 | $-1.19696$ | 0.8213 | $-1.24913$ | 0.8408 | $-1.19151$ | 0.7994 | -1.253493 | 0.8409 |
| 8 | 1 | ARITHMETIC | $-1.2817$ | 0.7781 | $-1.1977$ | 0.7631 | -1.15263 | 0.7532 | -1.15541 | 0.7712 | $-1.17347$ | 0.7877 | -1.14656 | 0.7525 | -1.182986 | 0.7877 |
| 9 | 1 | ARITHMETIC | $-1.1899$ | 0.7418 | $-1.1101$ | 0.7266 | $-1.06393$ | 0.7173 | $-1,07919$ | 0.7330 | $-1.06415$ | 0.7470 | $-1.06065$ | 0.7164 | -1.079357 | 0.7469 |
| 10 | 1 | ARITHMETIC | $-1.0453$ | 0.7126 | -0.9619 | 0.6970 | -0.90858 | 0.6882 | $-0.94640$ | 0.7028 | $-0.90325$ | 0.7152 | $-0.90753$ | 0.6871 | -0.924438 | 0.7150 |
| 11 | 1 | ARITHMETIC | $-0.8151$ | 0.6883 | $-0.7120$ | 0.6736 | -0.64015 | 0.6662 | $-0.71771$ | 0.6798 | -0.65848 | 0.6912 | -0.63979 | 0.6654 | -0.685503 | 0.6910 |
| 12 | 1 | ARITHMETIC | -0.4383 | 0.6726 | $-0.2871$ | 0.6623 | $-0.18242$ | 0.6572 | $-0.32434$ | 0.6683 | $-0.27087$ | 0.6796 | $-0.18059$ | 0.6566 | $-0.302887$ | 0.6792 |
| 13 | 1 | ARITHMETIC | 0.1660 | 0.6670 | 0.3730 | 0.6568 | 0.50326 | 0.6510 | 0.29986 | 0.6641 | 0.32880 | 0.6782 | 0.50615 | 0.6503 | 0.293606 | 0.6774 |
| 14 | 1 | ARITHMETIC | 0.9679 | 0.6525 | 1.1497 | 0.6391 | 1.24659 | 0.6320 | 1.07202 | 0.6488 | 1.08913 | 0.6671 | 1.24599 | 0.6312 | 1.055489 | 0.6660 |
| 15 | 1 | ARITHMETIC | 1.6257 | 0.6254 | 1.6449 | 0.6127 | 1.65040 | 0.6070 | 1.62078 | 0.6236 | 1.67036 | 0.6444 | 1.64471 | 0.6067 | 1.646939 | 0.6433 |
| 16 | 1 | ARITHMETIC | 1.7976 | 0.6055 | 1.6724 | 0.6031 | 1.60834 | 0.6017 | 1.69296 | 0.6102 | 1.79168 | 0.6301 | 1.60010 | 0.6018 | 1.778654 | 0.6288 |
| 17 | 1 | ARITHMETIC | 1.6191 | 0.6125 | 1.4503 | 0.6111 | 1.36579 | 0.6077 | 1.47538 | 0.6164 | 1.61024 | 0.6355 | 1.35947 | 0.6078 | 1.604503 | 0.6347 |
| 18 | 1 | ARITHMETIC | 1.3692 | 0.6187 | 1.2196 | 0.6105 | 1.13831 | 0.6030 | 1.22210 | 0.6170 | 1.37666 | 0.6359 | 1.13637 | 0.6025 | 1.379250 | 0.6358 |
| 19 | 1 | ARITHMETIC | 1.2090 | 0.6091 | 1.0895 | 0.5948 | 1.01064 | 0.5864 | 1.05644 | 0.6040 | 1.22342 | 0.6216 | 1.01261 | 0.5853 | 1.238470 | 0.6212 |
| 20 | 1 | ARITHMETIC | 1.1875 | 0.5888 | 1.0640 | 0.5771 | 0.96832 | 0.5732 | 0.99256 | 0.5871 | 1.17131 | 0.6028 | 0.96827 | 0.5718 | 1.201936 | 0.6004 |
| 21 | 1 | ARITHMETIC | 1.2546 | 0.5708 | 1.0303 | 0.5654 | 0.89237 | 0.5599 | 0.95314 | 0.5733 | 1.13925 | 0.5929 | 0.87968 | 0.5582 | 1.179838 | 0.5890 |
| 22 | 1 | ARITHMETIC | 1.2263 | 0.5399 | 0.8766 | 0.5246 | 0.75151 | 0.5121 | 0.86630 | 0.5379 | 0.99302 | 0.5659 | 0.73357 | 0.5109 | 1.027757 | 0.5635 |
| 23 | 1 | ARITHMETIC | 1.0762 | 0.4826 | 0.8502 | 0.4642 | 0.72676 | 0.4560 | 0.80345 | 0.4809 | 0.77987 | 0.5114 | 0.70422 | 0.4560 | 0.797451 | 0.5113 |
| 24 | 1 | ARITHMETIC | 0.6695 | 0.4841 | 0.4802 | 0.4726 | 0.32334 | 0.4703 | 0.43006 | 0.4935 | 0.37588 | 0.5176 | 0.28941 | 0.4715 | 0.382385 | 0.5187 |
| 25 | 1 | ARITHMETIC | 0.2508 | 0.4985 | 0.1363 | 0.4876 | 0.06273 | 0.4829 | 0.09607 | 0.5049 | -0.01844 | 0.5223 | 0.04545 | 0.4837 | -0.006806 | 0.5232 |
| 26 | 1 | ARITHMETIC | 0.1002 | 0.4463 | 0.1744 | 0.4312 | 0.23064 | 0.4218 | 0.15639 | 0.4445 | $-0.04595$ | 0.4561 | 0.23782 | 0.4209 | $-0.027946$ | 0.4570 |

Table 22. Current recruitment devs estimates (1975-2019) for all model scenarios.

| index |  |  | M19F00 |  | M19F00a |  | M19F01 |  | M19F02 |  | M19F03 |  | M19F04 |  | M19F05 |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | value | std dev | value | std dev | value | std dev | value | std dev | value | std dey | value | std dev | alue | std dev |
| 1 | 1 | ARITHMETIC | 1.325987 | 0.21947 | 1.26031 | 0.25829 | 1.1945 | 0.30320 | 1.15328 | 0.30225 | 0.86809 | 0.32553 | 1.11503 | 0.31661 | 0.873803 | 0,33236 |
| 2 | 1 | ARITHMETIC | 1.999511 | 0.14193 | 2.10527 | 0.14112 | 2.1559 | 0.14562 | 2.11267 | 0.15067 | 1.92607 | 0.15117 | 2.11699 | 0.14489 | 1.948271 | 0.15060 |
| 3 | 1 | ARITHMETIC | 1.741158 | 0.15634 | 1.79450 | 0.15903 | 1.7409 | 0.17475 | 1.74449 | 0.17805 | 1.64245 | 0.17117 | 1.67740 | 0.17631 | 1.647896 | 0.17221 |
| 4 | 1 | ARITHMETIC | 0.914330 | 0.25716 | 0.91906 | 0.26315 | 0.8178 | 0.29703 | 0.87446 | 0.30204 | 0.94441 | 0.26110 | 0.73860 | 0.30411 | 0.938347 | 0.26352 |
| 5 | 1 | ARITHMETIC | 0.060257 | 0.39606 | 0.15632 | 0.37163 | 0.1218 | 0.40566 | 0.07721 | 0.44154 | -0.06675 | 0.43056 | 0.06231 | 0.41081 | -0.073395 | 0.43698 |
| 6 | 1 | ARITHMETIC | -0.436228 | 0.47983 | $-0.29754$ | 0.43368 | -0.2936 | 0.46888 | $-0.41781$ | 0.53348 | $-0.57721$ | 0.51730 | $-0.33556$ | 0.47194 | -0.565462 | 0.51549 |
| 7 | 1 | ARITHMETIC | 0.057849 | 0.25068 | 0.08108 | 0.25329 | 0.1607 | 0.25902 | 0.07293 | 0.27445 | -0.10833 | 0.26401 | 0.11128 | 0.26485 | -0.121861 | 0.26766 |
| 8 | I | ARITHMETIC | -0.518498 | 0.33685 | -0.40743 | 0.32326 | $-0.4401$ | 0.36189 | $-0.54060$ | 0.37438 | -0.25303 | 0.24297 | -0.45249 | 0.35993 | -0.258598 | 0.24387 |
| 9 | 1 | ARITHMETIC | 1.068860 | 0.10832 | 1.06916 | 0.10705 | 1.1744 | 0.10699 | 1.09664 | 0.10918 | 0.84628 | 0.11017 | 1.15444 | 0.10700 | 0.837807 | 0.11057 |
| 10 | 1 | ARITHMETIC | 0.874922 | 0.14305 | 0.91568 | 0.13217 | 0.9568 | 0.13703 | 0.86325 | 0.14717 | 0.75097 | 0.14269 | 0.92987 | 0.13732 | 0.744673 | 0.14369 |
| 11 | 1 | ARITHMETIC | 1.172213 | 0.13044 | 0.85682 | 0.14854 | 0.7990 | 0.16062 | 0.93231 | 0.15232 | 0.95242 | 0.13655 | 0.76898 | 0.16024 | 0.962625 | 0.13501 |
| 12 | 1 | ARITHMETIC | 1.134704 | 0.13537 | 0.95193 | 0.13239 | 1.0451 | 0.12651 | 0.98791 | 0.14234 | 0.94857 | 0.14102 | 1.03449 | 0.12446 | 0.961474 | 0.14092 |
| 13 | 1 | ARITHMETIC | 1.127278 | 0.13076 | 0.94278 | 0.12646 | 0.8736 | 0.14097 | 0.97327 | 0.13736 | 0.98944 | 0.13308 | 0.83020 | 0.14230 | 0.993998 | 0.13253 |
| 14 | 1 | ARITHMETIC | 0.737460 | 0.15415 | 0.41544 | 0.15996 | 0.3241 | 0.17331 | 0.47560 | 0.17379 | 0.69924 | 0.15395 | 0.29543 | 0.17444 | 0.694339 | 0.15454 |
| 15 | 1 | ARITHMETIC | 0.004827 | 0.18563 | -0.13141 | 0.18008 | -0.1839 | 0.19135 | -0.14532 | 0.20310 | $-0.17150$ | 0.21099 | $-0.20957$ | 0.19309 | -0.187352 | 0.21377 |
| 16 | 1 | ARITHMETIC | -1.180975 | 0.38180 | -1.40841 | 0.40897 | $-1.5941$ | 0.48663 | -1.52964 | 0.50361 | -1.32340 | 0.40960 | -1.63103 | 0.49750 | $-1.339677$ | 0.41479 |
| 17 | 1 | ARITHMETIC | -1.404156 | 0.33220 | $-1.51656$ | 0.33544 | -1.5957 | 0.35933 | $-1.53360$ | 0.36485 | -1.42398 | 0.32064 | -1.62032 | 0.36386 | -1.435798 | 0.32304 |
| 18 | 1 | ARITHMETIC | -1.526055 | 0.29294 | $-1.53753$ | 0.28809 | -1.5417 | 0.29511 | $-1.53361$ | 0.30441 | -1.39119 | 0.25789 | $-1.55755$ | 0.29720 | -1.397713 | 0.25815 |
| 19 | 1 | ARITHMETIC | $-1.529904$ | 0.27300 | -1.48781 | 0.27121 | $-1.4885$ | 0.28543 | -1.53198 | 0.29714 | -1.48169 | 0.27427 | -1.50814 | 0.28884 | $-1.500872$ | 0.27647 |
| 20 | 1 | ARITHMETIC | -1.259841 | 0.22362 | -1.19852 | 0.22551 | $-1.1435$ | 0.23061 | -1.19402 | 0.23688 | -1.25593 | 0.24584 | $-1.15282$ | 0.23121 | $-1.269587$ | 0.24563 |
| 21 | 1 | ARITHMETIC | $-1.004762$ | 0.19234 | $-0.91708$ | 0.19233 | $-0.8269$ | 0.19186 | -0.87836 | 0.19708 | $-0.72265$ | 0.17359 | -0.84062 | 0.19226 | $-0.740183$ | 0.17330 |
| 22 | 1 | ARITHMETIC | -1.078268 | 0.22288 | $-1.01880$ | 0.22794 | $-1.0485$ | 0.24793 | $-1.05749$ | 0.24727 | $-1.01225$ | 0.23320 | $-1.07560$ | 0.25020 | $-1.036970$ | 0.23392 |
| 23 | 1 | ARITHMETIC | $-0.008056$ | 0.10696 | 0.05202 | 0.10829 | 0.1272 | 0.10884 | 0.11232 | 0.10896 | 0.02733 | 0.11155 | 0.11215 | 0.10864 | 0.009917 | 0.11119 |
| 24 | 1 | ARITHMETIC | $-0.920764$ | 0.20125 | -0.85936 | 0.20485 | $-0.8502$ | 0.21716 | $-0.90117$ | 0.22133 | $-0.84470$ | 0.20866 | $-0.86471$ | 0.21773 | -0.860841 | 0.20859 |
| 25 | 1 | ARITHMETIC | 0.289723 | 0,10549 | 0.38974 | 0.10445 | 0.4616 | 0.10480 | 0.41001 | 0.10665 | 0.41959 | 0.10446 | 0.44562 | 0.10465 | 0.396694 | 0.10416 |
| 26 | 1 | ARITHMETIC | $-0.366276$ | 0.20085 | $-0.29864$ | 0.20562 | $-0.2978$ | 0.21995 | $-0.34294$ | 0.22357 | -0.29201 | 0.21269 | $-0.30902$ | 0.22019 | $-0.306770$ | 0.21209 |
| 27 | 1 | ARITHMETIC | 0.815665 | 0.09646 | 0.86132 | 0.09686 | 0.9013 | 0.09881 | 0.90922 | 0.09894 | 0.93503 | 0.09762 | 0.89129 | 0.09891 | 0.914377 | 0.09759 |
| 28 | 1 | ARITHMETIC | -0.324659 | 0.23741 | $-0.26477$ | 0.23954 | $-0.2525$ | 0.25385 | -0.30239 | 0.26489 | -0.24681 | 0.25659 | $-0.25004$ | 0.25371 | -0.252331 | 0.25580 |
| 29 | 1 | ARITHMETIC | 0.782948 | 0.11247 | 0.85807 | 0.10883 | 0.8867 | 0.11053 | 0.90219 | 0.11281 | 1.03049 | 0.10797 | 0.88050 | 0.11109 | 1.011588 | 0.10834 |
| 30 | I | ARITHMETIC | 0.754406 | 0.10830 | 0.72155 | 0.11167 | 0.6930 | 0.11698 | 0.76029 | 0.11716 | 0.84219 | 0,11706 | 0.69958 | 0.11696 | 0.839696 | 0.11671 |
| 31 | 1 | ARITHMETIC | -0.561822 | 0.24624 | $-0.52333$ | 0.24261 | $-0.6105$ | 0.26630 | $-0.62183$ | 0.28186 | -0.46522 | 0.25987 | -0.60852 | 0.26952 | -0.477087 | 0.26272 |
| 32 | 1 | ARITHMETIC | $-0.827576$ | 0.27219 | -0.88981 | 0.29428 | $-0.9527$ | 0.31647 | -0.90211 | 0.31438 | $\sim 0.84391$ | 0.30278 | $-0.94964$ | 0.32136 | $-0.845306$ | 0.30424 |
| 33 | 1 | ARITHMETIC | $-1.079504$ | 0.31525 | $-0.98223$ | 0.31350 | $-0.9470$ | 0.32000 | -0.99731 | 0.33015 | -0.97928 | 0.31748 | -0.93565 | 0.32438 | -0.983900 | 0.31895 |
| 34 | 1 | ARITHMETIC | -0.651014 | 0.26285 | -0.45404 | 0.25695 | $-0.3760$ | 0.25660 | $-0.50319$ | 0.26709 | -0.50286 | 0.26369 | $-0.30953$ | 0.25406 | $-0.506445$ | 0.26382 |
| 35 | 1 | ARITHMETIC | 1.219595 | 0.10001 | 1.32960 | 0.09379 | 1.3675 | 0.09161 | 1.36945 | 0.09620 | 1.34668 | 0.10032 | 1.41816 | 0.08972 | 1.378909 | 0.09736 |
| 36 | 1 | ARITHMETIC | 1.079563 | 0.10839 | 1.00232 | 0.11536 | 0.8413 | 0.12839 | 0.96347 | 0.12734 | 1.07831 | 0.12005 | 0.81370 | 0.13253 | 1.059373 | 0.12192 |
| 37 | 1 | ARITHMETIC | 0.165279 | 0.17803 | 0.04304 | 0.18429 | -0.1110 | 0.19518 | -0.01554 | 0.19961 | 0.01705 | 0.19485 | -0.01670 | 0.18271 | 0.052942 | 0.18644 |
| 38 | 1 | ARITHMETIC | -1.429645 | 0.46100 | $-1.35238$ | 0.43217 | $-1.5650$ | 0.48335 | $-1.58832$ | 0.52578 | -1.55216 | 0.46028 | -1.62522 | 0.50258 | -1.629746 | 0.47288 |
| 39 | 1 | ARITHMETIC | -0.447087 | 0.18059 | -0.46539 | 0.18595 | $-0.5666$ | 0.18716 | -0.48835 | 0.18585 | $-0.53477$ | 0.17530 | $-0.46516$ | 0.15891 | -0.459241 | 0.15041 |
| 40 | 1 | ARITHMETIC | -0.834498 | 0.21397 | -0.83214 | 0.21754 | $-1.0070$ | 0.23089 | $-0.94203$ | 0.23099 | -1.01823 | 0.22108 | $-1.03765$ | 0.21588 | -1.077281 | 0.20752 |
| 41 | 1 | ARITHMETIC | -1.244014 | 0.27292 | -1.22138 | 0,27541 | $-1.3175$ | 0.28034 | -1.31779 | 0.28752 | -1.30915 | 0.25684 | -1.31331 | 0.24850 | $-1.357546$ | 0.23090 |
| 42 | 1 | ARITHMETIC | $-0.893146$ | 0.24196 | $-0.86516$ | 0.24412 | -0.8876 | 0.23938 | -0.86864 | 0.23975 | $-0.92572$ | 0.23069 | -0.73199 | 0.18381 | -0.840550 | 0.17699 |
| 43 | 1 | ARITHMETIC | 0.959708 | 0.14287 | 0.99193 | 0.14244 | 0.8948 | 0.13040 | 0.91736 | 0.13034 | 0.78189 | 0.12072 | 1.08909 | 0.11217 | 0.928495 | 0.10095 |
| 44 | 1 | ARITHMETIC | 1.240503 | 0.22169 | 1.21181 | 0.22408 | 0.8241 | 0.19598 | 0.86635 | 0.19511 | 0.82803 | 0.17921 | 0.91640 | 0.19794 | 0.823768 | 0.18082 |

Table 23. Logit-scale parameters for the probability of terminal molt for males for all model scenarios. The (arithmetic) probability of terminal molt was fixed at 0 for males less than 60 mm CW and at 1 for males greater than 145 mm CW in Scenarios M19F03 and M19F05.

| scenario: | M19F00 |  | M19F00a |  | M19F01 |  | M19F02 |  | M19F03 |  | M19F04 |  | M19F05 |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| size bin | value | std err | value | std err | value | std err | value | std err | value | std err | value | std err | value | std err |
| 27.5 | -12.03 | 7.44 | -12.06 | 7.42 | -12.04 | 7.39 | -12.48 | 7.59 | -- | -- | -11.92 | 7.22 | -- | -- |
| 32.5 | -10.85 | 5.62 | -10.84 | 5.60 | -10.82 | 5.57 | -11.15 | 5.73 | -- | -- | -10.70 | 5.43 | -- | -- |
| 37.5 | -9.66 | 4.02 | -9.62 | 4.00 | -9.61 | 3.99 | -9.82 | 4.11 | -- | -- | -9.48 | 3.86 | -- | -- |
| 42.5 | -8.48 | 2.68 | -8.40 | 2.67 | -8.39 | 2.66 | -8.49 | 2.74 | -- | -- | -8.27 | 2.56 | -- | -- |
| 47.5 | -7.31 | 1.63 | -7.19 | 1.61 | -7.19 | 1.61 | -7.17 | 1.64 | -- | -- | -7.06 | 1.53 | -- | -- |
| 52.5 | -6.16 | 0.91 | -6.01 | 0.88 | -6.01 | 0.88 | -5.87 | 0.87 | -- | -- | -5.89 | 0.84 | -- | -- |
| 57.5 | -5.11 | 0.54 | -4.92 | 0.52 | -4.92 | 0.51 | -4.65 | 0.46 | -- | -- | -4.81 | 0.49 | -- | -- |
| 62.5 | -4.49 | 0.36 | -4.27 | 0.36 | -4.24 | 0.35 | -3.82 | 0.28 | -2.91 | 0.28 | -4.20 | 0.35 | -2.95 | 0.28 |
| 67.5 | -4.10 | 0.29 | -3.95 | 0.29 | -3.92 | 0.29 | -3.48 | 0.22 | -3.29 | 0.29 | -3.98 | 0.29 | -3.37 | 0.30 |
| 72.5 | -3.46 | 0.22 | -3.34 | 0.23 | -3.39 | 0.22 | -3.04 | 0.19 | -2.86 | 0.25 | -3.40 | 0.22 | -2.86 | 0.25 |
| 77.5 | -2.93 | 0.17 | -2.71 | 0.17 | -2.76 | 0.17 | -2.50 | 0.14 | -2.17 | 0.16 | -2.68 | 0.17 | -2.14 | 0.15 |
| 82.5 | -2.50 | 0.14 | -2.26 | 0.14 | -2.24 | 0.14 | -1.92 | 0.12 | -1.66 | 0.14 | -2.19 | 0.13 | -1.67 | 0.13 |
| 87.5 | -2.03 | 0.12 | -1.90 | 0.12 | -1.90 | 0.12 | -1.53 | 0.10 | -1.41 | 0.12 | -1.91 | 0.11 | -1.43 | 0.12 |
| 92.5 | -1.44 | 0.11 | -1.42 | 0.11 | -1.48 | 0.10 | -1.00 | 0.09 | -0.86 | 0.11 | -1.50 | 0.10 | -0.86 | 0.10 |
| 97.5 | -0.95 | 0.09 | -0.95 | 0.10 | -1.07 | 0.10 | -0.57 | 0.08 | -0.47 | 0.10 | -1.07 | 0.10 | -0.47 | 0.09 |
| 102.5 | -0.68 | 0.09 | -0.59 | 0.09 | -0.67 | 0.09 | -0.39 | 0.08 | -0.32 | 0.10 | -0.65 | 0.09 | -0.34 | 0.09 |
| 107.5 | -0.53 | 0.09 | -0.47 | 0.09 | -0.45 | 0.08 | -0.23 | 0.08 | -0.15 | 0.10 | -0.41 | 0.08 | -0.14 | 0.09 |
| 112.5 | -0.06 | 0.10 | -0.12 | 0.09 | -0.15 | 0.09 | 0.20 | 0.09 | 0.30 | 0.11 | -0.13 | 0.08 | 0.30 | 0.10 |
| 117.5 | 0.56 | 0.13 | 0.41 | 0.11 | 0.34 | 0.10 | 0.77 | 0.11 | 0.90 | 0.13 | 0.37 | 0.10 | 0.95 | 0.14 |
| 122.5 | 1.44 | 0.20 | 1.09 | 0.14 | 0.98 | 0.13 | 1.55 | 0.16 | 1.76 | 0.19 | 0.99 | 0.13 | 1.80 | 0.19 |
| 127.5 | 2.81 | 0.36 | 1.88 | 0.29 | 1.55 | 0.20 | 2.81 | 0.30 | 3.11 | 0.31 | 1.54 | 0.20 | 3.16 | 0.30 |
| 132.5 | 5.06 | 0.59 | 3.95 | 0.61 | 3.17 | 0.55 | 4.17 | 0.34 | 4.35 | 0.34 | 3.16 | 0.54 | 4.40 | 0.35 |
| 137.5 | 7.20 | 1.06 | 6.22 | 0.91 | 5.44 | 0.79 | 6.01 | 0.65 | 6.12 | 0.73 | 5.45 | 0.77 | 6.15 | 0.74 |
| 142.5 | 9.01 | 1.68 | 8.21 | 1.42 | 7.54 | 1.21 | 7.80 | 1.17 | 8.03 | 1.54 | 7.55 | 1.20 | 8.05 | 1.55 |
| 147.5 | 10.50 | 2.32 | 9.85 | 2.03 | 9.29 | 1.79 | 9.35 | 1.77 | -- | -- | 9.31 | 1.78 | -- | -- |
| 152.5 | 11.69 | 2.85 | 11.18 | 2.57 | 10.73 | 2.35 | 10.64 | 2.33 | -- | -- | 10.75 | 2.34 | -- | -- |
| 157.5 | 12.63 | 3.19 | 12.24 | 2.95 | 11.87 | 2.75 | 11.71 | 2.73 | -- | -- | 11.89 | 2.75 | -- | -- |
| 162.5 | 13.36 | 3.26 | 13.06 | 3.07 | 12.78 | 2.92 | 12.59 | 2.90 | -- | -- | 12.80 | 2.91 | -- | -- |
| 167.5 | 13.91 | 3.01 | 13.71 | 2.88 | 13.50 | 2.77 | 13.32 | 2.76 | -- | -- | 13.52 | 2.77 | -- | -- |
| 172.5 | 14.35 | 2.42 | 14.21 | 2.33 | 14.08 | 2.27 | 13.95 | 2.26 | -- | -- | 14.09 | 2.26 | -- | -- |
| 177.5 | 14.69 | 1.44 | 14.63 | 1.40 | 14.56 | 1.37 | 14.49 | 1.36 | -- | -- | 14.57 | 1.37 | -- | -- |
| 182.5 | 15.00 | 0.00 | 15.00 | 0.00 | 15.00 | 0.00 | 15.00 | 0.00 | -- | -- | 15.00 | 0.00 | -- | -- |

Table 24. Logit-scale parameters for the probability of terminal molt for females for all model scenarios. The (arithmetic) probability of terminal molt was fixed at 0 for females less than 50 mm CW in Scenarios M19F03 and M19F05 and at 1 for females greater than 105 mm CW for all scenarios.

| scenario: | M19F00 |  | M19F00a |  | M19F01 |  | M19F02 |  | M19F03 |  | M19F04 |  | M19F05 |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| size bin | value | std err | value | std err | value | std err | value | std err | value | std err | value | std err | value | std err |
| 27.5 | -15.00 | 0.00 | -15.00 | 0.00 | -15.00 | 0.00 | -15.00 | 0.00 | -- | -- | -15.00 | 0.00 | -- | -- |
| 32.5 | -13.77 | 0.78 | -13.78 | 0.78 | -13.79 | 0.78 | -13.79 | 0.78 | -- | -- | -13.81 | 0.78 | -- | -- |
| 37.5 | -12.48 | 1.18 | -12.50 | 1.18 | -12.53 | 1.18 | -12.52 | 1.18 | - -- | -- | -12.56 | 1.18 | -- | -- |
| 42.5 | -11.09 | 1.29 | -11.12 | 1.29 | -11.15 | 1.28 | -11.13 | 1.28 | -- | -- | -11.20 | 1.28 | -- | -- |
| 47.5 | -9.53 | 1.15 | -9.56 | 1.15 | -9.60 | 1.15 | -9.58 | 1.15 | -- | -- | -9.66 | 1.14 | -- | -- |
| 52.5 | -7.76 | 0.86 | -7.79 | 0.86 | -7.83 | 0.86 | -7.80 | 0.86 | -6.82 | 0.99 | -7.90 | 0.86 | -6.89 | 1.00 |
| 57.5 | -5.75 | 0.52 | -5.78 | 0.53 | -5.81 | 0.52 | -5.78 | 0.52 | -5.05 | 0.45 | -5.88 | 0.52 | -5.11 | 0.45 |
| 62.5 | -3.58 | 0.24 | -3.60 | 0.24 | -3.63 | 0.24 | -3.60 | 0.24 | -3.34 | 0.21 | -3.70 | 0.24 | -3.39 | 0.20 |
| 67.5 | -1.77 | 0.11 | -1.78 | 0.11 | -1.81 | 0.11 | -1.78 | 0.11 | -1.79 | 0.11 | -1.87 | 0.11 | -1.85 | 0.11 |
| 72.5 | -0.43 | 0.09 | -0.44 | 0.08 | -0.48 | 0.08 | -0.44 | 0.08 | -0.51 | 0.09 | -0.52 | 0.08 | -0.54 | 0.09 |
| 77.5 | 0.31 | 0.09 | 0.28 | 0.09 | 0.24 | 0.08 | 0.29 | 0.09 | 0.22 | 0.09 | 0.26 | 0.08 | 0.27 | 0.09 |
| 82.5 | 0.59 | 0.10 | 0.59 | 0.10 | 0.56 | 0.09 | 0.59 | 0.10 | 0.55 | 0.10 | 0.55 | 0.09 | 0.56 | 0.09 |
| 87.5 | 1.28 | 0.16 | 1.23 | 0.15 | 1.20 | 0.14 | 1.26 | 0.15 | 1.18 | 0.14 | 1.03 | 0.12 | 1.04 | 0.12 |
| 92.5 | 2.58 | 0.35 | 2.36 | 0.29 | 2.36 | 0.27 | 2.53 | 0.31 | 2.26 | 0.25 | 2.08 | 0.22 | 2.12 | 0.22 |
| 97.5 | 4.03 | 0.67 | 3.61 | 0.50 | 3.67 | 0.49 | 3.96 | 0.60 | 3.48 | 0.47 | 3.50 | 0.41 | 3.53 | 0.43 |
| 102.5 | 5.52 | 1.27 | 4.91 | 0.99 | 5.03 | 1.00 | 5.42 | 1.18 | 4.78 | 0.99 | 5.02 | 0.87 | 5.06 | 0.93 |

Table 25. Log-scale NMFS survey catchability and selectivity parameters for all model scenarios.

| name | label | phase | scale | M19F00 value | std err | M19FOOa value | std err | M19F01 value | std err | M19F02 value | std err | M19F03 value | std err | M19F04 value | std err | M19F05 value | std err |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| ĐpQ[1] | ENMFS trawl survey: males, 1975-1981 | ®5 | OG | -0.693 | 0.000 | -0.693 | 0.000 | -0.693 | 0.000 | -0.693 | 0.000 | -0.693 | 0.000 | -0.693 | 0.000 | -0.693 | 0.000 |
| ${ }^{\text {bpQ[2] }}$ | - NMFS trawl survey: males, 1982+ | $\square 5$ | LOG | -0.45 | 0.05 | -0.635 | 0.06 | -0.70 | 0.06 | -0.75 | 0.06 | -0.84 | 0.06 | -0.70 | 0.053 | -0.76 | 0.055 |
| ĐpQ[3] | ENMFS trawl survey: females, 1975-1981 | 5 | LOG | -0.693 | 0.000 | -0.693 | 0.000 | -0.693 | 0.000 | -0.693 | 0.000 | -0.693 | 0.001 | -0.693 | . 000 | -0.693 | . 001 |
| $\pm \mathrm{pQ} 4$ ] | $\square$ NMFS trawl survey: females, 1982+ | $\boxminus 5$ | LOG | . 922 | . 073 | 85 | . 086 | -1.268 | . 84 | . 432 | . 88 | -1.437 | 105 | . 291 | 076 | 1.357 | . 099 |
| $\square \mathrm{pS} 111]$ | - 250 for NMFS survey selectivity (males, pre-1982) | $\square 1$ | ARITHMETIC | 52.441 | 25 | 90.000 | 0.001 | 90.000 | 0.000 | 90.000 | 0.000 | 90.000 | 0.000 | 90.000 | 0.000 | 90.000 | 0.000 |
| EpS1[2] | $\mathrm{E}_{250}$ for NMFS survey selectivity (males, 1982+) | 1 | ARITHMETIC | 34.262 | 4.137 | 40.160 | 6.282 | 40.369 | . 64 | 48.332 | 5.27 | 46.97 | 5.617 | 51.811 | 4.53 | 55.73 | . 69 |
| $\square \mathrm{ps} 13$ ] | $\mathrm{E}_{250}$ for NMFS survey selectivity (females, pre-1982) | $\square 1$ | ARITHMET | 5.408 | 854 | 6.838 | . 071 | 7.775 | . 96 | 2.30 | 3.32 | 2.15 | 4.94 | .50 | 96 | 2.97 | 4.820 |
| $\square \mathrm{pS1}$ [4] | - 250 for NMFS survey selectivity (females, 1982+) | $\square 1$ | ARITHMETIC | -35.492 | 30.433 | -33.961 | 30.933 | -36.975 | 32.573 | -47.549 | 41.643 | -0.042 | 18.679 | 4.632 | 15.305 | 18.651 | 14.067 |
| $\exists_{\text {pS2[1] }}$ | $\square_{\text {z95-250 for }}$ NMFS survey selectivity (males, pre-1982) | $\square 1$ | ARITHMETİ | 23.612 | 14 | 86.141 | 6.981 | 84.091 | 6.598 | 81.019 | 8 | 92.61 | 7.614 | 80.670 | 6.020 | 89.255 | 7.01 |
| EpS2[2] | Ez95-250 for NMFS survey selectivity (males, 1982+) | $\exists 1$ | ARITHMET | 75.23 | 10.334 | 99.001 | 17.736 | 100.000 | 0.003 | 100.000 | 0.00 | 100.000 | 0.000 | 100.000 | 0.00 | 100.000 | 0.0 |
| EpS2[3] | $\Xi_{\text {z95-z50 for NMFS survey selectivity (females, pre-1982) }}$ | E1 | ARITHMETIC | 40.090 | 5.841 | 59.809 | 6.261 | 59.360 | 5.973 | 65.786 | 6.834 | 68.015 | 8.994 | 60.215 | 5.987 | 67.834 | 8.860 |
| $\square \mathrm{pS2} 24]$ | $\square$ z95-z50 for NMFS survey selectivity (females, 1982+) | ®1 | ARITHMETIC | 100.000 | 0.002 | 100.000 | 0.002 | 100.000 | 0.002 | 100.000 | 0.003 | 100.000 | 0.001 | 100.000 | 0.001 | 100.000 | 0.00 |

Table 26. BSFRF SBS (side-by-side) male availability parameters for all model scenarios in which they were estimated.

|  | $\square$ BSFRF <br> M19F04 value | std err | (males, 20 | std err | $\square$ BSFRF availability (males, 2014) |  |  |  | $\square$ BSFRF availability (males, 2015) |  |  |  | $\square$ BSFRF availability (males, 2016) |  |  |  | $\square$ BSFRF availability (males, 2017) |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| index |  |  | M19F05 value |  | M19F04 <br> value | std err | M19F05 value | std err | M19F04 value | std err | M19F05 <br> value | std err | M19F04 <br> value | std err | M19F05 <br> value | std err | M19F04 <br> value | std err | M19F05 <br> value | std err |
| 1 | -3.297 | 0.615 | -3.255 | 0.614 | -3.608 | 0.708 | -3.590 | 0.700 | -3.084 | 0.588 | -3.030 | 0.578 | -0.591 | 0.477 | -0.438 | 0.488 | 0.113 | 0.340 | 0.321 | 0.352 |
| 2 | -3.463 | 0.509 | -3.428 | 0.510 | -3.558 | 0.569 | -3.558 | 0.563 | -3.057 | 0.465 | -3.021 | 0.457 | -0.848 | 0.390 | -0.716 | 0.399 | -0.433 | 0.298 | -0.234 | 0.308 |
| 3 | -3.606 | 0.443 | -3.578 | 0.446 | -3.505 | 0.466 | -3.523 | 0.463 | -3.032 | 0.381 | -3.015 | 0.376 | -1.082 | 0.344 | -0.971 | 0.351 | -0.901 | 0.292 | -0.717 | 0.301 |
| 4 | -3.709 | 0.407 | -3.690 | 0.412 | -3.445 | 0.401 | -3.481 | 0.400 | -3.002 | 0.331 | -3.006 | 0.330 | -1.271 | 0.326 | -1.185 | 0.332 | -1.204 | 0.299 | -1.046 | 0.306 |
| 5 | -3.764 | 0.388 | -3.753 | 0.394 | -3.374 | 0.366 | -3.430 | 0.365 | -2.970 | 0.308 | -2.999 | 0.308 | -1.400 | 0.324 | -1.347 | 0.327 | -1.356 | 0.311 | -1.235 | 0.316 |
| 6 | -3.764 | 0.375 | -3.763 | 0.381 | -3.292 | 0.347 | -3.368 | 0.347 | -2.932 | 0.298 | -2.992 | 0.297 | -1.458 | 0.326 | -1.444 | 0.326 | -1.394 | 0.326 | -1.319 | 0.328 |
| 7 | -3.713 | 0.363 | -3.722 | 0.368 | -3.205 | 0.334 | -3.302 | 0.333 | -2.872 | 0.292 | -2.968 | 0.291 | -1.440 | 0.327 | -1.476 | 0.323 | -1.347 | 0.342 | -1.328 | 0.340 |
| 8 | -3.618 | 0.349 | -3.638 | 0.354 | -3.107 | 0.320 | -3.225 | 0.318 | -2.775 | 0.287 | -2.910 | 0.283 | -1.350 | 0.327 | -1.443 | 0.318 | -1.241 | 0.357 | -1.283 | 0.350 |
| 9 | -3.476 | 0.334 | -3.505 | 0.339 | -2.987 | 0.302 | -3.125 | 0.300 | -2.632 | 0.279 | -2.803 | 0.274 | -1.193 | 0.326 | -1.346 | 0.312 | -1.091 | 0.369 | -1.197 | 0.356 |
| 10 | -3.283 | 0.321 | -3.320 | 0.326 | -2.829 | 0.283 | -2.984 | 0.280 | -2.446 | 0.267 | -2.647 | 0.262 | -0.974 | 0.323 | -1.189 | 0.305 | -0.907 | 0.379 | -1.076 | 0.360 |
| 11 | -3.048 | 0.310 | -3.092 | 0.316 | -2.623 | 0.263 | -2.794 | 0.260 | -2.242 | 0.254 | -2.465 | 0.248 | -0.706 | 0.321 | -0.979 | 0.297 | -0.699 | 0.386 | -0.925 | 0.360 |
| 12 | -2.792 | 0.304 | -2.843 | 0.309 | -2.367 | 0.245 | -2.551 | 0.241 | -2.061 | 0.241 | -2.298 | 0.235 | -0.415 | 0.321 | -0.740 | 0.292 | -0.471 | 0.392 | -0.749 | 0.360 |
| 13 | -2.529 | 0.303 | -2.589 | 0.307 | -2.083 | 0.232 | -2.276 | 0.227 | -1.924 | 0.230 | -2.164 | 0.224 | -0.125 | 0.322 | -0.485 | 0.290 | -0.228 | 0.396 | -0.544 | 0.358 |
| 14 | -2.244 | 0.306 | -2.315 | 0.310 | -1.774 | 0.224 | -1.969 | 0.217 | -1.828 | 0.222 | -2.060 | 0.215 | 0.141 | 0.325 | -0.235 | 0.291 | 0.023 | 0.398 | -0.317 | 0.356 |
| 15 | -1.899 | 0.312 | -1.979 | 0.315 | -1.437 | 0.218 | -1.625 | 0.211 | -1.756 | 0.215 | -1.965 | 0.208 | 0.362 | 0.327 | -0.002 | 0.292 | 0.270 | 0.396 | -0.074 | 0.353 |
| 16 | -1.484 | 0.320 | -1.574 | 0.322 | -1.097 | 0.216 | -1.267 | 0.208 | -1.684 | 0.209 | -1.858 | 0.204 | 0.527 | 0.326 | 0.199 | 0.292 | 0.498 | 0.390 | 0.170 | 0.348 |
| 17 | -1.044 | 0.327 | -1.143 | 0.329 | -0.824 | 0.216 | -0.977 | 0.208 | -1.605 | 0.203 | -1.745 | 0.199 | 0.624 | 0.320 | 0.348 | 0.290 | 0.689 | 0.380 | 0.391 | 0.341 |
| 18 | -0.668 | 0.334 | -0.776 | 0.337 | -0.660 | 0.219 | -0.806 | 0.210 | -1.557 | 0.199 | -1.672 | 0.195 | 0.642 | 0.309 | 0.422 | 0.283 | 0.820 | 0.366 | 0.563 | 0.333 |
| 19 | -0.445 | 0.343 | -0.560 | 0.345 | -0.618 | 0.224 | -0.763 | 0.215 | -1.540 | 0.197 | -1.637 | 0.193 | 0.573 | 0.295 | 0.411 | 0.274 | 0.877 | 0.350 | 0.669 | 0.323 |
| 20 | -0.419 | 0.353 | -0.533 | 0.355 | -0.648 | 0.230 | -0.797 | 0.220 | -1.540 | 0.196 | -1.623 | 0.192 | 0.413 | 0.280 | 0.305 | 0.265 | 0.850 | 0.332 | 0.696 | 0.311 |
| 21 | -0.566 | 0.362 | -0.671 | 0.364 | -0.683 | 0.234 | -0.836 | 0.222 | -1.514 | 0.195 | -1.586 | 0.192 | 0.174 | 0.265 | 0.116 | 0.256 | 0.734 | 0.315 | 0.637 | 0.300 |
| 22 | -0.807 | 0.367 | -0.895 | 0.368 | -0.693 | 0.234 | -0.851 | 0.223 | -1.453 | 0.193 | -1.516 | 0.191 | -0.119 | 0.252 | -0.134 | 0.247 | 0.533 | 0.302 | 0.492 | 0.293 |
| 23 | -1.086 | 0.368 | -1.155 | 0.370 | -0.670 | 0.233 | -0.835 | 0.222 | -1.395 | 0.193 | -1.454 | 0.191 | -0.451 | 0.245 | -0.432 | 0.244 | 0.258 | 0.300 | 0.271 | 0.297 |
| 24 | -1.385 | 0.374 | -1.432 | 0.376 | -0.655 | 0.236 | -0.831 | 0.224 | -1.369 | 0.196 | -1.430 | 0.194 | -0.814 | 0.256 | -0.768 | 0.256 | -0.073 | 0.318 | -0.010 | 0.320 |
| 25 | -1.703 | 0.395 | -1.731 | 0.397 | -0.672 | 0.249 | -0.862 | 0.236 | -1.391 | 0.206 | -1.458 | 0.203 | -1.198 | 0.296 | -1.132 | 0.294 | -0.441 | 0.368 | -0.332 | 0.372 |
| 26 | -2.037 | 0.445 | -2.044 | 0.447 | -0.735 | 0.283 | -0.941 | 0.268 | -1.461 | 0.231 | -1.538 | 0.227 | -1.589 | 0.371 | -1.507 | 0.366 | -0.831 | 0.453 | -0.678 | 0.458 |
| 27 | -2.381 | 0.530 | -2.369 | 0.532 | -0.831 | 0.347 | -1.055 | 0.328 | -1.576 | 0.281 | -1.664 | 0.276 | -1.987 | 0.481 | -1.889 | 0.473 | -1.230 | 0.573 | -1.035 | 0.576 |
| 28 | -2.730 | 0.650 | -2.698 | 0.652 | -0.941 | 0.444 | -1.183 | 0.421 | -1.726 | 0.364 | -1.825 | 0.357 | -2.388 | 0.624 | -2.276 | 0.612 | -1.633 | 0.725 | -1.397 | 0.725 |
| 29 | -3.080 | 0.803 | -3.029 | 0.803 | -1.055 | 0.572 | -1.315 | 0.546 | -1.886 | 0.482 | -1.999 | 0.472 | -2.792 | 0.794 | -2.666 | 0.779 | -2.038 | 0.904 | -1.761 | 0.902 |
| 30 | -3.431 | 0.983 | -3.360 | 0.983 | -1.171 | 0.730 | -1.450 | 0.700 | -2.049 | 0.630 | -2.175 | 0.618 | -3.195 | 0.987 | -3.055 | 0.969 | -2.442 | 1.106 | -2.125 | 1.101 |
| 31 | -3.782 | 1.187 | -3.692 | 1.186 | -1.288 | 0.913 | -1.586 | 0.880 | -2.213 | 0.805 | -2.353 | 0.790 | -3.599 | 1.202 | -3.445 | 1.181 | -2.847 | 1.329 | -2.489 | 1.321 |
| 32 | -4.132 | 1.411 | -4.024 | 1.409 | -1.406 | 1.118 | -1.722 | 1.081 | -2.377 | 1.002 | -2.531 | 0.985 | -4.002 | 1.434 | -3.835 | 1.411 | -3.252 | 1.569 | -2.853 | 1.560 |

Table 27. BSFRF SBS (side-by-side) female availability parameters for all model scenarios. in which they were estimated.


Table 28．Mean capture rate，selectivity and retention parameter estimates for the directed fishery（TCF）for all model scenarios．

| name | label | index | phase | M19F00 value | std err | M19F00a value | std err | M19F01 value | std err | M19F02 value | std err | M19F03 value | std err | M19F04 value | std err | M19F05 value | std err |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $\square \mathrm{pDC2}$［1］ | $\square$ TCF：female offset | 曰1 | 1 | －2．351 | 0.300 | －1．968 | 0.269 | －2．002 | 0.265 | －2．121 | 0.260 | －2．202 | 0.225 | －2．242 | 0.224 | －2．365 | 0.209 |
| ■plgtRet［1］ | $\square$ TCF：logit－scale max retention（pre－1997） | 曰1 | 3 | 14.999 | 2.211 | 14.999 | 4.414 | 14.999 | 4.355 | 14.999 | 4.757 | 14.999 | 5.155 | 14.999 | 4.336 | 14.999 | 4.561 |
| ＠pLgtRet［2］ | ETCF：logit－scale max retention（2005－2009） | ®1 | 3 | 2.101 | 1.305 | 14.210 | 996.790 | 14.919 | 337.810 | 14.863 | 526.220 | 14.993 | 39.551 | 14.868 | 479.580 | 14.928 | 322.800 |
| $\square \mathrm{pLgtRet}[3]$ | $\square$ TCF：logit－scale max retention（2013＋） | 日1 | 3 | 4.031 | 2.222 | 14.633 | 619.480 | 14.990 | 45.731 | 14.980 | 78.614 | 14.987 | 57.716 | 14.977 | 88.452 | 14.984 | 66.741 |
| $\square \mathrm{pLnC}[1]$ | $\square$ TCF：base capture rate，pre－1965（ $=0.05$ ） | 曰1 | －1 | －2．996 | 0.000 | －2．996 | 0.000 | －2．996 | 0.000 | －2．996 | 0.000 | －2．996 | 0.000 | －2．996 | 0.000 | －2．996 | 0.000 |
| $\square \mathrm{pLnC}[2]$ | ■TCF：base capture rate，1965＋ | ®1 | 1 | －1．418 | 0.083 | －1．580 | 0.085 | －1．679 | 0.083 | －1．634 | 0.083 | －1．818 | 0.086 | －1．678 | 0.081 | －1．767 | 0.082 |
| $\square$ pDevsS1［1］ | $\square \ln (250$ devs）for TCF selectivity（males，1991＋） | $\square 1$ | 2 | 0.037 | 0.018 | 0.086 | 0.011 | 0.100 | 0.012 | 0.092 | 0.011 | 0.090 | 0.010 | 0.099 | 0.012 | 0.090 | 0.010 |
|  |  | ®2 | 2 | 0.124 | 0.012 | 0.040 | 0.011 | 0.050 | 0.011 | 0.044 | 0.010 | 0.038 | 0.010 | 0.049 | 0.011 | 0.041 | 0.010 |
|  |  | ■ 3 | 2 | 0.107 | 0.014 | 0.118 | 0.013 | 0.130 | 0.014 | 0.121 | 0.013 | 0.112 | 0.012 | 0.128 | 0.013 | 0.113 | 0.012 |
|  |  | ®4 | 2 | 0.088 | 0.021 | 0.070 | 0.018 | 0.081 | 0.018 | 0.075 | 0.017 | 0.066 | 0.017 | 0.079 | 0.018 | 0.068 | 0.017 |
|  |  | ■5 | 2 | 0.001 | 0.027 | －0．002 | 0.026 | 0.002 | 0.027 | 0.009 | 0.025 | 0.005 | 0.024 | 0.001 | 0.026 | 0.008 | 0.023 |
|  |  | ＠ 6 | 2 | 0.130 | 0.040 | 0.153 | 0.038 | 0.167 | 0.038 | 0.161 | 0.037 | 0.161 | 0.036 | 0.164 | 0.038 | 0.159 | 0.035 |
|  |  | 日 7 | 2 | －0．079 | 0.017 | －0．076 | 0.016 | －0．067 | 0.016 | －0．064 | 0.015 | －0．061 | 0.015 | －0．066 | 0.016 | －0．062 | 0.015 |
|  |  | －8 | 2 | －0．087 | 0.018 | －0．080 | 0.016 | －0．068 | 0.016 | －0．067 | 0.015 | －0．062 | 0.015 | －0．066 | 0.016 | －0．062 | 0.015 |
|  |  | $\square 9$ | 2 | －0．124 | 0.016 | －0．122 | 0.015 | －0．114 | 0.015 | －0．108 | 0.014 | －0．103 | 0.014 | －0．113 | 0.015 | －0．103 | 0.014 |
|  |  | 曰10 | 2 | 0.019 | 0.014 | 0.018 | 0.014 | 0.029 | 0.013 | 0.027 | 0.013 | 0.030 | 0.013 | 0.029 | 0.013 | 0.029 | 0.013 |
|  |  | ®11 | 2 | 0.189 | 0.016 | 0.189 | 0.015 | 0.198 | 0.015 | 0.192 | 0.014 | 0.195 | 0.014 | 0.197 | 0.015 | 0.193 | 0.014 |
|  |  | $\square_{12}$ | 2 | －0．040 | 0.017 | －0．035 | 0.015 | －0．027 | 0.015 | －0．022 | 0.015 | －0．020 | 0.015 | －0．025 | 0.015 | －0．020 | 0.015 |
|  |  | $\boxminus 13$ | 2 | －0．100 | 0.014 | －0．096 | 0.013 | －0．083 | 0.012 | －0．085 | 0.012 | －0．085 | 0.012 | －0．082 | 0.012 | －0．086 | 0.012 |
|  |  | $\boxminus 14$ | 2 | －0．138 | 0.016 | －0．142 | 0.014 | －0．128 | 0.013 | －0．124 | 0.013 | －0．124 | 0.013 | －0．126 | 0.013 | －0．124 | 0.013 |
|  |  | ■15 | 2 | －0．125 | 0.021 | －0．122 | 0.019 | －0．113 | 0.019 | －0．103 | 0.017 | －0．098 | 0.017 | －0．112 | 0.019 | －0．099 | 0.017 |
|  |  | $\boxminus 16$ | 2 |  |  |  |  | －0．157 | 0.017 | －0．147 | 0.016 | －0．145 | 0.016 | －0．156 | 0.017 | －0．146 | 0.016 |
| ■ps1［28］ | E 250 for TCF retention（2005－2009） | ®1 | 1 | 138.799 | 1.573 | 137.700 | 0.303 | 137.716 | 0.337 | 137.711 | 0.331 | 137.711 | 0.329 | 137.716 | 0.348 | 137.711 | 0.328 |
| $\boxminus p S 1[29]$ | $\mathrm{E}_{250}$ for TCF retention（2013＋） | －1 | 1 | 125.230 | 0.725 | 125.170 | 0.566 | 125.216 | 0.544 | 125.269 | 0.539 | 125.254 | 0.538 | 125.189 | 0.543 | 125.249 | 0.538 |
| $\square \mathrm{pS} 155]$ | E $\mathbf{5 5 0}$ for TCF retention（pre－1991） | ®1 | 1 | 138.043 | 0.420 | 138.527 | 0.448 | 138.635 | 0.452 | 138.545 | 0.441 | 138.638 | 0.446 | 138.591 | 0.444 | 138.577 | 0.438 |
| $\square \mathrm{pS} 16]$ | Ez50 for TCF retention（1991－1996） | 曰1 | 1 | 137.483 | 0.250 | 138.337 | 0.331 | 138.378 | 0.340 | 138.418 | 0.347 | 138.475 | 0.357 | 138.380 | 0.339 | 138.438 | 0.352 |
| $\square \mathrm{pS} 1[8]$ | $\square \ln (250)$ for TCF selectivity（males） | ®1 | 1 | 4.858 | 0.008 | 4.865 | 0.008 | 4.857 | 0.007 | 4.860 | 0.007 | 4.859 | 0.007 | 4.859 | 0.007 | 4.860 | 0.007 |
| $\square \mathrm{pS} 1[9]$ | $\square \mathrm{z} 50$ for TCF selectivity（females） | ®1 | 1 | 96.441 | 2.583 | 96.842 | 2.621 | 96.722 | 2.641 | 96.719 | 2.600 | 95.205 | 2.202 | 94.863 | 2.164 | 94.174 | 1.986 |
| ■pS2［28］ | $\square$ slope for TCF retention（2005－2009） | －1 | 1 | 0.865 | 0.634 | 2.000 | 0.507 | 2.000 | 0.649 | 2.000 | 0.628 | 2.000 | 0.618 | 1.999 | 0.691 | 2.000 | 0.614 |
| $\square \mathrm{pS2}[29]$ | $\square$ slope for TCF retention（2013＋） | $\square 1$ | 1 | 0.563 | 0.115 | 0.568 | 0.108 | 0.570 | 0.104 | 0.563 | 0.100 | 0.565 | 0.100 | 0.575 | 0.105 | 0.567 | 0.101 |
| $\square \mathrm{pS2}[5]$ | $\square$ slope for TCF retention（pre－1991） | 曰1 | 1 | 0.687 | 0.125 | 0.686 | 0.118 | 0.687 | 0.115 | 0.678 | 0.115 | 0.689 | 0.116 | 0.692 | 0.116 | 0.694 | 0.117 |
| $\square \mathrm{pS} 2[6]$ | $\square$ slope for TCF retention（1997＋） | －1 | 1 | 0.954 | 0.190 | 0.937 | 0.222 | 0.933 | 0.222 | 0.920 | 0.217 | 0.908 | 0.212 | 0.931 | 0.221 | 0.918 | 0.217 |
| EpS2［7］ | Eslope for TCF selectivity（males，pre－1997） | 曰1 | 1 | 0.118 | 0.006 | 0.112 | 0.006 | 0.110 | 0.006 | 0.114 | 0.006 | 0.116 | 0.006 | 0.111 | 0.006 | 0.117 | 0.006 |
| EpS2［8］ | $\square$ slope for TCF selectivity（males，1997＋） | ®1 | 1 | 0.155 | 0.008 | 0.156 | 0.008 | 0.158 | 0.008 | 0.160 | 0.007 | 0.159 | 0.007 | 0.158 | 0.007 | 0.159 | 0.007 |
| EpS2［9］ | $\square$ slope for TCF selectivity（females） | $\boxminus 1$ | 1 | 0.185 | 0.019 | 0.184 | 0.018 | 0.179 | 0.017 | 0.179 | 0.017 | 0.184 | 0.017 | 0.189 | 0.018 | 0.191 | 0.018 |

Table 29. Log-scale male capture rate dev parameter estimates for the directed fishery (TCF) for all model scenarios.

|  | M19F00 |  | M19F00a |  | M19F01 |  | M19F02 |  | M19F03 |  | M19F04 |  | M19F05 |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| year | value | std err | value | std err | value | std err | value | std err | value | std err | value | std err | value | std err |
| 1965 | -0.548 | 0.463 | -0.569 | 0.459 | -0.588 | 0.456 | -0.561 | 0.459 | -0.494 | 0.460 | -0.590 | 0.456 | -0.501 | 0.460 |
| 1966 | -0.773 | 0.369 | -0.775 | 0.365 | -0.785 | 0.362 | -0.769 | 0.364 | -0.724 | 0.365 | -0.786 | 0.362 | -0.729 | 0.365 |
| 1967 | 0.449 | 0.336 | 0.460 | 0.329 | 0.459 | 0.325 | 0.466 | 0.328 | 0.494 | 0.327 | 0.459 | 0.324 | 0.492 | 0.328 |
| 1968 | 0.294 | 0.315 | 0.322 | 0.308 | 0.333 | 0.304 | 0.330 | 0.306 | 0.353 | 0.307 | 0.334 | 0.304 | 0.350 | 0.308 |
| 1969 | 0.474 | 0.304 | 0.504 | 0.298 | 0.518 | 0.294 | 0.514 | 0.295 | 0.532 | 0.296 | 0.517 | 0.295 | 0.528 | 0.298 |
| 1970 | 0.332 | 0.303 | 0.356 | 0.295 | 0.369 | 0.290 | 0.370 | 0.290 | 0.375 | 0.291 | 0.366 | 0.291 | 0.370 | 0.294 |
| 1971 | 0.138 | 0.293 | 0.159 | 0.283 | 0.172 | 0.277 | 0.182 | 0.276 | 0.169 | 0.276 | 0.168 | 0.277 | 0.160 | 0.280 |
| 1972 | -0.033 | 0.261 | -0.005 | 0.251 | 0.016 | 0.244 | 0.036 | 0.245 | 0.001 | 0.242 | 0.010 | 0.244 | -0.015 | 0.246 |
| 1973 | -0.281 | 0.199 | -0.227 | 0.193 | -0.191 | 0.188 | -0.161 | 0.191 | -0.220 | 0.185 | -0.197 | 0.188 | -0.247 | 0.187 |
| 1974 | -0.094 | 0.136 | 0.002 | 0.136 | 0.055 | 0.134 | 0.090 | 0.137 | 0.008 | 0.130 | 0.052 | 0.133 | -0.028 | 0.130 |
| 1975 | 0.130 | 0.103 | 0.271 | 0.108 | 0.340 | 0.106 | 0.373 | 0.109 | 0.281 | 0.104 | 0.340 | 0.105 | 0.240 | 0.102 |
| 1976 | 0.908 | 0.096 | 1.064 | 0.103 | 1.142 | 0.103 | 1.187 | 0.105 | 1.085 | 0.101 | 1.148 | 0.100 | 1.046 | 0.098 |
| 1977 | 1.711 | 0.113 | 1.812 | 0.124 | 1.885 | 0.123 | 2.007 | 0.131 | 1.827 | 0.117 | 1.901 | 0.121 | 1.797 | 0.115 |
| 1978 | 2.041 | 0.150 | 1.966 | 0.166 | 2.017 | 0.161 | 2.250 | 0.175 | 1.996 | 0.152 | 2.042 | 0.159 | 1.980 | 0.151 |
| 1979 | 2.818 | 0.229 | 2.383 | 0.225 | 2.407 | 0.205 | 2.703 | 0.229 | 2.488 | 0.220 | 2.443 | 0.201 | 2.483 | 0.223 |
| 1980 | 2.015 | 0.178 | 2.066 | 0.172 | 2.242 | 0.175 | 2.133 | 0.167 | 2.073 | 0.162 | 2.290 | 0.175 | 2.071 | 0.161 |
| 1981 | 0.207 | 0.112 | 0.357 | 0.116 | 0.534 | 0.119 | 0.353 | 0.110 | 0.390 | 0.108 | 0.565 | 0.118 | 0.386 | 0.108 |
| 1982 | -0.791 | 0.123 | -0.750 | 0.123 | -0.667 | 0.123 | -0.705 | 0.122 | -0.641 | 0.122 | -0.652 | 0.123 | -0.637 | 0.122 |
| 1983 | -1.796 | 0.244 | -1.801 | 0.245 | -1.768 | 0.246 | -1.733 | 0.248 | -1.708 | 0.248 | -1.760 | 0.246 | -1.692 | 0.249 |
| 1984 | -0.779 | 0.174 | -0.771 | 0.176 | -0.759 | 0.175 | -0.675 | 0.176 | -0.715 | 0.176 | -0.752 | 0.175 | -0.680 | 0.177 |
| 1987 | -1.338 | 0.208 | -1.271 | 0.211 | -1.230 | 0.211 | -1.189 | 0.211 | -1.120 | 0.213 | -1.214 | 0.212 | -1.116 | 0.214 |
| 1988 | -0.527 | 0.105 | -0.407 | 0.105 | -0.336 | 0.103 | -0.361 | 0.103 | -0.224 | 0.103 | -0.320 | 0.103 | -0.224 | 0.104 |
| 1989 | 0.669 | 0.081 | 0.772 | 0.078 | 0.820 | 0.076 | 0.821 | 0.077 | 0.998 | 0.077 | 0.832 | 0.076 | 0.999 | 0.078 |
| 1990 | 1.347 | 0.087 | 1.498 | 0.082 | 1.529 | 0.079 | 1.519 | 0.081 | 1.669 | 0.082 | 1.540 | 0.079 | 1.680 | 0.082 |
| 1991 | 1.352 | 0.105 | 1.762 | 0.118 | 1.823 | 0.119 | 1.742 | 0.116 | 1.826 | 0.115 | 1.835 | 0.119 | 1.852 | 0.116 |
| 1992 | 2.049 | 0.142 | 1.933 | 0.115 | 1.938 | 0.113 | 1.856 | 0.110 | 1.875 | 0.108 | 1.944 | 0.112 | 1.910 | 0.109 |
| 1993 | 1.442 | 0.147 | 1.576 | 0.143 | 1.594 | 0.144 | 1.491 | 0.140 | 1.428 | 0.136 | 1.596 | 0.143 | 1.470 | 0.135 |
| 1994 | 0.932 | 0.193 | 0.800 | 0.160 | 0.794 | 0.160 | 0.748 | 0.157 | 0.696 | 0.150 | 0.790 | 0.158 | 0.743 | 0.150 |
| 1995 | 0.340 | 0.178 | 0.222 | 0.166 | 0.168 | 0.161 | 0.219 | 0.166 | 0.204 | 0.161 | 0.163 | 0.158 | 0.251 | 0.161 |
| 1996 | 0.055 | 0.378 | -0.370 | 0.409 | -0.381 | 0.411 | -0.356 | 0.407 | -0.381 | 0.402 | -0.383 | 0.409 | -0.339 | 0.402 |
| 2005 | -2.086 | 0.189 | -2.210 | 0.206 | -2.172 | 0.206 | -2.208 | 0.206 | -2.159 | 0.207 | -2.173 | 0.206 | -2.156 | 0.207 |
| 2006 | -1.490 | 0.123 | -1.715 | 0.138 | -1.660 | 0.137 | -1.704 | 0.137 | -1.650 | 0.137 | -1.660 | 0.137 | -1.645 | 0.137 |
| 2007 | -1.473 | 0.110 | -1.653 | 0.119 | -1.614 | 0.117 | -1.652 | 0.117 | -1.618 | 0.117 | -1.625 | 0.116 | -1.617 | 0.117 |
| 2008 | -1.826 | 0.154 | -1.819 | 0.155 | -1.743 | 0.154 | -1.800 | 0.154 | -1.786 | 0.154 | -1.746 | 0.154 | -1.782 | 0.154 |
| 2009 | -1.198 | 0.265 | -1.147 | 0.263 | -1.049 | 0.264 | -1.125 | 0.258 | -1.091 | 0.260 | -1.046 | 0.265 | -1.087 | 0.260 |
| 2013 | -1.821 | 0.136 | -1.705 | 0.137 | -1.621 | 0.135 | -1.652 | 0.136 | -1.647 | 0.136 | -1.638 | 0.135 | -1.656 | 0.136 |
| 2014 | -0.623 | 0.089 | -0.581 | 0.093 | -0.463 | 0.090 | -0.558 | 0.088 | -0.546 | 0.087 | -0.503 | 0.089 | -0.568 | 0.087 |
| 2015 | -0.363 | 0.088 | -0.359 | 0.090 | -0.249 | 0.087 | -0.309 | 0.085 | -0.278 | 0.084 | -0.300 | 0.085 | -0.295 | 0.084 |
| 2017 | -1.864 | 0.125 | -2.151 | 0.143 | -2.042 | 0.140 | -2.037 | 0.141 | -1.983 | 0.141 | -2.097 | 0.139 | -1.995 | 0.140 |
| 2018 | -- | -- | -- | -- | -1.836 | 0.135 | -1.833 | 0.135 | -1.784 | 0.134 | -1.894 | 0.133 | -1.798 | 0.133 |

Table 30. Comparison of mean capture rate, In-scale capture rate devs, and selectivity parameter estimates for the snow crab fishery (SCF) for all model scenarios.

| name | label | index | phase | M19F00 value | std err | M19F00a value | std err | M19F01 value | std err | M19F02 value | std err | M19F03 value | std err | M19F04 value | std err | M19F05 value | std err |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| -pDC2[2] | ESCF: female offset | $\boxminus 1$ | 2 | -1.749 | 0.150 | -3.361 | 0.619 | -3.388 | 0.621 | -3.521 | 0.622 | -3.394 | 0.616 | -3.403 | 0.618 | -3.415 | 0.611 |
| $\square$ pDevsLnc[2] | -SCF: 1992+ | $\square 1$ | 2 | 1.945 | 0.091 | 0.572 | 0.105 | 0.586 | 0.105 | 0.571 | 0.105 | 0.512 | 0.104 | 0.583 | 0.105 | 0.516 | 0.104 |
|  |  | $\pm 2$ | 2 | 1.641 | 0.093 | 0.897 | 0.098 | 0.904 | 0.098 | 0.889 | 0.098 | 0.807 | 0.097 | 0.900 | 0.097 | 0.816 | 0.097 |
|  |  | -3 | 2 | 1.241 | 0.097 | 0.331 | 0.181 | 0.330 | 0.180 | 0.332 | 0.181 | 0.242 | 0.179 | 0.325 | 0.180 | 0.255 | 0.179 |
|  |  | $\square 4$ | 2 | 1.170 | 0.104 | 0.290 | 0.237 | 0.285 | 0.236 | 0.303 | 0.238 | 0.199 | 0.234 | 0.280 | 0.237 | 0.218 | 0.234 |
|  |  | $\square 5$ | 2 | -0.271 | 0.244 | 1.203 | 0.144 | 1.194 | 0.143 | 1.229 | 0.144 | 1.099 | 0.140 | 1.194 | 0.142 | 1.123 | 0.140 |
|  |  | ®6 | 2 | 0.784 | 0.211 | 0.919 | 0.160 | 0.895 | 0.163 | 0.909 | 0.162 | 0.901 | 0.161 | 0.915 | 0.160 | 0.912 | 0.159 |
|  |  | Đ7 | 2 | 0.999 | 0.203 | -0.136 | 0.354 | -0.147 | 0.352 | -0.122 | 0.351 | -0.134 | 0.351 | -0.133 | 0.353 | -0.123 | 0.351 |
|  |  | $\square 8$ | 2 | -0.035 | 0.336 | -0.995 | 0.551 | -1.000 | 0.548 | -0.976 | 0.550 | -0.982 | 0.548 | -0.991 | 0.551 | -0.975 | 0.549 |
|  |  | $\square 9$ | 2 | -0.982 | 0.513 | -0.731 | 0.495 | -0.733 | 0.493 | -0.717 | 0.491 | -0.718 | 0.492 | -0.724 | 0.496 | -0.711 | 0.492 |
|  |  | ®10 | 2 | -0.834 | 0.441 | -0.447 | 0.384 | -0.448 | 0.382 | -0.447 | 0.379 | -0.419 | 0.384 | -0.437 | 0.384 | -0.413 | 0.384 |
|  |  | ®11 | 2 | -0.614 | 0.358 | -1.148 | 0.497 | -1.155 | 0.495 | -1.152 | 0.493 | -1.115 | 0.500 | -1.148 | 0.496 | -1.112 | 0.500 |
|  |  | $\boxminus 12$ | 2 | -1.311 | 0.453 | -1.422 | 0.497 | -1.426 | 0.496 | -1.428 | 0.494 | -1.390 | 0.501 | -1.420 | 0.497 | -1.389 | 0.500 |
|  |  | $\square 13$ | 2 | -1.652 | 0.465 | -1.470 | 0.467 | -1.472 | 0.467 | -1.476 | 0.465 | -1.435 | 0.470 | -1.466 | 0.467 | -1.435 | 0.469 |
|  |  | ®14 | 2 | -0.540 | 0.229 | -0.107 | 0.205 | -0.109 | 0.204 | -0.118 | 0.204 | -0.079 | 0.204 | -0.090 | 0.204 | -0.076 | 0.204 |
|  |  | $\square_{15}$ | 2 | -0.251 | 0.170 | 0.041 | 0.164 | 0.030 | 0.163 | 0.030 | 0.163 | 0.069 | 0.163 | 0.046 | 0.163 | 0.070 | 0.163 |
|  |  | $\mathrm{E}_{16}$ | 2 | -0.144 | 0.145 | 0.136 | 0.142 | 0.138 | 0.141 | 0.116 | 0.141 | 0.124 | 0.141 | 0.151 | 0.141 | 0.124 | 0.140 |
|  |  | ®17 | 2 | -0.729 | 0.202 | -0.490 | 0.207 | -0.483 | 0.207 | -0.499 | 0.206 | -0.494 | 0.206 | -0.473 | 0.206 | -0.495 | 0.206 |
|  |  | ®18 | 2 | -0.504 | 0.181 | -0.105 | 0.160 | -0.107 | 0.159 | -0.103 | 0.159 | -0.085 | 0.159 | -0.102 | 0.159 | -0.087 | 0.159 |
|  |  | $\boxminus 19$ | 2 | -0.330 | 0.181 | -0.017 | 0.170 | -0.022 | 0.169 | -0.001 | 0.169 | 0.014 | 0.169 | -0.017 | 0.169 | 0.013 | 0.169 |
|  |  | $\square 20$ | 2 | 0.256 | 0.136 | 0.530 | 0.130 | 0.531 | 0.129 | 0.559 | 0.129 | 0.568 | 0.128 | 0.537 | 0.129 | 0.568 | 0.128 |
|  |  | -21 | 2 | -0.386 | 0.200 | 0.189 | 0.163 | 0.210 | 0.162 | 0.212 | 0.162 | 0.215 | 0.161 | 0.210 | 0.161 | 0.206 | 0.161 |
|  |  | - 22 | 2 | -0.269 | 0.148 | 0.081 | 0.144 | 0.130 | 0.143 | 0.089 | 0.143 | 0.101 | 0.142 | 0.121 | 0.142 | 0.082 | 0.142 |
|  |  | $\square_{23}$ | 2 | 0.646 | 0.099 | 0.953 | 0.093 | 1.005 | 0.091 | 0.971 | 0.091 | 1.005 | 0.089 | 0.986 | 0.090 | 0.984 | 0.088 |
|  |  | - 24 | 2 | 0.423 | 0.107 | 0.702 | 0.100 | 0.753 | 0.098 | 0.729 | 0.098 | 0.773 | 0.096 | 0.728 | 0.097 | 0.754 | 0.096 |
|  |  | - 25 | 2 | 0.202 | 0.125 | 0.463 | 0.119 | 0.514 | 0.117 | 0.502 | 0.117 | 0.548 | 0.115 | 0.483 | 0.116 | 0.530 | 0.115 |
|  |  | - 26 | 2 | -0.457 | 0.208 | -0.240 | 0.216 | -0.182 | 0.217 | -0.187 | 0.216 | -0.148 | 0.217 | -0.210 | 0.216 | -0.163 | 0.216 |
|  |  | $\pm 27$ | 2 |  |  |  |  | -0.221 | 0.258 | -0.215 | 0.258 | -0.177 | 0.260 | -0.248 | 0.257 | -0.189 | 0.259 |
| $\square \mathrm{pLnC}[3]$ | $\square$ SCF: base capture rate, pre-1978 ( $=0.01$ ) | $\pm 1$ | -2 | -4.605 | 0.000 | -4.605 | 0.000 | -4.605 | 0.000 | -4.605 | 0.000 | -4.605 | 0.000 | -4.605 | 0.000 | -4.605 | 0.000 |
| $\square \mathrm{pLnc}[4]$ | $\square$ SCF: base capture rate, 1992+ | 曰1 | 2 | -2.862 | 0.102 | -3.428 | 0.123 | -3.505 | 0.124 | -3.526 | 0.122 | -3.732 | 0.116 | -3.557 | 0.113 | -3.693 | 0.106 |
| - psi [10] | Eascending $\mathbf{z 5 0}$ for SCF selectivity (males, pre-1997) | $\Xi 1$ | 2 | 87.648 | 1.558 | 113.170 | 2.039 | 113.617 | 1.991 | 115.440 | 1.911 | 113.499 | 1.864 | 113.489 | 1.947 | 113.866 | 1.836 |
| -ps1[11] | Elascending $\mathbf{z 5 0}$ for SCF selectivity (males, 1997-2004) | $\square 1$ | 2 | 95.647 | 3.832 | 94.504 | 3.036 | 94.623 | 3.059 | 95.940 | 3.096 | 95.758 | 3.008 | 94.609 | 2.975 | 95.734 | 2.961 |
| $\square \mathrm{pS} 1$ [12] | $\square$ ascending 250 for SCF selectivity (males, 2005+) | -1 | 2 | 105.452 | 1.410 | 105.556 | 1.188 | 105.657 | 1.180 | 106.572 | 1.149 | 106.295 | 1.103 | 105.846 | 1.165 | 106.315 | 1.097 |
| $\square \mathrm{pS} 1$ [13] | $\boxminus$ ascending 250 for SCF selectivity (females, pre-1997) | ®1 | 2 | 70.333 | 4.978 | 74.138 | 4.872 | 74.155 | 4.844 | 74.154 | 4.851 | 73.422 | 4.650 | 74.086 | 4.752 | 73.547 | 4.635 |
| $\square \mathrm{pS} 1$ [14] | $\square$ ascending $\mathbf{z 5 0}$ for SCF selectivity (females, 1997-2004) | $\boxminus 1$ | 2 | 76.365 | 4.529 | 76.921 | 4.483 | 76.928 | 4.439 | 76.865 | 4.458 | 76.348 | 4.447 | 76.990 | 4.394 | 76.484 | 4.427 |
| $\square \mathrm{pS} 115 \mathrm{5}$ | $\square$ ascending $\mathbf{z 5 0}$ for SCF selectivity (females, 2005+) | $\boxminus 1$ | 2 | 84.942 | 5.484 | 81.126 | 4.013 | 80.715 | 4.017 | 80.706 | 4.030 | 79.972 | 3.937 | 80.666 | 3.847 | 80.056 | 3.826 |
| -ps2[10] | $\square$ ascending slope for SCF selectivity (males, pre-1997) | $\boxminus 1$ | 2 | 0.376 | 0.131 | 0.100 | 0.000 | 0.100 | 0.000 | 0.100 | 0.000 | 0.100 | 0.000 | 0.100 | 0.000 | 0.100 | 0.000 |
| -ps2[11] | Eascending slope for SCF selectivity (males, 1997-2004) | $\boxminus 1$ | 2 | 0.209 | 0.064 | 0.224 | 0.065 | 0.225 | 0.066 | 0.209 | 0.056 | 0.211 | 0.056 | 0.227 | 0.066 | 0.212 | 0.056 |
| $\square \mathrm{bs} 2[12]$ | $\square$ ascending slope for SCF selectivity (males, 2005+) | $\square 1$ | 2 | 0.175 | 0.015 | 0.182 | 0.013 | 0.181 | 0.013 | 0.180 | 0.013 | 0.182 | 0.013 | 0.180 | 0.013 | 0.182 | 0.013 |
| $\square \mathrm{bs} 2[13]$ | Eslope for SCF selectivity (females, pre-1997) | $\square 1$ | 2 | 0.221 | 0.127 | 0.162 | 0.065 | 0.163 | 0.065 | 0.162 | 0.065 | 0.170 | 0.068 | 0.164 | 0.064 | 0.170 | 0.066 |
| $\square \mathrm{pS} 2[14]$ | Eslope for SCF selectivity (females, 1997-2004) | $\boxminus 1$ | 2 | 0.263 | 0.128 | 0.257 | 0.119 | 0.259 | 0.119 | 0.259 | 0.120 | 0.264 | 0.126 | 0.259 | 0.117 | 0.262 | 0.123 |
| -ps2[15] | $\square$ slope for SCF selectivity (females, 2005+) | Đ1 | 2 | 0.157 | 0.049 | 0.193 | 0.057 | 0.190 | 0.055 | 0.190 | 0.055 | 0.193 | 0.058 | 0.193 | 0.054 | 0.194 | 0.056 |
| $\square \mathrm{pS4} 41]$ | $\square$ descending slope for SCF selectivity (males, pre-1997) | $\boxminus 1$ | 2 | 0.500 | 0.001 | 0.100 | 0.000 | 0.100 | 0.000 | 0.100 | 0.000 | 0.100 | 0.000 | 0.100 | 0.000 | 0.100 | 0.000 |
| $\square \mathrm{pS4} 42]$ | $\square$ descending slope for SCF selectivity (males, 1997-2004) | $\boxminus 1$ | 2 | 0.126 | 0.081 | 0.164 | 0.093 | 0.157 | 0.090 | 0.167 | 0.103 | 0.168 | 0.103 | 0.164 | 0.096 | 0.171 | 0.107 |
| $\square \mathrm{pS4} 43]$ | $\square$ descending slope for SCF selectivity (males, 2005+) | $\boxminus 1$ | 2 | 0.182 | 0.024 | 0.191 | 0.024 | 0.189 | 0.023 | 0.193 | 0.025 | 0.196 | 0.025 | 0.192 | 0.024 | 0.197 | 0.025 |

Table 31. Comparison of mean capture rate, $\ln$-scale capture rate devs, and selectivity parameters estimates for the BBRKC fishery (RKF) for all model scenarios.

| name | label | index | phase | M19F00 value | std err | M19F00a value | std err | M19F01 value | std err | M19F02 value | std err | M19F03 value | std err | M19F04 value | std err | M19F05 value | std err |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| -pDC2[4] | -RKF: female offset | $\square 1$ | 2 | -0.834 | 3.018 | -1.153 | 3.735 | -1.278 | 3.713 | -1.430 | 3.589 | -1.832 | 2.062 | -1.686 | 2.318 | -1.950 | 1.864 |
| $\square \mathrm{pDevsLnC}[4]$ | -RKF: 1992+ | $\exists 1$ | 2 | 0.845 | 0.275 | 0.565 | 0.191 | 0.548 | 0.190 | 0.522 | 0.189 | 0.466 | 0.185 | 0.548 | 0.190 | 0.474 | 0.185 |
|  |  | ®2 | 2 | 2.220 | 0.213 | 1.589 | 0.129 | 1.566 | 0.126 | 1.539 | 0.125 | 1.433 | 0.121 | 1.564 | 0.125 | 1.447 | 0.122 |
|  |  | -3 | 2 | -0.044 | 0.381 | 0.162 | 0.344 | 0.120 | 0.338 | 0.143 | 0.346 | 0.088 | 0.330 | 0.121 | 0.338 | 0.105 | 0.333 |
|  |  | $\boxminus 4$ | 2 | 0.010 | 0.396 | 0.294 | 0.425 | 0.260 | 0.418 | 0.268 | 0.424 | 0.282 | 0.421 | 0.265 | 0.419 | 0.287 | 0.424 |
|  |  | $\square 5$ | 2 | -0.012 | 0.392 | 0.256 | 0.425 | 0.227 | 0.419 | 0.236 | 0.424 | 0.255 | 0.424 | 0.233 | 0.421 | 0.258 | 0.426 |
|  |  | $\boxminus 6$ | 2 | -0.028 | 0.389 | 0.219 | 0.419 | 0.194 | 0.415 | 0.202 | 0.419 | 0.222 | 0.419 | 0.200 | 0.417 | 0.224 | 0.421 |
|  |  | $\boxminus 7$ | 2 | -0.035 | 0.387 | 0.195 | 0.412 | 0.173 | 0.409 | 0.179 | 0.412 | 0.196 | 0.412 | 0.178 | 0.411 | 0.197 | 0.413 |
|  |  | $\square 8$ | 2 | -0.052 | 0.381 | 0.145 | 0.398 | 0.129 | 0.397 | 0.128 | 0.397 | 0.145 | 0.397 | 0.135 | 0.398 | 0.145 | 0.398 |
|  |  | $\square 9$ | 2 | -0.060 | 0.376 | 0.098 | 0.379 | 0.089 | 0.380 | 0.082 | 0.378 | 0.106 | 0.381 | 0.095 | 0.382 | 0.106 | 0.382 |
|  |  | $\boxminus 10$ | 2 | -0.083 | 0.368 | 0.027 | 0.361 | 0.019 | 0.362 | 0.015 | 0.361 | 0.041 | 0.364 | 0.024 | 0.363 | 0.041 | 0.365 |
|  |  | ®11 | 2 | -0.126 | 0.355 | -0.070 | 0.341 | -0.069 | 0.343 | -0.076 | 0.342 | -0.053 | 0.344 | -0.064 | 0.344 | -0.053 | 0.344 |
|  |  | $\square 12$ | 2 | -0.161 | 0.345 | -0.146 | 0.323 | -0.140 | 0.326 | -0.146 | 0.325 | -0.128 | 0.326 | -0.132 | 0.327 | -0.127 | 0.326 |
|  |  | $\boxminus 13$ | 2 | -0.218 | 0.333 | -0.250 | 0.306 | -0.236 | 0.310 | -0.247 | 0.309 | -0.233 | 0.309 | -0.227 | 0.311 | -0.233 | 0.309 |
|  |  | 曰14 | 2 | -0.229 | 0.326 | -0.292 | 0.296 | -0.280 | 0.299 | -0.284 | 0.299 | -0.278 | 0.299 | -0.274 | 0.300 | -0.277 | 0.299 |
|  |  | ■15 | 2 | -0.125 | 0.319 | -0.191 | 0.282 | -0.167 | 0.287 | -0.184 | 0.285 | -0.195 | 0.282 | -0.161 | 0.287 | -0.195 | 0.282 |
|  |  | $\square_{16}$ | 2 | -0.236 | 0.314 | -0.320 | 0.278 | -0.295 | 0.282 | -0.298 | 0.282 | -0.306 | 0.280 | -0.291 | 0.282 | -0.306 | 0.280 |
|  |  | ®17 | 2 | -0.280 | 0.318 | -0.399 | 0.285 | -0.385 | 0.288 | -0.362 | 0.291 | -0.361 | 0.290 | -0.383 | 0.288 | -0.360 | 0.290 |
|  |  | ®18 | 2 | -0.239 | 0.328 | -0.323 | 0.296 | -0.312 | 0.299 | -0.277 | 0.304 | -0.274 | 0.304 | -0.310 | 0.299 | -0.272 | 0.304 |
|  |  | $\boxminus 19$ | 2 | -0.196 | 0.336 | -0.235 | 0.307 | -0.221 | 0.310 | -0.190 | 0.316 | -0.189 | 0.314 | -0.218 | 0.311 | -0.188 | 0.314 |
|  |  | $\boxminus 20$ | 2 | -0.189 | 0.329 | -0.193 | 0.303 | -0.160 | 0.309 | -0.166 | 0.309 | -0.174 | 0.306 | -0.161 | 0.309 | -0.181 | 0.305 |
|  |  | E21 | 2 | -0.139 | 0.311 | -0.188 | 0.278 | -0.125 | 0.288 | -0.167 | 0.282 | -0.175 | 0.280 | -0.134 | 0.286 | -0.188 | 0.278 |
|  |  | - 22 | 2 | -0.233 | 0.307 | -0.345 | 0.273 | -0.280 | 0.281 | -0.304 | 0.278 | -0.302 | 0.277 | -0.297 | 0.279 | -0.312 | 0.276 |
|  |  | $\mathrm{E}_{23}$ | 2 | -0.221 | 0.314 | -0.343 | 0.277 | -0.280 | 0.285 | -0.279 | 0.285 | -0.266 | 0.286 | -0.299 | 0.282 | -0.274 | 0.285 |
|  |  | - 24 | 2 | -0.168 | 0.327 | -0.255 | 0.288 | -0.193 | 0.296 | -0.174 | 0.299 | -0.156 | 0.301 | -0.212 | 0.293 | -0.164 | 0.299 |
|  |  | ®25 | 2 |  |  |  |  | -0.183 | 0.313 | -0.162 | 0.317 | -0.145 | 0.319 | -0.200 | 0.310 | -0.152 | 0.318 |
| - plnc[7] | $\square$ RKF: base capture rate, pre-1953 ( $=0.02$ ) | ®1 | -2 | -3.912 | 0.000 | -3.912 | 0.000 | -3.912 | 0.000 | -3.912 | 0.000 | -3.912 | 0.000 | -3.912 | 0.000 | -3.912 | 0.000 |
| $\square \mathrm{pLnC}[8]$ | -RKF: base capture rate, 1992+ | $\boxminus 1$ | 2 | -4.012 | 0.159 | -3.595 | 0.117 | -3.663 | 0.118 | -3.626 | 0.120 | -3.758 | 0.120 | -3.665 | 0.115 | -3.702 | 0.115 |
| $\square \mathrm{pS} 1[22]$ | $\square$ z95 for RKF selectivity (males, pre-1997) | $\boxminus 1$ | 3 | 157.784 | 6.506 | 151.838 | 4.156 | 152.477 | 4.147 | 152.695 | 4.047 | 151.025 | 4.078 | 152.312 | 4.110 | 151.034 | 4.047 |
| $\square \mathrm{pS} 1[23]$ | @z95 for RKF selectivity (males, 1997-2004) | $\boxminus 1$ | 3 | 180.000 | 0.005 | 180.000 | 0.001 | 180.000 | 0.001 | 180.000 | 0.001 | 180.000 | 0.001 | 180.000 | 0.001 | 180.000 | 0.001 |
| $\square \mathrm{pS1}$ [24] | Ez95 for RKF selectivity (males, 2005+) | $\boxminus 1$ | 3 | 180.000 | 0.000 | 180.000 | 0.000 | 180.000 | 0.000 | 180.000 | 0.000 | 180.000 | 0.000 | 180.000 | 0.000 | 180.000 | 0.000 |
| $\square \mathrm{ps} 1[25]$ | $\mathrm{E}_{\text {z95 }}$ for RKF selectivity (females, pre-1997) | $\square 1$ | 3 | 121.870 | 39.215 | 125.661 | 42.789 | 125.322 | 41.346 | 125.189 | 40.260 | 118.660 | 23.645 | 119.762 | 27.833 | 116.967 | 22.375 |
| $\square \mathrm{ps} 126]$ | $\mathrm{E}_{\text {z95 }}$ for RKF selectivity (females, 1997-2004) | $\Xi 1$ | 3 | 122.103 | 56.686 | 123.545 | 50.295 | 124.984 | 56.212 | 125.454 | 57.613 | 121.229 | 48.066 | 120.998 | 50.369 | 119.121 | 45.518 |
| $\square \mathrm{pS} 1[27]$ | Ez95 for RKF selectivity (females, 2005+) | ®1 | 3 | 140.000 | 0.037 | 140.000 | 0.034 | 140.000 | 0.036 | 140.000 | 0.035 | 140.000 | 0.103 | 140.000 | 0.040 | 140.000 | 0.107 |
| ■pS2[22] | Eln(z95-250) for RKF selectivity (males, pre-1997) | ®1 | 3 | 3.070 | 0.163 | 2.930 | 0.134 | 2.943 | 0.133 | 2.931 | 0.129 | 2.914 | 0.133 | 2.935 | 0.132 | 2.908 | 0.133 |
| $\square \mathrm{lpS} 2[23]$ | Eln(z95-250) for RKF selectivity (males, 1997-2004) | $\triangle 1$ | 3 | 3.550 | 0.086 | 3.458 | 0.074 | 3.452 | 0.074 | 3.439 | 0.072 | 3.433 | 0.072 | 3.447 | 0.073 | 3.431 | 0.071 |
| $\square \mathrm{bS} 2[24]$ | $\square \ln (295-250)$ for RKF selectivity (males, 2005+) | $\square 1$ | 3 | 3.516 | 0.041 | 3.435 | 0.038 | 3.428 | 0.036 | 3.413 | 0.035 | 3.408 | 0.035 | 3.418 | 0.036 | 3.405 | 0.035 |
| - pS2[25] | Eln(z95-250) for RKF selectivity (males, pre-1997) | $\boxminus 1$ | 3 | 2.789 | 0.697 | 2.830 | 0.607 | 2.825 | 0.599 | 2.823 | 0.590 | 2.743 | 0.529 | 2.759 | 0.565 | 2.719 | 0.544 |
| $\square \mathrm{pS} 2[26]$ | Eln(z95-250) for RKF selectivity (males, 1997-2004) | 曰1 | 3 | 2.859 | 0.909 | 2.866 | 0.781 | 2.883 | 0.795 | 2.892 | 0.799 | 2.865 | 0.860 | 2.849 | 0.872 | 2.840 | 0.895 |
| $\boxminus p S 2[27]$ | Eln(z95-250) for RKF selectivity (males, 2005+) | $\boxminus 1$ | 3 | 2.985 | 0.212 | 2.970 | 0.205 | 2.971 | 0.204 | 2.972 | 0.204 | 3.026 | 0.201 | 2.999 | 0.203 | 3.039 | 0.201 |

Table 32. Comparison of mean capture rate and selectivity parameters estimates for the groundfish fisheries (GTF).

| name | label | M19F00 value | std err | M19F00a value | std err | M19F01 value | std err | M19F02 value | std err | M19F03 value | std err | M19F04 value | std err | M19F05 value | std err |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| EpDC2[3] | GTF: female offset | -0.957 | 0.072 | -0.981 | 0.087 | -1.081 | 0.079 | -1.231 | 0.081 | -1.002 | 0.083 | -1.137 | 0.078 | -1.048 | 0.082 |
| EpLnC[6] | GTF: base capture rate, ALL YEARS | -4.408 | 0.067 | -4.611 | 0.073 | -4.830 | 0.066 | -4.804 | 0.068 | -4.992 | 0.069 | -4.843 | 0.060 | -4.948 | 0.061 |
| \#pS1[16] | z50 for GF.AllGear selectivity (males, pre-1987) | 55.070 | 1.852 | 57.435 | 2.141 | 59.312 | 2.273 | 61.798 | 2.619 | 57.543 | 2.499 | 60.814 | 2.316 | 59.036 | 2.555 |
| EpS1[17] | z50 for GF.AllGear selectivity (males, 1987-1996) | 59.005 | 4.849 | 60.873 | 6.670 | 68.343 | 6.466 | 78.079 | 6.766 | 68.399 | 5.326 | 71.569 | 6.318 | 70.533 | 5.218 |
| EpS1[18] | z50 for GF.AllGear selectivity (males, 1997+) | 80.710 | 2.123 | 86.047 | 2.398 | 87.470 | 2.298 | 93.086 | 2.404 | 92.847 | 2.489 | 89.264 | 2.235 | 93.451 | 2.401 |
| \#pS1[19] | z50 for GF.AllGear selectivity (males, pre-1987) | 41.206 | 1.659 | 41.987 | 1.914 | 41.429 | 1.744 | 40.562 | 1.645 | 41.453 | 1.663 | 41.795 | 1.711 | 41.970 | 1.654 |
| EpS1[20] | z50 for GF.AllGear selectivity (males, 1987-1996) | 40.000 | 0.000 | 40.000 | 0.000 | 40.000 | 0.000 | 40.000 | 0.000 | 40.000 | 0.000 | 40.000 | 0.000 | 40.000 | 0.002 |
| EpS1[21] | z50 for GF.AllGear selectivity (males, 1997+) | 76.232 | 2.497 | 79.614 | 2.733 | 77.468 | 2.549 | 78.569 | 2.695 | 85.086 | 3.036 | 77.762 | 2.492 | 84.499 | 2.955 |
| EpS2[16] | slope for GF.AllGear selectivity (males, pre-1987) | 0.104 | 0.010 | 0.094 | 0.009 | 0.088 | 0.008 | 0.084 | 0.008 | 0.093 | 0.010 | 0.087 | 0.008 | 0.091 | 0.009 |
| EpS2[17] | slope for GF.AllGear selectivity (males, 1987-1996) | 0.057 | 0.012 | 0.049 | 0.013 | 0.041 | 0.007 | 0.037 | 0.005 | 0.046 | 0.007 | 0.040 | 0.006 | 0.046 | 0.006 |
| EpS2[18] | slope for GF.AllGear selectivity (males, 1997+) | 0.075 | 0.004 | 0.069 | 0.004 | 0.067 | 0.003 | 0.063 | 0.003 | 0.061 | 0.003 | 0.068 | 0.003 | 0.062 | 0.003 |
| EpS2[19] | slope for GF.AllGear selectivity (females, pre-1987) | 0.137 | 0.022 | 0.124 | 0.021 | 0.130 | 0.021 | 0.139 | 0.022 | 0.138 | 0.020 | 0.130 | 0.020 | 0.138 | 0.020 |
| $\square \mathrm{pS} 2[20]$ | slope for GF.AllGear selectivity (females, 1987-1996) | 0.185 | 0.038 | 0.189 | 0.038 | 0.184 | 0.039 | 0.182 | 0.039 | 0.168 | 0.038 | 0.182 | 0.039 | 0.167 | 0.038 |
| ■pS2[21] | slope for GF.AllGear selectivity (females, 1997+) | 0.073 | 0.006 | 0.070 | 0.005 | 0.073 | 0.005 | 0.071 | 0.005 | 0.063 | 0.005 | 0.075 | 0.005 | 0.064 | 0.005 |

Table 33. Log-scale capture rate dev parameter estimates for the groundfish fisheries (GTF) for all model scenarios.

| scenario: | M19F00 |  | M19F00a |  | M19F01 |  | M19F02 |  | M19F03 |  | M19F04 |  | M19F05 |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| year | value | std err | value | std err | value | std err | value | std err | value | std err | value | std err | value | std err |
| 1973 | 1.312 | 0.098 | 1.352 | 0.099 | 1.524 | 0.092 | 1.496 | 0.097 | 1.428 | 0.097 | 1.519 | 0.091 | 1.397 | 0.097 |
| 1974 | 1.703 | 0.077 | 1.764 | 0.081 | 1.945 | 0.073 | 1.916 | 0.076 | 1.853 | 0.077 | 1.943 | 0.071 | 1.819 | 0.075 |
| 1975 | 0.856 | 0.072 | 0.924 | 0.077 | 1.110 | 0.069 | 1.081 | 0.071 | 1.036 | 0.072 | 1.111 | 0.066 | 1.002 | 0.069 |
| 1976 | 0.319 | 0.079 | 0.376 | 0.084 | 0.562 | 0.077 | 0.542 | 0.078 | 0.519 | 0.080 | 0.565 | 0.074 | 0.487 | 0.077 |
| 1977 | 0.009 | 0.100 | 0.033 | 0.104 | 0.217 | 0.098 | 0.210 | 0.099 | 0.203 | 0.100 | 0.223 | 0.097 | 0.175 | 0.098 |
| 1978 | -0.254 | 0.130 | -0.270 | 0.134 | -0.085 | 0.130 | -0.087 | 0.130 | -0.070 | 0.131 | -0.075 | 0.129 | -0.094 | 0.130 |
| 1979 | 0.369 | 0.098 | 0.289 | 0.101 | 0.482 | 0.094 | 0.467 | 0.095 | 0.520 | 0.095 | 0.497 | 0.092 | 0.497 | 0.093 |
| 1980 | -0.009 | 0.126 | -0.094 | 0.126 | 0.099 | 0.121 | 0.063 | 0.121 | 0.132 | 0.121 | 0.115 | 0.120 | 0.118 | 0.120 |
| 1981 | -0.201 | 0.158 | -0.276 | 0.158 | -0.103 | 0.156 | -0.129 | 0.156 | -0.058 | 0.156 | -0.089 | 0.155 | -0.059 | 0.156 |
| 1982 | -0.969 | 0.353 | -1.029 | 0.350 | -0.893 | 0.357 | -0.895 | 0.358 | -0.836 | 0.361 | -0.883 | 0.358 | -0.826 | 0.362 |
| 1983 | -0.418 | 0.302 | -0.457 | 0.300 | -0.328 | 0.303 | -0.306 | 0.305 | -0.260 | 0.306 | -0.320 | 0.303 | -0.239 | 0.307 |
| 1984 | -0.203 | 0.326 | -0.211 | 0.325 | -0.093 | 0.329 | -0.056 | 0.331 | -0.030 | 0.334 | -0.088 | 0.329 | -0.001 | 0.336 |
| 1985 | -0.615 | 0.425 | -0.605 | 0.423 | -0.495 | 0.434 | -0.462 | 0.435 | -0.452 | 0.446 | -0.489 | 0.433 | -0.432 | 0.450 |
| 1986 | -0.461 | 0.321 | -0.439 | 0.318 | -0.308 | 0.323 | -0.295 | 0.321 | -0.254 | 0.331 | -0.303 | 0.322 | -0.243 | 0.332 |
| 1987 | -0.659 | 0.319 | -0.606 | 0.319 | -0.421 | 0.325 | -0.381 | 0.324 | -0.363 | 0.329 | -0.407 | 0.325 | -0.356 | 0.330 |
| 1988 | -1.066 | 0.363 | -0.997 | 0.364 | -0.818 | 0.375 | -0.796 | 0.374 | -0.769 | 0.379 | -0.806 | 0.375 | -0.768 | 0.380 |
| 1989 | -0.874 | 0.295 | -0.782 | 0.295 | -0.594 | 0.300 | -0.584 | 0.299 | -0.560 | 0.301 | -0.583 | 0.300 | -0.562 | 0.301 |
| 1990 | -0.532 | 0.231 | -0.448 | 0.233 | -0.250 | 0.235 | -0.258 | 0.234 | -0.252 | 0.233 | -0.240 | 0.235 | -0.254 | 0.233 |
| 1991 | 0.569 | 0.104 | 0.630 | 0.108 | 0.518 | 0.070 | 0.477 | 0.071 | 0.405 | 0.069 | 0.524 | 0.070 | 0.406 | 0.069 |
| 1992 | 0.869 | 0.097 | 0.893 | 0.101 | 0.798 | 0.067 | 0.752 | 0.069 | 0.666 | 0.066 | 0.803 | 0.067 | 0.670 | 0.066 |
| 1993 | 0.704 | 0.134 | 0.668 | 0.136 | 0.431 | 0.082 | 0.391 | 0.084 | 0.291 | 0.082 | 0.435 | 0.082 | 0.300 | 0.082 |
| 1994 | 1.168 | 0.117 | 1.072 | 0.120 | 0.952 | 0.071 | 0.927 | 0.073 | 0.821 | 0.071 | 0.956 | 0.071 | 0.833 | 0.071 |
| 1995 | 1.155 | 0.150 | 1.015 | 0.152 | 0.886 | 0.080 | 0.881 | 0.081 | 0.758 | 0.080 | 0.889 | 0.079 | 0.773 | 0.079 |
| 1996 | 1.440 | 0.145 | 1.287 | 0.147 | 1.002 | 0.083 | 1.012 | 0.084 | 0.877 | 0.083 | 1.006 | 0.082 | 0.896 | 0.082 |
| 1997 | 1.415 | 0.190 | 1.311 | 0.190 | 1.429 | 0.079 | 1.459 | 0.080 | 1.445 | 0.080 | 1.447 | 0.079 | 1.463 | 0.080 |
| 1998 | 1.186 | 0.252 | 1.110 | 0.250 | 1.321 | 0.089 | 1.360 | 0.090 | 1.348 | 0.089 | 1.340 | 0.089 | 1.367 | 0.089 |
| 1999 | 0.740 | 0.370 | 0.684 | 0.367 | 0.689 | 0.136 | 0.729 | 0.137 | 0.729 | 0.136 | 0.709 | 0.136 | 0.748 | 0.136 |
| 2000 | 0.816 | 0.308 | 0.785 | 0.312 | 0.674 | 0.127 | 0.711 | 0.128 | 0.729 | 0.127 | 0.694 | 0.127 | 0.747 | 0.127 |
| 2001 | 1.129 | 0.197 | 1.129 | 0.199 | 0.801 | 0.100 | 0.832 | 0.101 | 0.863 | 0.100 | 0.819 | 0.100 | 0.879 | 0.100 |
| 2002 | 0.455 | 0.304 | 0.459 | 0.308 | 0.073 | 0.159 | 0.104 | 0.160 | 0.140 | 0.160 | 0.088 | 0.159 | 0.155 | 0.159 |
| 2003 | -0.162 | 0.418 | -0.157 | 0.425 | -0.318 | 0.190 | -0.293 | 0.190 | -0.267 | 0.190 | -0.305 | 0.189 | -0.253 | 0.190 |
| 2004 | 0.001 | 0.310 | 0.026 | 0.314 | -0.011 | 0.127 | 0.003 | 0.128 | 0.025 | 0.127 | -0.002 | 0.127 | 0.036 | 0.127 |
| 2005 | -0.246 | 0.323 | -0.213 | 0.327 | -0.367 | 0.155 | -0.359 | 0.156 | -0.343 | 0.156 | -0.361 | 0.155 | -0.333 | 0.156 |
| 2006 | -0.232 | 0.285 | -0.204 | 0.288 | -0.386 | 0.148 | -0.385 | 0.149 | -0.378 | 0.149 | -0.383 | 0.148 | -0.370 | 0.149 |
| 2007 | -0.344 | 0.287 | -0.316 | 0.290 | -0.110 | 0.116 | -0.116 | 0.116 | -0.116 | 0.116 | -0.112 | 0.115 | -0.110 | 0.116 |
| 2008 | -0.593 | 0.333 | -0.572 | 0.337 | -0.434 | 0.155 | -0.440 | 0.155 | -0.439 | 0.155 | -0.441 | 0.154 | -0.436 | 0.155 |
| 2009 | -0.780 | 0.394 | -0.776 | 0.398 | -0.821 | 0.223 | -0.817 | 0.223 | -0.812 | 0.223 | -0.834 | 0.222 | -0.811 | 0.223 |
| 2010 | -0.900 | 0.455 | -0.904 | 0.459 | -1.113 | 0.294 | -1.101 | 0.294 | -1.100 | 0.294 | -1.125 | 0.293 | -1.098 | 0.294 |
| 2011 | -0.905 | 0.476 | -0.895 | 0.482 | -0.682 | 0.204 | -0.667 | 0.205 | -0.662 | 0.204 | -0.696 | 0.204 | -0.664 | 0.204 |
| 2012 | -1.074 | 0.486 | -1.032 | 0.495 | -1.234 | 0.295 | -1.232 | 0.295 | -1.219 | 0.295 | -1.251 | 0.294 | -1.226 | 0.295 |
| 2013 | -0.944 | 0.405 | -0.877 | 0.413 | -0.863 | 0.201 | -0.881 | 0.201 | -0.858 | 0.201 | -0.888 | 0.200 | -0.871 | 0.201 |
| 2014 | -0.908 | 0.366 | -0.850 | 0.371 | -0.811 | 0.193 | -0.841 | 0.194 | -0.809 | 0.193 | -0.844 | 0.193 | -0.824 | 0.193 |
| 2015 | -0.953 | 0.399 | -0.922 | 0.401 | -0.946 | 0.244 | -0.974 | 0.244 | -0.938 | 0.244 | -0.985 | 0.243 | -0.952 | 0.244 |
| 2016 | -0.891 | 0.428 | -0.873 | 0.430 | -0.800 | 0.253 | -0.818 | 0.253 | -0.782 | 0.253 | -0.843 | 0.252 | -0.794 | 0.253 |
| 2017 | -1.021 | 0.496 | -1.003 | 0.499 | -1.213 | 0.374 | -1.222 | 0.375 | -1.186 | 0.377 | -1.259 | 0.371 | -1.201 | 0.376 |
| 2018 | 0.000 | 0.000 | 0.000 | 0.000 | -1.015 | 0.346 | -1.019 | 0.347 | -0.97 | 0.349 | . 07 | 0.343 | .995 | 0.348 |

Table 34. (Unweighted) negative log-likelihoods and (weighted) objective function values for fishery-related data components from the model scenarios. TCF: directed Tanner crab fishery; SCF: snow crab fishery; RKF: BBRKC fishery; GTF: groundfish fisheries.

| fleet | catch.type | data.type | x | NLLs <br> M19F01 | M19F02 | M19F03 | M19F04 | M19F05 | Objective funct M19F01 | tion values <br> M19F02 | M19F03 | M19F04 | M19F05 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $\square \mathrm{GTF}$ | छtotal catch | ■abundance | all sexes | 0.15 | 0.15 | 0.16 | 0.15 | 0.16 | 2.99 | 3.03 | 3.19 | 3.04 | 3.23 |
|  |  | $\square$ biomass | all sexes | 1.24 | 1.28 | 1.48 | 1.26 | 1.49 | 24.71 | 25.60 | 29.69 | 25.11 | 29.78 |
|  |  | En.at.z | female | 293.88 | 293.59 | 274.46 | 290.59 | 273.66 | 293.88 | 293.59 | 274.46 | 290.59 | 273.66 |
|  |  |  | male | 288.00 | 294.90 | 285.09 | 291.02 | 287.46 | 288.00 | 294.90 | 285.09 | 291.02 | 287.46 |
| $\square \mathrm{RKF}$ | 曰total catch | Gabundance | female | 21,939.76 | 25,053.62 | 28,472.19 | 18,960.49 | 30,238.99 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  |  | male | 8,919.09 | 8,498.83 | 8,786.52 | 8,885.16 | 8,754.89 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  | $\square$ biomass | female | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.06 | 0.06 | 0.07 | 0.07 | 0.07 |
|  |  |  | male | 1.20 | 1.23 | 1.36 | 1.22 | 1.36 | 23.99 | 24.60 | 27.22 | 24.34 | 27.28 |
|  |  | En.at.z | female | 3.16 | 3.16 | 3.06 | 3.24 | 3.13 | 3.16 | 3.16 | 3.06 | 3.24 | 3.13 |
|  |  |  | male | 75.34 | 72.80 | 74.43 | 75.72 | 75.27 | 75.34 | 72.80 | 74.43 | 75.72 | 75.27 |
| $\square$ SCF | 日total catch | $\boxminus$ abundance | female | 267.01 | 271.11 | 367.66 | 280.76 | 363.74 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  |  | male | 100.24 | 101.77 | 96.25 | 98.78 | 96.11 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  | $\square$ biomass | female | 0.10 | 0.10 | 0.10 | 0.10 | 0.10 | 1.92 | 1.92 | 1.92 | 1.92 | 1.92 |
|  |  |  | male | 1.12 | 1.04 | 0.89 | 1.12 | 0.89 | 22.35 | 20.89 | 17.75 | 22.37 | 17.76 |
|  |  | En.at.z | female | 15.07 | 15.10 | 15.69 | 15.88 | 16.30 | 15.07 | 15.10 | 15.69 | 15.88 | 16.30 |
|  |  |  | male | 133.09 | 129.50 | 124.76 | 132.85 | 125.92 | 133.09 | 129.50 | 124.76 | 132.85 | 125.92 |
| $\square$ TCF | $\square$ retained cal | Gabundance | female | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  |  | male | 226.68 | 224.71 | 218.46 | 227.25 | 219.40 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  | Ebiomass | female | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  |  | male | 0.44 | 0.41 | 0.37 | 0.44 | 0.39 | 8.75 | 8.19 | 7.35 | 8.72 | 7.74 |
|  |  | En.at.z | male | 54.24 | 54.37 | 51.98 | 54.69 | 52.92 | 54.24 | 54.37 | 51.98 | 54.69 | 52.92 |
|  | $\square$ total catch | ■abundance | female | 13,346.33 | 13,369.84 | 17,446.24 | 13,835.53 | 15,990.44 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  |  | male | 10.78 | 10.19 | 9.31 | 10.65 | 9.37 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  | Ebiomass | female | 0.53 | 0.50 | 0.50 | 0.53 | 0.50 | 10.63 | 10.07 | 9.96 | 10.65 | 9.92 |
|  |  |  | male | 0.21 | 0.20 | 0.19 | 0.21 | 0.19 | 4.19 | 3.99 | 3.77 | 4.18 | 3.80 |
|  |  | En.at.z | female | 18.19 | 18.11 | 18.16 | 17.86 | 17.97 | 18.19 | 18.11 | 18.16 | 17.86 | 17.97 |
|  |  |  | male | 89.01 | 88.69 | 88.13 | 87.57 | 87.80 | 89.01 | 88.69 | 88.13 | 87.57 | 87.80 |

Table 35. (Unweighted) negative log-likelihoods and (weighted) objective function values for survey-related data components from the model scenarios. Rows consisting of all zero values indicate a data component which was not included in any of the models. Blank cells indicate a data component (row) that was not included in the associated scenario (column).

| fleet | catch.type | data.type | NLLs <br> M19F01 | M19F02 | M19F03 | M19F04 | M19F05 | Objective funct M19F01 | tion values M19F02 | M19F03 | M19F04 | M19F05 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| - NMFS 0 | $\square$ index catch | abundance | 838.19 | 792.09 | 757.00 | 818.86 | 748.14 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  | biomass | 250.88 | 234.17 | 267.37 | 251.37 | 264.97 | 250.88 | 234.17 | 0.00 | 251.37 | 0.00 |
|  |  | n.at.z | 991.65 | 982.94 | 1,126.75 | 993.77 | 1,135.21 | 991.65 | 982.94 | 0.00 | 993.77 | 0.00 |
| $\square$ NMFS F | Bindex catch | abundance | 388.10 | 372.88 | 329.04 | 383.67 | 328.60 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  | biomass | 320.69 | 307.61 | 278.83 | 324.17 | 278.68 | 0.00 | 0.00 | 278.83 | 0.00 | 278.68 |
|  |  | n.at.z | 336.03 | 338.50 | 343.70 | 334.71 | 347.17 | 0.00 | 0.00 | 343.70 | 0.00 | 347.17 |
| $\square$ NMFS M | $\square$ index catch | abundance | 289.94 | 261.72 | 240.01 | 280.35 | 234.33 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  | biomass | 151.12 | 131.45 | 141.07 | 147.12 | 136.47 | 0.00 | 0.00 | 141.07 | 0.00 | 136.47 |
|  |  | n.at.z | 465.57 | 460.96 | 449.00 | 466.11 | 455.33 | 0.00 | 0.00 | 449.00 | 0.00 | 455.33 |
| $\square$ SBS BSFRF females | $\square$ index catch | abundance |  |  |  | 9.12 | 7.70 |  |  |  | 0.00 | 0.00 |
|  |  | biomass |  |  |  | 2.86 | 2.88 |  |  |  | 2.86 | 2.88 |
|  |  | n.at.z |  |  |  | 45.54 | 45.09 |  |  |  | 45.54 | 45.09 |
| GSBS BSFRF males | ■index catch | abundance |  |  |  | 10.44 | 8.65 |  |  |  | 0.00 | 0.00 |
|  |  | biomass |  |  |  | 1.99 | 2.24 |  |  |  | 1.99 | 2.24 |
|  |  | n.at.z |  |  |  | 48.87 | 48.93 |  |  |  | 48.87 | 48.93 |
| $\square$ SBS NMFS females | $\square$ index catch | abundance |  |  |  | 7.92 | 8.40 |  |  |  | 0.00 | 0.00 |
|  |  | biomass |  |  |  | 9.72 | 9.86 |  |  |  | 9.72 | 9.86 |
|  |  | n.at.z |  |  |  | 23.26 | 22.53 |  |  |  | 23.26 | 22.53 |
| $\square$ SBS NMFS males | $\square$ index catch | abundance |  |  |  | 2.91 | 3.76 |  |  |  | 0.00 | 0.00 |
|  |  | biomass |  |  |  | 3.67 | 4.62 |  |  |  | 3.67 | 4.62 |
|  |  | n.at.z |  |  |  | 30.00 | 30.15 |  |  |  | 30.00 | 30.15 |

Table 36. (Unweighted) negative log-likelihoods and (weighted) objective function values for fits to growth (molt increment) and male maturity ogive data components from the model scenarios.

| category | NLLs |  | M19F03 | objective function values |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | M19F01 | M19F02 |  | M19F04 | M19F05 | M19F01 | M19F02 | M19F03 | M19F04 | M19F05 |
| growth data | 493.63 | 524.81 | 539.86 | 490.63 | 540.15 | 493.63 | 524.81 | 539.86 | 490.63 | 540.15 |
| maturity ogive data | 215.21 | 116.53 | 95.42 | 210.40 | 95.61 | 0.00 | 116.53 | 95.42 | 0.00 | 95.61 |

Table 37. Root mean square errors (RMSE) for fishery-related data components from the model scenarios. TCF: directed Tanner crab fishery; SCF: snow crab fishery; RKF: BBRKC fishery; GTF: groundfish fisheries. Rows consisting of all zero values indicate a data component which was not included in any of the models.

| fleet | catch.type | data.type | x | M19F01 | M19F02 | M19F03 | M19F04 | M19F05 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| - GTF | $\square$ total catch | Eabundance | all sexes | 0.10 | 0.10 | 0.11 | 0.10 | 0.11 |
|  |  | $\square$ biomass | all sexes | 0.23 | 0.24 | 0.25 | 0.23 | 0.25 |
|  |  | En.at.z | female | 371.61 | 384.39 | 388.75 | 372.54 | 391.04 |
|  |  |  | male | 373.16 | 358.55 | 369.45 | 371.28 | 366.79 |
| - RKF | $\square$ total catch | Eabundance | female | 40.31 | 43.08 | 45.92 | 37.48 | 47.33 |
|  |  |  | male | 25.70 | 25.09 | 25.51 | 25.65 | 25.47 |
|  |  | $\square$ biomass | female | 0.02 | 0.02 | 0.02 | 0.02 | 0.02 |
|  |  |  | male | 0.30 | 0.30 | 0.32 | 0.30 | 0.32 |
|  |  | \#n.at.z | female | 48.67 | 48.79 | 54.10 | 48.20 | 52.53 |
|  |  |  | male | 56.41 | 56.51 | 55.96 | 56.25 | 55.74 |
| ■SCF | $\square$ total catch | Eabundance | female | 4.29 | 4.32 | 5.04 | 4.40 | 5.01 |
|  |  |  | male | 2.63 | 2.65 | 2.58 | 2.61 | 2.57 |
|  |  | Ebiomass | female | 0.08 | 0.08 | 0.08 | 0.08 | 0.08 |
|  |  |  | male | 0.28 | 0.27 | 0.25 | 0.28 | 0.25 |
|  |  | En.at.z | female | 54.82 | 55.60 | 54.83 | 50.76 | 51.28 |
|  |  |  | male | 227.18 | 232.72 | 235.32 | 224.44 | 235.23 |
| $\square$ TCF | - retained catch | Elabundance | female | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  |  | male | 3.37 | 3.35 | 3.30 | 3.37 | 3.31 |
|  |  | $\square$ biomass | female | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  |  | male | 0.15 | 0.14 | 0.14 | 0.15 | 0.14 |
|  |  | \#n.at.z | male | 355.25 | 367.67 | 384.72 | 354.36 | 381.60 |
|  | - total catch | Eabundance | female | 40.84 | 40.88 | 46.70 | 41.59 | 44.71 |
|  |  |  | male | 1.16 | 1.13 | 1.08 | 1.15 | 1.08 |
|  |  | Ebiomass | female | 0.26 | 0.25 | 0.25 | 0.26 | 0.25 |
|  |  |  | male | 0.16 | 0.16 | 0.15 | 0.16 | 0.15 |
|  |  | En.at.z | female | 182.37 | 188.63 | 208.21 | 194.75 | 231.22 |
|  |  |  | male | 499.38 | 511.41 | 513.56 | 514.17 | 520.01 |

Table 38. Average root mean square errors (RMSE) for survey-related data components from the model scenarios. Rows consisting of all zero values indicate a data component which was not included in any of the models. Blank cells indicate a data component (row) that was not included in the likelihood in the associated scenario (column).

| fleet | catch.type | data.type | x | M19F01 | M19F02 | M19F03 | M19F04 | M19F05 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $\square$ NMFS 0 | $\square$ index catch | ■abundance | female | 2.92 | 2.86 | 2.69 | 2.90 | 2.69 |
|  |  |  | male | 3.12 | 2.99 | 3.05 | 3.06 | 3.02 |
|  |  | $\square$ biomass | female | 2.32 | 2.29 | 2.30 | 2.34 | 2.31 |
|  |  |  | male | 2.40 | 2.27 | 2.56 | 2.38 | 2.54 |
|  |  | - n.at.z | female | 373.93 | 385.98 | 356.45 | 364.22 | 344.26 |
|  |  |  | male | 487.07 | 499.49 | 490.64 | 477.53 | 480.54 |
| $\square$ NMFS F | $\square$ index catch | Eabundance | female | 2.92 | 2.86 | 2.69 | 2.90 | 2.69 |
|  |  |  | male | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  | $\square$ biomass | female | 2.65 | 2.60 | 2.48 | 2.66 | 2.48 |
|  |  |  | male | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  | $\square$ n.at.z | female | 125.86 | 125.69 | 136.02 | 125.48 | 133.02 |
| $\square$ NMFS M | $\square$ index catch | $\square$ abundance | female | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  |  | male | 3.59 | 3.41 | 3.27 | 3.53 | 3.23 |
|  |  | $\square$ biomass | female | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
|  |  |  | male | 2.59 | 2.42 | 2.50 | 2.56 | 2.46 |
|  |  | En.at.z | male | 195.51 | 188.89 | 185.98 | 192.03 | 183.02 |
| -SBS BSFRF females | Bindex catch | Elabundance | female |  |  |  | 1.31 | 1.23 |
|  |  |  | male |  |  |  | 0.00 | 0.00 |
|  |  | $\square$ biomass | female |  |  |  | 0.76 | 0.75 |
|  |  |  | male |  |  |  | 0.00 | 0.00 |
|  |  | - n.at.z | female |  |  |  | 44.41 | 46.72 |
| $\square$ SBS BSFRF males | ■index catch | Eabundance | female |  |  |  | 0.00 | 0.00 |
|  |  |  | male |  |  |  | 2.04 | 1.86 |
|  |  | $\square$ biomass | female |  |  |  | 0.00 | 0.00 |
|  |  |  | male |  |  |  | 0.89 | 0.95 |
|  |  | Gn.at.z | male |  |  |  | 191.47 | 198.80 |
| $\square$ SBS NMFS females | $\square$ index catch | $\square$ abundance | female |  |  |  | 1.17 | 1.22 |
|  |  |  | male |  |  |  | 0.00 | 0.00 |
|  |  | $\square$ biomass | female |  |  |  | 1.25 | 1.28 |
|  |  |  | male |  |  |  | 0.00 | 0.00 |
|  |  | $\square \mathrm{n} .9 \mathrm{t} . \mathrm{z}$ | female |  |  |  | 48.92 | 53.02 |
| $\square$ SBS NMFS males | $\square$ index catch | Eabundance | female |  |  |  | 0.00 | 0.00 |
|  |  |  | male |  |  |  | 1.08 | 1.23 |
|  |  | $\square$ biomass | female |  |  |  | 0.00 | 0.00 |
|  |  |  | male |  |  |  | 1.21 | 1.36 |
|  |  | En.at.z | male |  |  |  | 316.65 | 303.44 |

Table 39. Root mean square errors (RMSE) for fits to growth (molt increment) and male maturity ogive data components from the model scenarios.

| category | M19F01 | M19F02 | M19F03 | M19F04 | M19F05 |
| :--- | ---: | ---: | ---: | ---: | ---: |
| growth data | 0.42 | 0.42 | 0.43 | 0.42 | 0.43 |
| maturity ogive data | 2.51 | 1.92 | 1.77 | 2.48 | 1.80 |

Table 40. Effective sample sizes used for NMFS 0 EBS trawl survey size composition data for the 2018 assessment model (M19F00) and the author's preferred model (M19F03). Effective sample sizes were estimated using the McAllister-Ianelli approach. Note that, while effective N's were calculated for this dataset in MF 1903, it was not included in the model objective function (the weight in the likelihood was set to 0 ). Input sample sizes were set at 200.

| Sum of val year | Column Labels M19F00 | M19F03 | Sum of val year | Column Labels M19F00 | M19F03 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1975 | 700.312702 | 479.8666701 | 1996 | 1935.626324 | 1479.261951 |
| 1976 | 835.8906704 | 679.480725 | 1997 | 1891.598819 | 1523.133367 |
| 1977 | 874.9223597 | 775.5274286 | 1998 | 2046.203568 | 1345.547918 |
| 1978 | 892.441064 | 1415.818871 | 1999 | 1158.80325 | 1541.437445 |
| 1979 | 1130.270061 | 802.0601696 | 2000 | 1589.385175 | 2628.510003 |
| 1980 | 1441.838602 | 1103.810405 | 2001 | 2168.1765 | 1789.809452 |
| 1981 | 1138.908258 | 723.7818753 | 2002 | 1943.798287 | 2242.190544 |
| 1982 | 518.8477363 | 480.801695 | 2003 | 1488.112154 | 2703.663841 |
| 1983 | 1067.859284 | 866.8946961 | 2004 | 978.9173627 | 1103.316885 |
| 1984 | 572.9407661 | 790.9737623 | 2005 | 3262.607163 | 4249.657685 |
| 1985 | 326.2645986 | 386.0794348 | 2006 | 1505.176736 | 2452.948118 |
| 1986 | 676.7917083 | 818.0047904 | 2007 | 1294.785121 | 1506.294676 |
| 1987 | 789.3102243 | 1520.471983 | 2008 | 2318.550309 | 2770.433117 |
| 1988 | 1107.233577 | 1722.63342 | 2009 | 1414.661594 | 2372.096875 |
| 1989 | 2579.165029 | 1673.448147 | 2010 | 12011.00017 | 4232.237577 |
| 1990 | 2756.786708 | 2063.757876 | 2011 | 1806.553577 | 2278.879216 |
| 1991 | 3162.992353 | 1499.878515 | 2012 | 1476.147611 | 1820.248354 |
| 1992 | 2697.685485 | 2936.358538 | 2013 | 2662.685394 | 2493.220045 |
| 1993 | 1972.898268 | 1429.015184 | 2014 | 1191.826672 | 1135.930166 |
| 1994 | 1603.111983 | 1219.109801 | 2015 | 2445.230566 | 1933.403136 |
| 1995 | 1758.283681 | 1208.444231 | 2016 | 1168.110952 | 1004.934831 |
|  |  |  | 2017 | 1151.365149 | 1026.282821 |
|  |  |  | 2018 | 2277.011147 | 1905.751709 |
|  |  |  | 2019 |  | 4102.775173 |

Table 41. Effective sample sizes used for retained catch size composition data from the directed fishery for the 2018 assessment model (M19F00) and the author's preferred model (M19F03). Effective sample sizes were estimated using the McAllister-Ianelli approach.

| year | $\square \text { M19F00 }$ <br> effective $\mathbf{N}$ | input ss | $\square \text { M19F03 }$ <br> effective $\mathbf{N}$ | input ss |
| :---: | :---: | :---: | :---: | :---: |
| 1980 | 24.8 | 97.8 | 24.8 | 104.6 |
| 1981 | 1466.9 | 83.1 | 476.1 | 88.9 |
| 1982 | 1992.2 | 99.3 | 1097.8 | 106.2 |
| 1983 | 52.4 | 12.3 | 55.6 | 13.2 |
| 1984 | 426.3 | 18.7 | 203.8 | 20.0 |
| 1988 | 128.0 | 91.0 | 142.5 | 97.3 |
| 1989 | 1429.7 | 30.3 | 413.9 | 32.4 |
| 1990 | 256.1 | 200.0 | 242.1 | 200.0 |
| 1991 | 144.3 | 200.0 | 903.1 | 200.0 |
| 1992 | 99.0 | 200.0 | 313.2 | 200.0 |
| 1993 | 131.3 | 200.0 | 599.2 | 200.0 |
| 1994 | 145.3 | 200.0 | 273.7 | 200.0 |
| 1995 | 175.6 | 11.2 | 307.6 | 12.0 |
| 1996 | 172.8 | 32.6 | 1951.4 | 34.8 |
| 2005 | 14.4 | 5.2 | 18.3 | 5.5 |
| 2006 | 301.0 | 21.6 | 120.6 | 23.1 |
| 2007 | 1641.2 | 51.0 | 224.3 | 45.2 |
| 2008 | 972.8 | 25.6 | 402.8 | 27.4 |
| 2009 | 128.9 | 17.8 | 126.5 | 19.0 |
| 2013 | 770.9 | 35.0 | 581.1 | 35.8 |
| 2014 | 219.2 | 103.3 | 285.1 | 113.7 |
| 2015 | 164.3 | 200.0 | 263.6 | 190.3 |
| 2017 | 104.0 | 25.5 | 132.4 | 27.3 |
| 2018 |  |  | 73.8 | 26.0 |

Table 42. Effective sample sizes used for total catch size composition data from the directed fishery for the 2018 assessment model (M19F00) and the author's preferred model (M19F03). Effective sample sizes were estimated using the McAllister-Ianelli approach.

| year | $\begin{aligned} & \text { M19F00 } \\ & \text { female } \\ & \text { effective } \mathrm{N} \end{aligned}$ | input ss | $\square$ male <br> effective $\mathbf{N}$ | input ss | $\begin{aligned} & \square \text { M19F03 } \\ & \text { female } \\ & \text { effective } \mathbf{N} \end{aligned}$ | input ss | $\square$ male <br> effective $\mathbf{N}$ | input ss |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1991 | 421.1 | 41.2 | 1343.4 | 200.0 | 245.5 | 44.0 | 499.5 | 200.0 |
| 1992 | 555.0 | 64.3 | 121.7 | 200.0 | 1450.9 | 68.8 | 363.3 | 200.0 |
| 1993 | 307.9 | 76.9 | 267.9 | 200.0 | 232.8 | 82.3 | 270.1 | 200.0 |
| 1994 | 62.7 | 15.7 | 549.0 | 42.6 | 81.4 | 16.7 | 1044.4 | 45.5 |
| 1995 | 100.7 | 22.9 | 310.4 | 41.1 | 136.0 | 24.5 | 285.2 | 43.9 |
| 1996 | 249.3 | 2.5 | 31.3 | 5.0 | 171.1 | 1.3 | 22.3 | 2.8 |
| 2005 | 41.7 | 8.1 | 99.4 | 144.9 | 48.2 | 8.7 | 118.1 | 154.9 |
| 2006 | 442.5 | 32.6 | 285.3 | 178.0 | 341.9 | 34.8 | 330.7 | 190.3 |
| 2007 | 302.4 | 24.4 | 394.4 | 200.0 | 231.5 | 26.1 | 560.6 | 200.0 |
| 2008 | 46.3 | 4.7 | 1149.5 | 200.0 | 45.1 | 5.1 | 1250.6 | 200.0 |
| 2009 | 23.6 | 1.1 | 162.5 | 127.0 | 21.5 | 1.2 | 168.1 | 135.8 |
| 2013 | 59.7 | 5.2 | 1475.0 | 127.0 | 44.7 | 5.6 | 2528.6 | 135.8 |
| 2014 | 175.6 | 8.8 | 210.5 | 200.0 | 126.6 | 9.4 | 248.3 | 200.0 |
| 2015 | 75.3 | 11.9 | 133.0 | 200.0 | 81.9 | 12.8 | 189.4 | 200.0 |
| 2017 | 52.1 | 12.6 | 168.4 | 138.0 | 58.5 | 13.5 | 243.4 | 147.6 |
| 2018 |  |  |  |  | 13.7 | 16.0 | 94.3 | 200.0 |

Table 43. Effective sample sizes used for bycatch size composition data from the snow crab fishery for the 2018 assessment model (M19F00) and the author's preferred model (M19F03). Effective sample sizes were estimated using the McAllister-Ianelli approach.

| year | ```GM19F00 Gemale effective N``` | input ss | $\square$ male effective $\mathbf{N}$ | input ss | ```GM19F03 female effective N``` | input ss | $\square$ male effective $\mathbf{N}$ | input ss |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1990 |  |  |  |  | 38.9 | 3.8 | 42.9 | 110.2 |
| 1991 |  |  |  |  | 22.7 | 5.4 | 86.1 | 92.0 |
| 1992 | 18.3 | 6.3 | 186.3 | 46.1 | 24.0 | 6.7 | 29.7 | 49.3 |
| 1993 | 30.7 | 11.3 | 117.4 | 51.2 | 38.0 | 12.1 | 28.6 | 54.7 |
| 1994 | 40.7 | 11.2 | 37.3 | 21.9 | 30.0 | 12.0 | 13.9 | 23.4 |
| 1995 | 42.1 | 3.1 | 86.7 | 13.9 | 40.5 | 3.4 | 26.2 | 14.9 |
| 1996 | 46.2 | 4.9 | 289.1 | 24.0 | 73.2 | 5.2 | 104.5 | 25.6 |
| 1997 | 111.8 | 4.8 | 449.8 | 29.2 | 106.4 | 5.2 | 390.0 | 31.2 |
| 1998 | 21.5 | 2.4 | 1131.3 | 14.0 | 21.7 | 2.5 | 546.4 | 15.0 |
| 1999 | 30.3 | 0.6 | 132.8 | 7.2 | 32.1 | 0.6 | 128.9 | 7.7 |
| 2000 | 30.6 | 0.5 | 285.3 | 9.1 | 34.3 | 0.6 | 253.4 | 9.7 |
| 2001 | 121.8 | 1.2 | 565.8 | 22.9 | 132.8 | 1.3 | 436.8 | 24.5 |
| 2002 | 45.6 | 0.9 | 59.8 | 7.2 | 47.2 | 0.9 | 66.1 | 7.7 |
| 2003 | 45.1 | 1.1 | 110.1 | 5.1 | 45.7 | 1.2 | 130.5 | 5.4 |
| 2004 | 30.7 | 5.2 | 23.1 | 6.2 | 30.5 | 5.6 | 23.7 | 6.5 |
| 2005 | 154.2 | 2.7 | 123.0 | 72.0 | 75.2 | 2.9 | 134.4 | 77.0 |
| 2006 | 49.9 | 9.2 | 76.5 | 76.4 | 30.4 | 9.9 | 77.1 | 81.6 |
| 2007 | 44.2 | 5.3 | 384.9 | 101.4 | 27.3 | 5.7 | 421.7 | 108.4 |
| 2008 | 15.0 | 5.3 | 97.0 | 62.1 | 20.2 | 5.7 | 102.0 | 66.4 |
| 2009 | 21.2 | 3.5 | 470.9 | 81.2 | 33.1 | 3.7 | 449.5 | 86.9 |
| 2010 | 76.4 | 1.8 | 382.8 | 88.7 | 91.7 | 2.0 | 279.0 | 94.8 |
| 2011 | 62.1 | 1.4 | 228.2 | 69.5 | 58.9 | 1.5 | 183.2 | 74.3 |
| 2012 | 47.3 | 1.4 | 209.1 | 53.9 | 78.5 | 2.1 | 153.3 | 86.4 |
| 2013 | 203.9 | 2.6 | 248.0 | 95.0 | 117.9 | 2.8 | 216.7 | 101.6 |
| 2014 | 67.5 | 5.9 | 532.0 | 182.8 | 141.0 | 6.3 | 402.6 | 195.4 |
| 2015 | 107.8 | 1.7 | 520.2 | 146.5 | 56.6 | 1.8 | 354.9 | 155.9 |
| 2016 | 112.9 | 1.7 | 468.7 | 142.8 | 28.9 | 2.1 | 844.4 | 128.6 |
| 2017 | 63.6 | 0.8 | 709.0 | 41.1 | 96.2 | 0.9 | 491.4 | 44.0 |
| 2018 |  |  |  |  | 16.2 | 1.8 | 406.7 | 48.3 |

Table 44. Effective sample sizes used for bycatch size composition data from the BBRKC fishery for the 2018 assessment model (M19F00) and the author's preferred model (M19F03. Effective sample sizes were estimated using the McAllister-Ianelli approach.

| year | $\begin{aligned} & \text { M19F00 } \\ & \text { female } \\ & \text { effective } \mathrm{N} \end{aligned}$ | input ss | $\square$ male <br> effective $\mathbf{N}$ | input ss | $\begin{aligned} & \square \text { M19F03 } \\ & \text { female } \\ & \text { effective } \mathbf{N} \end{aligned}$ | input ss | $\square$ male effective $\mathbf{N}$ | input ss |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1990 |  |  |  |  | 42.5 | 0.3 | 12.6 | 12.4 |
| 1991 |  |  |  |  | 91.0 | 0.7 | 16.0 | 17.9 |
| 1992 | 83.1 | 0.8 | 33.2 | 15.1 | 82.5 | 0.8 | 23.1 | 16.2 |
| 1993 | 275.0 | 8.8 | 32.9 | 54.1 | 319.6 | 9.4 | 24.5 | 57.8 |
| 1996 | 3.4 | 0.0 | 12.6 | 0.8 |  |  |  |  |
| 1997 | 25.2 | 0.3 | 19.6 | 7.6 | 27.8 | 0.3 | 27.4 | 8.1 |
| 1998 | 21.0 | 0.1 | 55.7 | 3.4 | 21.6 | 0.2 | 83.0 | 3.6 |
| 1999 | 17.5 | 0.1 | 51.2 | 1.5 | 18.3 | 0.1 | 41.9 | 1.6 |
| 2000 | 40.5 | 0.3 | 134.6 | 6.2 | 41.1 | 0.3 | 92.2 | 6.6 |
| 2001 | 51.1 | 0.3 | 113.6 | 3.4 | 50.0 | 0.3 | 69.8 | 3.6 |
| 2002 | 35.5 | 0.4 | 87.3 | 5.5 | 36.7 | 0.4 | 60.3 | 5.9 |
| 2003 | 53.3 | 0.3 | 58.2 | 4.1 | 52.7 | 0.4 | 42.5 | 4.4 |
| 2004 | 20.3 | 0.3 | 31.5 | 3.6 | 21.0 | 0.3 | 24.4 | 3.8 |
| 2005 | 12.6 | 0.5 | 44.3 | 7.2 | 14.1 | 0.6 | 34.5 | 7.7 |
| 2006 | 23.8 | 0.6 | 22.6 | 5.9 | 28.4 | 0.5 | 19.0 | 5.9 |
| 2007 | 102.5 | 0.7 | 91.4 | 10.3 | 91.7 | 0.7 | 71.7 | 10.7 |
| 2008 | 91.8 | 0.9 | 62.5 | 27.9 | 108.5 | 1.0 | 81.2 | 29.8 |
| 2009 | 109.0 | 0.5 | 19.3 | 24.9 | 116.8 | 0.6 | 22.5 | 22.6 |
| 2010 | 35.9 | 0.2 | 51.3 | 4.4 | 52.2 | 0.2 | 43.1 | 4.6 |
| 2011 | 6.0 | 0.0 | 68.6 | 2.5 | 5.8 | 0.0 | 50.9 | 2.5 |
| 2012 | 6.9 | 0.4 | 66.0 | 4.5 | 7.2 | 0.4 | 48.6 | 4.9 |
| 2013 | 9.7 | 0.4 | 86.1 | 15.5 | 9.6 | 0.5 | 110.8 | 16.6 |
| 2014 | 19.3 | 0.2 | 155.1 | 22.9 | 19.8 | 0.3 | 169.1 | 24.4 |
| 2015 | 86.3 | 1.3 | 195.1 | 16.1 | 89.9 | 1.5 | 119.9 | 17.1 |
| 2016 | 18.9 | 1.8 | 25.1 | 22.5 | 19.6 | 1.9 | 21.4 | 25.3 |
| 2017 | 34.0 | 0.6 | 76.0 | 27.8 | 32.7 | 0.7 | 55.7 | 29.7 |
| 2018 |  |  |  |  | 5.5 | 0.0 | 89.1 | 10.1 |

Table 45. Effective sample sizes used for bycatch size composition data from the groundfish fisheries for the 2018 assessment model (M19F00) and the author's preferred model (M19F03). Effective sample sizes were estimated using the McAllister-Ianelli approach.

| year | M19F00 <br> female effective $\mathbf{N}$ | input ss | male effective $\mathbf{N}$ | input ss | M19F03 <br> female effective $\mathbf{N}$ | input ss | male effective $\mathbf{N}$ | input ss |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1973 | 226.8 | 39.9 | 357.5 | 39.9 | 220.4 | 39.9 | 269.0 | 39.9 |
| 1974 | 209.7 | 30.1 | 726.4 | 30.1 | 220.2 | 30.1 | 470.6 | 30.1 |
| 1975 | 195.0 | 15.4 | 334.1 | 15.4 | 230.7 | 15.4 | 254.9 | 15.4 |
| 1976 | 107.3 | 100.2 | 178.4 | 100.2 | 114.1 | 100.2 | 125.7 | 100.2 |
| 1977 | 327.3 | 140.1 | 233.1 | 140.1 | 312.4 | 140.1 | 210.2 | 140.1 |
| 1978 | 193.1 | 237.1 | 249.6 | 237.1 | 175.7 | 237.1 | 239.9 | 237.1 |
| 1979 | 889.5 | 223.5 | 594.2 | 223.5 | 776.5 | 223.5 | 763.7 | 223.5 |
| 1980 | 419.3 | 137.6 | 1045.8 | 137.6 | 817.6 | 137.6 | 704.2 | 137.6 |
| 1981 | 56.1 | 74.7 | 1050.0 | 74.7 | 67.3 | 74.7 | 791.0 | 74.7 |
| 1982 | 62.1 | 157.6 | 529.6 | 157.6 | 90.6 | 157.6 | 509.7 | 157.6 |
| 1983 | 134.6 | 196.0 | 345.6 | 196.0 | 293.9 | 196.0 | 401.5 | 196.0 |
| 1984 | 235.0 | 301.2 | 354.6 | 301.2 | 482.9 | 301.2 | 679.8 | 301.2 |
| 1985 | 278.0 | 263.5 | 169.9 | 263.5 | 274.3 | 263.5 | 239.7 | 263.5 |
| 1986 | 193.5 | 165.2 | 281.7 | 165.2 | 155.0 | 165.2 | 405.1 | 165.2 |
| 1987 | 671.1 | 289.3 | 266.3 | 289.3 | 718.7 | 289.3 | 282.6 | 289.3 |
| 1988 | 224.1 | 130.2 | 404.9 | 130.2 | 218.5 | 130.2 | 339.5 | 130.2 |
| 1989 | 595.1 | 400.0 | 810.5 | 400.0 | 906.0 | 400.0 | 747.2 | 400.0 |
| 1990 | 308.5 | 255.4 | 997.0 | 255.4 | 349.5 | 255.4 | 953.5 | 255.4 |
| 1991 | 186.1 | 75.7 | 330.4 | 75.7 | 213.5 | 80.9 | 316.1 | 80.9 |
| 1992 | 63.6 | 31.6 | 177.7 | 31.6 | 68.4 | 33.8 | 166.4 | 33.8 |
| 1993 | 93.8 | 11.6 | 77.8 | 11.6 | 108.2 | 12.4 | 72.8 | 12.4 |
| 1994 | 429.9 | 40.0 | 238.3 | 40.0 | 442.7 | 42.8 | 236.5 | 42.8 |
| 1995 | 60.2 | 48.3 | 58.2 | 48.3 | 65.5 | 51.6 | 52.2 | 51.6 |
| 1996 | 597.2 | 86.0 | 176.8 | 86.0 | 512.7 | 92.0 | 158.0 | 92.0 |
| 1997 | 184.6 | 101.8 | 49.5 | 101.8 | 137.2 | 108.8 | 43.0 | 108.8 |
| 1998 | 303.0 | 121.6 | 119.1 | 121.6 | 182.3 | 130.0 | 93.1 | 130.0 |
| 1999 | 1011.6 | 114.4 | 441.8 | 114.4 | 569.3 | 122.4 | 288.5 | 122.4 |
| 2000 | 899.8 | 117.4 | 556.9 | 117.4 | 638.7 | 125.6 | 338.9 | 125.6 |
| 2001 | 1246.6 | 138.7 | 775.7 | 138.7 | 1297.3 | 148.2 | 523.1 | 148.2 |
| 2002 | 891.3 | 137.0 | 429.6 | 137.0 | 736.2 | 146.5 | 391.5 | 146.5 |
| 2003 | 300.1 | 90.4 | 196.9 | 90.4 | 307.9 | 96.7 | 196.2 | 96.7 |
| 2004 | 30.3 | 134.5 | 110.2 | 134.5 | 32.8 | 143.8 | 119.3 | 143.8 |
| 2005 | 1814.9 | 157.9 | 1545.9 | 157.9 | 1652.0 | 168.9 | 1226.6 | 168.9 |
| 2006 | 134.6 | 139.2 | 182.0 | 139.2 | 151.8 | 148.9 | 197.8 | 148.9 |
| 2007 | 106.0 | 146.7 | 187.6 | 146.7 | 117.7 | 156.9 | 199.2 | 156.9 |
| 2008 | 164.5 | 223.4 | 184.2 | 223.4 | 182.4 | 233.3 | 172.7 | 233.3 |
| 2009 | 536.6 | 160.0 | 313.3 | 160.0 | 524.9 | 171.1 | 345.8 | 171.1 |
| 2010 | 2097.5 | 127.9 | 628.5 | 127.9 | 1362.2 | 136.7 | 719.2 | 136.7 |
| 2011 | 66.8 | 149.6 | 83.2 | 149.6 | 61.7 | 160.0 | 71.3 | 160.0 |
| 2012 | 102.6 | 118.1 | 412.4 | 118.1 | 112.1 | 126.2 | 426.6 | 126.2 |
| 2013 | 433.7 | 244.6 | 359.5 | 244.6 | 567.0 | 247.6 | 314.9 | 247.6 |
| 2014 | 794.9 | 231.0 | 1037.7 | 231.0 | 741.6 | 233.1 | 1105.9 | 233.1 |
| 2015 | 203.2 | 242.1 | 219.3 | 242.1 | 250.8 | 245.1 | 232.4 | 245.1 |
| 2016 | 56.9 | 166.2 | 229.2 | 166.2 | 57.4 | 177.6 | 244.0 | 177.6 |
| 2017 | 173.8 | 98.6 | 80.6 | 98.6 | 149.8 | 108.2 | 75.1 | 108.2 |
| 2018 |  |  |  |  | 214.1 | 64.7 | 279.5 | 64.7 |

Table 46. Comparison of fits to mature survey biomass by sex (in 1000's t) from the 2018 assessment model (M19F00) and the author's preferred model (M19F03).

| year | $\begin{aligned} & \square \text { M19F00 } \\ & \square \text { female } \\ & \text { observed } \end{aligned}$ | predicted | $\square$ male observed | predicted | $\begin{aligned} & \text { M19F03 } \\ & \text { - female } \\ & \text { observed } \end{aligned}$ | predicted | Emale observed | predicted |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1975 | 72.4 | 102.8 | 540.9 | 333.2 | 72.4 | 91.5 | 540.9 | 381.9 |
| 1976 | 68.7 | 91.6 | 283.3 | 295.7 | 68.7 | 80.3 | 283.3 | 328.2 |
| 1977 | 91.6 | 82.8 | 249.8 | 241.9 | 91.6 | 70.6 | 249.8 | 261.2 |
| 1978 | 58.3 | 80.2 | 176.2 | 192.1 | 58.3 | 64.8 | 176.2 | 201.2 |
| 1979 | 42.5 | 84.2 | 82.7 | 183.6 | 42.5 | 65.2 | 82.7 | 187.0 |
| 1980 | 141.1 | 87.3 | 239.3 | 186.2 | 141.1 | 68.5 | 239.3 | 201.0 |
| 1981 | 86.7 | 77.5 | 130.2 | 156.7 | 86.7 | 56.4 | 130.2 | 180.4 |
| 1982 | 130.0 | 56.1 | 117.5 | 160.0 | 130.0 | 43.7 | 117.5 | 161.6 |
| 1983 | 43.1 | 43.9 | 67.9 | 117.2 | 43.1 | 31.7 | 67.9 | 120.2 |
| 1984 | 32.1 | 35.4 | 56.3 | 83.5 | 32.1 | 23.8 | 56.3 | 85.0 |
| 1985 | 12.1 | 31.4 | 26.8 | 65.2 | 12.1 | 20.5 | 26.8 | 65.0 |
| 1986 | 9.4 | 33.5 | 34.9 | 81.9 | 9.4 | 23.9 | 34.9 | 77.6 |
| 1987 | 25.3 | 38.0 | 70.1 | 108.6 | 25.3 | 28.7 | 70.1 | 98.2 |
| 1988 | 60.9 | 42.4 | 160.2 | 139.6 | 60.9 | 33.5 | 160.2 | 122.5 |
| 1989 | 50.6 | 45.6 | 226.1 | 165.9 | 50.6 | 37.0 | 226.1 | 143.3 |
| 1990 | 85.2 | 46.2 | 230.3 | 172.2 | 85.2 | 38.8 | 230.3 | 151.2 |
| 1991 | 96.5 | 43.5 | 258.4 | 153.9 | 96.5 | 38.0 | 258.4 | 144.0 |
| 1992 | 54.4 | 37.7 | 233.1 | 132.8 | 54.4 | 34.5 | 233.1 | 134.1 |
| 1993 | 24.3 | 30.1 | 135.3 | 99.0 | 24.3 | 29.0 | 135.3 | 108.8 |
| 1994 | 20.9 | 23.2 | 92.2 | 72.9 | 20.9 | 23.3 | 92.2 | 86.7 |
| 1995 | 25.8 | 17.9 | 67.7 | 53.9 | 25.8 | 18.6 | 67.7 | 67.4 |
| 1996 | 20.6 | 14.1 | 58.3 | 40.7 | 20.6 | 15.1 | 58.3 | 53.3 |
| 1997 | 8.2 | 11.8 | 25.9 | 34.7 | 8.2 | 13.0 | 25.9 | 44.9 |
| 1998 | 6.5 | 10.5 | 25.9 | 32.0 | 6.5 | 11.7 | 25.9 | 40.0 |
| 1999 | 10.5 | 10.4 | 34.5 | 32.2 | 10.5 | 11.6 | 34.5 | 39.4 |
| 2000 | 10.7 | 11.0 | 40.2 | 35.9 | 10.7 | 12.2 | 40.2 | 42.1 |
| 2001 | 15.4 | 12.7 | 47.9 | 43.4 | 15.4 | 13.9 | 47.9 | 48.3 |
| 2002 | 14.4 | 14.4 | 46.4 | 51.8 | 14.4 | 15.7 | 46.4 | 56.3 |
| 2003 | 21.5 | 17.0 | 62.4 | 62.7 | 21.5 | 18.7 | 62.4 | 67.7 |
| 2004 | 13.5 | 20.2 | 65.9 | 77.0 | 13.5 | 22.3 | 65.9 | 82.4 |
| 2005 | 33.5 | 22.7 | 108.9 | 92.9 | 33.5 | 25.3 | 108.9 | 98.7 |
| 2006 | 43.1 | 24.5 | 169.4 | 106.8 | 43.1 | 27.9 | 169.4 | 114.2 |
| 2007 | 32.5 | 25.5 | 173.7 | 117.0 | 32.5 | 29.4 | 173.7 | 127.1 |
| 2008 | 26.2 | 24.6 | 150.0 | 124.6 | 26.2 | 28.8 | 150.0 | 135.2 |
| 2009 | 19.5 | 22.9 | 85.6 | 122.2 | 19.5 | 27.0 | 85.6 | 130.3 |
| 2010 | 14.7 | 21.8 | 88.1 | 110.2 | 14.7 | 25.3 | 88.1 | 115.4 |
| 2011 | 21.2 | 22.4 | 105.9 | 101.8 | 21.2 | 25.2 | 105.9 | 104.6 |
| 2012 | 36.4 | 25.5 | 122.3 | 105.2 | 36.4 | 27.8 | 122.3 | 107.7 |
| 2013 | 42.1 | 29.4 | 170.4 | 126.9 | 42.1 | 31.5 | 170.4 | 128.6 |
| 2014 | 32.2 | 30.1 | 191.3 | 150.1 | 32.2 | 32.2 | 191.3 | 148.9 |
| 2015 | 24.1 | 26.9 | 137.2 | 144.4 | 24.1 | 29.0 | 137.2 | 142.2 |
| 2016 | 16.4 | 22.5 | 131.2 | 118.1 | 16.4 | 24.4 | 131.2 | 116.2 |
| 2017 | 15.6 | 19.8 | 104.5 | 102.7 | 15.6 | 20.9 | 104.5 | 97.4 |
| 2018 | 14.9 | 19.0 | 86.8 | 90.4 | 14.9 | 19.0 | 86.8 | 82.8 |
| 2019 |  |  |  |  | 14.6 | 19.7 | 48.8 | 75.0 |

Table 47. Comparison of estimates of mature biomass-at-mating by sex (in 1000's t) from the 2018 assessment model (M19F00) and the author's preferred model (M19F03).

|  | $\begin{aligned} & \text { M19F00 } \\ & \text { female } \end{aligned}$ | male | $\begin{aligned} & \text { M19F03 } \\ & \text { female } \end{aligned}$ | male | year | $\square \text { M19F00 }$ |  | $\square \text { M19F03 }$ |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| year |  |  |  |  |  | female | male | female | male |
| 1948 | 0 | 0 | 0 | 0 | 1981 | 62.61178291 | 75.56595728 | 82.26932357 | 131.5457245 |
| 1949 | 0 | 0 | 0 | 0 | 1982 | 51.87527796 | 70.87388133 | 63.4376148 | 120.0710505 |
| 1950 | 0.029290529 | 0.010148921 | 0.052876667 | 0.032672463 | 1983 | 39.72084346 | 54.03546272 | 44.82566346 | 91.63937955 |
| 1951 | 0.248247159 | 0.137859662 | 0.426265689 | 0.358680293 | 1984 | 29.98144143 | 35.0648082 | 31.06340546 | 60.86077393 |
| 1952 | 1.009797269 | 0.996117696 | 1.729951628 | 2.05994497 | 1985 | 25.60570341 | 33.03218917 | 27.90313734 | 55.15366568 |
| 1953 | 2.27389353 | 3.799354119 | 4.034889901 | 6.798217159 | 1986 | 26.02824674 | 39.80612176 | 31.91702343 | 64.05134402 |
| 1954 | 3.530051511 | 8.111411502 | 6.516424806 | 13.71206184 | 1987 | 29.57524959 | 52.15248396 | 39.27885587 | 80.65351304 |
| 1955 | 4.505292273 | 11.95069256 | 8.572618804 | 19.91123245 | 1988 | 34.25326965 | 69.06617841 | 47.76469389 | 102.1912584 |
| 1956 | 5.23017135 | 14.86119389 | 10.17885791 | 24.64490259 | 1989 | 38.49425689 | 75.18367028 | 55.51587713 | 112.8729649 |
| 1957 | 5.78636166 | 17.08319575 | 11.47760673 | 28.32425004 | 1990 | 40.93261177 | 69.26230859 | 61.27541543 | 111.8311327 |
| 1958 | 6.245965522 | 18.83845512 | 12.60903413 | 31.34932668 | 1991 | 40.45010837 | 66.70096787 | 63.29311168 | 116.5835973 |
| 1959 | 6.678762971 | 20.34192753 | 13.71131601 | 34.10406971 | 1992 | 36.03354495 | 57.41112499 | 59.78533715 | 108.7895289 |
| 1960 | 7.166650145 | 21.80054365 | 14.95040896 | 36.96211501 | 1993 | 29.65188903 | 49.3061425 | 51.56526726 | 100.0945072 |
| 1961 | 7.836129047 | 23.456348 | 16.58795884 | 40.38361165 | 1994 | 23.0563003 | 39.76180942 | 41.6729754 | 83.5553754 |
| 1962 | 8.952929857 | 25.7583941 | 19.17276358 | 45.27954043 | 1995 | 17.59936166 | 29.9848262 | 32.93880923 | 65.64733696 |
| 1963 | 11.20855208 | 29.68301023 | 24.08692299 | 53.71943121 | 1996 | 13.60743973 | 24.14996006 | 26.2383916 | 52.55986347 |
| 1964 | 16.36886723 | 37.82739217 | 34.63784649 | 71.00574611 | 1997 | 10.89988773 | 20.4377007 | 21.53158493 | 43.25608304 |
| 1965 | 27.65874111 | 55.00374857 | 56.44912529 | 107.3336525 | 1998 | 9.235893888 | 18.20281208 | 18.67026921 | 37.79694288 |
| 1966 | 47.58206403 | 93.89734099 | 93.53874708 | 180.1369655 | 1999 | 8.542129551 | 17.98680288 | 17.4372762 | 36.17624799 |
| 1967 | 72.62268113 | 148.279836 | 139.8721418 | 279.6484942 | 2000 | 8.83666402 | 19.51660887 | 17.74578088 | 37.62178453 |
| 1968 | 93.82954052 | 214.5323711 | 180.3751698 | 388.7986146 | 2001 | 9.692410669 | 23.12988283 | 19.18716563 | 42.12837696 |
| 1969 | 104.9105219 | 255.7631836 | 203.5673987 | 456.8869593 | 2002 | 11.02624145 | 28.07170871 | 21.7207241 | 49.14444027 |
| 1970 | 107.109204 | 271.41282 | 210.1384986 | 481.6364664 | 2003 | 12.9610676 | 34.12756344 | 25.48135684 | 58.66512211 |
| 1971 | 105.2731072 | 271.6575832 | 207.2362958 | 478.9748353 | 2004 | 15.62280239 | 42.27234237 | 30.63971212 | 71.66628579 |
| 1972 | 103.0811121 | 267.6417929 | 200.5408889 | 464.2391531 | 2005 | 18.32526828 | 51.63285385 | 36.34643336 | 86.95724602 |
| 1973 | 100.1838676 | 261.581255 | 190.297984 | 440.9478977 | 2006 | 20.8324607 | 60.09081306 | 42.1541279 | 102.2607715 |
| 1974 | 95.18617647 | 246.8464928 | 175.7955327 | 402.2135648 | 2007 | 23.30097298 | 67.36561163 | 47.78906901 | 116.9103724 |
| 1975 | 87.98656329 | 230.3186616 | 158.2999936 | 358.4925464 | 2008 | 23.64643945 | 76.38178815 | 49.26304162 | 130.6858117 |
| 1976 | 77.82978018 | 188.5578534 | 137.8142501 | 289.4207129 | 2009 | 21.09129729 | 76.86605777 | 44.74294322 | 128.1643334 |
| 1977 | 67.70984017 | 130.9737959 | 118.1804075 | 209.8222867 | 2010 | 17.86953558 | 68.49052925 | 38.25400266 | 111.6231267 |
| 1978 | 63.0060915 | 96.16056729 | 106.342965 | 163.6950948 | 2011 | 16.63246429 | 59.23689734 | 35.08159051 | 95.47813585 |
| 1979 | 65.72347908 | 74.32736881 | 107.0998424 | 142.6686239 | 2012 | 19.85647861 | 57.81165696 | 39.75256727 | 94.20587388 |
| 1980 | 67.71303871 | 70.16135116 | 98.82641083 | 131.1216495 | 2013 | 25.76424691 | 70.26742291 | 49.64969507 | 114.7756218 |
|  |  |  |  |  | 2014 | 28.58151286 | 83.75361085 | 54.83740078 | 135.7892866 |
|  |  |  |  |  | 2015 | 26.38078068 | 82.0122724 | 51.08625555 | 131.8573143 |
|  |  |  |  |  | 2016 | 22.15777388 | 75.99847076 | 43.11401688 | 117.1288802 |
|  |  |  |  |  | 2017 | 18.40263189 | 64.09196727 | 35.57943126 | 96.36555861 |
|  |  |  |  |  | 2018 |  |  | 29.66231471 | 79.45494853 |

Table 48. Estimated population size (millions) for females on July 1 of year. from the author's preferred model, Model M19F03.
$\ll$ Table too large: available online in the zip file "TannerCrab.PopSizeStructure.csvs.zip".>>
Table 49. Estimated population size (millions) for males on July 1 of year. from the author's preferred mode, Model M19F03.
$\ll$ Table too large: available online as a zipped csv file "TannerCrab.PopSizeStructure.csvs.zip".>>

Table 50. Comparison of estimates of recruitment (in millions) from the 2018 assessment model (M19F00) and the author's preferred model (M19F03).

| year | M19F00 | M19F03 | year | M19F00 | M19F03 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1948 | 70.09251687 | 132.6537449 | 1981 | 101.416713 | 229.8722417 |
| 1949 | 70.09557738 | 133.2203853 | 1982 | 496.0095549 | 690.0962537 |
| 1950 | 70.19921793 | 134.4944439 | 1983 | 408.5677129 | 627.3626472 |
| 1951 | 70.53781202 | 136.6898133 | 1984 | 550.0166431 | 767.3734676 |
| 1952 | 71.30143949 | 140.1373441 | 1985 | 529.7681206 | 764.4249404 |
| 1953 | 72.76979101 | 145.363002 | 1986 | 525.8487112 | 796.315064 |
| 1954 | 75.37788738 | 153.237184 | 1987 | 356.0941501 | 595.7349193 |
| 1955 | 79.85259349 | 165.2823008 | 1988 | 171.1538503 | 249.3996718 |
| 1956 | 87.52520379 | 184.3743392 | 1989 | 52.28767878 | 78.81849366 |
| 1957 | 101.1405733 | 216.5609486 | 1990 | 41.82858297 | 71.27724516 |
| 1958 | 127.3271887 | 276.6196651 | 1991 | 37.02824035 | 73.65268281 |
| 1959 | 185.5896413 | 407.587403 | 1992 | 36.8859791 | 67.27971079 |
| 1960 | 339.6142312 | 742.4221286 | 1993 | 48.32235441 | 84.32008419 |
| 1961 | 757.2893156 | 1588.034161 | 1994 | 62.36311147 | 143.7247786 |
| 1962 | 1462.061345 | 2839.791915 | 1995 | 57.94345627 | 107.5877226 |
| 1963 | 1736.132801 | 3206.061429 | 1996 | 168.9628999 | 304.259962 |
| 1964 | 1452.379666 | 2674.087231 | 1997 | 67.82772625 | 127.2116239 |
| 1965 | 1131.170889 | 2117.061574 | 1998 | 227.5701775 | 450.4022196 |
| 1966 | 963.730419 | 1816.266851 | 1999 | 118.091505 | 221.0847323 |
| 1967 | 943.2576586 | 1724.0517 | 2000 | 385.0604766 | 754.1423021 |
| 1968 | 1008.697227 | 1669.653411 | 2001 | 123.1097967 | 231.3054485 |
| 1969 | 980.6227068 | 1442.516104 | 2002 | 372.6665098 | 829.6787085 |
| 1970 | 843.9469644 | 1165.600708 | 2003 | 362.1799899 | 687.2774432 |
| 1971 | 561.9043515 | 778.2159348 | 2004 | 97.11673458 | 185.9225205 |
| 1972 | 369.6842268 | 524.6240813 | 2005 | 74.45238686 | 127.3120944 |
| 1973 | 318.0087047 | 510.3866376 | 2006 | 57.8718913 | 111.1939148 |
| 1974 | 641.4445935 | 705.3119393 | 2007 | 88.82972036 | 179.0548001 |
| 1975 | 1257.959539 | 2031.682265 | 2008 | 576.7044896 | 1138.228615 |
| 1976 | 971.5504765 | 1529.970097 | 2009 | 501.346918 | 870.3214169 |
| 1977 | 424.9897994 | 761.2503091 | 2010 | 200.9415568 | 301.1484309 |
| 1978 | 180.9087231 | 276.9401217 | 2011 | 40.77585148 | 62.70194572 |
| 1979 | 110.113046 | 166.2250105 | 2012 | 108.9237585 | 173.4310903 |
| 1980 | 180.4735272 | 265.6626468 | 2013 | 73.93881264 | 106.9458916 |
|  |  |  | 2014 | 49.09325854 | 79.9499305 |
|  |  |  | 2015 | 69.72714155 | 117.3112392 |
|  |  |  | 2016 | 444.7192575 | 647.0591576 |
|  |  |  | 2017 | 588.8895622 | 677.6176162 |
|  |  |  | 2018 |  | 1234.937393 |

Table 51. Comparison of exploitation rates (i.e., catch divided by biomass) from the 2018 assessment model (M19F00) and the author's preferred model (M19F03).

| year | M19F00 | M19F03 | year | M19F00 | M19F03 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1949 | 0.001622787 | 0.000851594 | 1981 | 0.046786646 | 0.026757506 |
| 1950 | 0.002688423 | 0.001373949 | 1982 | 0.025203452 | 0.014146239 |
| 1951 | 0.004152346 | 0.00210409 | 1983 | 0.013099622 | 0.007026487 |
| 1952 | 0.006247527 | 0.003273033 | 1984 | 0.026017138 | 0.015184677 |
| 1953 | 0.009322445 | 0.00529258 | 1985 | 0.015433162 | 0.005593393 |
| 1954 | 0.01260007 | 0.007679072 | 1986 | 0.019326482 | 0.007141423 |
| 1955 | 0.014753304 | 0.009385377 | 1987 | 0.031682959 | 0.013068075 |
| 1956 | 0.015980555 | 0.010368409 | 1988 | 0.040555945 | 0.020050353 |
| 1957 | 0.016287354 | 0.010527445 | 1989 | 0.091529287 | 0.054189398 |
| 1958 | 0.016548995 | 0.010701976 | 1990 | 0.152834055 | 0.091491752 |
| 1959 | 0.016393883 | 0.0105286 | 1991 | 0.14575004 | 0.075020873 |
| 1960 | 0.01602232 | 0.010226125 | 1992 | 0.173127894 | 0.095630499 |
| 1961 | 0.015550666 | 0.009976282 | 1993 | 0.130835171 | 0.054998198 |
| 1962 | 0.014008919 | 0.009008863 | 1994 | 0.098005158 | 0.038802861 |
| 1963 | 0.01190419 | 0.007751795 | 1995 | 0.085254294 | 0.031793233 |
| 1964 | 0.010409007 | 0.006771504 | 1996 | 0.047280956 | 0.019457438 |
| 1965 | 0.015993162 | 0.009020895 | 1997 | 0.033563022 | 0.01697017 |
| 1966 | 0.015931948 | 0.009115647 | 1998 | 0.031137612 | 0.01149889 |
| 1967 | 0.043643679 | 0.025403963 | 1999 | 0.01512733 | 0.005898124 |
| 1968 | 0.048268751 | 0.028737808 | 2000 | 0.012987827 | 0.006044593 |
| 1969 | 0.063683821 | 0.038160681 | 2001 | 0.016821106 | 0.006725774 |
| 1970 | 0.059569187 | 0.035828691 | 2002 | 0.010714727 | 0.003631334 |
| 1971 | 0.050880748 | 0.030686751 | 2003 | 0.006018027 | 0.002625986 |
| 1972 | 0.045502754 | 0.028496504 | 2004 | 0.006466766 | 0.003153082 |
| 1973 | 0.055554121 | 0.035566735 | 2005 | 0.012287384 | 0.006120349 |
| 1974 | 0.074143668 | 0.048631197 | 2006 | 0.018752949 | 0.008653291 |
| 1975 | 0.064643017 | 0.04403133 | 2007 | 0.020865591 | 0.010646254 |
| 1976 | 0.100923862 | 0.070635583 | 2008 | 0.014201418 | 0.007946933 |
| 1977 | 0.140735249 | 0.098008396 | 2009 | 0.012001593 | 0.006769104 |
| 1978 | 0.118938682 | 0.075778039 | 2010 | 0.006272852 | 0.00328305 |
| 1979 | 0.152736347 | 0.085590706 | 2011 | 0.007820264 | 0.004469626 |
| 1980 | 0.093896849 | 0.05814116 | 2012 | 0.004964941 | 0.00300838 |
|  |  |  | 2013 | 0.015086706 | 0.008840832 |
|  |  |  | 2014 | 0.052987808 | 0.031389129 |
|  |  |  | 2015 | 0.072375017 | 0.044605374 |
|  |  |  | 2016 | 0.009963209 | 0.005834419 |
|  |  |  | 2017 | 0.020021174 | 0.010205414 |
|  |  |  | 2018 |  | 0.01100967 |

Table 52. Values required to determine Tier level and OFL for the models considered here. These values are presented only to illustrate the effect of incremental changes in the model scenarios. Results from the author's preferred model (M19F03) are highlighted in green.

| Model <br> Scenario | average <br> recruitment <br> millions | Final MMB | BO | Bmsy | Fmsy | MSY | Fofl | OFL | projected <br> MMB | projected MMB <br> /Bmsy |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| M19F00 | 223.63 | 66.64 | 86.55 | 30.29 | 0.74 | 12.75 | 0.74 | 20.87 | 35.95 | 1.19 |
| M19F00a | 284.28 | 82.05 | 94.24 | 32.99 | 0.89 | 14.58 | 0.89 | 27.90 | 41.52 | 1.26 |
| M19F01 | 316.79 | 68.79 | 100.85 | 35.30 | 0.81 | 15.58 | 0.81 | 22.54 | 35.66 | 1.01 |
| M19F02 | 367.48 | 71.54 | 105.59 | 36.96 | 1.11 | 17.89 | 1.03 | 24.75 | 34.63 | 0.94 |
| M19F03 | 393.84 | 82.61 | 118.96 | 41.64 | 1.18 | 19.49 | 1.12 | 29.48 | 39.68 | 0.95 |
| M19F04 | 377.28 | 74.03 | 106.76 | 37.37 | 0.87 | 16.87 | 0.87 | 24.87 | 37.50 | 1.00 |
| M19F05 | 418.73 | 80.33 | 116.44 | 40.75 | 1.21 | 19.40 | 1.14 | 28.58 | 38.42 | 0.94 |

## Figures



Figure 1. Eastern Bering Sea District of Tanner crab Registration Area J including sub-districts and sections (from Bowers et al. 2008).


Figure 2. Upper: retained catch (males, 1000's $t$ ) in the directed fisheries (US pot fishery [green bars], Russian tangle net fishery [red bars], and Japanese tangle net fisheries [blue bars]) for Tanner crab since 1965/66. Lower: Retained catch (males, 1000's t) in directed fishery since 2001/02. The directed fishery was closed from 1996/97 to 2004/05, from 2010/11 to 2012/13, and in 2016/17.


Figure 3. Time series of retained catch biomass (1000's t) in the directed Tanner crab (TCF: red; eastern area: triangles; western area: circles; all EBS: squares), snow crab (SCF: green), and BBRKC (RFF: blue) fisheries since 2005. The directed fishery was closed from 2010/11 to 2012/13, and in 2016/17. Legalsized Tanner crab can be incidentally-retained in the snow crab and BBRKC fisheries up to a cap of $5 \%$ the target catch.


Figure 4. Upper: total catch (retained + discards) of Tanner crab (males and females, 1000's $t$ ) in the directed Tanner crab, snow crab, Bristol Bay red king crab, and groundfish fisheries. Bycatch reporting began in 1973 for the groundfish fisheries and in 1992 for the crab fisheries. Lower: detail since 2005.


Figure 5. Retained catch size compositions in the directed Tanner crab fisheries since the fishery reopened in 2013/14 (red: western area, green: eastern area; blue: all EBS).


Figure 6. Total catch (retained + discards) size compositions for males, normalized by fleet, during 1990/91-1999/2000 in the directed Tanner crab (aggregated across areas, TCF: blue), snow crab (SCF: green), and BBRKC (RKF: red) fisheries (solid line: new shell crab; dotted line: old shell crab).


Figure 7. Total catch (retained + discards) size compositions for males, normalized by fleet, during 2000/01-2009/10 in the directed Tanner crab (aggregated across areas, TCF: blue), snow crab (SCF: green), and BBRKC (RKF: red) fisheries (solid line: new shell crab; dotted line: old shell crab). The directed fishery was closed in 2000/01-2004/05 and was open only in the western area in 2005/06 and in the eastern area in 2009/10.


Figure 8. Total catch (retained + discards) size compositions for males, normalized by fleet, during 2010/11-2018/19 in the directed Tanner crab (aggregated across areas, TCF: blue), snow crab (SCF: green), and BBRKC (RKF: red) fisheries (solid line: new shell crab; dotted line: old shell crab). The directed fishery was closed in 2010/11-2012/13 and 206/17, and was open only in the western area in 2017/18 and 2018/19.


Figure 9. Bycatch size compositions for females, normalized by fleet, during 1990/91-1999/2000 in the directed Tanner crab (aggregated across areas, TCF: blue), snow crab (SCF: green), and BBRKC (RKF: red) fisheries (solid line: new shell crab; dotted line: old shell crab).


Figure 10. Bycatch size compositions for females, normalized by fleet, during 2000/01-2009/10 in the directed Tanner crab (aggregated across areas, TCF: blue), snow crab (SCF: green), and BBRKC (RKF: red) fisheries (solid line: new shell crab; dotted line: old shell crab).


Figure 11. Bycatch size compositions for females, normalized by fleet, during 2010/11-2018/19 in the directed Tanner crab (aggregated across areas, TCF: blue), snow crab (SCF: green), and BBRKC (RKF: red) fisheries (solid line: new shell crab; dotted line: old shell crab).


Figure 12. Annual bycatch size compositions in the groundfish fisheries by sex, expanded to total bycatch, during 1991/92-2006/07. Red lines: females; green lines: males.


Figure 13. Annual bycatch size compositions in the groundfish fisheries by sex, expanded to total bycatch, during 2007/08-2018/19. Red lines: females; green lines: males.


Figure 14. Annual estimates of area-swept biomass from the NMFS EBS bottom trawl survey, by sex, maturity state, and management area. Red lines: total biomass; green lines: biomass in the eastern area; blue: biomass in the western area.


Figure 15. Annual estimates of area-swept biomass from the NMFS EBS bottom trawl survey for preferred-size ( $>125 \mathrm{~mm}$ CW) legal males. Red lines: total biomass; green lines: biomass in the eastern area; blue: biomass in the western area.


Figure 16. Spatial footprints (stations occupied in green) during the BSFRF-NMFS cooperative side-by-side (SBS) catchability studies in 20132017. Squares and circles represent stations in the standard NMFS EBS bottom trawl survey (which extends beyond the area shown in the maps).


Figure 17. Annual estimates of area-swept biomass from the BSFRF-NMFS cooperative side-by-side (SBS) catchability studies in 2013-2017. The SBS studies had different spatial footprints each year, so annual changes in biomass do not necessarily reflect underlying population trends. Red lines: BSFRF; green lines: NMFS.


Figure 18. Size compositions from the NMFS EBS bottom trawl survey for 1975-2019.


Figure 19. Annual size compositions of area-swept abundance by sex from the BSFRF-NMFS cooperative side-by-side (SBS) catchability studies in 2013-2015. Red lines: BSFRF; green lines: NMFS.


Figure 19 (cont.). Annual size compositions of area-swept abundance by sex from the BSFRF-NMFS cooperative side-by-side (SBS) catchability studies in 2017. Red lines: BSFRF; green lines: NMFS


Figure 20. Annual estimates of area-swept abundance (blue circles) from the NMFS EBS bottom trawl survey, by sex and maturity state for 2014 and 2015. Local abundance scales with symbol area. The background "heatmap" represents bottom water temperatures at the time of the survey.


Figure 21. Annual estimates of area-swept abundance (blue circles) from the NMFS EBS bottom trawl survey, by sex and maturity state for 2016 and 2017. Local abundance scales with symbol area. The background "heatmap" represents bottom water temperatures at the time of the survey.


Figure 22. Annual estimates of area-swept abundance (blue circles) from the NMFS EBS bottom trawl survey, by sex and maturity state for 2018 and 2019. Local abundance scales with symbol area. The background "heatmap" represents bottom water temperatures at the time of the survey.


Figure 23. Male maturity ogives (the fraction of new shell mature males, relative to all new shell males) as determined from chela height:carapace width ratios from the NMFS EBS bottom trawl survey for years when chela heights were collected with 0.1 mm precision..


Figure 24. Molt increment data collected collaboratively by NMFS, BSFRF, and ADFG.


Figure 25. Size-weight relationships developed from NMFS EBS summer trawl survey data.


Figure 26. Assumed size distribution for recruits entering the population.


Figure 27. Fits to retained and total catch biomass in the directed fishery from all model scenarios.


Figure 28. Fits to total catch biomass in the snow crab fishery from all scenarios.


Figure 29. Fits to total catch biomass in the BBRKC fishery from all scenarios.


Figure 30. Fits to total catch biomass in the groundfish fisheries for all scenarios.


Figure 31. Fits to mature biomass from the NMFS " 0 " EBS bottom trawl survey data for all. Note that scenarios M19F03 and M19F05 do not include the mature male component in the likelihood (they fit total male biomass) and fit both mature and immature biomass for females.


Figure 32. Fits to mature biomass from the NMFS "M" and NMFS "F" EBS bottom trawl survey data for scenarios M19F00a, M19F01, M19F02, M19F02, M19F03, M19F04, and M19F05. Note that only scenarios M19F03 and M19F05 include these data components in the model objective function.


Figure 33. Fits to survey biomass from the NMFS SBS bottom trawl survey data for scenarios M19F04 and M19F05.

SBS BSFRF males


Figure 34. Fits to survey biomass from the BSFRF SBS bottom trawl survey data for scenarios M19F04 and M19F05.


Figure 35. Fits to molt increment data for scenarios M19F00a, M19F01, M19F02, M19F02, M19F03, M19F04, and M19F05.


Figure 36. Fits to male maturity ogive data for scenarios M19F00a, M19F01, M19F02, M19F02, M19F03, M19F04, and M19F05. Note that only scenarios M1902, M19F03, and M19F05 include the data in the likelihood.


Figure 37. Directed fishery catchability (capture rates) from all model scenarios.


Figure 38. Directed fishery selectivity curves from all scenarios for the pre-1991 time period and 1991-1994. The $50 \%$-selected parameter varies annually for $1991+$.


Figure 39. Directed fishery selectivity curves from all scenarios for 1995-1996 and 2005-2007. The $50 \%$-selected parameter varies annually for 1991+.


Figure 40. Directed fishery selectivity curves from all scenarios for 2008-2009 and 2013-2015. The $50 \%$-selected parameter varies annually for 1991+.


Figure 41 . Directed fishery selectivity curves from all scenarios for 2008-2009 and 2013-2015. The $50 \%$-selected parameter varies annually for 1991+.


Figure 42. Directed fishery retention curves from all scenarios for the pre-1991, 1991-1996, and post-2004 time periods


Figure 43. Snow crab fishery catchability (capture rates) from all scenarios.


Figure 44. Snow crab fishery selectivity curves from all scenarios for 3 time periods: pre-1997, 1997-2004, 2005+.


Figure 45. BBRKC fishery catchability (capture rates) from all scenarios.


Figure 46. BBRKC fishery selectivity curves from all scenarios for 3 time periods: pre-1997, 1997-2004, 2005+.


Figure 47. Catchability (capture rates) in the groundfish fisheries from all scenarios.


Figure 48. Groundfish fisheries selectivity curves from all scenarios estimated for 3 time periods: pre-1987, 1987-1996, 1997+


Figure 49. NMFS "0" survey catchabilities for all scenarios for the 1975-1981 and 1982+ time periods.


Figure 50. NMFS "0" survey selectivity functions for all scenarios for the 1975-1981 and 1982+ time periods.


Figure 51. NMFS " 0 " survey capture probabilities (i.e., catchability $x$ selectivity) for all scenarios for the 1975-1981 and 1982+ time periods.


Figure 52. Survey availabilities from scenarios M19F04 and M19F05 for the 2013-2017 SBS studies.


Figure 53. Comparison of empirical "observed" and predicted availability in the 2013-2017 SBS studies from scenario M19F04. The "observed" availability is the ratio of abundance in the NMFS SBS survey to that in the full NMFS survey by size bin. Observed: red points, lines. Red fills are from loess smoothing of the observed availability. Predicted: green points, lines.


Figure 54 . Estimates of natural mortality from all scenarios.


Figure 55. Estimates of the probability of terminal molt from all scenarios.


Figure 56. Estimates of mean growth from all scenarios. Dashed line is 1:1.



- M19F00a
- M19F01
- M19F02
- M19F03
$\div$ M19F04
- M19F05
- M19F00a
- M19F01
- M19F02
+ M19F03
- M19F04
* M19F05

Figure 57. Estimated recruitment time series from all scenarios.


Figure 58. Estimated recent recruitment time series from all scenarios.


Figure 59. Estimated (Feb. 15) mature biomass time series from all scenarios.


Figure 60. Estimated recent (Feb. 15) mature biomass time series from all scenarios.


Figure 61. Estimated (July 1) biomass time series by population category for all scenarios.


Figure 62. MCMC results from scenario M19F03, the author's preferred model, for OFL-related quantities.


Figure 63. The Fofs harvest control rule.


Figure 64. The OFL and ABC from the author's preferred model, scenario M19F03.


Figure 65. Quad plot for the author's preferred model, scenario M19F03.


Figure 66. The ratio of estimated abundance by size from the NMFS and BSFRF side-by-side catchability studies. The heavy green line is the size-specific mean over the 5 years. These represent simple empirical estimates of the size-specific catchability of the NMFS survey gear relative to the BSFRF gear. If the BSFRF survey gear is assumed to capture all crab within the area swept, these curves represent empirical estimates of the size-specific NMFS survey gear catchability (i.e., fully selected catchability [q] $x$ selectivity).

# Appendix A: <br> Description of the Tanner Crab Stock Assessment Model, Version 2 

September, 2019

## Introduction

The "TCSAM02" (Tanner Crab Stock Assessment Model, version 2) modeling framework was developed "from scratch" to eliminate many of the constraints imposed on potential future assessment models by TCSAM2013, the previous assessment model framework (Stockhausen, 2016). Like TCSAM2013, TCSAM02 uses AD Model Builder libraries as the basis for model optimization using a maximum likelihood (or Bayesian) approach. The model code for TCSAM02 is available on GitHub (the 2019 assessment model code is available at " 201909 CPTdoRetro"). TCSAM02 was first used for the Tanner crab assessment in 2017 (Stockhausen, 2017) and will be used until a transition is made to Gmacs (the Generalized Model for Alaska Crab $\underline{S}$ tocks). Gmacs is intended to be used for all crab stock assessments conducted for the North Pacific Fisheries Management Council (NPFMC), including both lithodid (king crab) and Chionoecetes (Tanner and snow crab) stocks, while TCSAM02 is specific to Chionoecetes biology (i.e., terminal molt).

TCSAM02 is referred to here as a "modeling framework" because, somewhat similar to Stock Synthesis (Methot and Wetzel, 2013), model structure and parameters are defined "on-the-fly" using control filesrather than editing and re-compiling the underlying code. In particular, the number of fisheries and surveys, as well as their associated data types (abundance, biomass, and /or size compositions) and the number and types of time blocks defined for every model parameter, are defined using control files in TCSAM02 and have not been pre-determined. Priors can be placed on any model parameter. New data types (e.g., growth data) can also be included in the model optimization that could not be fit with TCSAM2013. Additionally, status determination and OFL calculations can be done directly within a TCSAM02 model run, rather having to run a separate "projection model".

## Model Description

## A. General population dynamics

TCSAM02 is a stage/size-based population dynamics model. Population abundance at the start (July 1) of year $y$ in the model, $n_{y, x, m, s, z}$, is characterized by sex $x$ (male, female), maturity state $m$ (immature, mature), shell condition $s$ (new shell, old shell), and size $z$ (carapace width, CW). Changes in abundance due to natural mortality, molting and growth, maturation, shell aging, fishing mortality and recruitment are tracked on an annual basis. Because the principal crab fisheries occur during the winter, the model year runs from July 1 to June 30 of the following calendar year.

The order of calculation steps to project population abundance from year $y$ to $y+1$ depends on the assumed timing of the fisheries $\left(\delta t_{y}^{F}\right)$ relative to molting/growth/mating $\left(\delta t_{y}^{m}\right)$ in year $y$. The steps when the fisheries occur before molting/growth/mating ( $\delta t_{y}^{F} \leq \delta t_{y}^{m}$ ) are outlined below first (Steps A1.1-A1.4), followed by the steps when molting/growth/mating occurs after the fisheries ( $\delta t_{y}^{m}<\delta t_{y}^{F}$;


Fig. 1. Timing of annual events in TCSAM02 when fisheries occur before molting/growth/mating. Steps A2.1-A2.4).

## A1. Calculation sequence when $\boldsymbol{\delta} \boldsymbol{t}_{\boldsymbol{y}}^{\boldsymbol{F}} \leq \boldsymbol{\delta} \boldsymbol{t}_{\boldsymbol{y}}^{\boldsymbol{m}}$

## Step A1.1: Survival prior to fisheries

Natural mortality is applied to the population from the start of the model year (July 1) until just prior to prosecution of pulse fisheries for year $y$ at $\delta t_{y}^{F}$. The numbers surviving to $\delta t_{y}^{F}$ in year $y$ are given by:

| $n_{y, x, m, s, z}^{1}=e^{-M_{y, x, m, s, z} \cdot \delta t_{y}^{F}} \cdot n_{y, x, m, s, z}$ | A 1.1 |
| :--- | :---: |

where $M$ represents the annual rate of natural mortality in year $y$ on crab classified as $x, m, s, z$.

## Step A1.2: Prosecution of the fisheries

The directed and bycatch fisheries are modeled as simultaneous pulse fisheries occurring at $\delta t_{y}^{F}$ in year $y$. The numbers that remain after the fisheries are prosecuted are given by:

| $n_{y, x, m, s, z}^{2}=e^{-F_{y, x, m, s, z}^{T} \cdot n_{y, x, m, s, z}^{1}}$ | A1.2 |
| :--- | :---: |

where $F_{y, x, m, s, z}^{T}$ represents the total fishing mortality (over all fisheries) on crab classified as $x, m, s, z$ in year $y$.

## Step A1.3: Survival after fisheries to time of molting/growth/mating

Natural mortality is again applied to the population from just after the fisheries to the time just before molting/growth/mating occurs for year $y$ at $\delta t_{y}^{m}$ (generally Feb. 15). The numbers surviving to $\delta t_{y}^{m}$ in year $y$ are given by:

| $n_{y, x, m, s, z}^{3}=e^{-M_{y, x, m, s, z} \cdot\left(\delta t_{y}^{m}-\delta t_{y}^{F}\right)} \cdot n_{y, x, m, s, z}^{2}$ | A1.3 |
| :--- | :---: |

where, as above, $M$ represents the annual rate of natural mortality in year $y$ on crab classified as $x, m, s, z$.

Step A1.4: Molting, growth, and maturation
The changes in population structure due to molting, growth and maturation of immature (new shell) crab, as well as the change in shell condition for mature new shell (MAT, NS) crab to mature old shell (MAT, OS) crab due to aging, are given by:

| $n_{y, x, M A T, N S, z}^{4}=\phi_{y, x, z} \cdot \sum_{z^{\prime}} \Theta_{y, x, z, z^{\prime}} \cdot n_{y, x, I M M, N S, z^{\prime}}^{3}$ | A1.4a |
| :--- | :---: |
| $n_{y, x, I M M, N S, z}^{4}=\left(1-\phi_{y, x, z}\right) \cdot \sum_{z^{\prime}} \Theta_{y, x, Z, z^{\prime}} \cdot n_{y, x, I M M, N S, z^{\prime}}^{3}$ | A1.4b |
| $n_{y, x, M A T, O S, z}^{4}=n_{y, x, M A T, O S, z}^{3}+n_{y, x, M A T, N S, z}^{3}$ | A1.4c |

where $\Theta_{y, x, z, z^{\prime}}$ is the growth transition matrix in year $y$ for an immature new shell (IMM, NS) crab of sex $x$ and pre-molt size $z$ ' to post-molt size $z$ and $\phi_{y, x, z}$ is the probability that a just-molted crab of sex $x$ and post-molt size $z$ has undergone its terminal molt to maturity (MAT). All crab that molted remain new shell (NS) crab. Additionally, all mature crab that underwent terminal molt to maturity the previous year are assumed to change shell condition from new shell to old shell (A1.4c). Note that the numbers of immature old shell (IMM, OS) crab are identically zero in the current model because immature crab are assumed to molt each year until they undergo the terminal molt to maturity; consequently, the "missing" equation for $m=I M M, s=O S$ is unnecessary.

Step A1.5: Survival to end of year, recruitment, and update to start of next year
Finally, the population abundance at the start of year $y+1$, due to natural mortality on crab from just after the time of molting/growth/mating in year $y$ until the end of the model year (June 30) and recruitment $\left(R_{y, x, z}\right)$ at the end of year $y$ of immature new shell (IMM, NS) crab by sex $x$ and size $z$, is given by:
$n_{y+1, x, m, s, z}=\left\{\begin{array}{ll|l}e^{-M_{y, x, I M M, N S, z} \cdot\left(1-\delta t_{y}^{m}\right)} \cdot n_{y, x, I M M, N S, z}^{4}+R_{y, x, Z} & m=I M M, s=N S \\ e^{-M_{y, x, m, s, z} \cdot\left(1-\delta t_{y}^{m}\right)} \cdot n_{y, x, m, s, z}^{4} & \text { otherwise }\end{array} \quad\right.$ A1.5

## A2. Calculation sequence when $\delta t_{\boldsymbol{y}}^{\boldsymbol{m}}<\boldsymbol{\delta} \boldsymbol{t}_{\boldsymbol{y}}^{\boldsymbol{F}}$

Step A2.1: Survival prior to molting/growth/mating
As in the previous sequence, natural mortality is first applied to the population from the start of the model year (July 1), but this time until just prior to molting/growth/mating in year $y$ at $\delta t_{y}^{m}$ (generally Feb. 15). The numbers surviving at $\delta t_{y}^{m}$ in year $y$ are given by:

| $n_{y, x, m, s, z}^{1}=e^{-M_{y, x, m, s, z} \cdot \delta t_{y}^{m}} \cdot n_{y, x, m, s, z}$ | A2.1 |
| :--- | :--- |

where $M$ represents the annual rate of natural mortality in year $y$ on crab classified as $x, m, s, z$.
Step A2.2: Molting, growth, and maturation
The changes in population structure due to molting, growth and maturation of immature new shell (IMM, NS) crab, as well as the change in shell condition for mature new shell (MAT, NS) crab to mature old shell (MAT, OS) crab due to aging, are given by:

| $n_{y, x, M A T, N S, z}^{2}=\phi_{y, x, z} \cdot \sum_{z^{\prime}} \Theta_{y, x, z, z^{\prime}} \cdot n_{y, x, I M M, N S, z^{\prime}}^{1}$ | A2.2a |
| :--- | :---: |
| $n_{y, x, I M M, N S, z}^{2}=\left(1-\phi_{y, x, z}\right) \cdot \sum_{z^{\prime}} \Theta_{y, x, z, z^{\prime}} \cdot n_{y, x, I M M, N S, z^{\prime}}^{1}$ | A2.2b |
| $n_{y, x, M A T, O S, z}^{2}=n_{y, x, M A T, O S, z}^{1}+n_{y, x, M A T, N S, z}^{1}$ | A2.2c |

where $\Theta_{y, x, z, z^{\prime}}$ is the growth transition matrix in year $y$ for an immature new shell (IMM, NS) crab of sex $x$ and pre-molt size $z$ ' to post-molt size $z$ and $\phi_{y, x, z}$ is the probability that a just-molted crab of $\operatorname{sex} x$ and post-molt size $z$ has undergone its terminal molt to maturity. Additionally, mature new shell (MAT, NS) crab that underwent their terminal molt to maturity the previous year are assumed to change shell condition from new shell to old shell (A2.2c). Again, the numbers of immature old shell crab are identically zero because immature crab are assumed to molt each year until they undergo the terminal molt to maturity.

Step A2.3: Survival after molting/growth/mating to prosecution of fisheries
Natural mortality is again applied to the population from just after molting/growth/mating to the time at which the fisheries occur for year $y$ (at $\delta t_{y}^{F}$ ). The numbers surviving at $\delta t_{y}^{F}$ in year $y$ are then given by:

| $n_{y, x, m, s, z}^{3}=e^{-M_{y, x, m, s, z} \cdot\left(\delta t_{y}^{F}-\delta t_{y}^{m}\right)} \cdot n_{y, x, m, s, z}^{2}$ | A 2.3 |
| :--- | :---: |

where, as above, $M$ represents the annual rate of natural mortality in year $y$ on crab classified as $x, m, s, z$.
Step A2.4: Prosecution of the fisheries
The directed fishery and bycatch fisheries are modeled as pulse fisheries occurring at $\delta t_{y}^{F}$ in year $y$. The numbers that remain after the fisheries are prosecuted are given by:

| $n_{y, x, m, s, z}^{4}=e^{-F_{y, x, m, s, z}^{T} \cdot n_{y, x, m, s, z}^{3}}$ | A2.4 |
| :--- | :---: |

where $F_{y, x, m, s, z}^{T}$ represents the total fishing mortality (over all fisheries) on crab classified as $x, m, s, z$ in year $y$.

Step A2.5: Survival to end of year, recruitment, and update to start of next year Finally, population abundance at the start of year $y+1$ due to natural mortality on crab from just after prosecution of the fisheries in year $y$ until the end of the model year (June 30) and recruitment of immature new (IMM, NS) shell crab at the end of year $y\left(R_{y, x, z}\right)$ and are given by:

$$
\begin{array}{|ll|l|}
\hline n_{y+1, x, m, s, z}= \begin{cases}e^{-M_{y, x, I M M, N S, z} \cdot\left(1-\delta t_{y}^{F}\right)} \cdot n_{y, x, I M M, N S, z}^{4}+R_{y, x, z} & m=I M M, s=N S \\
e^{-M_{y, x, m, S, z} \cdot\left(1-\delta t_{y}^{F}\right)} \cdot n_{y, x, m, s, z}^{4} & \text { otherwise }\end{cases} & \text { A2.5 } \\
\hline
\end{array}
$$

## B. Parameter specification

Because parameterization of many model processes (e.g., natural mortality, fishing mortality) in TCSAM02 is fairly flexible, it is worthwhile discussing how model processes and their associated parameters are configured in TCSAM02 before discussing details of the model processes themselves. Each type of model process has a set of (potentially estimable) model parameters and other information associated with it, but different "elements" of a model process can be defined that apply, for example, to different segments of the population and/or during different time blocks. In turn, several "elements" of a model parameter associated with a model process may also be defined (and applied to different elements of the process). At least one combination of model parameters and other information associated with a model process must be defined-i.e., one process element must be defined.

Model processes and parameters are configured in a "ModelParametersInfo" file, one of the three control files required for a model run (the others are the "ModelConfiguration" file and the "ModelOptions" file). As an example of the model processes and parameter specification syntax, Text Box 1 presents the part of a "ModelParametersInfo" file concerned with specifying fishing processes in the directed Tanner crab fishery.

In Text Box 1, the keyword "fisheries" identifies the model process in question. The first section, following the "PARAMETER_COMBINATIONS" keyword (up to the first set of triple blue dots), specifies the indices associated with fishing process parameters ( $\mathrm{pHM}, \mathrm{pLnC}, \mathrm{pDC} 1, \mathrm{pDC} 2, \mathrm{pDC} 3$, $\mathrm{pDC4}$, pDevsLnC, pLnEffX, pLgtRet), selectivity and retention functions (idxSelFcn, idxRetFcn), and effort averaging time period (effAvgID) that apply to a single fishing process element. In this example, the indices for the selectivity and retention functions, as well as those for the effort averaging time period, constitute the "other information" specified for each fishing process element. Each fishing process element in turn applies to a specific fishery (FISHERY $=1$ indicates the directed fishery, in this case), time block (specified by YEAR_BLOCK), and components of the model population (specified by SEX, MATURITY STATE, and SHELL CONDITION). Using indices to identify which parameters and selectivity and retention functions apply to a given combination of fishery/time block/sex/maturity state/shell condition allows one to "share" individual parameters and selectivity and retention functions across different fishery/time block/sex/maturity state/shell condition combinations.

The second section (following the "PARAMETERS" keyword) determines the characteristics for each of the fishing process parameters, organized by parameter name (note: the parameters associated with the different selectivity and retention functions are specified in a different section of the ModelParametersInfo file). Here, each parameter name corresponds to an ADMB "param_init_bounded_number_vector" in the model code-the exception being pDevsLnC, which corresponds to an ADMB "param_init_bounded_vector_vector".

Each row under a "non-devs" parameter name in the fisheries section (e.g., pLnC) specifies the index used to associate an element of the parameter with the fishing processes defined in the PARAMETER_COMBINATIONS section, as well as characteristics of the element in the associated ADMB number_vector (upper and lower bounds, initial value, and initial estimation phase), various flags for initialization ("jitter", "resample"), definition of an associated prior probability distribution, and a label. Each row under a "devs" parameter name (e.g., pDevsLnC) specifies much the same information for the associated ADMB devs vector, with the "read" flag replacing the "initial value" entry. If "read?" is TRUE, then a vector of initial values is read from the file after all "info" rows for the devs parameter have
been read. The "jitter" flag (if set to TRUE) provides the ability to change the initial value for an element of a non-devs parameter using a randomly selected value based on the element's upper and lower bounds. For a devs parameter, an element with jitter set to TRUE is initialized using a vector of randomlygenerated numbers (subject to being a devs vector within the upper and lower bounds). The "resample" flag was intended to specify an alternative method to providing randomly-generated initial values (based on an element's prior probability distribution, rather than its upper and lower bounds), but this has not yet been fully implemented.

Some model processes apply only to specific segments of the population (e.g., growth only applies to immature, new shell crab). In general, though, a model process element can be defined to apply to any segment of the population (by specifying SEX, MATURITY STATE, and SHELL CONDITION appropriately) and range of years (by specifying YEAR_BLOCK). In turn, an element of a parameter may be "shared" across multiple processes by specifying the element's index in multiple rows of a PARAMETERS_COMBINATION block.


```
PARAMETERS
pHM #handling mortality (0-1)
3 #number of parameters
pLnC #base (ln-scale) capture rate (mature males)
9 #number of parameters
\begin{tabular}{|c|c|c|c|c|c|c|c|c|c|c|c|}
\hline 9 & \multicolumn{2}{|l|}{\multirow[t]{2}{*}{limits}} & eters & \multirow[b]{2}{*}{initial value} & \multirow[b]{2}{*}{\begin{tabular}{l}
start \\
phase
\end{tabular}} & \multirow[b]{2}{*}{|resample? \(\mid\)} & \multirow[t]{2}{*}{} & \multicolumn{2}{|l|}{\multirow[t]{2}{*}{priors}} & \multirow[b]{2}{*}{\[
-1
\]} & \multirow[b]{2}{*}{label} \\
\hline \#id & & & & & & & & & & & \\
\hline 1 & -15 & 15 & OFF & -2.995732274 & -1 & OFF & 1 & none & none & none & TCF:_base_capture_rate,_pre-1965_(=0.05) \\
\hline 2 & -15 & 15 & ON & -1.164816291 & 1 & OFF & 1 & none & none & none & TCF:_base_capture_rate,_1965+ \\
\hline
\end{tabular}
pDC1 #main temporal ln-scale capture rate offset
pDC1 #main temporal 1n-sca
pDC2 #ln-scale capture rate offset for female crabs
6 #number of parameters
## m
pDevsLnC #annual ln-scale capture rate deviations
pDevsLnC #annual in-scale capture rate
*)
```

Text Box 1. Abbreviated example of process and parameter specifications in a "ModelParametersInfo" file for fishing mortality in TCSAM02. Only parameter combinations and parameters relevant to the directed fishery are shown. Input values are in black text, comments are in green, triple blue dots indicate additional input lines not shown.

## C. Model processes: natural mortality

The natural mortality rate applied to crab of sex $x$, maturity state $m$, shell condition $s$, and size $z$ in year $y$, $M_{y, x, m, s, z}$, can be specified using one of two parameterizations. The first parameterization option uses a $\ln$-scale parameterization with an option to include an inverse- size dependence using Lorenzen's approach:

| $\ln M_{y, x, m, s}=\mu_{y, x, m, s^{-}}^{0^{-}} \sum^{\frac{4^{-}}{-}} \delta \mu^{-} \mu_{y, x, m, s^{-}}^{i^{-}}$ | C.1a |
| :---: | :---: |
| $M_{y, x, m, s, z^{-}}=\left\{\begin{array}{lc} \exp \left(\ln M_{y, x, m, s}\right)^{-} & \text {if Lorenzenoption } \bar{\imath} \text { s not selected }{ }^{-} \\ -\exp \left(\ln M_{y, x, m, s}\right) \cdot \frac{z_{\text {base }}}{z^{-}} & \text {if Lorenzenoption } \bar{\imath} \text { s selected } \end{array}\right.$ | $\begin{aligned} & \text { C.1b } \\ & \text { C.1c } \end{aligned}$ |

where the $\mu^{0^{-}}$and the $\delta \mu^{i}$ 's are (potentially) estimable parameters defined for time block $T$, sex $S$ (MALE, FEMALE, or ANY), maturity $M$ (IMMATURE, MATURE, or ANY), and shell condition $S$ (NEWSHELL, OLDSHELL, or ANY), and $\{y, x, m, s\}$ falls into the set $\{T, X, M, S\}$. In Eq. C.1c, $z_{\text {base }}{ }^{-}$ denotes the specified reference size ( mm CW ) for the inverse-size dependence.

The second parameterization option uses an arithmetic parameterization in order to provide backward compatibility with the 2016 assessment model based on TCSAM2013. In TCSAM2013, the natural mortality rate $M_{y, x, m, s, z}$-was parameterized using:

| $M_{y, x, m=I M M, s, z^{-}}={ }^{\text {b }}{ }^{\text {base }}{ }^{-} \cdot \delta M_{I M M^{-}}$ | C.2a |
| :---: | :---: |
| $M_{y, x, m=M A T, s, Z^{-}}=\left\{\begin{array}{lc} M^{\text {base }^{-}} \cdot \delta M_{x, M A T^{-}} & \text {otherwise }^{-} \\ M^{\text {base }^{-}} \cdot \delta M_{x, M A T^{-}} \delta M_{x, M A T^{-}}^{T} & 1980^{-} \leq y \leq 1984^{-} \end{array}\right.$ | C.2b |

where $M^{\text {base }}$ was a fixed value ( $0.23 \mathrm{yr}^{-1}$ ), $\delta M_{I M M^{-} \text {-was a multiplicative factor applied for all immature }}$ crab, the $\delta M_{x, M A T}$-were sex-specific multiplicative factors for mature crab, and the $\delta M_{x, M A T}^{T}$-were additional sex-specific multiplicative factors for mature crab during the 1980-1984 time block (which has been identified as a period of enhanced natural mortality on mature crab, the mechanisms for which are not understood). While it would be possible to replicate Eq.s C.2a and C.2b using ln-scale parameters, TCSAM2013 also placed informative arithmetic-scale priors on some of these parameters-and this could not be duplicated on the ln -scale. Consequently, the second option uses the following parameterization, where the parameters (and associated priors) are defined on the arithmetic-scale:

$$
\ln M_{y, x, m, s}=\ln \left[\mu_{y, x, m, s}^{0^{-}}\right]+\sum_{i=1^{-}}^{4^{-}} \ln \left[\delta \mu_{y, x, m, s}^{i^{-}}\right]^{-} \quad \quad \text { C. } 3 \mathrm{a}
$$

A system of equations identical to C.2a-b can be achieved under the following assignments:

| $\mu_{\{y, x, m, S\} \in\{T=A L L, X=A L L, M=A L L, S=A L L\}^{-}}^{0^{-}}=^{-} M^{\text {base }}$ | C. 4 a |
| :--- | :---: |
| $\delta \mu_{\{y, x, m, s\} \in\{T=A L L, X=A L L, M=I M M, S=A L L\}^{-}}^{1^{-}}=^{-} \delta M_{I M M^{-}}$ | C.4e |
| $\delta \mu_{\{y, x, m, s\} \in\{T=A L L, X=x, M=M A T, S=A L L\}^{-}}^{1^{-}}=^{-} \delta M_{x, M A T^{-}}$ | C.4f |
| $\delta \mu_{\{y, x, m, s\} \in\{T=1980-1984, X=x, M=M A T, S=A L L\}^{-}}^{2^{-}}=^{-} \delta M_{x, M A T^{-}}^{T}$ | C. 4 g |

where unassigned $\delta \mu_{y, x, m, s}^{i}$-are set equal to 1 . Pending further model testing using alternative model configurations, the TCSAM2013 option is standard.

It is worth noting explicitly that, given the number of potential parameters above that could be used, extreme care must be taken when defining a model to achieve a set of parameters that are not confounded and are, at least potentially, estimable.

## D. Model processes: growth

Because Tanner crab are assumed to undergo a terminal molt to maturity, in TCSAM02 only immature crab experience growth. Annual growth of immature crab is implemented as using two options, the first based on a formulation used in Gmacs and the second (mainly for purposes of backward compatibility) based on that used in TCSAM2013. In TCSAM02, growth can vary by time block and sex, so it is expressed by sex-specific transition matrices for time block $t, \Theta_{t, x, z, z^{\prime}}$, that specify the probability that crab of sex $x$ in pre-molt size bin $z^{\prime}$ grow to post-molt size bin $z$ at molting.

In the Gmacs-like approach (the standard approach as of May, 2017), the sex-specific growth matrices are given by:

| $\Theta_{t, x, z, z^{\prime}}=c_{t, x, z^{\prime}} . \int_{z-b i n / 2^{-}}^{+b i n / 2^{-}} \int_{\beta_{t, x^{-}}}^{-} \Gamma\left(\frac{z^{\prime \prime-}-\bar{z}_{t, x, z^{\prime}}}{\beta^{-}}\right) d z^{\prime \prime-}$ | Sex-specific $(x)$ transition matrix for growth from pre-molt $z^{\prime}$ to post-molt $z$, with $z \geq z^{\prime}$ | D.1a |
| :---: | :---: | :---: |
| $c_{t, x, z^{\prime}}=-\left[\int_{z^{\prime}}^{\infty-} \Gamma\left(\frac{z^{\prime \prime-}-\bar{z}_{t, x, z^{\prime}}}{\beta_{t, x^{-}}}\right) d z^{\prime \prime}\right]^{-1-}$ | Normalization constant so $1=\sum_{Z^{-}} \Theta_{t, x, z, z^{-}}$ | D.1b |
| $\bar{z}_{t, x, z^{\prime}}=^{-} e^{a_{t, x}} \cdot z^{\prime} b_{t, x^{-}}$ | Mean size after molt, given pre-molt size $z^{\prime^{-}}$ | D.1c |

where the integral represents a cumulative gamma distribution across the post-molt ( $z$ ) size bin. This approach may have better numerical stability properties than the TCSAM2013 approach below.

The TCSAM2013 approach is an approximation to the Gmacs approach, where the sex-specific growth matrices $\Theta_{t, x, z, z^{\prime}}$ are given by

| $\Theta_{t, x, z, z^{\prime}}=\bar{c}_{t, x, z^{\prime}} \cdot \Delta_{z, z^{\prime}} \alpha_{t, x, z^{\prime}}-1 \cdot e^{-\frac{\Delta_{z, z^{\prime}}}{\beta_{t, x}-}}$ | Sex-specific ( $x$ ) transition matrix for growth from pre-molt $z^{\prime}$ to post-molt $z$, with $z \geq z^{\prime}$ | D. 2 a |
| :---: | :---: | :---: |
| $c_{t, x, z^{\prime}}=\left[\sum_{z^{\prime}} \Delta_{z, z^{\prime}} \alpha_{t, x, z^{\prime}}-1 \cdot e^{-\frac{\Delta_{z, z^{\prime}} \beta_{t, x^{\prime}}}{-1}}\right]^{-1}$ | Normalization constant so $1=\sum_{z^{-}} \Theta_{t, x, z, z^{-}}$ | D. 2 b |
| $\Delta_{z, z^{\prime}}=^{-} z-z^{\prime}$ | Actual growth increment | D.2c |
| $\alpha_{t, x, z^{\prime}}=\left[\bar{z}_{t, x, z^{\prime}}-z^{\prime}\right] / \beta_{t, x^{-}}$ | Mean molt increment, scaled by $\beta_{t, x^{-}}$ | D.2d |
| $\bar{z}_{t, x, z^{\prime}}=e^{-} e^{a_{t, x}} \cdot z^{\prime \prime}{ }^{b_{t, x^{-}}}$ | Mean size after molt, given pre-molt size $z^{\prime^{-}}$ | D.2e |

In both approaches, the $a_{t, x}, b_{t, x,}$, and $\beta_{t, x}$ are arithmetic-scale parameters with imposed bounds. $\Theta_{t, x, z, z^{\prime}}$ is used to update the numbers-at-size for immature crab, $n_{y, x, z}$, from pre-molt size $z^{\prime}$ to post-molt size $z^{-}$ using:

| $n_{y, x, z^{-}}^{+-}=\sum_{z^{\prime}} \Theta_{t, x, z, z^{\prime}} \cdot n_{y, x, z^{\prime}}$ | numbers at size of immature crab after <br> growth | D. 3 |
| :--- | :--- | :--- |

where $y$ falls within time block $t$ (see also Eq.s A1.4a-b and A2.2a-b).
Priors using normal distributions are imposed on $a_{t, x}$ and $b_{t, x}$ in TCSAM2013, with the values of the hyper-parameters hard-wired in the model code. While priors may be defined for the associated parameters here, these are identified by the user in the model input files and are not hard-wired in the model code.

## E. Model processes: maturity (terminal molt)

Maturation of immature crab in TCSAM02 is based on a similar approach to that taken in TCSAM2013, except that the sex- and size-specific probabilities of terminal molt for immature crab, $\phi_{t, x, z^{-}}$-(where size $z$ is post-molt size), can vary by time block. After molting and growth, the numbers of (new shell) crab at post-molt size $z$ remaining immature, $n_{y, x, I M M, N S, Z}^{+}$, and those maturing, $n_{x, M A T, N S, Z}^{+-}$, are given by:

| $n_{y, \chi, I M M, N S, z^{-}}^{+-}=$ | $\left(1-\phi_{t, x, z}\right) \cdot n_{y, x, I M M, N S, z^{-}}$ | crab remaining immature |
| :--- | :--- | :--- |
| $n_{y, x, M A T, N S, z^{-}}^{+-} \quad \phi_{t, x, z^{-}} \cdot n_{y, x, I M M, N S, z^{-}}$ | E.1a |  |

where $y$ falls in time block $t$ and $n_{y, x, I M M, N S, z}$-is the number of immature, new shell crab of sex $x$ at postmolt size $z$.

The sex- and size-specific probabilities of terminal molt, $\phi_{t, x, z}$, are related to logit-scale model parameters $p_{t, x, Z}^{m a t}$ by:

| $\phi_{t, F E M, z^{-}}= \begin{cases}\frac{1^{-}}{\overline{1}+e^{p_{t, F E M, Z^{-}}^{\text {mat- }}}} & z \leq z_{t, F E M}^{\text {mat }} \\ 1^{-} & z>z_{t, F E M}^{\text {mat }}\end{cases}$ | female probabilities of maturing at <br> post-molt size $z$ | E.2a |
| :--- | :--- | :--- | :--- |
| $\phi_{t, M A L E, z^{-}}=\left\{\begin{array}{lll}\frac{1^{-}}{\overline{1}+e^{p_{t, M A L E, z^{-}}^{\text {mat- }}}} & z \leq z_{t, M A L E}^{\text {mat }} \\ 1^{-} & z>z_{t, M A L E}^{\text {mat }}\end{array}\right.$ | male probabilities of maturing at <br> post-molt size $z$ | E. 2 b |

where the $z_{t, x}^{m a t}$ are constants specifying the minimum pre-molt size at which to assume all immature crab will mature upon molting. The $z_{t, x}^{m a t^{-}}$are used here pedagogically; in actuality, the user specifies the number of logit-scale parameters to estimate (one per size bin starting with the first bin) for each sex, and this determines the $z_{t, x}^{\text {mat }}$ - used above. This parameterization is similar to that implemented in TCSAM2013 for the 2016 assessment model.

Second difference penalties are applied to the parameter estimates in TCSAM2013's objective function to promote relatively smooth changes in these parameters with size. Similar penalties (smoothness, nondecreasing) can be applied in TCSAM02.

## F. Model processes: recruitment

Recruitment in TCSAM02 consists of immature new shell crab entering the population at the end of the model year (June 30). Recruitment in TCSAM02 has a similar functional form to that used in TCSAM2013, except that the sex ratio at recruitment is not fixed at 1:1 and multiple time blocks can be specified. In TCSAM2013, two time blocks were defined: "historical" (model start to 1974) and "current" (1975-present), with "current" recruitment starting in the first year of NMFS survey data. In TCSAM02, recruitment in year $y$ of immature new shell crab of sex $x$ at size $z$ is specified as

| $R_{y, x z^{-}}=\dot{R}_{y}-\ddot{R}_{y, x} \cdot \dddot{R}_{y, z^{-}}$ | recruitment of immature, new shell crab <br> by sex and size bin | F. 1 |
| :--- | :--- | :--- |

where $\dot{R}_{y}$-represents total recruitment in year $y$ and $\ddot{R}_{y, x}$ represents the fraction of sex $x$ crab recruiting, and $\dddot{R}_{y, z}$ is the size distribution of recruits, which is assumed identical for males and females.

Total recruitment in year $y, \dot{R}_{y}$, is parameterized as

| $\dot{R}_{y^{-}}=^{-} e^{p L n R_{t}+\delta R_{t, y^{-}}} y \in t^{-}$ | total recruitment in year $y$ | F. 2 |
| :--- | :--- | :--- |

where $y$ falls within time block $t, p \operatorname{Ln} R_{t}$ is the $\ln$-scale mean recruitment parameter for $t$, and $\delta R_{t, y}$ is an element of a "devs" parameter vector for $t$ (constrained such that the elements of the vector sum to zero over the time block).

The fraction of crab recruiting as sex $x$ in year $y$ in time block $t$ is parameterized using the logistic model

| $\ddot{R}_{y, x}=\left\{\begin{array}{l} \frac{1^{-}}{1+e^{p L g t R x_{t}}} \quad x={ }^{-} M A L E^{-} \\ 1-\ddot{R}_{y, M A L E^{-}} \quad x=\text { FEMALE }^{-} \end{array} \quad y \in t^{-}\right.$ | sex-specific fraction recruiting in year $y$ |
| :---: | :---: |

where $p L g t R x_{t}$ is a logit-scale parameter determining the sex ratio in time block $t$.
The size distribution for recruits in time block $t, \dddot{R}_{t, Z}$, is assumed to be a gamma distribution and is parameterized as

| $\dddot{R}_{t, z^{-}}=^{-} c^{-1} \cdot \Delta_{z} \frac{\alpha_{t}}{\beta_{t}-1} \cdot e^{-\frac{\Delta_{z}^{-}}{\beta_{t^{-}}}}$ | size distribution of recruiting crab | F .4 |
| :--- | :--- | :--- |
| $c_{t}={ }^{-} \sum_{z^{-}} \Delta_{z}^{\frac{\alpha_{t}}{\beta_{t}-1}} \cdot e^{-\frac{\Delta_{z^{-}}}{\beta_{t^{-}}}}$ | normalization constant so that $1=\sum_{z} \dddot{R_{t, z^{-}}}$ | F .5 |
| $\Delta_{z}={ }^{-} z+\delta z / 2-z_{\text {min }^{-}}$ | offset from minimum size bin | F .6 |
| $\alpha_{t}=e^{\text {pLnRa }_{t^{-}}}$ | gamma distribution location parameter | F .7 |
| $\beta_{t}==^{\text {pLnRb }_{t^{-}}}$ | gamma distribution shape parameter | F .8 |

where $p L n R a_{t}$ and $p L n R b_{t}$ are the $\ln$-scale location and shape parameters and the constant $\delta z$ is the size bin spacing.

A final time-blocked parameter, $p \operatorname{LnRCV}_{t}$, is associated with the recruitment process representing the $\ln$ scale coefficient of variation (cv) in recruitment variability in time block $t$. These parameters are used to apply priors on the recruitment "devs" in the model likelihood function.

## G. Selectivity and retention functions

Selectivity and retention functions in TCSAM02 are specified independently from the fisheries and surveys to which they are subsequently applied. This allows a single selectivity function to be "shared" among multiple fisheries and/or surveys, as well as among multiple time block/sex/maturity state/shell condition categories, if so desired.

Currently, the following functions are available for use as selectivity or retention curves in a model:

| $S_{z^{-}}=\left\{1+e^{-\beta \cdot\left(z-z_{50}\right)}\right\}^{-1^{-}}$ | standard logistic | G. 1 |
| :---: | :---: | :---: |
| $S_{z^{-}}=\left\{1+e^{-\beta \cdot\left(z-\exp \left(l n Z_{50}\right)\right)}\right\}^{-1^{-}}$ | logistic w/ alternative parameterization | G. 2 |
| $S_{z^{-}}=\left\{1+e^{-\ln (19) \cdot \frac{\left(z-Z_{50}\right)}{\Delta q_{955}-50}}\right\}^{-1^{-}}$ | logistic w/ alternative parameterization | G. 3 |
| $S_{z^{-}}=\left\{1+e^{\left.-\ln (19) \cdot \frac{-\left(z-z_{50}\right)^{-}}{\exp \left(\ln \Delta Z_{95}-50\right)}\right)^{-1-}}\right.$ | logistic w/ alternative parameterization | G. 4 |
| $S_{z^{-}}=\left\{1+e^{\left.-\ln (19) \cdot \frac{\left(z-\exp \left(\ln Z_{50}\right)\right)}{\exp \left(\ln \Delta z_{95}-50\right)}\right)}\right\}^{-1^{-}}$ | logistic w/ alternative parameterization | G. 5 |
| $S_{z^{-}}=-\frac{1^{-}}{1+e^{-\beta_{a} \cdot\left(z-z_{a 50}\right.}} \cdot \frac{-\frac{1^{-}}{1+e^{\beta_{d} \cdot\left(z-z_{d 50}\right)^{-}}}}{}$ | double logistic | G. 6 |
| $S_{z^{-}}=\frac{1^{-}}{1+e^{-\ln (19) \cdot \frac{\left(z-z_{a 50}\right)^{-}}{\Delta z_{a(95-50)^{-}}}} \cdot-\frac{1^{-}}{1+e^{\ln (19) \cdot \frac{\left(z-z_{d 50}\right)^{-}}{\Delta z_{d(95-50)^{-}}}}} \text {. }}$ | double logistic with alt. parameterization | G. 7 |
|  | double logistic with alt. parameterization | G. 8 |
|  | double logistic with alt. parameterization | G. 9 |
| $S_{z^{-}}=-\frac{1^{-}}{1+e^{-\beta_{a} \cdot\left(z-z_{a 50}\right)}} \cdot \frac{-}{1+e^{\beta_{d} \cdot\left(z-\left[z_{a 50}+\exp \left(\ln Z_{d 50-a 50}\right)\right]\right)^{-}}}$ | double logistic with alt. parameterization | G. 10 |

A double normal selectivity function (requiring 6 parameters to specify) has also been implemented as an alternative to the double logistic functions. In the above functions, all symbols (e.g., $\beta, z_{50}, \Delta z_{95-50}$ ) represent parameter values, except " $z$ " which represents crab size.

Selectivity parameters are defined independently of the functions themselves, and subsequently assigned. It is thus possible to "share" parameters across multiple functions. The "parameters" used in selectivity functions are further divided into mean parameters across a time block and annual deviations within a time block. To accommodate the 6-parameter double normal equation, six "mean" parameter sets ( $p S 1$, $p S 2, \ldots, p S 6$ ) and six associated sets of "devs" parameter vectors ( $p D e v s S 1, p D e v s S 2, \ldots, p D e v s S 6$ ) are defined to specify the parameterization of individual selectivity/retention functions. Thus, for example, $z_{50}$ in eq. F 1 is actually expressed as $z_{50, y^{-}}=\bar{z}_{50}+\delta z_{50, y^{-}}$in terms of model parameters $p S 1$ and $p D e v s S 1_{y}$, where $\bar{z}_{50}=p S 1$ is the mean size-at- $50 \%$-selected over the time period and $\delta z_{50, y^{-}}=^{-}$ $p \operatorname{DevsS} 1_{y}$-is the annual deviation.

Finally, three different options to normalize individual selectivity curves are provided: 1) no normalization, 2 ) specifying a fully-selected size, and 3 ) re-scaling such that the maximum value of the
re-scaled function is 1. A normalization option must be specified in the model input files for each defined selectivity/retention curve.

## H. Fisheries

Unlike TCSAM2013, which explicitly models 4 fisheries that catch Tanner crab (one as a directed fishery, three as bycatch), there is no constraint in TCSAM02 on the number of fisheries that can be incorporated in the model. All fisheries are modeled as "pulse" fisheries occurring at the same time.

TCSAM02 uses the Gmacs approach to modeling fishing mortality (also implemented in TCSAM2013). The total (retained + discards) fishing mortality rate, $F_{f, y, x, m, s, z}$, in fishery $f$ during year $y$ on crab in state $x, m, s$, and $z$ (i.e., sex, maturity state, shell condition, and size) is related to the associated fishery capture rate $\phi_{f, y, x, m, s, z}$-by

| $F_{f, y, x, m, s, z^{-}}=$ | $\left.h_{f, t} \cdot\left(1-\rho_{f, y, x, m, s, z}\right)+\rho_{f, y, x, m, s, z}\right] \cdot \phi_{f, y, x, m, s, z^{-}}$ | fishing mortality rate |
| :--- | :--- | :--- |
| H. 1 |  |  |

where $h_{f, t}$ is the handling (discard) mortality for fishery $f$ in time block t (which includes year $y$ ) and $\rho_{f, y, x, m, s, z}-$ is the fraction of crabs in state $x, m, s, z$ that were caught and retained (i.e., the retention function). The retention function is assumed to be identically 0 for females in a directed fishery and for both sexes in a bycatch fishery.

In TCSAM2013, the same retention function (in each of two time blocks) was applied to male crab regardless of maturity state or shell condition. Additionally, full retention of large males was assumed, such that the retention function essentially reached 1 at large sizes. In TCSAM02, different retention functions can be applied based on maturity state and/or shell condition, and "max retention" is now an (potentially) estimable logit-scale parameter. Thus, in TCSAM02, the retention function $\rho_{f, y, x, m, s, z}$-is given by

| $\rho_{f, y, x, m, s, z^{-}}=-\frac{1^{-}}{1+e^{\rho_{f, t, x, m, s}}} \cdot R_{f, y, x, m, s, z^{-}}$ | retention function | H. 2 |
| :--- | :--- | :--- |

where $f$ corresponds to the directed fishery, $y$ is in time block $t, x=$ MALE, $\rho_{f, t, x, m, s}$ is the corresponding logit-scale "max retention" parameter, and $R_{f, y, x, m, s, z}$-is the associated selectivity/retention curve.

If $n_{y, x, m, s, z}$ is the number of crab classified as $x, m, s, z$ in year $y$ just prior to the prosecution of the fisheries, then

| $c_{f, y, x, m, s, z^{-}}=\frac{\phi_{f, y, x, m, s, z^{-}}}{F_{y, x, m, s, z^{-}}^{T}} \cdot\left[1-e^{-F_{y, x, m, z}^{T}}\right] \cdot n_{y, x, m, s, z^{-}}$ | number of crab <br> captured | H.3 |
| :--- | :--- | :--- |

is the number of crab classified in that state that were captured by fishery $f$, where $F_{y, x, m, s, z^{-}}^{T}=-$ $\sum_{f} F_{f, y, x, m, s, z}$-represents the total (across all fisheries) fishing mortality on those crab. The number of crab retained in fishery $f$ classified as $x, m, s, z$ in year $y$ is given by

| $r_{f, y, x, m, s, z^{-}}=\frac{\rho_{f, y, x, m, s, z^{-}} \phi_{f, y, x, m, s, z^{-}}}{F_{y, x, m, s, z^{-}}^{T}} \cdot\left[1-e^{-F_{y, x, m, s, z}^{T}}\right] \cdot n_{y, x, m, s, z^{-}}$ | number of <br> retained crab | H. 4 |
| :--- | :--- | :---: |

while the number of discarded crab, $d_{f, y, x, m, s, z}$, is given by

$$
\begin{array}{|l|l|l|}
\hline d_{f, y, x, m, s, z^{-}}=\frac{\left(1-\rho_{f, y, x, m, s, z}\right) \cdot \phi_{f, y, x, m, s, z^{-}}}{F_{y, x, m, s, z^{-}}^{T}} \cdot\left[1-e^{-F_{y, x, m, s, z}^{T}}\right] \cdot n_{y, x, m, s, z^{-}} & \begin{array}{l}
\text { number of } \\
\text { discarded crab }
\end{array} & \text { H. } 5 \\
\hline
\end{array}
$$

and the discard mortality, $d m_{f, y, x, m, s, z}$, is

$$
\begin{array}{|l|l|l|}
\hline d m_{f, y, x, m, s, z^{-}}=\frac{h_{f, y^{-}}\left(1-\rho_{f, y, x, m, s, z}\right) \cdot \phi_{f, y, x, m, s, z^{-}}}{F_{y, x, m, s, z^{-}}^{T}} \cdot\left[1-e^{\left.-F_{y, x, m, z}^{T}\right]} \cdot n_{y, x, m, s, z^{-}}\right. & \begin{array}{l}
\text { discard } \\
\text { mortality } \\
\text { (numbers) }
\end{array} & \text { H. } 6 \\
\hline
\end{array}
$$

The capture rate $\phi_{f, y, x, m, s, z}$ - not the fishing mortality rate $\left.F_{f, y, x, m, s, z}\right)$ is modeled as a function separable into separate year and size components such that

| $\phi_{f, y, x, m, s, z^{-}}={ }^{-} \phi_{f, y, x, m, s} \cdot S_{f, y, x, m, s, z^{-}}$ | fishing capture <br> rate | H. 7 |
| :--- | :--- | :--- |

where $\phi_{f, y, x, m, s}$ is the fully-selected capture rate in year $y$ and $S_{f, y, x, m, s, z}$-is the size-specific selectivity.
The fully-selected capture rate $\phi_{f, y, x, m, s}$ for $y$ in time block $t$ is parameterized in the following manner:

$$
\begin{array}{|l|l}
\hline \phi_{f, y, x, m, s}=-\exp \left(\ln C_{f, t, x, m, s}+p \operatorname{Devs} C_{f, y, x, m, s}\right)^{-} & \text {H. } 8
\end{array}
$$

where the $p \operatorname{Devs} C_{f, y, x, m, s}$ are elements for year $y$ in time block $t$ of a "devs" vectors representing annual variations from the $\ln$-scale mean fully-selected capture rate $\ln C_{f, t, x, m, s}$. The latter is expressed in terms of model parameters as

| $\ln C_{f, t, x, m, s}=p \operatorname{Ln} C_{f, t, x, m, s}+\sum_{i=1}^{4} \delta C_{f, t, x, m, s^{-}}^{i}$ | H. 9 |
| :--- | :---: |

where the $p L n C_{f, t, x, m, s}$ is the mean $\ln$-scale capture rate (e.g., for mature males) and the $\delta C_{f, t, x, m, s}^{i}$-are $\ln -$ scale offsets.

## I. Surveys

If $n_{y, x, m, s, z}$ is the number of crab classified as $x, m, s, z$ in year $y$ just prior to the prosecution of a survey, then the survey abundance, $a_{v, y, x, m, s, z}$, of crab classified in that state by survey $v$ is given by

| $v, y, x, m, s, z^{-}={ }^{-} q_{v, y, x, m, s, z^{-}} n_{y, x, m, s, z^{-}}$ | survey abundance | I.1 |
| :--- | :--- | :---: |

where $q_{v, y, x, m, s, z}$-is the size-specific survey catchability on this component of the population.
The survey catchability $q_{v, y, x, m, s, z}$-is decomposed in the usual fashion into separate time block and size components such that, for $y$ in time block $t$ :

| $q_{v, y, x, m, s, z^{-}}=q_{v, t, x, m, s} \cdot S_{v, t, x, m, s, z^{-}} \cdot A_{v, t, x, m, s, z^{-}}$ | survey catchability | I. 2 |
| :--- | :--- | :---: |

where $q_{v, t, x, m, s}$ is the fully-selected catchability in time block $t, S_{v, t, x, m, s, z}-$ is the size-specific survey selectivity, and $A_{v, t, x, m, s, z}$-is the size-specific availability of the population to the survey. If the survey covers the complete stock area (as the standard NMFS EBS bottom trawl is assumed to do for Tanner
crab), then $A_{v, t, x, m, s, z} \equiv 1$. However, if the survey does not cover the complete stock, as is the case with the BSFRF/NMFS side-by-side catchability studies, then $A_{v, t, x, m, s, z}$-needs to be estimated or assumed.

The fully-selected catchability $q_{v, t, x, m, s}$ is parameterized in a fashion similar to that for fully-selected fishery capture rates (except that annual "devs" are not included) in the following manner:

| $q_{v, t, x, m, s}=\exp ^{-}\left(p L n Q_{v, t, x, m, s}+\sum_{i=1^{-}}^{4^{-}} \delta Q_{v, t, x, m, s}^{i^{-}}\right)^{-}$ | I. 3 |
| :--- | :---: |

where the $p L n Q_{v, t, x, m, s}$ is the mean $\ln$-scale catchability (e.g., for mature males) and the $\delta Q_{v, t, x, m, s}^{i}$-are $\ln$ scale offsets.

## J. Model fitting: objective function equations

The TCSAM02 model is fit by minimizing an objective function, $\sigma$, with additive components consisting of: 1) negative log-likelihood functions based on specified prior probability distributions associated with user-specified model parameters, and 2) several negative log-likelihood functions based on input data components, of the form:

| $\sigma=^{-}-2 \sum_{p^{-}} \lambda_{p}^{-} \cdot \ln \left(\wp_{p}\right)^{-}-2 \sum_{l^{-}} \lambda_{l}^{-} \cdot \ln \left(\mathcal{L}_{l}\right)^{-}$ | model objective function | J. 1 |
| :--- | :--- | :--- |

where $\wp_{p}$ represents the $p$ th prior probability function, $\mathcal{L}_{l}$ represents the $l$ th likelihood function, and the $\lambda$ 's represent user-adjustable weights for each component.

## Prior Probability Functions

Prior probability functions can be associated with each model parameter or parameter vector by the user in the model input files (see Section L below for examples on specifying priors).

## Likelihood Functions

The likelihood components included in the model's objective function are based on normalized size frequencies and time series of abundance or biomass from fishery or survey data. Survey data optionally consists of abundance and/or biomass time series for males, females, and/or all crab (with associated survey cv's), as well as size frequencies by sex, maturity state, and shell condition. Fishery data consists of similar data types for optional retained, discard, and total catch components.

## Size frequency components

Likelihood components involving size frequencies are based on multinomial sampling:

| $\ln (\mathcal{L}))^{-} \sum_{y^{-}} n_{y, c^{-}} \sum_{z^{-}}\left\{p_{y, c, z^{-}}^{o b s^{-}} \cdot \ln \left(p_{y, c, z}^{m o d^{-}}+\delta\right)-p_{y, c, z^{-}}^{o b s^{-}} \cdot \ln \left(p_{y, c, z^{-}}^{o b s^{-}}+\delta\right)\right\}^{-}$ | multinomial <br> log-likelihood | J. 2 |
| :--- | :--- | :---: |

where the $y$ 's are years for which data exists, " $c$ " indicates the population component classifiers (i.e., sex, maturity state, shell condition) the size frequency refers to, $n_{y, c}$-is the classifier-specific effective sample size for year $\mathrm{y}, p_{y, c, z}^{o b s}$-is the observed size composition in size bin $z$ (i.e., the size frequency normalized to sum to 1 across size bins for each year), $p_{y, c, z}^{m o{ }^{-}}$is the corresponding model-estimated size composition, and $\delta$ is a small constant. The manner in which the observed and estimated size frequencies for each data component are aggregated (e.g., over shell condition) prior to normalization is specified by the user in the model input files. Data can be entered in input files at less-aggregated levels of than will be used in the model; it will be aggregated in the model to the requested level before fitting occurs.

## Aggregated abundance/biomass components

Likelihood components involving aggregated (over size, at least) abundance and or biomass time series can be computed using one of three potential likelihood functions: the normal, the lognormal, and the "norm2". The likelihood function used for each data component is user-specified in the model input files.

The $\ln$-scale normal likelihood function is

| $\ln \left(\mathcal{L}^{N}\right)_{c^{-}}=-\frac{1}{2} \sum_{y^{-}}\left\{\frac{\left[a_{y, c}^{\theta b s^{-}}-a_{y, c}^{\bmod }\right]^{2}}{\sigma_{y, c^{-}}^{2-}}\right\}^{-}$ | normal log- <br> likelihood | J. 3 |
| :--- | :--- | :---: |

where $a_{y, c}^{o b s^{-}}$is the observed abundance/biomass value in year $y$ for aggregation level $c, a_{y, c}^{m o d}$ is the associated model estimate, and $\sigma_{y, c}^{2}$-is the variance associated with the observation.

The $\ln$-scale lognormal likelihood function is

| $\ln \left(\mathcal{L}^{L N}\right)_{c^{-}}=-\frac{1}{2} \sum_{y^{-}}\left\{\frac{\left[\ln \left(a_{y, c}^{\text {®bs }}+\delta\right)-\ln \left(a_{y, c}^{\text {mod }}+\delta\right)\right]^{-}}{\sigma_{y, c^{-}}^{2-}}\right\}^{-}$ | lognormal log- <br> likelihood | J. 4 |
| :--- | :--- | :--- |

where $a_{y, c}^{o b s^{-}}$is the observed abundance/biomass value in year $y$ for aggregation level $c, a_{y, c}^{m o d}$ is the associated model estimate, and $\sigma_{y, c}^{2}$-is the ln -scale variance associated with the observation.

For consistency with TCSAM2013, a third type, the "norm2", may also be specified

| $\ln \left(\mathcal{L}^{N 2}\right)_{x^{-}}=^{-}-\frac{1}{2} \sum_{y^{-}}\left[a_{y, x}^{* b s^{-}}-a_{y, x}^{m o d}\right]^{2^{-}}$ | "norm2" log-likelihood | J.5 |
| :--- | :--- | :--- |

This is equivalent to specifying a normal $\log$-likelihood with $\sigma_{y, x}^{2} \equiv 1.0$. This is the standard likelihood function applied in TCSAM2013 to fishery catch time series.

## Growth data

Growth (molt increment) data can be fit as part of a TCSAM02 model. Multiple datasets can be fit at the same time. The likelihood for each dataset $\left(\mathrm{L}_{d}\right)$ is based on the same gamma distribution used in the growth model:

| $\mathrm{L}_{d}=--\sum_{i \in d^{-}} \ln \left\{\Gamma\left(\frac{\tilde{z}_{i}-\bar{z}_{y_{\bar{i}}, x_{i}, z_{\bar{i}}^{-}}}{\beta_{y_{\bar{i}}, x_{i}-}}\right)\right\}$ | gamma log-likelihood | J. 6 |
| :--- | :--- | :--- |

where $z_{i}$ and $\tilde{z}_{i}$ are the pre-molt and post-molt sizes for individual $i$ (of sex $x_{i}$ collected in year $y_{i}$ ) in dataset $d$, respectively, $\bar{z}_{y_{i}, x_{i}, z_{i}}$ is the predicted mean post-molt size for individual $i$, and $\beta_{y_{i}, x_{i}}$ is the scale factor for the gamma distribution corresponding to individual $i$.

## Maturity ogive data

Annual maturity ogive data, the observed proportions-at-size of mature crab in a given year, can also be fit as part of a TCSAM02 model. This data consists of proportions of mature crab observed within a size bin, as well as the total number of observations for that size bin. The proportions are assumed to represent the fraction of new shell mature crab (i.e., having gone through terminal molt within the previous growth season) to all new shell crab within the size bin in that year. Multiple datasets can be fit at the same time. The likelihood for each observation is based on a binomial distribution with sample size equal to the
number of observations within the corresponding size bin, so the likelihood for each dataset $\left(L_{m}\right)$ is given by:

$$
\begin{array}{|l|l|l|}
\hline \mathrm{L}_{m}==_{y, Z^{-}}^{-} n_{y, z^{-}}\left\{p_{y, z^{-}}^{o b s^{-}} \cdot \ln \left(p_{y, z}^{\text {mod }}+\delta\right)+\left(1-p_{y, z}^{o b s}\right) \cdot \ln \left(1-p_{y, z}^{\text {mod }}+\delta\right)\right\}^{-} & \begin{array}{l}
\text { binomial log- } \\
\text { likelihood }
\end{array} & \text { J. } 7 \\
\hline
\end{array}
$$

where $y$ is a year, $z$ is a size bin, $n_{y, z}-$ is the total number of classified crab in size bin $z$ in year $y, p_{y, z}^{o b s}$ is the observed ratio of mature, new shell males to total new shell males in size bin z in year $\mathrm{y}, p_{y, z}^{o b s^{-}}$is the corresponding model-predicted ratio, and $\delta$ is a small constant to prevent trying to calculate $\ln (0)$.

## Effort data

In both TCSAM2013 and TCSAM02, fishery-specific effort data is used to predict annual fully-selected fishery capture rates for Tanner crab bycatch in the snow crab and Bristol Bay red king crab fisheries in the period before at-sea observer data is available (i.e., prior to 1991), based on the assumed relationship

$$
F_{f, y^{-}}=q_{f} \cdot E_{f, y^{-}}
$$

where $F_{f, y}$-is the fully-selected capture rate in fishery $f$ in year $y, q_{f}$ is the estimated catchability in fishery f, and $E_{f, y}$-is the reported annual, fishery-specific effort (in pots). In TCAM2013, the fishery $q$ 's are estimated directly from the ratio of fishery mean $F$ to mean $E$ over the time period ( $t_{f}$ ) when at-sea observer data is available from which to estimate the $F_{f, y}$ 's as parameters:

$$
q_{f}=\frac{\sum_{y \in t_{f}} F_{f, y^{-}}}{\sum_{y \in t_{f}} E_{f, y^{\prime}}}
$$

Note that, in this formulation, the fishery $q$ 's are not parameters (i.e., estimated via maximizing the likelihood) in the model. In TCSAM2013, the time period over which $q$ is estimated for each fishery is hard-wired. This approach is also available as an option in TCSAM02, although different time periods for the averaging can be specified in the model options file.

A second approach to effort extrapolation in which the fishery $q$ 's are fully-fledged parameters estimated as part of maximizing the likelihood is provided in TCSAM02 as an option, as well. In this case, the effort data is assumed to have a lognormal error distribution and the following negative log-likelihood components are included in the overall model objective function:

$$
L_{f}=\sum_{y^{-}} \frac{\left(\ln \left(E_{f, y}+\delta\right)-\ln \left(\frac{F_{f, y}}{q_{f}}+\delta\right)\right)^{2^{-}}}{2 \cdot \sigma_{f}^{2^{-}}}
$$

where $\sigma_{f}^{2}$ is the assumed $\ln$-scale variance associated with the effort data and $\delta$ is a small value so that the arguments of the $\ln$ functions do not go to zero.

## Aggregation fitting levels

A number of different ways to aggregate input data and model estimates prior to fitting likelihood functions have been implemented in TCSAM02. These include:

| Abundance/Biomass | Size Conpositions |  |
| :---: | :---: | :---: |
| by | by | extended by |
| total | total | X |
| x |  | x, m |
| $x$, mature only | x | -- |
| $x, m$ |  | m |
| $x, \mathrm{~s}$ |  | s |
| $x, m, s$ | $\mathrm{x}, \mathrm{m}$ | -- |
|  |  | S |
|  | $x, \mathrm{~s}$ |  |
|  | $\mathrm{x}, \mathrm{m}, \mathrm{s}$ |  |

where $x, m, s$ refer to sex, maturity state and shell condition and missing levels are aggregated over. For size compositions that are "extended by" $x, m, s$, or $\{x, m\}$, this involves appending the size compositions corresponding to each combination of "extended by" factor levels, renormalizing the extended composition to sum to 1 , and then fitting the extended composition using a multinomial likelihood.

## K. Devs vectors

For TCSAM02 to accommodate arbitrary numbers of fisheries and time blocks, it is necessary to be able to define arbitrary numbers of "devs" vectors. This is currently not possible using the ADMB C++ libraries, so TCSAM02 uses an alternative implementation of devs vectors from that implemented in ADMB. For the 2017 assessment, an $n$-element "devs" vector was implemented using an $n$-element bounded parameter vector. with the final element of the "devs" vector defined as $-\sum_{n-1} v_{i}$, where $v_{i}$ was the ith value of the parameter (or devs) vector, so that the sum over all elements of the devs vector was identically 0 . Penalties were placed on the final element of the devs vector to ensure it was bounded in the same manner as the parameter vector. However, this approach was problematic when initializing the model with the values for the $n$-1 elements that defined the $n$-element devs vector, the value of the $n$-th element ( $-\sum_{n-1} v_{i}$ ) was not guaranteed to satisfy the bounds placed on the vector. Thus, this approach was revised to allow specification of all n element values (the $v_{n}=-\sum_{n-1} v_{i}$-constraint was removed) while the likelihood penalty was changed to ensure the sum of the elements was 0 . The new approach also has the advantage that it more closely follows the one used in ADMB to define "devs" vectors. Test runs with both approaches showed no effect on convergence to the MLE solution.

## L. Priors for model parameters

A prior probability distribution can be specified for any element of model parameter. The following distributions are available for use as priors:

| indicator | parameters | constants | description |
| :--- | :--- | :--- | :--- |
| none | none | none | no prior applied |
| ar1_normal | $\mu, \sigma^{-}$ | none | random walk with normal deviates |
| cauchy | $x_{0}, \gamma^{-}$ | none | Cauchy pdf |
| chisquare | $v^{-}$ | none | $\chi^{2}$ pdf |
| constant | $\min , \max$ | none | uniform pdf |
| exponential |  | none | exponential pdf |
| gamma | $r, \mu^{-}$ | none | gamma pdf |
| invchisquare | $v^{-}$ | none | inverse $\chi^{2^{-}}$pdf |


| invgamma | $r, \mu^{-}$ | none | inverse gamma pdf |
| :--- | :--- | :--- | :--- |
| invgaussian | $\mu_{,}^{-}$ | none | inverse Gaussian pdf |
| lognormal | median, CV | none | lognormal pdf |
| logscale_normal | median, CV | none | normal pdf on ln-scale |
| normal | $\mu, \sigma^{-}$ | none | normal pdf |
| scaled_invchisquare | $v, s^{-}$ | none | inverse $\chi^{2^{-}}$scaled pdf |
| scaledCV_invchisquare | $v, C V^{-}$ | none | inverse $\chi^{2}$ pdf, scaled by CV |
| t | $v^{-}$ | none | t distribution |
| truncated_normal | $\mu, \sigma^{-}$ | min, max | truncated normal pdf |

## M. Parameters and other information determined outside the model

Several nominal model parameters are not estimated in the model, rather they are fixed to values determined outside the model. These include Tanner crab handling mortality rates for discards in the crab fisheries ( $32.1 \%$ ), the groundfish trawl fisheries ( $80 \%$ ), and the groundfish pot fisheries ( $50 \%$ ), as well the base rate for natural mortality ( $0.23 \mathrm{yr}^{-1}$ ). Sex- and maturity-state-specific parameters for individual weight-at-size have also been determined outside the model, based on fits to data collected on the NMFS EBS bottom trawl survey (Daly et al., 2016). Weight-at-size, $w_{x, m, z}$, is given by

$$
w_{x, m, z^{-}}=a_{x, m} \cdot z^{b_{x, m^{-}}}
$$

where

| sex | maturity state | $\boldsymbol{a}_{\boldsymbol{x}, \boldsymbol{m}^{-}}$ | $\boldsymbol{b}_{\boldsymbol{x}, \boldsymbol{m}^{-}}$ |
| :--- | :--- | :--- | :--- |
| male | all states | 0.000270 | 3.022134 |
|  | immature | 0.000562 | 2.816928 |
|  | mature | 0.000441 | 2.898686 |

and size is in mm CW and weight is in kg .
N. OFL calculations and stock status determination

Overfishing level (OFL) calculations and stock status determination for Tanner crab are based on Tier 3 considerations for crab stocks as defined by the North Pacific Fishery Management Council (NPFMC; NPFMC 2016). Tier 3 considerations require life history information such as natural mortality rates, growth, and maturity but use proxies based on a spawner-per-recruit approach for $\mathrm{F}_{\text {MSY }}, \mathrm{B}_{\mathrm{MSY}}$, and MSY because there is no reliable stock-recruit relationship.


Fig. 2. The Fofl harvest control rule. Equilibrium recruitment is assumed to be equal to the average recruitment over a selected time period (1982-present for Tanner crab). For Tier 3 stocks, the proxy for $\mathrm{B}_{\text {MSY }}$ is defined as $35 \%$ of longterm (equilibrium) mature male biomass (MMB) for the unfished stock $\left(\mathrm{B}_{0}\right)$. The proxy $\mathrm{F}_{\text {MSY }}$ for Tier 3 stocks is then the directed fishing mortality rate that results in $\mathrm{B}_{35 \%}$ (i.e., $\mathrm{F}_{35 \%}$ ), while the MSY proxy is the longterm total (retained plus discard) catch mortality resulting from fishing at $\mathrm{F}_{\text {MSY }}$. The OFL calculation for the upcoming year is based on a sloping
harvest control rule for $\mathrm{F}_{\text {OFL }}$ (Fig. 2), the directed fishing mortality rate that results in the OFL. If the "current" MMB (projected to Feb. 15 of the upcoming year under the $\mathrm{F}_{\text {OFL }}$ ) is above $\mathrm{B}_{\text {MSY }}\left(\mathrm{B}_{35 \%}\right)$, then $\mathrm{F}_{\text {OFL }}=\mathrm{F}_{\mathrm{MSY}}=\mathrm{F}_{35 \%}$. If the current MMB is between $\beta \cdot B_{M S Y}$-and $\mathrm{B}_{\mathrm{MSY}}$, then $\mathrm{F}_{\text {OFL }}$ is determined from the slope of the control rule. In either of these cases, the OFL is simply the projected total catch mortality under directed fishing at $\mathrm{F}_{\text {OFL }}$. If current MMB is less than $\beta \cdot B_{M S Y}$, then no directed fishing is allowed $\left(\mathrm{F}_{\mathrm{OFL}}=0\right)$ and the OFL is set to provide for stock rebuilding with bycatch in non-directed fisheries. Note that if current MMB is less than $\mathrm{B}_{\mathrm{MSY}}$, then the process of determining $\mathrm{F}_{\mathrm{OFL}}$ is generally an iterative one.

Stock status is determined by comparing "current" MMB with the Minimum Stock Size Threshold (MSST), which is defined as $0.5 x \mathrm{~B}_{\text {MSY: }}$ if "current" MMB is below the MSST, then the stock is overfished-otherwise, it is not overfished.

## N. 1 Equilibrium conditions

Both OFL calculations and stock status determination utilize equilibrium considerations, both equilibrium under unfished conditions (to determine $\mathrm{B}_{0}$ and $\mathrm{B}_{35 \%}$ ) and under fished conditions (to determine $\mathrm{F}_{35 \%}$ ). For Tier 3 stocks, because there is no reliable stock-recruit relationship, analytical solutions can be found for equilibrium conditions for any fishing mortality conditions. These solutions are described below (the notation differs somewhat from that used in previous sections).

## N.1.1 Population states

The Tanner crab population on July 1 can be characterized by abundance-at-size in four population states:
in- immature new shell crab
io- immature old shell crab
$m n$ - mature new shell crab
mo - mature old shell crab
where each of these states represents a vector of abundance-at-size (i.e., a vector subscripted by size).

## N.1.2 Population processes

The following processes then describe the dynamics of the population over a year:
$S_{l}$ - survival from start of year to time of molting/growth of immature crab, possibly including fishing mortality (a diagonal matrix)
$S_{2}$ - survival after time of molting/growth of immature crab to end of year, possibly including fishing mortality (a diagonal matrix)
$\Phi$ - probability of an immature crab molting $(\operatorname{pr}(\operatorname{molt} \mid z)$, where $z$ is pre-molt size; a diagonal matrix) ( $\operatorname{pr}(\operatorname{molt} \mid z)$ is assumed to be 1 in TCSAM02).
$\Theta$ - probability that a molt was terminal ( $\operatorname{pr}($ molt to maturity $\mid z$, molt), where $z$ is post-molt size; a diagonal matrix)
$T$ - size transition matrix (a non-diagonal matrix)
1 - identity matrix
$R$-number of recruits by size (a vector)
The matrices above are doubly-subscripted, and $R$ is singly-subscripted, by size. Additionally, the matrices above (except for the identity matrix) can also be subscripted by population state (in, io, mn, mo) for generality. For example, survival of immature crab may differ between those that molted and those that skipped.

## N.1.3 Population dynamics

The following equations then describe the development of the population from the beginning of one year to the beginning of the next:

$$
\begin{align*}
& i n^{+}=R+S_{2 i n} \cdot\left\{\left(1-\Theta_{i n}\right) \cdot T_{i n} \cdot \Phi_{i n} \cdot S_{1 i n} \cdot i n+T_{i o} \cdot\left(1-\Theta_{i o}\right) \cdot \Phi_{i o} \cdot S_{1 i o} \cdot i o\right\}^{-}  \tag{N.1}\\
& i o^{+}=S_{2 i o} \cdot\left\{\left(1-\Phi_{i n}\right) \cdot S_{1 i n} \cdot i n+\left(1-\Phi_{i o}\right) \cdot S_{1 i o} \cdot i o\right\}^{-}  \tag{N.2}\\
& m n^{+}=S_{2 m n} \cdot\left\{\Theta_{i n} \cdot T_{i n} \cdot \Phi_{i n} \cdot S_{1 i n} \cdot i n+\Theta_{i o} \cdot T_{i o} \cdot \Phi_{i o} \cdot S_{1 i o} \cdot i o\right\}^{-}  \tag{N.3}\\
& m o^{+}=S_{2 m o} \cdot\left\{S_{1 m n} \cdot m n+S_{1 m o} \cdot m o\right\}^{-} \tag{N.4}
\end{align*}
$$

where " + " indicates year +1 and all recruits $(R)$ are assumed to be new shell.

## N.1.4 Equilibrium equations

The equations reflecting equilibrium conditions (i.e., in $^{+}=i n$, etc.) are simply:

$$
\begin{align*}
& \text { in }=R+S_{2 i n} \cdot\left\{\left(1-\Theta_{i n}\right) \cdot T_{i n} \cdot \Phi_{i n} \cdot S_{1 i n} \cdot i n+\left(1-\Theta_{i o}\right) \cdot T_{i o} \cdot \Phi_{i o} \cdot S_{1 i o} \cdot i o\right\}^{-}  \tag{N.5}\\
& i o=S_{2 i o} \cdot\left\{\left(1-\Phi_{i n}\right) \cdot S_{1 i n} \cdot i n+\left(1-\Phi_{i o}\right) \cdot S_{1 i o} \cdot i o\right\}^{-}  \tag{N.6}\\
& m n=S_{2 m n} \cdot\left\{\Theta_{i n} \cdot T_{i n} \cdot \Phi_{i n} \cdot S_{1 i n} \cdot i n+\Theta_{i o} \cdot T_{i o} \cdot \Phi_{i o} \cdot S_{1 i o} \cdot i o\right\}^{-}  \tag{N.7}\\
& m o=S_{2 m o} \cdot\left\{S_{1 m n} \cdot m n+S_{1 m o} \cdot m o\right\}^{-} \tag{N.8}
\end{align*}
$$

where $R$ above is now the equilibrium (longterm average) number of recruits-at-size vector.

## N.1.5 Equilibrium solution

The equilibrium solution can be obtained by rewriting the above equilibrium equations as:

$$
\begin{align*}
& \text { in }=R+A \cdot i n+B \cdot i o^{-}  \tag{N.9}\\
& i o=C \cdot i n+D \cdot i o^{-}  \tag{N.10}\\
& m n=E \cdot i n+F \cdot i o^{-}  \tag{N.11}\\
& m o=G \cdot m n+H \cdot \mathrm{mo}^{-} \tag{N.12}
\end{align*}
$$

where $A, B, C, D, E, F, G$, and $H$ are square matrices. Solving for $i o$ in terms of in in eq. 10 , one obtains

$$
\begin{equation*}
i o=\{1-D\}^{-1} \cdot C \cdot i n^{-} \tag{N.13}
\end{equation*}
$$

Plugging eq. 13 into 9 and solving for in yields

$$
\begin{equation*}
\text { in }=\left\{1-A-B \cdot[1-D]^{-1} \cdot C\right\}^{-1} \cdot R^{-} \tag{N.14}
\end{equation*}
$$

Equations 13 for io and 14 for in can simply be plugged into eq. 11 to yield $m n$ :

$$
\begin{equation*}
m n=E \cdot i n+F \cdot i o^{-} \tag{N.15}
\end{equation*}
$$

while eq. 12 can then be solved for $m o$, yielding:

$$
\begin{equation*}
m o=\{1-H\}^{-1} \cdot G \cdot m n^{-} \tag{N.16}
\end{equation*}
$$

where (for completeness):

$$
\begin{align*}
& A=S_{2 i n} \cdot\left(1-\Theta_{i n}\right) \cdot T_{i n} \cdot \Phi_{i n} \cdot S_{1 i n^{-}}  \tag{N.17}\\
& B=S_{2 i n} \cdot\left(1-\Theta_{i o}\right) \cdot T_{i o} \cdot \Phi_{i o} \cdot S_{1 i o^{-}}  \tag{N.18}\\
& C=S_{2 i o} \cdot\left(1-\Phi_{i n}\right) \cdot S_{1 i n^{-}}  \tag{N.19}\\
& D=S_{2 i o} \cdot\left(1-\Phi_{i o}\right) \cdot S_{1 i o^{-}}  \tag{N.20}\\
& E=S_{2 m n} \cdot \Theta_{i n} \cdot T_{i n} \cdot \Phi_{i n} \cdot S_{1 i n^{-}} \tag{N.21}
\end{align*}
$$

$$
\begin{align*}
& F=S_{2 m n} \cdot \Theta_{i o} \cdot T_{i o} \cdot \Phi_{i o} \cdot S_{1 i o^{-}}  \tag{N.22}\\
& G=S_{2 m o} \cdot S_{1 m n^{-}}  \tag{N.23}\\
& H=S_{2 m o} \cdot S_{1 m o^{-}} \tag{N.24}
\end{align*}
$$

Note that $\Theta$, the size-specific conditional probability of a molt being the terminal molt-to-maturity, is defined above on the basis of post-molt, not pre-molt, size. This implies that whether or not a molt is terminal depends on the size a crab grows into, not the size it at which it molted. An alternative approach would be to assume that the conditional probability of terminal molt is determined by pre-molt size. This would result in an alternative set of equations, but these can be easily obtained from the ones above by simply reversing the order of the terms involving $T$ and $\Theta$ (e.g., the term $\left(1-\Theta_{i n}\right) \cdot T_{i n}$-becomes $T_{i n} .^{-}$ $\left(1-\Theta_{i n}\right)$ ).

## N. 2 OFL calculations

Because a number of the calculations involved in determining the OFL are iterative in nature, the OFL calculations do not involve automatically-differentiated (AD) variables. Additionally, they are only done after model convergence or when evaluating an MCMC chain. The steps involved in calculating the OFL are outlined as follows:

1. The initial population numbers-at-sex/maturity state/shell condition/size for the upcoming year are copied to a non-AD array.
2. Mean recruitment is estimated over a pre-determined time frame (currently 1982-present).
3. The arrays associated with all population rates in the final year are copied to non-AD arrays for use in the upcoming year.
4. Calculate the average selectivity and retention functions for all fisheries over the most recent 5year period.
5. Determine the average maximum capture rates for all fisheries over the most recent 5-year period.
6. Using the equilibrium equations, calculate $\mathrm{B}_{0}$ for unfished stock $\left(\mathrm{B} 35 \%=0.35 * \mathrm{~B}_{0}\right)$.
7. Using the equilibrium equations, iterate on the maximum capture rate for males in the directed fishery to find the one $\left(\mathrm{F}_{35 \%}\right)$ that results in the equilibrium $\mathrm{MMB}=\mathrm{B}_{35 \%}$.
8. Calculate "current" MMB under directed fishing at $\mathrm{F}=\mathrm{F}_{35 \%}$ by projecting initial population (1) to Feb. 15.
a. If current $\mathrm{MMB}>\mathrm{B}_{35 \%}, \mathrm{~F}_{\mathrm{OFL}}=\mathrm{F}_{35 \%}$. The associated total catch mortality is OFL.
b. Otherwise
i. set directed F based on the harvest control rule and the ratio of the calculated current MMB to $\mathrm{B}_{35 \%}$
ii. recalculate current MMB
iii. iterate i-iii until current MMB doesn't change between iterations. Then $F_{O F L}={ }^{-}$ $F\left(<F_{35 \%}\right)$ and the OFL is the associated total (retained plus discard) catch mortality.

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# Model Comparisons: Fits to Size Composition Data M19F00a vs M19F01 vs M19F02 vs M19F03 vs M19F04 vs M19F05 

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07 September, 2019

## Contents

Model fits to size compositions, by year 1
Survey size compositions 2
Fishery retained catch size compositions 43
Fishery total catch size compositions 47

## Model fits to size compositions, by year

Fits to the size composition data available to the model(s) are presented in this section as line plots by year. Not all of the fits presented are necessarily included in the parameter optimization for each model; some fits to datasets for a particular model may be included for comparison purposes with other models which include those data in their optimization. The reader should consult the main assessment document to determine which fits are included in the optimization for any particular model.

## Survey size compositions



Figure 1: Comparison of observed and predicted male, immature, all shell survey size comps for NMFS 0 . Page 1 of 5 .

NMFS 0: male, immature, all shell


Figure 2: Comparison of observed and predicted male, immature, all shell survey size comps for NMFS 0 . Page 2 of 5.

NMFS 0: male, immature, all shell


Figure 3: Comparison of observed and predicted male, immature, all shell survey size comps for NMFS 0 . Page 3 of 5 .

NMFS 0: male, immature, all shell


Figure 4: Comparison of observed and predicted male, immature, all shell survey size comps for NMFS 0 . Page 4 of 5.

NMFS 0: male, immature, all shell


Figure 5: Comparison of observed and predicted male, immature, all shell survey size comps for NMFS 0 . Page 5 of 5 .

NMFS 0: male, mature, all shell


Figure 6: Comparison of observed and predicted male, mature, all shell survey size comps for NMFS 0 . Page 1 of 5 .

NMFS 0: male, mature, all shell


Figure 7: Comparison of observed and predicted male, mature, all shell survey size comps for NMFS 0 . Page 2 of 5.

NMFS 0: male, mature, all shell


Figure 8: Comparison of observed and predicted male, mature, all shell survey size comps for NMFS 0 . Page 3 of 5 .

NMFS 0: male, mature, all shell


Figure 9: Comparison of observed and predicted male, mature, all shell survey size comps for NMFS 0 . Page 4 of 5.

NMFS 0: male, mature, all shell


Figure 10: Comparison of observed and predicted male, mature, all shell survey size comps for NMFS 0. Page 5 of 5.

NMFS 0: female, immature, all shell


Figure 11: Comparison of observed and predicted female, immature, all shell survey size comps for NMFS 0. Page 1 of 5.

NMFS 0: female, immature, all shell


Figure 12: Comparison of observed and predicted female, immature, all shell survey size comps for NMFS 0. Page 2 of 5.

NMFS 0: female, immature, all shell


Figure 13: Comparison of observed and predicted female, immature, all shell survey size comps for NMFS 0 . Page 3 of 5 .

NMFS 0: female, immature, all shell


Figure 14: Comparison of observed and predicted female, immature, all shell survey size comps for NMFS 0. Page 4 of 5.

NMFS 0: female, immature, all shell


Figure 15: Comparison of observed and predicted female, immature, all shell survey size comps for NMFS 0. Page 5 of 5.

NMFS 0: female, mature, all shell


Figure 16: Comparison of observed and predicted female, mature, all shell survey size comps for NMFS 0 . Page 1 of 5 .

NMFS 0: female, mature, all shell


Figure 17: Comparison of observed and predicted female, mature, all shell survey size comps for NMFS 0 . Page 2 of 5.

NMFS 0: female, mature, all shell


Figure 18: Comparison of observed and predicted female, mature, all shell survey size comps for NMFS 0 . Page 3 of 5 .

NMFS 0: female, mature, all shell


Figure 19: Comparison of observed and predicted female, mature, all shell survey size comps for NMFS 0 . Page 4 of 5.

NMFS 0: female, mature, all shell


Figure 20: Comparison of observed and predicted female, mature, all shell survey size comps for NMFS 0 . Page 5 of 5 .

NMFS M: male, all maturity, all shell


Figure 21: Comparison of observed and predicted male, all maturity, all shell survey size comps for NMFS M. Page 1 of 5 .

NMFS M: male, all maturity, all shell


Figure 22: Comparison of observed and predicted male, all maturity, all shell survey size comps for NMFS M. Page 2 of 5 .

NMFS M: male, all maturity, all shell


Figure 23: Comparison of observed and predicted male, all maturity, all shell survey size comps for NMFS M. Page 3 of 5 .

NMFS M: male, all maturity, all shell


Figure 24: Comparison of observed and predicted male, all maturity, all shell survey size comps for NMFS M. Page 4 of 5 .

NMFS M: male, all maturity, all shell


Figure 25: Comparison of observed and predicted male, all maturity, all shell survey size comps for NMFS M. Page 5 of 5 .

NMFS F: female, immature, all shell


Figure 26: Comparison of observed and predicted female, immature, all shell survey size comps for NMFS F. Page 1 of 5 .

NMFS F: female, immature, all shell


Figure 27: Comparison of observed and predicted female, immature, all shell survey size comps for NMFS F. Page 2 of 5.

NMFS F: female, immature, all shell


Figure 28: Comparison of observed and predicted female, immature, all shell survey size comps for NMFS F. Page 3 of 5 .

NMFS F: female, immature, all shell


Figure 29: Comparison of observed and predicted female, immature, all shell survey size comps for NMFS F. Page 4 of 5.

NMFS F: female, immature, all shell


Figure 30: Comparison of observed and predicted female, immature, all shell survey size comps for NMFS F. Page 5 of 5 .

NMFS F: female, mature, all shell


Figure 31: Comparison of observed and predicted female, mature, all shell survey size comps for NMFS F. Page 1 of 5 .

NMFS F: female, mature, all shell


Figure 32: Comparison of observed and predicted female, mature, all shell survey size comps for NMFS F. Page 2 of 5 .

NMFS F: female, mature, all shell


Figure 33: Comparison of observed and predicted female, mature, all shell survey size comps for NMFS F. Page 3 of 5 .

NMFS F: female, mature, all shell


Figure 34: Comparison of observed and predicted female, mature, all shell survey size comps for NMFS F. Page 4 of 5.

NMFS F: female, mature, all shell


Figure 35: Comparison of observed and predicted female, mature, all shell survey size comps for NMFS F. Page 5 of 5 .

SBS NMFS males: male, all maturity, all shell


Figure 36: Comparison of observed and predicted male, all maturity, all shell survey size comps for SBS NMFS males. Page 1 of 1.

SBS NMFS females: female, immature, all shell


Figure 37: Comparison of observed and predicted female, immature, all shell survey size comps for SBS NMFS females. Page 1 of 1.

SBS NMFS females: female, mature, all shell


Figure 38: Comparison of observed and predicted female, mature, all shell survey size comps for SBS NMFS females. Page 1 of 1.

SBS BSFRF males: male, all maturity, all shell


Figure 39: Comparison of observed and predicted male, all maturity, all shell survey size comps for SBS BSFRF males. Page 1 of 1.

SBS BSFRF females: female, immature, all shell


Figure 40: Comparison of observed and predicted female, immature, all shell survey size comps for SBS BSFRF females. Page 1 of 1 .

SBS BSFRF females: female, mature, all shell


Figure 41: Comparison of observed and predicted female, mature, all shell survey size comps for SBS BSFRF females. Page 1 of 1.

Fishery retained catch size compositions

TCF: male, all maturity, all shell


Figure 42: Comparison of observed and predicted male, all maturity, all shell retained catch size comps for TCF. Page 1 of 4 .

TCF: male, all maturity, all shell


Figure 43: Comparison of observed and predicted male, all maturity, all shell retained catch size comps for TCF. Page 2 of 4 .

TCF: male, all maturity, all shell


Figure 44: Comparison of observed and predicted male, all maturity, all shell retained catch size comps for TCF. Page 3 of 4 .

TCF: male, all maturity, all shell


Figure 45: Comparison of observed and predicted male, all maturity, all shell retained catch size comps for TCF. Page 4 of 4.

Fishery total catch size compositions

TCF: male, all maturity, all shell


Figure 46: Comparison of observed and predicted male, all maturity, all shell total catch size comps for TCF. Page 1 of 3 .

TCF: male, all maturity, all shell


Figure 47: Comparison of observed and predicted male, all maturity, all shell total catch size comps for TCF. Page 2 of 3 .

TCF: male, all maturity, all shell


Figure 48: Comparison of observed and predicted male, all maturity, all shell total catch size comps for TCF. Page 3 of 3 .

TCF: female, all maturity, all shell


Figure 49: Comparison of observed and predicted female, all maturity, all shell total catch size comps for TCF. Page 1 of 3 .

TCF: female, all maturity, all shell


Figure 50: Comparison of observed and predicted female, all maturity, all shell total catch size comps for TCF. Page 2 of 3 .

TCF: female, all maturity, all shell


Figure 51: Comparison of observed and predicted female, all maturity, all shell total catch size comps for TCF. Page 3 of 3 .

SCF: male, all maturity, all shell


Figure 52: Comparison of observed and predicted male, all maturity, all shell total catch size comps for SCF. Page 1 of 3 .

SCF: male, all maturity, all shell


Figure 53: Comparison of observed and predicted male, all maturity, all shell total catch size comps for SCF. Page 2 of 3 .

SCF: male, all maturity, all shell


Figure 54: Comparison of observed and predicted male, all maturity, all shell total catch size comps for SCF. Page 3 of 3 .

SCF: female, all maturity, all shell


Figure 55: Comparison of observed and predicted female, all maturity, all shell total catch size comps for SCF. Page 1 of 3 .

SCF: female, all maturity, all shell


Figure 56: Comparison of observed and predicted female, all maturity, all shell total catch size comps for SCF. Page 2 of 3 .

SCF: female, all maturity, all shell


Figure 57: Comparison of observed and predicted female, all maturity, all shell total catch size comps for SCF. Page 3 of 3 .

GTF: male, all maturity, all shell


Figure 58: Comparison of observed and predicted male, all maturity, all shell total catch size comps for GTF. Page 1 of 5 .

GTF: male, all maturity, all shell


Figure 59: Comparison of observed and predicted male, all maturity, all shell total catch size comps for GTF. Page 2 of 5 .

GTF: male, all maturity, all shell


Figure 60: Comparison of observed and predicted male, all maturity, all shell total catch size comps for GTF. Page 3 of 5 .

GTF: male, all maturity, all shell


Figure 61: Comparison of observed and predicted male, all maturity, all shell total catch size comps for GTF. Page 4 of 5 .

GTF: male, all maturity, all shell


Figure 62: Comparison of observed and predicted male, all maturity, all shell total catch size comps for GTF. Page 5 of 5 .

GTF: female, all maturity, all shell


Figure 63: Comparison of observed and predicted female, all maturity, all shell total catch size comps for GTF. Page 1 of 5 .

GTF: female, all maturity, all shell


Figure 64: Comparison of observed and predicted female, all maturity, all shell total catch size comps for GTF. Page 2 of 5 .

## GTF: female, all maturity, all shell



Figure 65: Comparison of observed and predicted female, all maturity, all shell total catch size comps for GTF. Page 3 of 5 .

GTF: female, all maturity, all shell


Figure 66: Comparison of observed and predicted female, all maturity, all shell total catch size comps for GTF. Page 4 of 5 .

GTF: female, all maturity, all shell


Figure 67: Comparison of observed and predicted female, all maturity, all shell total catch size comps for GTF. Page 5 of 5 .

RKF: male, all maturity, all shell


Figure 68: Comparison of observed and predicted male, all maturity, all shell total catch size comps for RKF. Page 1 of 3 .

RKF: male, all maturity, all shell


Figure 69: Comparison of observed and predicted male, all maturity, all shell total catch size comps for RKF. Page 2 of 3 .

RKF: male, all maturity, all shell


Figure 70: Comparison of observed and predicted male, all maturity, all shell total catch size comps for RKF. Page 3 of 3 .

RKF: female, all maturity, all shell


Figure 71: Comparison of observed and predicted female, all maturity, all shell total catch size comps for RKF. Page 1 of 3 .

RKF: female, all maturity, all shell


Figure 72: Comparison of observed and predicted female, all maturity, all shell total catch size comps for RKF. Page 2 of 3 .

RKF: female, all maturity, all shell


Figure 73: Comparison of observed and predicted female, all maturity, all shell total catch size comps for RKF. Page 3 of 3 .

# Model Comparisons: Fits to Fisheries Size Composition Data - M19F00a vs M19F01 vs M19F02 vs M19F03 vs M19F04 vs M19F05 

William Stockhausen

07 September, 2019

## Contents

Introduction ..... 1
Retained catch mean size compositions ..... 2
Total catch mean size compositions ..... 3
Fishery retained catch size composition residuals ..... 8
Effective Ns for retained catch size compositions ..... 14
Total catch size composition residuals ..... 15
Effective Ns for total catch size compositions ..... 63

## Introduction

Fits to fishery retained catch and total catch size composition data available to the model(s) are presented in this section. Included are plots of mean fits to size compositions, Pearson's residuals as bubble plots, and effective sample sizes. Not all of the fits presented are necessarily included in the parameter optimization for each model; some fits to datasets for a particular model may be included for comparison purposes with other models which include those data in their optimization. The reader should consult the main assessment document to determine which fits are included in the optimization for any particular model.

## Retained catch mean size compositions



Figure 1: Comparison of observed and predicted mean retained catch size comps for TCF.

Total catch mean size compositions


Figure 2: Comparison of observed and predicted mean total catch size comps for GTF.


Figure 3: Comparison of observed and predicted mean total catch size comps for RKF.


Figure 4: Comparison of observed and predicted mean total catch size comps for SCF.


Figure 5: Comparison of observed and predicted mean total catch size comps for TCF.

Fishery retained catch size composition residuals


Figure 6: Pearson's residuals for male proportions-at-size from the TCF for scenario M19F00a.


Figure 7: Pearson's residuals for male proportions-at-size from the TCF for scenario M19F01.


Figure 8: Pearson's residuals for male proportions-at-size from the TCF for scenario M19F02.


Figure 9: Pearson's residuals for male proportions-at-size from the TCF for scenario M19F03.


Figure 10: Pearson's residuals for male proportions-at-size from the TCF for scenario M19F04.


Figure 11: Pearson's residuals for male proportions-at-size from the TCF for scenario M19F05.

## Effective Ns for retained catch size compositions



Figure 12: Input and effective sample sizes from retained catch size compositions from the TCF fishery.

## Total catch size composition residuals



Figure 13: Pearson's residuals for male proportions-at-size from the TCF for scenario M19F00a.


Figure 14: Pearson's residuals for male proportions-at-size from the TCF for scenario M19F01.


Figure 15: Pearson's residuals for male proportions-at-size from the TCF for scenario M19F02.


Figure 16: Pearson's residuals for male proportions-at-size from the TCF for scenario M19F03.


Figure 17: Pearson's residuals for male proportions-at-size from the TCF for scenario M19F04.


Figure 18: Pearson's residuals for male proportions-at-size from the TCF for scenario M19F05.


Figure 19: Pearson's residuals for female proportions-at-size from the TCF for scenario M19F00a.


Figure 20: Pearson's residuals for female proportions-at-size from the TCF for scenario M19F01.


Figure 21: Pearson's residuals for female proportions-at-size from the TCF for scenario M19F02.


Figure 22: Pearson's residuals for female proportions-at-size from the TCF for scenario M19F03.


Figure 23: Pearson's residuals for female proportions-at-size from the TCF for scenario M19F04.


Figure 24: Pearson's residuals for female proportions-at-size from the TCF for scenario M19F05.


Figure 25: Pearson's residuals for male proportions-at-size from the SCF for scenario M19F00a.


Figure 26: Pearson's residuals for male proportions-at-size from the SCF for scenario M19F01.


Figure 27: Pearson's residuals for male proportions-at-size from the SCF for scenario M19F02.


Figure 28: Pearson's residuals for male proportions-at-size from the SCF for scenario M19F03.


Figure 29: Pearson's residuals for male proportions-at-size from the SCF for scenario M19F04.


Figure 30: Pearson's residuals for male proportions-at-size from the SCF for scenario M19F05.


Figure 31: Pearson's residuals for female proportions-at-size from the SCF for scenario M19F00a.


Figure 32: Pearson's residuals for female proportions-at-size from the SCF for scenario M19F01.


Figure 33: Pearson's residuals for female proportions-at-size from the SCF for scenario M19F02.


Figure 34: Pearson's residuals for female proportions-at-size from the SCF for scenario M19F03.


Figure 35: Pearson's residuals for female proportions-at-size from the SCF for scenario M19F04.


Figure 36: Pearson's residuals for female proportions-at-size from the SCF for scenario M19F05.


Figure 37: Pearson's residuals for male proportions-at-size from the GTF for scenario M19F00a.


Figure 38: Pearson's residuals for male proportions-at-size from the GTF for scenario M19F01.


Figure 39: Pearson's residuals for male proportions-at-size from the GTF for scenario M19F02.


Figure 40: Pearson's residuals for male proportions-at-size from the GTF for scenario M19F03.


Figure 41: Pearson's residuals for male proportions-at-size from the GTF for scenario M19F04.


Figure 42: Pearson's residuals for male proportions-at-size from the GTF for scenario M19F05.


Figure 43: Pearson's residuals for female proportions-at-size from the GTF for scenario M19F00a.


Figure 44: Pearson's residuals for female proportions-at-size from the GTF for scenario M19F01.


Figure 45: Pearson's residuals for female proportions-at-size from the GTF for scenario M19F02.


Figure 46: Pearson's residuals for female proportions-at-size from the GTF for scenario M19F03.


Figure 47: Pearson's residuals for female proportions-at-size from the GTF for scenario M19F04.


Figure 48: Pearson's residuals for female proportions-at-size from the GTF for scenario M19F05.


Figure 49: Pearson's residuals for male proportions-at-size from the RKF for scenario M19F00a.


Figure 50: Pearson's residuals for male proportions-at-size from the RKF for scenario M19F01.


Figure 51: Pearson's residuals for male proportions-at-size from the RKF for scenario M19F02.


Figure 52: Pearson's residuals for male proportions-at-size from the RKF for scenario M19F03.


Figure 53: Pearson's residuals for male proportions-at-size from the RKF for scenario M19F04.


Figure 54: Pearson's residuals for male proportions-at-size from the RKF for scenario M19F05.


Figure 55: Pearson's residuals for female proportions-at-size from the RKF for scenario M19F00a.


Figure 56: Pearson's residuals for female proportions-at-size from the RKF for scenario M19F01.


Figure 57: Pearson's residuals for female proportions-at-size from the RKF for scenario M19F02.


Figure 58: Pearson's residuals for female proportions-at-size from the RKF for scenario M19F03.


Figure 59: Pearson's residuals for female proportions-at-size from the RKF for scenario M19F04.


Figure 60: Pearson's residuals for female proportions-at-size from the RKF for scenario M19F05.

## Effective Ns for total catch size compositions



Figure 61: Input and effective sample sizes from total catch size compositions from the TCF fishery.


Figure 62: Input and effective sample sizes from total catch size compositions from the SCF fishery.


Figure 63: Input and effective sample sizes from total catch size compositions from the GTF fishery.


Figure 64: Input and effective sample sizes from total catch size compositions from the RKF fishery.

# Model Comparisons: Fits to Surveys Size Composition Data - M19F00a vs M19F01 vs M19F02 vs M19F03 vs M19F04 vs M19F05 

William Stockhausen

07 September, 2019

## Contents

Introduction ..... 1
Mean survey size compositions ..... 2
Residuals to survey size composition data ..... 10
Effective sample sizes for survey size compositions ..... 42

## Introduction

Fits to survey size composition data available to the model(s) are presented in this section. Included are plots of mean fits to size compositions, Pearson's residuals as bubble plots, and effective sample sizes. Not all of the fits presented are necessarily included in the parameter optimization for each model; some fits to datasets for a particular model may be included for comparison purposes with other models which include those data in their optimization. The reader should consult the main assessment document to determine which fits are included in the optimization for any particular model.

Mean survey size compositions


Figure 1: Comparison of observed and predicted mean survey size comps for NMFS 0.


Figure 2: Comparison of observed and predicted mean survey size comps for NMFS F.


Figure 3: Comparison of observed and predicted mean survey size comps for NMFS M.

## SBS BSFRF females



Figure 4: Comparison of observed and predicted mean survey size comps for SBS BSFRF females.

## SBS BSFRF males



Figure 5: Comparison of observed and predicted mean survey size comps for SBS BSFRF males.

## SBS NMFS females



Figure 6: Comparison of observed and predicted mean survey size comps for SBS NMFS females.

## SBS NMFS males



Figure 7: Comparison of observed and predicted mean survey size comps for SBS NMFS males.

## Residuals to survey size composition data



Figure 8: Pearson's residuals for male proportions-at-size from the NMFS 0 for scenario M19F00a.


Figure 9: Pearson's residuals for male proportions-at-size from the NMFS 0 for scenario M19F01.


Figure 10: Pearson's residuals for male proportions-at-size from the NMFS 0 for scenario M19F02.


Figure 11: Pearson's residuals for male proportions-at-size from the NMFS 0 for scenario M19F03.


Figure 12: Pearson's residuals for male proportions-at-size from the NMFS 0 for scenario M19F04.


Figure 13: Pearson's residuals for male proportions-at-size from the NMFS 0 for scenario M19F05.


Figure 14: Pearson's residuals for male proportions-at-size from the NMFS M for scenario M19F00a.


Figure 15: Pearson's residuals for male proportions-at-size from the NMFS M for scenario M19F01.


Figure 16: Pearson's residuals for male proportions-at-size from the NMFS M for scenario M19F02.


Figure 17: Pearson's residuals for male proportions-at-size from the NMFS M for scenario M19F03.


Figure 18: Pearson's residuals for male proportions-at-size from the NMFS M for scenario M19F04.


Figure 19: Pearson's residuals for male proportions-at-size from the NMFS M for scenario M19F05.


Figure 20: Pearson's residuals for male proportions-at-size from the SBS NMFS males for scenario M19F04.


Figure 21: Pearson's residuals for male proportions-at-size from the SBS NMFS males for scenario M19F05.


Figure 22: Pearson's residuals for male proportions-at-size from the SBS BSFRF males for scenario M19F04.


Figure 23: Pearson's residuals for male proportions-at-size from the SBS BSFRF males for scenario M19F05.


Figure 24: Pearson's residuals for female proportions-at-size from the NMFS 0 for scenario M19F00a.


Figure 25: Pearson's residuals for female proportions-at-size from the NMFS 0 for scenario M19F01.


Figure 26: Pearson's residuals for female proportions-at-size from the NMFS 0 for scenario M19F02.


Figure 27: Pearson's residuals for female proportions-at-size from the NMFS 0 for scenario M19F03.


Figure 28: Pearson's residuals for female proportions-at-size from the NMFS 0 for scenario M19F04.


Figure 29: Pearson's residuals for female proportions-at-size from the NMFS 0 for scenario M19F05.


Figure 30: Pearson's residuals for female proportions-at-size from the NMFS F for scenario M19F00a.


Figure 31: Pearson's residuals for female proportions-at-size from the NMFS F for scenario M19F01.


Figure 32: Pearson's residuals for female proportions-at-size from the NMFS F for scenario M19F02.


Figure 33: Pearson's residuals for female proportions-at-size from the NMFS F for scenario M19F03.


Figure 34: Pearson's residuals for female proportions-at-size from the NMFS F for scenario M19F04.


Figure 35: Pearson's residuals for female proportions-at-size from the NMFS F for scenario M19F05.


Figure 36: Pearson's residuals for female proportions-at-size from the SBS NMFS females for scenario M19F04.


Figure 37: Pearson's residuals for female proportions-at-size from the SBS NMFS females for scenario M19F05.


Figure 38: Pearson's residuals for female proportions-at-size from the SBS BSFRF females for scenario M19F04.


Figure 39: Pearson's residuals for female proportions-at-size from the SBS BSFRF females for scenario M19F05.

Effective sample sizes for survey size compositions


Figure 40: Input and effective sample sizes from retained catch size compositions from the NMFS 0.

NMFS M


Figure 41: Input and effective sample sizes from retained catch size compositions from the NMFS M.


Figure 42: Input and effective sample sizes from retained catch size compositions from the NMFS F.


Figure 43: Input and effective sample sizes from retained catch size compositions from the SBS NMFS males.


Figure 44: Input and effective sample sizes from retained catch size compositions from the SBS NMFS females.

## SBS BSFRF males



Figure 45: Input and effective sample sizes from retained catch size compositions from the SBS BSFRF males.

SBS BSFRF females


Figure 46: Input and effective sample sizes from retained catch size compositions from the SBS BSFRF females.


[^0]:    ${ }^{1}$ https://aws.state.ak.us/OnlinePublicNotices/Notices/Attachment.aspx?id=100244

[^1]:    ${ }^{2}$ https://github.com/wStockhausen/wtsTCSAM2013.git
    ${ }^{3} \mathrm{https}: / /$ github.com/wStockhausen/wtsTCSAM02.git

