# 2016 Stock Assessment and Fishery Evaluation Report for the Tanner Crab Fisheries of the Bering Sea and Aleutian Islands Regions 

William T. Stockhausen<br>Alaska Fisheries Science Center<br>6 September 2016<br>\title{ THIS INFORMATION IS DISTRIBUTED SOLELY FOR THE PURPOSE OF PREDISSEMINATION PEER REVIEW UNDER<br><br>APPLICABLE INFORMATION QUALITY GUIDELINES. IT HAS NOT BEEN FORMALLY DISSEMINATED BY NOAA<br><br>FISHERIES/ALASKA FISHERIES SCIENCE CENTER AND SHOULD NOT BE CONSTRUED TO REPRESENT ANY AGENCY<br><br>DETERMINATION OR POLICY }

## 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 directed fishery was opened in 2013/14 for the first time since 2009/10 because the stock was not overfished in 2012/13 (Stockhausen et al., 2013) and stock metrics met the State of Alaska (SOA) criteria for opening the fishery in 2013/14. TAC was set at $1,645,000 \mathrm{lbs}\left(746 \mathrm{t}\right.$ ) for the area west of $166^{\circ}$ W and at $1,463,000 \mathrm{lbs}(664 \mathrm{t})$ for the area east of $166^{\circ} \mathrm{W}$ in the SOA's Eastern Subdistrict of the Bering Sea District 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 was taken in the western area while $98.6 \%$ ( 654 t ) was taken in the eastern area. Prior to the closures, the retained catch averaged 770 t per year between 2005/062009/10.

Following the 2014 assessment (Stockhausen, 2014), 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.

Following last year's assessment (Stockhausen, 2015), TAC was set at $11,272,000 \mathrm{lbs}(5,113 \mathrm{t})$ for the eastern area and $8,396,000 \mathrm{lbs}$ ( 3808 t ) for the western area. On closing, essentially $100 \%$ of the TAC was taken in both areas $(11,268,885 \mathrm{lbs}[5,111 \mathrm{t}]$ in the eastern area, $8,373,493 \mathrm{lbs}[3,798 \mathrm{t}]$ in the western area based on the 5/20/2016 in-season catch report).

Non-retained females and sub-legal males are caught in the directed fishery as bycatch and discarded. Total bycatch (not discounted for assumed handling mortality) in the directed fishery was 3,104 t. Tanner crab are also caught as bycatch in the snow crab and Bristol Bay red king crab fisheries, in the groundfish fisheries and, to a 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 $1,414 \mathrm{t}$ for the 5 -year period 2011/12-2015/16. Bycatch in the snow crab fishery in 2015/16 was $3,536 \mathrm{t}$. The groundfish fisheries have been the next major source of Tanner crab bycatch over the same five year time period, averaging 296 t . Bycatch in the groundfish fisheries in 2015/16 was 352 t . The Bristol Bay red king crab fishery has typically been the smallest source of Tanner crab bycatch among these fisheries, averaging 61 t over the 5 -year time period, although 297 t caught and discarded in 2014/15. In 2015/16, this fishery accounted for $180 t$ 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 and $80 \%$ for Tanner crab discarded in the groundfish fisheries 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 (Model C), estimated MMB for 2015/16 was 73.9 thousand t (Table 30, Fig. 48). This was slightly smaller than that for 2014/15 (75.4 thousand t), but larger than that for 2013/14 ( 61.2 thousand t ). MMB has generally been rising since 2011/12. It remains above the very low levels seen in the mid-1990s to early 2000s (1990 to 2005 average: 29 thousand t ) and the 2014/15 estimate is the largest since 1978/79. However, it is considerably below model-estimated historic levels in the early 1970s when MMB peaked at $\sim 241$ thousand t (1971).

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

From the author's preferred model (Model C), the estimated total recruitment in 2016/17 (number of crab entering the population on July 1) is 120 million crab (Table 33, Fig. 45). Recruitment recently peaked in 2013 at 124 million crab, then declined in 2014 and 2015 below 100 million.
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 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2012 / 13$ | 16.77 | $59.35^{\mathrm{A}}$ | 0.00 | 0.00 | 0.71 | 19.02 | 8.17 |
| $2013 / 14$ | 16.98 | $72.70^{\mathrm{A}}$ | 1.41 | 1.26 | 2.78 | 25.35 | 17.82 |
| $2014 / 15$ | 13.40 | $71.57^{\mathrm{A}}$ | 6.85 | 6.16 | 9.16 | 31.48 | 25.18 |
| $2015 / 16$ | $12.82^{\mathrm{C}}$ | $73.93^{\mathrm{A}}$ | 8.92 | 8.91 | 11.38 | 27.19 | 21.75 |
| $2016 / 17$ |  | $45.34^{\mathrm{B}}$ |  |  |  | $25.61^{\mathrm{C}}$ | $20.49^{\mathrm{C}}$ |

(b) in millions lbs.

| Year | MSST | Biomass <br> (MMB) | TAC <br> (East + West) | Retained <br> Catch | Total Catch <br> Mortality | OFL | ABC |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2012 / 13$ | 36.97 | $130.84^{\mathrm{A}}$ | 0.00 | 0.00 | 1.57 | 41.93 | 18.01 |
| $2013 / 14$ | 37.43 | $160.28^{\mathrm{A}}$ | 3.11 | 2.78 | 6.14 | 55.89 | 39.29 |
| $2014 / 15$ | 29.53 | $157.78^{\mathrm{A}}$ | 15.10 | 13.58 | 20.19 | 69.40 | 55.51 |
| $2015 / 16$ | $28.27^{\mathrm{C}}$ | $162.99^{\mathrm{A}}$ | 19.67 | 19.64 | 25.09 | 59.94 | 47.95 |
| $2016 / 17$ |  | $99.95^{\mathrm{B}}$ |  |  |  | $56.46^{\mathrm{C}}$ | $45.17^{\mathrm{C}}$ |

A-Estimated biomass at the time of mating for the year concerned. Note this represents a revised estimate, based on the subsequent assessment, from the projection the previous year.
B-Projected biomass from the current stock assessment. This value will be updated next year.
C-Based on the author's preferred model (Model C).
6. Basis for the OFL
a) in 1000's t.

| Year | Tier ${ }^{\text {a }}$ | $\mathrm{B}_{\mathrm{MSY}}{ }^{\text {a }}$ | Current MMB $^{\mathbf{A}}$ | B/B $\mathrm{MSY}^{\text {a }}$ | $\mathrm{F}_{\text {OFL }}{ }^{\text {a }}$ | $\begin{gathered} \hline \text { Years to } \\ \text { define } \\ \mathbf{B}_{\mathrm{MSY}^{A}}{ }^{\mathbf{4}} \\ \hline \end{gathered}$ | $\underset{\text { Mortality }{ }^{\text {A,B }}}{\text { Natura }}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2012/13 | 3 a | 33.45 | 58.59 | 1.75 | $0.61 \mathrm{yr}^{-1}$ | 1982-2012 | $0.23 \mathrm{yr}^{-1}$ |
| 2013/14 | 3 a | 33.54 | 59.35 | 1.77 | $0.73 \mathrm{yr}^{-1}$ | 1982-2013 | $0.23 \mathrm{yr}^{-1}$ |
| 2014/15 | 3 a | 29.82 | 63.80 | 2.14 | 0.61 yr-1 | 1982-2014 | 0.23 yr-1 |
| 2015/16 | 3 a | 26.79 | 53.70 | 2.00 | $0.58 \mathrm{yr}-1$ | 1982-2015 | 0.23 yr-1 |
| 2016/17 | 3 a | 25.65 | 45.34 | 1.77 | $0.79 \mathrm{yr}^{-1}$ | 1982-2016 | $0.23 \mathrm{yr}^{-1}$ |

b) in millions lbs.

| Year | Tier ${ }^{\text {a }}$ | $\mathrm{B}_{\mathrm{MSY}}{ }^{\text {a }}$ | Current <br> MMB $^{\text {A }}$ | B/B MSY $^{\text {a }}$ | $\mathrm{F}_{\text {OFL }}{ }^{\text {a }}$ | Years to define B $_{\text {MSY }}{ }^{\text {a }}$ | Natural Mortality ${ }^{\text {A }}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2012/13 | 3a | 73.74 | 129.17 | 1.75 | $0.61 \mathrm{yr}^{-1}$ | 1982-2012 | $0.23 \mathrm{yr}^{-1}$ |
| 2013/14 | 3a | 73.94 | 130.84 | 1.77 | $0.73 \mathrm{yr}^{-1}$ | 1982-2013 | $0.23 \mathrm{yr}^{-1}$ |
| 2014/15 | 3 a | 65.74 | 140.66 | 2.14 | 0.61 yr-1 | 1982-2014 | 0.23 yr-1 |
| 2015/16 | 3 a | 59.06 | 118.38 | 2.00 | $0.58 \mathrm{yr}-1$ | 1982-2015 | $0.23 \mathrm{yr}-1$ |
| 2016/17 | 3a | 56.54 | 99.95 | 1.77 | $0.58 \mathrm{yr}^{-1}$ | 1982-2016 | $0.23 \mathrm{yr}^{-1}$ |

A-Calculated from the assessment reviewed by the Crab Plan Team in 20XX of 20XX/YY or based on the author's preferred model for 2016/17.
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 2016/17, is estimated at 45.34 thousand t . $\mathrm{B}_{\text {MSY }}$ for this stock is calculated to be 25.65 thousand t , so MSST is 12.82 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 2015/16 was 11.38 thousand t , which was less than the OFL for 2015/16 (27.19 thousand t ); consequently overfishing did not occur. The OFL for 2016/17 based on the author's preferred model (Model C) is 25.61 thousand $t$. The $\mathrm{ABC}_{\text {max }}$ for 2016/17, based on the $\mathrm{p}^{*} \mathrm{ABC}$, is 25.57 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 20.49 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. 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 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).

Based on the 2015 assessment (Stockhausen, 2015) and the new harvest strategy, TAC was set at $11,272,000 \mathrm{lbs}(5,113 \mathrm{t}$ ) for the eastern area and $8,396,000 \mathrm{lbs}(3,808 \mathrm{t})$ for the western area. On closing, essentially $100 \%$ of the TAC was taken in both areas ( $11,268,885 \mathrm{lbs}$ [ $5,111 \mathrm{t}]$ in the eastern area, $8,373,493 \mathrm{lbs}[3,798 \mathrm{t}]$ in the western area based on the 5/20/2016 in-season catch report).

## 2. Changes to the input data

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

| Data source | Data types | Time frame | Notes | Agency |
| :--- | :--- | :---: | :--- | :---: |
| NMFS EBS Bottom Trawl Survey | abundance, biomass, size compositions | 2016 | new | NMFS |
| NMFS EBS Bottom Trawl Survey | biomass cv's | $1975-2015$ | new calculation | NMFS |
| Directed fishery | retained catch (numbers, biomass) | $2015 / 16$ | new | ADFG |
|  | retained catch size compositions | $2015 / 16$ | new | ADFG |
|  | effort | $2015 / 16$ | new | ADFG |
|  | total catch, discards (biomass) | $2015 / 16$ | new | ADFG |
|  | total catch, discards size compositions | $2015 / 16$ | new | ADFG |
| Snow Crab Fishery | effort | $2015 / 16$ | new | ADFG |
|  | total catch, discards (biomass) | $2015 / 16$ | new | ADFG |
| Bristol Bay Red King Crab Fishery effort | $2015 / 16$ | new | ADFG | ADFG |
|  | size compositions | $2015 / 16$ | new | ADFG |
| Groundfish Fisheries | total catch, discards (biomass) | $2015 / 16$ | new | new |
|  | size compositions | $2015 / 16$ | new | new |
|  | total catch, discards (biomass) | $2015 / 16$ | new | NMFS/AKFIN |
|  | size compositions | $2015 / 16$ | NMFS/AKFIN |  |

## 3. Changes to the assessment methodology.

A number of potential changes to the model were reviewed by the CPT at its May 2016 meeting. The author's preferred model (Model C) embodies a number of the changes endorsed by the CPT, including: $1)$ using the Gmacs fishing mortality model; 2) estimating ln-scale female offsets to male fishing mortality in all fisheries; 3) estimating annual F-devs for 1992-present for bycatch in the BBRKC fishery; 4) eliminating constraints on minimum F's for bycatch in the BBRKC fishery; 5) requiring logistic selectivity curves to reach 1 in the largest model size bin; 5) using a logit scale, rather than a log scale, to estimate size-specific probabilities of terminal molt-to-maturity, 6) weighting sex-specific size composition by observed, rather than input, sample sizes when combining size compositions for bycatch in the groundfish fisheries, and 7) starting "current" recruitment estimates in 1975 (coincident with the NMFS EBS bottom trawl survey data), rather than in 1974. Model scenarios were also evaluated using 200 model runs using jittered initial parameter values to better achieve model convergence to the global minimum value for the model objective function. Additionally, CV's for estimates of mature survey biomass were recalculated using an approach that calculated CPUE across size classes at the haul level, then scaled to the regional (EBS) level using a standard approach for a stratified sampling design, as

[^0]opposed to the approach used last year which calculated CPUE in 1-mm CW size bins, scaled to the EBS, and then aggregated across size bins assuming independence of "errors" across size bins.

## 4. Changes to the assessment results

Results from the author's preferred model this year (Model C) are reasonably similar to those from the previous assessment, considering the large number of changes in the model. Average recruitment (1982present) was estimated at 179 million in last year's model, whereas it was estimated at 182 million in the author's preferred model this year. $\mathrm{B}_{\text {MSY }}$ was estimated at 26.79 thousand t last year and 25.65 thousand t this year. The largest difference was in $\mathrm{F}_{\mathrm{MSY}}$, which last year was estimated at $0.58 \mathrm{yr}^{-1}$ and $0.79 \mathrm{yr}^{-1}$ this year. This is partly due to the change this year to the Gmacs fishing mortality model which, although it assumes that fishery capture rates have a logistic size structure, imposes a somewhat different sizespecific mortality pattern for males in the directed fishery vis-à-vis the old model (which assumes fishing mortality has a logistic size dependence).

## B. Responses to SSC and CPT Comments

1. Responses to the most recent two sets of SSC and CPT comments on assessments in general. [Note: for continuity with the previous assessment, the following includes unaddressed comments prior to the most recent two sets of comments.]

June 2016 SSC Meeting
No general comments.
May2016 Crab Plan Team Meeting
No general comments.
October 2015 SSC Meeting
No general comments.
September 2015 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 2016 SSC Meeting
The SSC endorsed the CPT suggestions from its May meeting.
May2016 Crab Plan Team Meeting
The CPT outlined the base model to be used for this assessment, based on results presented by the author for a suite of models.
Response: The base model recommended by the CPT is the base model used here (Model B).
The CPT outlined a number of alternative models built on its recommended base model to be evaluated. Response: These models were evaluated for the assessment.

## October 2015 SSC Meeting

Comment: "The SSC endorses all of the CPT recommendations with respect to the poor fits to some of the retained catch time series, poor fits to the size composition data for retained catch and survey data, and issues with the total directed fishery selectivity curve for males (in particular the 1996 'outlier')." Response: See responses to CPT comments below.

Comment: "The SSC was unable to fully compare models, as the summary tables in the assessment did not include the number of model parameters for evaluating differences in likelihoods."
Response: A good point, and an oversight on my part. The number of model parameters will be included in at least one summary table.

Comment: "The SSC would have liked to have seen residual diagnostic plots for models assuming a lognormal likelihood (B and D) to assess more fully the rationale for not further considering these models." Response: Residual diagnostic output (z-scores) have been added to model output, and z -score plots are now included in the standard plots produced following a converged model run.

Comment: "There are continuing concerns about the most appropriate weights to use for different data components (CVs, effective N, etc.), and the SSC looks forward to recommendations from the dataweighting workshop."
Response: The CPT endorsed using an iterative approach to weighting composition data (the "Francis method"), but it has not yet been implemented for this model.

Comment: "Strong residual patterns in numbers at size remain a concern and suggest model misspecification with respect to growth."
Response: Growth increment data for Tanner crab in the Bering Sea was collected in 2015 for sub-adults and April-June, 2016 for smaller crab. This data was made available to the author this summer, but time did not permit substantive results to include in this assessment. The data appears to be very consistent with previous growth data collected near Kodiak Island, and is plotted against mean growth as estimated in last year's assessment in Fig. 2.

Comment: "The period with elevated M differs between male (1981-1985) and female crab (1980-84)." Response: This was a mistake (now corrected) in the code that produced the plot. The periods are the same (1980).

Comment: "The model overestimates female bycatch mortality in the snow crab fishery." Response: One factor responsible for this observation was that the estimated male fishing mortality rate in each fishery was equally applied to females, with only changes in selectivity available to better fit female bycatch. The option to estimate female-specific offsets to (log-scale mean) male fishing mortality rates has been added to the model and reduces this problem. Fits were also improved using a lognormal likelihood (with assumed cv's), rather than the standard normal likelihood.

## September 2015 CPT Meeting

Comment: "The model fits total catch well, but does a poorer job in fitting retained catch, catch of females, and catch in the bycatch fisheries."
Response: There appears to be a conflict in the model between fitting total (male) catch and retained catch in the directed fishery. Fitting discard catch rather than total catch improves the fit to retained catch. This may be an issue related to treating retained and total catch with equal uncertainty in the standard model likelihood. Fits to female bycatch are improved when estimating a female-specific offset to (log-scale male) mean fishing mortality. Fits to bycatch improved, in general, using a lognormal likelihood assumption for fishery catch data, but it is unclear whether the cv's assumed are reasonable.

Comment: "Strong residual patterns exist in fits of male survey and retained-catch size composition..." Response: See response to SSC comment regarding collection of growth increment data.

Comment: "It was not clear why the model estimates full selection [for males in the directed fishery] in 1996 at roughly $100 \mathrm{~cm} . . . "$
Response: This occurs due to a combination of two factors: 1) the sample size for male size comps from the directed fishery in 1996 is quite small, meaning that a poor fit to this size frequency has little effect on
the overall likelihood, and 2) the size-at-50\% selected in the directed fishery prior to 1992 is based on the mean size-at-50\% selected in the directed fishery after 1991 (size-at- $50 \%$ selected in the directed fishery is allowed to vary annually after 1991). Although it has cascading effects through many likelihood components because of its influence on underling population structure, the size-at- $50 \%$ selected in the directed fishery prior to 1992 most directly influences (I think) fits to retained catch size compositions prior to 1992. If the fit to the pre-1992 retained catch size compositions can be improved by changing the size-at- $50 \%$ selected in the pre-1992 directed fishery, there is little "cost" to doing so even by making the size- $50 \%$-selected in 1996 any value whatsoever.

Comment: "The poor fit of the models with lognormal fishery catch likelihoods (Models B and D [in the 2015 assessment] ... was surprising to some CPT members."
Response: These models exhibited questionable convergence in the 2015 assessment. From results obtained in May using similar models, it is clear those models had not converged and the results were spurious (as was suggested by the author at the time). For this assessment, I ran each model scenario 200 times with randomly-selected (jittered) initial parameter values to improve confidence in obtaining a "converged" model result. The models with lognormal fishery likelihoods (models including changes L0 and L1 in the report) now fit the data well-perhaps too well, in some cases.

Comment: "The author should consider fitting retained catch exactly."
Response: Time did not allow exploring this possibility.

## June 2015 SSC Meeting

No specific comments.

## 3. Older comments that were addressed this year or remain to be addressed:

Comment: "Future exploration...should consider the impact of handling mortality on the estimate of natural mortality and how the model behaves if Q for the most recent years is assumed known rather than being estimated."
Response: Not yet addressed.
Comment: "The CPT reiterates its suggestions from the September 2014 meeting, in particular that the sensitivity of the results to the prior on Q should be explored."
Response: Not yet addressed.
Comment: "The SSC encourages authors to explore alternative models such as time-varying growth to help address retrospective bias and patterns in other residuals."
Response: This can be addressed in the future with the new model code (currently being tested), but not with the current model.

Comment: "The SSC also encourages authors to explore model alternatives without time-varying selectivity for the groundfish fishery."
Response: Not yet addressed.
Comment: "Examine issues related to misfits of the size composition residuals for retained males and total males in the directed fishery. Consider exploring alternative growth components, specification of sample sizes, or a combination of fishing selectivity and handling mortality is causing mis-fits."
Response: Not yet addressed.
Comment: "Examine retrospective patterns of models being brought forward."
Response: Retrospective patterns for the author's preferred model are examined here for the first time. Patterns for rejected models were similar (but are not presented here).

Comment: "Evaluate the feasibility of estimating $F_{M S Y}$ (and $B_{M S Y}$ ) for the stock using the estimates of recruitment and MMB during the post-1982 period, and compare to the $\mathrm{F}_{35 \%}$ MSY proxy." Response: Not yet addressed.

Comment: "If time permits, apply the groundfish plan team's stock structure template to Tanner crab to synthesize the available information on stock structure."
Response: Not yet addressed.
Comment: The CPT "recommends that all assessment authors document assumptions and simulate data under those assumptions to test the ability of the model to estimate key parameters in an unbiased manner."
Response: Not yet addressed.
Comment: "Plot the input effective sample sizes for the compositional data versus the effective sample sizes inferred by the fit of the model..."
Response: Done.
Comment: "Allow M for immature as well as mature males to change during 1980-83 (the data on changes in abundance do not suggest that only mature males declined substantially) and test whether it is necessary to allow female M to change over time."
Response: Not yet addressed.
Comment: "Consider fitting to total biomass (by sex?) and to the compositional data rather than to mature biomass (include the fit to mature biomass by sex as a diagnostic)."
Response: Not yet addressed.
Comment: "Do not fit to male compositional data by maturity state for the years for which chela heightmaturity relationships are not available."
Response: Not yet addressed.
Comment: "There is still a residual pattern in the fit to the size-composition data for the survey. This could be due to time-varying growth, which should be examined as an alternative model." Response: Not yet addressed.

## 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. Evidence is lacking that the EBS shelf is home to two distinct, non-intermixing, non-interbreeding stocks that should be assessed and managed separately.

## 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 <br> 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. |
| 4 | 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 25 mm CW proceeds by a series of annual molts, up to a final (terminal) molt to maturity (Tamone et al., 2007). Relationships between pre-molt and post-molt size specific to Tanner crab in the EBS have not been evaluated, although data on individual molt increments from 125 crab collected in the EBS in 2015 and 2016 (Fig. 2).

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; Fig. 2).

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.

## 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. 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 self-fertilize 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. 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). 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).

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.

## 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 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 ADF\&G 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, with federal oversight (Bowers et al. 2008). The State of Alaska 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 (Fig. 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, I use the terms "east region" and "west region" as shorthand to refer to the regions demarcated by $166^{\circ} \mathrm{W}$.

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 $5.5 "(140 \mathrm{~mm} \mathrm{CW})$ in the east and 5 " $(127 \mathrm{~mm} \mathrm{CW})$ 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 previous assessments, 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 this assessment, I use the term "legal males" to refer to crab 125 mm CW, the minimum "preferred" size used in both eastern and western areas 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; Fig. 3). Foreign fishing for Tanner crab ended in 1980.

The domestic Tanner crab pot fishery developed rapidly in the mid-1970s (Tables 1 and 2; Fig.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 18.19 thousand t , and then fell sharply through the mid-1990s. The domestic Tanner crab fishery was closed between 1996/97 and 2004/05 as a result of conservation concerns regarding depressed stock status. It re-opened in 2005/06 and averaged 0.77 thousand t retained catch between 2005/06-2009/10 (Tables 1 and 2). For the 2010/11-2012/13 seasons, the State of Alaska closed directed commercial fishing for Tanner crab due to estimated female stock metrics being 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 $1,645,000 \mathrm{lbs}(746 \mathrm{t})$ for the area west of $166^{\circ} \mathrm{W}$ and at $1,463,000 \mathrm{lbs}(664 \mathrm{t})$ for the area east of $166^{\circ} \mathrm{W}$ in the State of Alaska's 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 $99.6 \%$ ( $3,829 \mathrm{t}$ ) 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 t ) was the largest taken in the fishery since 1992/93 (Tables 1, 2; Fig. 3).

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 (Tables 4 and 5, Fig.s 5-7). 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 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

## 1. Summary of new information

Survey biomass and size composition data from the 2016 NMFS EBS Bottom Trawl Survey were added to the assessment dataset. Last year, coefficients of variation for annual mature male and female survey biomass were calculated based on survey biomass information (estimates and cv's) provided at 1 mm CW size bins for the EBS region by the NMFS Kodiak Lab (R. Foy, NMFS, pers. comm.). In this assessment, the cv's for mature survey biomass for the EBS were calculated by aggregating over sizes at the haul level, then scaling up to the EBS. Model runs with cv's calculated using both approaches were made to discern the impact of the change. This change is discussed in more detail in the section on survey biomass estimates below (Section D.2.d).

Estimates of total retained biomass and abundance, as well as retained size frequencies by shell condition, in the 2015/16 directed fishery were provided by ADFG (J. Webb, ADFG, pers. comm.) based on fish ticket data and dockside observer sampling. ADFG also provided estimates of Tanner crab bycatch (sexspecific numbers, biomass and size compositions) in the 2015/16 directed Tanner crab, snow crab, and Bristol Bay red king crab fisheries.

Tanner crab bycatch data in the groundfish fisheries (biomass, size compositions) were extracted for 2015/16 from the groundfish observer and AKFIN databases.

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

| Data source | Data types | Time frame | Notes | Agency |
| :---: | :---: | :---: | :---: | :---: |
| NMFS EBS Bottom Trawl Survey | abundance, biomass, size compositions | 2016 | new | NMFS |
| NMFS EBS Bottom Trawl Survey | biomass cv's | 1975-2015 | new calculation | NMFS |
| Directed fishery | retained catch (numbers, biomass) | 2015/16 | new | ADFG |
|  | retained catch size compositions | 2015/16 | new | ADFG |
|  | effort | 2015/16 | new | ADFG |
|  | total catch, discards (biomass) | 2015/16 | new | ADFG |
|  | total catch, discards size compositions | 2015/16 | new | ADFG |
| Snow Crab Fishery | effort | 2015/16 | new | ADFG |
|  | total catch, discards (biomass) | 2015/16 | new | ADFG |
|  | size compositions | 2015/16 | new | ADFG |
| Bristol Bay Red King Crab Fishery | effort | 2015/16 | new | ADFG |
|  | total catch, discards (biomass) | 2015/16 | new | ADFG |
|  | size compositions | 2015/16 | new | ADFG |
| Groundfish Fisheries | total catch, discards (biomass) | 2015/16 | new | NMFS/AKFIN |
|  | size compositions | 2015/16 | new | NMFS/AKFIN |

The following table summarizes the data coverage in the assessment model:


## 2. Data presented as time series

For the stock biomass and fishery 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. Total catch

Retained catch ( 1000 's $t$ ) 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 Fig. 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 has increased each of the last three years as TACs have increased (Fig.s 3, 6). The retained catch for 2015/16 (8,910 t) was the largest since 1992/1993 (15,920 t; Table 1).

## b. Information on bycatch and discards

Annual bycatch (discards) of Tanner crab are provided by sex in Tables 3 and 4 (and Fig.s 4-6) from ADFG crab observer sampling, starting in 1992/93 for the directed Tanner crab fishery, the snow crab
fishery, and the BBRKC fishery. Annual discards for the groundfish fisheries, based on NMFS groundfish observer programs, are also provided starting in 1973/74, but sex is undifferentiated. A value of 0.321 is used 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 for handling mortality is used to reflect differences in gear and on-deck operations with those of the crab fleets.

Estimated bycatch mortality in the groundfish fisheries was highest ( $\sim 15,000 \mathrm{t}$ ) in the early 1970s, but was substantially reduced by 1977 to $\sim 2,000 \mathrm{t}$ with the curtailment of foreign fishing fleets. It declined further in the 1980s (to $\sim 500 \mathrm{t}$ ) but increased somewhat in the late 1980s to a peak of $\sim 2,000 \mathrm{t}$ before undergoing a slow but rather steady decline to the present ( 282 t in 2015/16). Since reliable at-sea ADFG crab observer data has been available (1992), the snow crab fishery has consistently accounted for the fraction of bycatch mortality among the crab fisheries, followed by the directed fishery and the BBRKC fishery (Table 4, Fig. 5). Estimated bycatch mortality was highest for all crab fisheries in the early 1990s ( $\sim 12,000 \mathrm{t}$ total) but subsequently declined as (presumably) the stock declined and the directed fishery was curtailed. Since the directed fishery re-opened in 2013/14, bycatch mortality has averaged 325 t in the directed fishery, 579 t in the snow crab fishery, 32 t in the BBRKC fishery, and 300 t in the groundfish fisheries.

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 2014/14, 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 crab observer sampling is presented in Fig. 7 by fishery region (and total) for the two most recent periods the fishery was open (spanning 2005/06-2015/16). These appear to indicate a shift to retaining somewhat smaller minimum sizes since 2013/14, compared with 2005/06-2009/10.

Size compositions of estimated total catch (retained + discards) from at-sea crab fishery observer sampling in the directed fishery are presented by shell condition and fishery region in Fig. 8 for male crab and in Fig. 9 for female crab. The male size compositions suggest that about half the males caught in the directed fishery in 2015/16 were less than the minimum "preferred" size of 125 mm CW . If old shell males really are males at least one year past their terminal molt (as assumed in the assessment model), the size compositions for these crab suggest that $30-50 \%$ of these crab (which will not grow) are less than the preferred size.

Size compositions for Tanner crab bycatch by sex in the snow crab fishery from at-sea crab fishery observer sampling are presented by shell condition in Fig. 10. Fig. 11 presents similar information for the BBRKC fishery. Fig. 12 presents relative catch size composition information from groundfish observer sampling in the groundfish fisheries for males and females, respectively, from 1973/74 to the present. The male bycatch 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, the BBRKC fishery appears to catch mostly larger Tanner crab males, while the groundfish fisheries take a wide range of sizes as bycatch.

Raw and input sample sizes (number of individuals measured) for the various fisheries are presented in Tables 5-9.

## 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 (Table 10, Fig. 13). Estimated biomass of mature crab in the survey time series started at its maximum ( $281,000 \mathrm{t}$ ) in 1975, decreased rapidly to a low $(14,000 \mathrm{t}$ ) in 1986, and rebounded quickly to a smaller peak ( $134,000 \mathrm{t}$ ) in 1991. After 1991, mature survey biomass decreased again, reaching a minimum of $10,500 \mathrm{t}$ in 1998. Recovery following this decline was slow and mature survey biomass did not peak again until 2008 ( $67,000 \mathrm{t}$ ), after which it has fluctuated more rapidly-immediately decreasing the following year by almost $50 \%$ and reaching a minimum in 2012 ( $36,000 \mathrm{t}$ ), followed by an increase of almost $50 \%$ in 2013 and reaching a peak in $2014(82,000 \mathrm{t}$ ). The most recent trend (2014-2016) has been a declining one (Fig. 14). Trends in the male and female components of mature survey biomass, as well as legal male abundance, have primarily been in synchrony with one another (Fig. 13), as have changes in the eastern and western fishery regions (east and west of $166^{\circ} \mathrm{W}$ longitude; Fig.s 15, 16), although the magnitudes differ.

Survey biomass estimates are not direct inputs to the stock assessment model. Instead, survey size compositions and standardized sex-specific weight-at-size regressions from Daly et al. (2014) are used to calculate the corresponding sex-specific mature survey biomass on an annual basis. This approach has been used since the 2012 assessment (Rugolo and Turnock, 2012a), although the weight-at-size regressions were changed in 2015 to agree with the standardized versions used by the NMFS EBS Bottom Trawl Survey (Daly et al., 2014). These biomass estimates, while similar in scale, do not correspond exactly to corresponding time series published in recent survey technical memoranda. First, the minimum size of crab included in the assessment model is 25 mm CW, while the "tech memo" time series includes crab of all sizes. Second, maturity state for males in the assessment has been based on a maturity ogive developed by Rugolo and Turnock (2010), while size cut-points are used to classify male maturity for the tech memos.

Last year, coefficients of variation for annual mature male and female survey biomass were calculated based on survey biomass information (estimates and cv's) provided at 1 mm CW size bins for the EBS region by the NMFS Kodiak Lab (R. Foy, NMFS, pers. comm.). For this data, haul-level estimates of CPUE at $1-\mathrm{mm} \mathrm{CW}$ size bin widths were expanded to regional (east/west of $166^{\circ} \mathrm{W}$ longitude, entire EBS) scales using standard formulae. In order to obtain estimates of mature (or any other combination of sizes) survey biomass across the EBS for each sex , it was simply necessary to sum across sizes-which was the rationale for providing the data in this format. In order to obtain the associated cv's with the summed data, however, it was necessary to assume observation "errors" were uncorrelated between size bins. However, this approach tends to underestimate the "true" cv's one obtains by aggregating first across sizes at the haul level, then scaling up to the EBS (as opposed to aggregating to the EBS level for 1 mm CW size bins, then aggregating across size bins; Fig. 17). In this assessment, the cv's for mature survey biomass for the EBS were calculated by aggregating over sizes at the haul level, then scaling up to the EBS. Model runs with cv's calculated using both approaches were made to discern the impact of the change (discussed below).

## e. Survey catch-at-length

Plots of survey size compositions for male crab, expanded to total abundance by shell condition and fishery region, in Fig.s 18 and 19. The absence of small (new shell) crab in the eastern region since 2009 is notable, as is the progression of a possible cohort (with two size modes) through the new shell size classes in both regions starting in 2009 that starts to show up, but much reduced in amplitude, in the old shell crab size comps in 2014. Plots of survey size compositions for female crab, expanded to total abundance by maturity status (based on morphometric characteristics) and fishery region, are shown in Fig.s 20 and 21. Similar to males, a cohort progression of immature females starting in 2009 is evident in both regions, although it is much clearer in the eastern region. It can also be tracked into the old shell size size comps starting in 2013.

Observed sample sizes for the size compositions, aggregated to the EBS regional level used in the assessment, are presented in Table 11.

## f. Other time series data.

Spatial patterns of abundance in the 2013-2016 NMFS bottom trawl surveys are mapped in Fig.s 22-26 for immature males, mature males, legal males, immature females, and mature females, respectively. A decline in the abundance of immature crab over time in the middle shelf of the EBS and around the Pribilof Islands is evident in Fig. 22. A similar decline is apparent for mature and legal-sized males crab in the middle shelf (Fig.s 23 and 24), but it does not occur in the Pribilofs. Immature females (Fig. 25) do not extend as far into the middle shelf as males (compare distributions for 2013), and the distribution appears to recede from the middle shelf to the shelf edge over 2013-2016. A similar phenomenon occurs for mature females (Fig. 26), although these extended further into the middle shelf region than immature females in 2013 (more like mature males).

The decline in abundance of Tanner crab from the middle shelf region over the last four years has occurred as bottom temperatures in the EBS have risen since 2012 from the second-lowest value during the 1975-2015 annual NMFS EBS summer trawl surveys to the second-highest in 2016 (Fig. 27). Associated with these increased mean temperatures is a withdrawal of an extensive cold pool in summer 2012 to the northwest in subsequent years and a concomitant warming of the middle and inner shelf areas (Fig. 28). It is unknown, however, whether or not the increasingly-warm middle shelf in the summer is responsible for the increased absence of Tanner crab from the middle shelf during the survey and, if it is, whether this constitutes a survey-specific phenomenon (i.e., changes in catchability or availability without actual changes in population abundance) or a factor driving a true decline in the Tanner crab stock.

While of interest, it should be noted that these spatial patterns of survey abundance and bottom temperature, as well as the time series of average bottom temperature during the survey, do not play a role in the assessment model.

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 12).

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

## a. Growth-per-molt

Sex-specific growth curves derived by Rugolo and Turnock (2010) were shown in Fig. 2. These curves provide the basis for priors on sex-specific growth estimated within the assessment model.

## b. Weight-at size

Weight-at-size relationships used in the assessment model for males, immature females, and mature females is depicted in Fig. 29.

## c. Size distribution at recruitment

The assumed size distribution for recruits to the population in the assessment model is presented in Fig. 30.

## 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.
## 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 Rugolo's and Turnock's 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.

In December 2012, a new analyst (Stockhausen) was assigned as principal author for the Tanner crab assessment. 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 ${ }^{2}$.

## 2. Model Description

## a. Overall modeling approach

TCSAM 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 3 of the 2014 assessment (Stockhausen, 2014).

In brief, crab enter the modeled population as recruits following the size distribution in Fig. 30. An equal (50:50) sex ratio is assumed at recruitment, 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 calculated based on fishery-specific stage/size-

[^1]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 $1(\delta t=0.375$ 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 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 (Stockhausen, 2014).

## b. Changes since the previous assessment.

Model code is available on github (https://github.com/wStockhausen/wtsTCSAM2013; the current branch is '2016AssessmentModel'). A substantial amount of work has been done since Sept. 2015 to implement alternative approaches to model parameterization, data-fitting, and model output formats in the code. In addition, all model options can now be specified in a "control file", as can parameter estimation phases and initial parameter values, and are no longer "hard-wired" in the model code. The changes made up to May 2016 are summarized in the following table:

| Category | Description |
| :---: | :--- |
| recruitment | The beginning of the "historic" and "current" recruitment periods now inputs. <br> Initial parameter values and estimation phase set now inputs. |
| natural <br> mortality | linitial parameter values and estimation phase now inputs. <br> Time period for high natural mortality now an input. |
| fishing | Phase to estimate fishing mortality in BBRKC fishery now an input. <br> mortality <br> Lognormal likelihoods implemented for fishery catch data (assumed cv's are inputs). <br> Option to fit male discard (rather than total mortality) in directed fishery implemented. <br> Ln-scale offsets to mean fishing mortality/capture for female crab added as parameters. <br> Parameters added to estimate scalars to extrapolate fishing mortality using effort. <br> Methods to estrapolate fishing mortality using effort are set in control file. <br> Implemented alternative methods to normalize size comps from the groundfish fisheries. <br> Normalization method for size comps from the groundfish fisheries set in control file. |
| molt to <br> maturity | Implemented parameter estimation on logit scale. |
| control file | Added nominal legal size as input. Was hard-wired to 138 mm CW. <br> Survey Q: means, std devs now set in control file. |
| other | Model start year now an input. <br> Revised code to vectorize many calculations. <br> Added z-scores from likelihood calculations to output. <br> Added ability to jitter initial parameter values <br> R package revised to run multiple models, jittered parameter runs |

Models implementing many of these changes were reviewed by the CPT at its May 2016 meeting; the most substantial option not reviewed was the addition of using parameters to estimate the values used to extrapolate effort to fishing mortality in the snow crab and BBRKC bycatch fisheries. This option is addressed in models considered for this assessment.

Model changes made subsequent to May 2016 are summarized here:

| Category | Description |
| :---: | :--- |
| fishing <br> mortality | implemented phased reduction of penalties on F-devs as option <br> implemented option to remove penalties on F-devs in final estimation phase <br> implemented option to remove minimum F's for BBRKC bycatch fishery |
| control file | All parameter phases now inputs (no longer hardwired) <br> All initial parameter values now inputs (if not jittering) <br> legal/preferred size now an input (no longer hardwired) |
| other | Model output completely revised to facilitate model comparisons <br> R package revised to facilitate model comparisons |

The model changes above associated with fishing mortality were implemented to address CPT requests for alternative models to be considered for this assessment.

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

The model code has been previously reviewed by members of the CPT and the assessment author.

## 3. Model Selection and Evaluation

## a. Description of alternative model configurations

Based on analyses presented to the CPT at its May 2016 meeting, it was concluded that the 2015 assessment model ("2015AMO", with "O" for "original") had not converged to its global minimum objective function value; instead, it had converged to a local minimum. The model was re-evaluated using the 2015 data to determine its global minimum by making 200 runs with randomly-selected ("jittered") initial values. The run ("2015AMR", with "R" for "re-run") with the smallest objective function and smallest maximum gradient was selected as the run most likely to have arrived at the global minimum. The 2015AMR achieved a slightly lower objective function value (2048.68) than the 2015AMO assessment model (2049.07), conclusively indicating that the 2015AMO had not converged to the global minimum.

Two data configurations were considered in this assessment; the two configurations differed in how input cv's for regional (EBS) mature survey biomass estimates were calculated. In the "old" method, cv's were calculated assuming independence of errors across $1-\mathrm{mm}$ CW size bins:

$$
c v_{\text {mat }}=\frac{\sqrt{\sum_{z}\left(c v_{z} \cdot b_{z}\right)^{2}}}{\sum_{z} b_{z}}
$$

where $c v_{\text {mat }}$ is the cv associated with the estimate of mature biomass $\left(=\sum_{z} b_{z}\right)$ and $c v_{z}$ is the cv associated with $b_{z}$, the survey estimate of mature biomass for size bin $z$. In the "new" method, estimates of survey biomass at the individual haul level (i.e., summed across size bins for each individual haul) were expanded to the regional (EBS) level using the survey's stratified sampling design, with the regional level cv calculated based on this stratification. The impact of this change on the assessment was quantified using the new cv's for mature survey biomass, but without otherwise updating the 2015 datafiles to 2016, and evaluating the 2015 assessment model using the parameter jittering approach with 200 jittered runs. The resulting "best" model run is referred to here as 2015AMN (" N " for "new").

At the May CPT meeting, models with the following incremental changes to the 2015 assessment model were evaluated:

| Change | Description |
| :---: | :--- |
| O | 2015 assessment model |
| A | start "current" recruitment estimation in 1975, instead of 1974 |
| B | normalize groundfish fishery size comps using original sample sizes, not input sample sizes |
| C | estimate log-scale fishing mortality/capture rate offsets for female crab |
| D | fit to male discard mortality in directed fishery |
| E | turn on fishing mortality/capture rate estimation for BBRKC |
| F | set initial estimate for historic log-scale recruitment ( $=11.4$ ) |
| G | estimate probability of molt-to-maturity using logit-scale parameterization |
| H | change model start year to 1930, keep start year for "historic" recruitment deviations = 1949 |
| I | enforce logistic selectivity = 1 in largest size bin |
| J | use GMACS fishing mortality model |
| L0 | use lognormal NLL's with moderate cv's for fits to fishery catch data |
| L1 | use lognormal NLL's with small cv's for fits to fishery catch data |

Based on these the review of these models, the CPT requested the following configuration, referred to here as Model B ("B" for "base"), be used as the "base" model for evaluating additional alternative model configurations:

| Change | Description |
| :---: | :--- |
| A | start "current" recruitment estimation in 1975, instead of 1974 |
| B | normalize groundfish fishery size comps using original sample sizes, not input sample sizes |
| C | estimate log-scale fishing mortality/capture rate offsets for female crab |
| E | turn on fishing mortality/capture rate estimation for BBRKC |
| G | estimate probability of molt-to-maturity using logit-scale parameterization |
| I | enforce logistic selectivity $=1$ in largest size bin |
| J | use GMACS fishing mortality model |

Based on requested alternatives proposed by the CPT in May, the following alternative models were evaluated for this assessment:

| Scenario | Description |
| :---: | :--- |
| 2015AMO | 2015 assessment model and data |
| 2015AMR | $2015 A M O$ re-evaluated using parameter jittering |
| 2015AMN | $2015 A M O$ + new approach to calculate CVs for mature survey biomass |
| 2015AM | $2015 A M N+2016$ data (using new approach to calculate CVs for mature survey biomass) |
| Model A | Model B, but using old fishing mortality model |
| Model B | Model selected by CPT in May as 'base" model for 2016 assessment |
| Model C | Model B + no minimum F's imposed on BBRKC fishery bycatch |
| Model D | Model C + effort extrapolation parameters estimated |
| Model E | Model D + penalty on F-devs reduced to 0 in final estimation phase |
| Model F | Model D + lognormal likelihoods assumed for fishery catch data (change L0 from May) |
| Model G | Model E + lognormal likelihoods assumed for fishery catch data (change L0 from May) |

In implementing the lognormal fishery catch likelihoods (Models F and G), it was necessary to specify relative error sizes for each data source. The same set of values were used for both models, as documented in the following table:

| Fishery | Data Source | Likelihood <br> Component | Assumed <br> CV |
| :--- | :--- | :--- | ---: |
| Directed fishery | fish tickets | retained catch | $5 \%$ |
|  | at-sea observers | total catch/discards | $20 \%$ |
| snow crab | at-sea observers | total catch/discards | $20 \%$ |
| BBRKC | at-sea observers | total catch/discards | $20 \%$ |
| groundfish | at-sea observers | total catch/discards | $20 \%$ |

The values chosen were subjective, based on the author's experience with such data. It seems likely the chosen values can be refined in future work.

## b. Progression of results from the previous assessment to the preferred base model

The following table summarizes basic model results for the 11 model/data combinations considered here:

| Model Scenario |  | Data | $\begin{array}{\|c} \# \\ \text { params } \end{array}$ | \# of jitter runs | Objective Function |  | invertible hessian? | Mean Recruitment |  | MMB (1000's t) |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Year |  |  |  | value | max gradient |  | 1982+ | 2000+ | 1982+ | last 3 <br> years | final year |
| 2015AMO | 2015 | old cv's | 307 | -- | 2049.07 | 0.0000875 | yes | 179.4 | 164.9 | 36.5 | 59.6 | 71.6 |
| 2015AMR | 2015 | old cv's | 307 | 200 | 2048.68 | 0.0002388 | yes | 176.8 | 163.9 | 35.8 | 57.7 | 69.3 |
| 2015AMN | 2015 | new cv's | 307 | 200 | 1838.14 | 0.0003343 | yes | 193.4 | 188.1 | 42.7 | 68.7 | 83.3 |
| 2015AM | 2016 | new cv's | 312 | 200 | 1952.73 | 0.0002182 | yes | 183.5 | 174.1 | 41.8 | 71.3 | 74.3 |
| Model A | 2016 | new cv's | 341 | 200 | 2338.77 | 1.5256000 | yes | -- | -- | -- | -- | -- |
| Model B | 2016 | new cv's | 341 | 200 | 2406.67 | 0.0002237 | yes | 182.2 | 171.4 | 39.7 | 70.2 | 73.9 |
| Model C | 2016 | new cv's | 341 | 200 | 2406.75 | 0.0004336 | yes | 182.3 | 171.5 | 40.7 | 70.2 | 73.9 |
| Model D | 2016 | new cv's | 343 | 200 | 2391.11 | 0.0004838 | yes | 168.8 | 165.2 | 37.9 | 63.7 | 67.2 |
| Model E | 2016 | new cv's | 343 | 200 | 2286.11 | 0.0000145 | yes | 174.2 | 176.0 | 40.1 | 68.3 | 72.4 |
| Model F | 2016 | new cv's | 343 | 200 | 2997.88 | 0.0003812 | yes | 163.6 | 160.8 | 37.6 | 61.8 | 63.3 |
| Model G | 2016 | new cv's | 343 | 200 | 2672.99 | 0.0000301 | yes | 172.7 | 175.6 | 40.5 | 68.8 | 70.9 |

The first three models illustrate progress from the 2015 assessment model (2015AMO) to a converged version based on the same data but evaluated using 200 jittered parameter runs (2015AMR), and finally to a converged version using cv's for the NMFS trawl survey mature biomass time series based on the "new" cv calculation (2015AMN). The next three (2015AM, Model A, Model B) illustrate the progression from the 2015 assessment model configuration with 2016 data to the CPT's requested base model for this assessment (Model B). Models C through G illustrate incremental changes to Model B requested by the CPT in May.

All new model scenarios were evaluated using 200 runs with jittered initial parameter values to select the run with the smallest objective function value and smallest maximum gradient. For each model, the selected run was re-run to invert the hessian and obtain standard deviations for parameter estimates. While all models resulted in hessians that were invertible and provided uncertainty estimates associated with the parameter estimates, the "best" run for Model A had clearly not yet converged to a minimum because the maximum gradient value was far too large (1.5256). It is surprising that the hessian was invertible for this model, but the result is clearly not valid and Model A is dropped from further consideration (note: it was not a model requested by the CPT).

Results of the progression from the 2015 assessment model with 2015, model scenario 2015AMO, to the same model configuration but with 2016 data (including the "new" survey biomass cv's), model scenario 2015AM, are provided in Appendix A.

Results of the change from the 2015AM model scenario to the base model requested by the CPT for the 2016 assessment, Model B, are summarized in Appendix B.

Results of the change from Model B to Model C, relevant to model selection, are summarized in Appendix C.

Results of the progression from Model C: Model D: Model E: Model F: Model G, relevant to model selection, are summarized in Appendix D.

More complete comparisons are provided in the accompanying on-line material at the Council website.
c. Evidence of search for balance between realistic (but possibly overparameterized) and simpler (but not realistic) models.
All models considered were parameterized in substantially similar fashion, so no simpler or more realistic models were considered.

## d. Convergence status and convergence criteria

Convergence in all models was assessed by running each model 200 times with randomly-selected ("jittered") initial parameter values for each run. 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 achieving a minimum threshold for the max gradient or exceeding the maximum number of iterations.

## e. Sample sizes assumed for the compositional data

Input sample sizes used for compositional data are listed in Tables 5-9 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):

$$
S S_{y}^{\text {inp }}=\min \left(200, \frac{S S_{y}}{(\overline{S S} / 200)}\right)
$$

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

## f. Parameter sensibility

As noted in Appendix D, estimates for the ln-scale effort extrapolation (fishery q) parameters estimated for the snow crab and BBRKC fisheries in Models D, E, F and G are unreasonably small (on the order of -19) and consequently result in associated bycatch fishing mortality rates before 1992 in these fisheries that are essentially zero. Uncertainty estimates associated with these parameters were also very large (std. dev. $=\sim 800$ ). Consequently, these models were no longer considered as viable candidates for preferred model.

Most parameter estimates obtained for Model C appear to be reasonable, or at least consistent with the 2015 assessment (Tables 20-28). An exception was the estimated $1996 \ln$-scale deviation to $50 \%$-selected for total-catch of males in the directed fishery, which hit its lower bound in Model C. Other parameters that were limited by the bounds placed on them in Model C were also limited un the 2015 assessment, and those that did so hit their upper bounds. These included the female growth parameter "a" (Table 20), the offset from 50 -to- $95 \%$ selected for female selectivity in surveys 1982-present (Table 20), and the sizes at $50 \%$-selected for male bycatch in the BBRKC fishery before 1997 and after 2004 (Table 25). Another parameter in Model C that had a questionable value was the $\ln$-scale female offset to the fully-selected
male fishery capture rate in the BBRKC fishery, which had a value of 2.44 (Table 24)-implying female Tanner crab experienced 10 times the capture rate in the BBRKC fishery that males did. However, a similar value (2.44) was estimated in Model B.
g. Criteria used to evaluate the model or to choose among alternative models

Criteria used to evaluate the alternative models were based primarily on: 1) goodness of fit and likelihood criteria, 2) parameter sensibility, and 3) biological realism.

## h. Residual analysis

Residuals for the author's preferred model, Model C, are discussed below under the Results section.
i. Evaluation of the model(s)

Of the models evaluated with data for 2016, Models 2015AM and Model A were run to illustrate the progression of models (and data) from the 2015 assessment to the CPT's base model for this assessment (Model B), and thus were not considered as suitable for selection. Of the remaining models, Models B and $C$ yielded almost identical results, so Model $C$ was preferred relative to Model B because it removed a constraint on bycatch F rates in the BBRKC fishery that fixed minimum F's. Model D was eliminated from consideration because the estimated parameters converting effort to bycatch fishing mortality rates (i.e., fishery q's) in the snow crab and BBRKC fisheries were unreasonably small-resulting in predicted bycatches of almost 0 prior to the period when observations of bycatch were available (early 1990s). Models E, F, and G were also eliminated from further consideration for this reason, because each was "built" on Model D as a base model. It will be worthwhile, in future work, to reconsider the incremental changes embodied in Models E, F and G using Model C as a base rather than model D (i.e., eliminate estimating fishery q's as model parameters).

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

Model C was selected as the author's preferred model for the 2016 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 13-18 fro the 2015 assessment model and Model C. Weighting factors applied to the various components included in the overall model objective function, including likelihoods, penalties and priors, are listed in Table 19.

## 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 21-28.
ii. Abundance and biomass time series, including spawning biomass and MMB.

Estimates for mature survey biomass, by sex, are listed in Table 29 and for mature biomass at mating, by sex, in Table 30. Numbers at size for males and females are given by year in 5 mm CW size bins in Tables 31 and 32 , respectively.

## iii. Recruitment time series

The estimated recruitment time series from the 2015 assessment and Model C are listed in Table 33.
iv. Time series of catch divided by biomass.

A comparison of catch divided by biomass (i.e., exploitation rate) from the 2015 assessment and Model C is listed in Table 34.

## c. Graphs of estimates

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

Estimates of natural mortality by sex and maturity state are shown in Fig. 31. Mortality rates are assumed equal by sex for immature crab, but are allowed to differ by sex for mature crab. Mortality rates for mature crab are estimated by sex across two time periods: 1949-1979+1985-2013 and 1980-1984. The latter period has been identified as a period of high natural mortality in the BBRKC stock (Zheng et al., 2012) and was identified as a separate period for Tanner crab in the 2012 assessment. The following table summarizes the estimated rates by stock component:

| Stock component | Normal period |  | High Mortality |  |
| :--- | :--- | :--- | :--- | :--- |
|  | 2015 assessment | Model C | 2015 Assessment | Model C |
| immature crab | 0.24 | 0.24 | 0.24 | 0.24 |
| mature females | 0.35 | 0.33 | 0.37 | 0.44 |
| mature males | 0.26 | 0.27 | 0.92 | 0.76 |

While the rates are almost identical in the "normal" period, Model C's estimates for mature males and females are substantially smaller than those from the 2015 assessment. This is the reverse of what occurred moving from the 2014 assessment to the 2015 assessment with the adoption of the "standardized" trawl survey dataset that included the "old" mature survey biomass cv's. When these were replaced by the new cv 's, the natural mortality rates decreased.

Estimated sex- and size-specific probabilities of the terminal molt-to-maturity are quite similar for the 2015 assessment model and Model C, despite different parameterizations used in the two models (Fig. 32). Estimated sex-specific mean post-molt size, as a function of pre-molt size, is also quite similar for the two models (Fig. 33).

For both sexes, survey selectivity curves (Fig. 34) estimated by the 2015 assessment model and Model C are almost identical for the first survey time period (pre-1982) for both sexes, but have slightly larger slopes and reach higher asymptotes in the 2015 assessment model for the second and third time periods (1982-present). This is a result of Model C estimating a smaller survey q for females and a larger estimated size at $95 \%$-selected for males.

Retention curves in the directed fishery estimated by the 2015 assessment model and Model C are almost identical (Fig. 35). The estimated selectivity curve for males in the directed fishery prior to 1991 (Fig. 36) for Model C is slightly left-shifted to smaller sizes relative to that from the 2015 assessment; this is probably a result of the different fishing mortality models used (the 2015 assessment used the "standard" Tanner crab model used in prior assessments, while Model C uses the Gmacs model; see Stockhausen, 2015). Conversely, the estimated selectivity curve for female bycatch in the directed fishery (Fig. 36) for Model C is substantially left-shifted to smaller sizes relative to that from the 2015 assessment model. This is not a result of the two different fishing mortality models; rather, it is a result of estimating a femalespecific offset to the male capture rate in the directed fishery in Model C (none was estimated in the 2015 assessment).

Estimated selectivity curves in the period 1991-present from Model C are generally left-shifted to smaller sizes compared to those from the 2015 assessment model (Fig. 37). In part, this reflects the difference in fishing mortality models: the selectivity functions in Model C reflect annual size-dependence in fishery
capture rates in the directed fishery while those in the 2015 assessment model reflect the size dependence of fishery (retained + discard) mortality rates.

Separate curves are estimated for 3 different time periods for each bycatch fishery, corresponding to changes in available data and fishery activity. For the snow crab fishery, separate sex-specific curves are estimated for 1989/90-1996/97, 1997/98-2004/05, and 2005/06-present. The time periods are the same for the BBRKC fishery. The directed Tanner crab fishery was closed during 1997/98-2004/05, which may have encouraged changes in how the snow crab and BBRKC fisheries were prosecuted-with associated changes in bycatch selectivity on Tanner crab. For the groundfish fisheries, the three time periods corresponding to the selectivity curves are 1973-1987, 1988-1996, and 1997-present. These correspond to changes in the groundfish fleets and Tanner crab fishery, with the curtailment of foreign and joint-venture fishing by 1988, the expansion of domestic fisheries from 1988 to 1996, and the closure of the tanner crab fishery in 1996/97. Estimated male selectivity curves in the bycatch fisheries (Fig.s 38-40) from the two models are similar for each time period, whereas the female selectivity curves tend to be left-shifted to smaller sizes in Model A relative to the 2015 assessment model (Fig.s 38-40). Again, this latter phenomenon is due to estimating female-specific offsets to male capture rates in Model A.

## iii. Estimated full selection F over time

Estimated time series of fully-selected F on males in the directed fishery and as bycatch in the snow crab, BBRKC and groundfish fisheries are compared in Fig.s 41-44 between Model C and the 2015 assessment. It should be noted that fully-selected "capture rates" are estimated directly in Model C while mortality rates are derived after applying assumed handling mortality rates, whereas the 2015 assessment model estimates the mortality rates directly (and does not estimate capture rates at all). For males in the directed fishery (Fig. 41), rates in Model C are slightly higher early in the model period (pre-2000), but rates in both models are similar more recently (post 2000). Because these are "fully-selected" rates, there is no difference between capture rate, total mortality rate, and retained mortality rate as long as retention is $100 \%$ for large crab (as is the case for both models). In contrast, capture and (bycatch) mortality rates for females in the directed fishery in Model C are generally lower than for the 2015 assessment model because the same mortality rates are applied to males and females in the 2015 assessment model while a female-specific ln-scale offset to the male rate is estimated in Model C. Similar observations hold for comparisons of the results for the snow crab fishery (Fig. 42) and the groundfish fisheries (Fig. 44). Results for the BBRKC fishery show more contrast between the two models (Fig. 43), but this is partly because the F's were fixed (not estimated) in the 2015 assessment whereas they are estimated for 1992present in Model C. As noted previously, the estimated female-specific offset for this fishery in Model C is greater than 1 .
ii. Estimated male, female, mature male, total and effective mature biomass time series The time series of recruitment estimated in the 2015 assessment and by Model C are remarkably similar (Table 33, Fig. 45). Both indicate a peak in recruitment in 1964 (probably a model artifact reflecting the start of retained catch data in 1965) followed by a steady decline into the mid-1970s, another peak in1976 followed again by declining recruitment. This decline bottoms out in 1980-1982, recruitment increases to a 4 -year plateau in the mid-1980s, declines to low values in the early-to-mid 1990s, then undergoes a period of oscillations with increasing amplitude through 2005 followed by a 4 -year low to 2008. After 2008, both models estimate increased recruitment in 2009-2011, followed by a return to lower levels in 2012-present. In general, recruitment is estimated to be much lower since 1990 than prior to 1990.

Estimates of population abundance in the 2015 assessment and from Model C exhibit similar patterns of variability, although the magnitudes differ in some cases (Fig.s 46, 47). Abundance in both models builds to a maximum in 1965-66, although the 2015 assessment estimates a somewhat larger maximum than does Model C. Abundance then follows a declining trend, with superimposed fluctuations, to 1982-83, rebuilds to a much smaller peak in 1987, and declines into a broad "valley" extending from 1993 to 2001
or so. Since 2000, population abundance has exhibited (in both models) fairly large fluctuations, possibly superimposed on a (very) gradual upward trend. Model C estimates slightly higher abundance than the 2015 assessment, although the pattern of variability is the same.

Estimates of mature biomass from the 2015 assessment and Model C also (not surprisingly) exhibit similar patterns of variability (Fig. 48), being basically smoothed versions of the population abundance trajectories.

## iv. Estimated fishing mortality versus estimated spawning stock biomass

See Section F (Calculation of the OFL; Fig. 94).
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

Model fit to retained catch is shown Fig. 49. The fits are generally quite good in both the 2015 assessment and for Model C, except for the terminal model year, where both models underpredict actual retained catch. Similarly, fits to male total (retained+discard) mortality, based on at-sea observer data, are generally quite good for both models, although (in contrast to retained catch) both models overpredict total mortality in the terminal model year (Fig. 50). Similar observations hold for predictions of male discard mortality in the directed fishery (Fig. 51), although these data are not directly fit in the model. These opposing terminal year misfits may indicate a recently-introduced (post-2009) bias between the atsea observer data and the dockside observer data which the models can't resolve. Recent changes in retention practices not reflected in the models may also be a source of this tension.

Fits to bycatch data are also generally good for males in both the 2015 assessment and for Model C for the snow crab fishery (Fig. 52). Fits to males look poorer in both models in the BBRKC fishery (Fig. 53), although Model C captures the mean level slightly better than does the 2015 assessment. One reason for the "poor" fits to the BBRKC fishery bycatch is that the bycatch levels ( $<100 \mathrm{t}$ ) are smaller than the assumed uncertainty ( $\sim 500 \mathrm{t}$ ) in the likelihood, so the models think the fits are adequate. Improving the fits would require assuming smaller levels of uncertainty, but this may not be worthwhile in terms of overall model performance.

Fits to female bycatch data in all the crab fisheries (Fig. 51-53) are not really very good for either the 2015 assessment model or Model C, even though Model C includes female-specific offsets to male fishing mortality. The problem with both models is twofold: first) predicted female bycatch is constrained to follow a temporal pattern similar to that for males, but observed mortality des not; and second) female bycatch levels in all the crab fisheries are much smaller than the assumed uncertainty levels and consequently fitting female bycatch levels more closely has little leverage in minimizing the overall model objective functions.

Bycatch in the groundfish fisheries is not sex-specific. Fits to total bycatch mortality in the groundfish fisheries are very good both for Model C and in the 2015 assessment. Both models nicely capture the peak at the beginning of the time series, followed by the rapid decline and subsequent fluctuations. Since 2008/09, total bycatch mortality has been less than 500 t and both models have over-predicted it (although the predictions are essentially identical).

The "goodness of fit"s to the fishery catch data, as they influence the likelihoods in the 2015 assessment model and Model C, is also evident of plots of $z$-scores for the fishery catch data (Fig.s 55 and 56, males only). That almost all the z -scores are < 1 indicates that probably little improvement to the current fits in
terms of absolute (rather than relative) error will occur without changing the assumed uncertainty levels for the fishery data.

## ii. Graphs of model fits to survey numbers

Time series of observed biomass of mature crab in the NMFS bottom trawl surveys are compared by sex with model-predicted values for Model C and the 2015 assessment in Fig. 57. The difference in cv's for the observed data appears to have little direct impact on the trajectories of the model-predicted time series. Both the model and the assessment under-predict mature female survey biomass in the early 1980s and again in the early 1990s. They also under-predict mature male survey biomass in the early 1990s as well as in the mid-2000s. The scale of the standardized log-scale residuals (Fig. 58) indicates mediocre fits for (the standard deviation of the residuals is $\sim 2$, whereas $\sim 1$ would indicate a good fit). In almost all cases, though, Model C exhibits slightly smaller relative errors in comparison with the 2015 assessment results.

Model predictions for total survey numbers of preferred males ( $\geq 125 \mathrm{~mm} \mathrm{CW}$ ) are compared with observations from the survey in Fig. 59. These data are not fit in the models, and so provide a somewhat independent test of model fitting. Prior to 2000, both models tended to underpredict observed survey abundance when it was high, but overpredict it when it was low. In recent years, both models rather substantially over-predict numbers of large crab in the survey.
iii. Graphs of model fits to catch proportions by length

Model-predicted proportions at size from the 2015 assessment and Model A for retained males in the directed Tanner crab fishery are presented in Fig. 60. A plot of the Pearson's residuals for the fits is presented in Fig. 61. Both models appear to fit the observed proportions quite similarly, although Model C fits slightly better in 1991-1996 and 2005-2008 (the fishery was closed 1997-2004) because, although its shapes are similar to those from the 2015 assessment, they are slightly right-shifted to larger sizes (as the data tends to be). For $2014(2014 / 15)$, both models predict more retained crab at larger sizes than is seen in the data. This pattern extends to $2015(2015 / 16)$ for Model C. This is consistent with a recent shift in industry retention to smaller sizes not yet reflected in the models.

Model-predicted patterns from the 2015 assessment and Model C for the proportions caught-at-size in the directed fishery are shown in Fig. 62 for males, Fig. 63 for females, and as Pearson's residuals for both sexes in Fig. 64. General residual patterns indicate that the fishery catches a larger proportion of small male crab than predicted by the models (except in 1996), and catches fewer large male crab than predicted by the models. This is particularly true in $2009(2009 / 10)$, when the area west of $166^{\circ} \mathrm{W}$ longitude was closed to directed fishing. Conceivably, among other potential explanations, this pattern may indicate that an asymptotic selectivity curve is inappropriate for the male selection process or that the model overestimates growth into the largest size classes for males. 1996 is the exception to this, and exhibits extremely poor (though different) absolute fits to the data for the two models (Fig. 62), although the relative fits are good (as evidenced by the small values for the Pearson's residuals for males in 1996; Fig. 64). As previously noted, however, the relative weight (input sample size) put on fitting this data in the likelihood is quite small. It is notable that the fit to the 1996 bycatch size composition for females is much better, but in general the residuals for females are much smaller. This is somewhat surprising given that a single selectivity pattern is estimated for females while the male selectivity pattern (the $50 \%$-selected parameter of the logistic function) is allowed to vary from year-to-year after 1991.

Model-predicted patterns from the 2015 assessment and Model C for the proportions caught-at-size as bycatch in the snow crab fishery are shown in Fig. 65 for males, Fig. 66 for females, and as Pearson's residuals for both sexes in Fig. 67. Estimates from both models for males are almost identical. Estimates for females are quite similar, although some differences between the models can be seen at small sizes for 1992-1996.

Model-predicted patterns from the 2015 assessment and Model C for the proportions caught-at-size as bycatch in the BBRKC fishery are shown in Fig. 68 for males, Fig. 69 for females, and as Pearson's residuals for both sexes in Fig. 70. As with snow crab, estimates from both models for males are almost identical. Estimates for females are also almost identical.

Model-predicted patterns from the 2015 assessment and Model C for the proportions caught-at-size as bycatch in the groundfish fisheries are shown in Fig. 71 for males, Fig. 72 for females, and as Pearson’s residuals for both sexes in Fig. 73. These proportions-at-size are fit as extended size compositions, where the annual proportions sum to 1 over both sexes, in contrast to the proportions in the crab fisheries where the proportions sum to 1 over each sex individually. Extended size compositions are fit for the groundfish fisheries because the associated observed bycatch mortality is not sex-specific and the extended compositions allow the models to extract information on the relative abundance of males vs. females in these fisheries. The model-predicted size compositions in the groundfish fisheries are relatively similar for males, differing mainly in magnitude. For females, the patterns for 1973-1996 are similar and differ, like males, somewhat in overall magnitude rather than in shape. However, during the period 1997-present the magnitudes are substantially different (unfortunately, the model-predicted size compositions from the 2015 assessment blend into the data bars) - with the 2015 assessment size compositions of much smaller magnitude (and much worse fit) than those from Model C. The poor behavior of the 2015 assessment model was traced earlier this year to how the sex-specific size compositions were combined to form the extended composition. Previous to this year, the size compositions were combined using the input sample sizes to weight the size compositions. However, this approach did not always preserve the relative abundance scales inherent in the observed sample sizes. In Model C, the extended size compositions are created using the observed male and female sample sizes to weight the sex-specific size compositions, then fit using the input effective sample sizes. The new approach vastly improved the overall fits for the female size compositions (Fig. 73), as well as slightly improving the fits to the male size compositions.

## iv. Graphs of model fits to survey proportions by length

Model fits from the 2015 assessment and Model C to observed proportions-at-size in the annual NMFS trawl survey are shown for males in Fig. 74. The similarity in results between the two models is fairly remarkable. As with the 2015 assessment model, Model C appears to be suitably sensitive to relatively large cohorts recruiting to the model size range (e.g., 1997-2002), but appears to be less able to track strong cohorts through time (the mode in the model proportions at $\sim 100 \mathrm{~mm} \mathrm{CW}$ in 1982 disappears after two years, but appears to last until at least 1985 in the observed proportions. After 1982, the model tends to under-predict size proportions for males in the $70-120 \mathrm{~mm}$ range and over-predict the proportion of large (> 120 mm CW) males after 2000. Model fits to proportions at size in the survey for females are shown in Fig. 75. The model tends to over-predict proportions-at-size in the $65-85 \mathrm{~mm}$ CW range. The patterns of residuals for males and females evident in the bubble plots for Model A are almost identical to those obtained from the 2015 assessment (Fig. 76).
v. Marginal distributions for the fits to the compositional data.

Marginal fits for the Model C-predicted proportion of crab by size in the directed fishery catch are similar to those for the 2015 assessment model: the models somewhat over-predict proportions for retained males at sizes smaller than the peak and under-predict proportions at sizes larger than the peak (Fig. 77). Model C does a slightly poorer job in this respect than the 2015 assessment model. In contrast, the model underpredicts proportions near the peak and somewhat smaller for all males caught (retained and discarded) in the directed fishery, but over-estimates the proportions for crab larger than the peak (Fig. 78, lower plot). This may indicate an unresolved tension between the retained size comps and the total-catch size comps. Model C appears to reflect observed marginal female bycatch size composition pattern quite well, while the 2015 assessment model under-predicts proportions of crab just smaller than the peak and over-predicts proportions just larger (Fig. 78, upper plot).

The observed and predicted (Model A) marginal proportions for males taken as bycatch in the snow crab fishery are in good agreement at all sizes for both models (Fig. 79, lower plot), while both models tend to underestimate the proportion of females taken as bycatch near the peak proportions ( $\sim 80-90 \mathrm{~mm} \mathrm{CW})$ and over-estimate the proportions at larger sizes (Fig. 79, upper plot). The opposite pattern is true for both models regarding the proportion-at-size of females taken as bycatch in the BBRKC fishery, where intermediate-size females are over-represented in the model predictions and under-represented at larger sizes (Fig. 80). The patterns of model-predicted marginal proportions-at-size for males taken as bycatch in the BBRKC fishery are similar to that found for the snow crab fishery, but shifted to larger sizes by $\sim 20$ mm CW. Unfortunately, these result in poorer fits to the observations, overestimating proportions at larger sizes and underestimating them at smaller sizes, than those for the snow crab fishery. The patterns of marginal predicted proportions at size for males and females taken in the groundfish fishery (Fig. 81) obtained using Model C are much closer to the data than those obtained in the 2015 assessment. The improvement occurs Model C uses an improved approach to combining the male and female size compositions prior to fitting them (documented at the May 2016 CPT meeting).

Marginal fits of Model A-predicted proportion-at-sizes in the survey are presented in Fig. 82. The model's marginal survey proportions fit the data quite well, and in quite similar fashion to the 2014 assessment.

## vi. Plots of implied versus input effective sample sizes and time-series of implied effective sample sizes.

Time series of implied effective sample sizes, using the McAllister-Ianelli method, are shown in Fig.s 8385 for retained catch and total catch size compositions in the directed fishery (Fig. 83), bycatch size compositions in the snow crab, BBRKC and groundfish fisheries (Fig. 84), and the NMFS EBS bottom trawl survey (Fig. 85). 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).
Not available.
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.
Not available.

> 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).

Results from a 10 -year retrospective analysis for Model C, the author's preferred model, are shown in Fig.s 86-89 for mature biomass-at-mating, recruitment, mature survey biomass and retained catch biomass. The plots for mature biomass-at-mating and recruitment (Fig.s 86, 87) display strong retrospective patterns, such that models that are terminated earlier are biased high relative to models that are terminated later. The plot for mature survey biomass indicates the model is almost always biased high in the terminal year of the model run, particularly when the end-year observations are smaller than the previous year (Fig. 88). However, there does not seem to be a similar pattern for fitting retained catch biomass (Fig. 89).
ii. Historic analysis (plot of actual estimates from current and previous assessments). Many of the plots contained in this assessment feature comparisons between results from the 2015 assessment model and the author's preferred model for this assessment. Most of them indicate little difference between the two models, particularly for more recent periods (e.g., since 1990).

## g. Uncertainty and sensitivity analyses

Not available.

## 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 2015/16 was 27.19 thousand t while the total catch mortality for 2014/15 was 11.38 thousand $t$, based on applying discard 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 reported catch by fleet for 2015/16 (Tables 1 and 4). 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 (Fig. 90):

$$
\begin{array}{llc}
\hline B, F_{35 \%}, B_{35 \%} * & F_{\text {OFL }}=F_{35 \%} * & \\
& \text { a. } \frac{B}{B_{35 \%^{*}}}>1 & \\
& \text { b. } \beta<\frac{B}{B_{35 \%} *} \leq 1 & F_{\text {OFL }}=F^{*}{ }_{35 \%} \frac{\frac{B}{B_{35 \%}^{*}}-\alpha}{1-\alpha}
\end{array} \quad \text { ABC } \leq(1-\mathrm{by}) * \text { OFL }
$$

and is based on an estimate of "current" spawning biomass at mating ( $B$ above, taken as MMB at mating in the assessment year) and spawning biomass per recruit (SBPR)-based proxies for $\mathrm{F}_{\text {MSY }}$ and $\mathrm{B}_{\text {MSY }}$. In the above equations, $\alpha=0.1$ and $\beta=0.25$. For Tanner crab, the proxy for $\mathrm{F}_{\mathrm{MSY}}$ 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 $\mathrm{B}_{\text {MSY }}$ 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 2015/16 require estimates of $B=\mathrm{MMB}_{2016 / 17}$ (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 $\mathrm{F}_{\mathrm{OFL}}=\mathrm{F}_{35 \%}$. If the ratio is less than one but greater than $\beta$, then the stock falls into Tier 3 b and $\mathrm{F}_{\mathrm{OFL}}$ is reduced from $\mathrm{F}_{35 \%}$ following the descending limb of the control rule (Fig. 90). If the ratio is less than $\beta$, then the stock falls into Tier 3 c 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 previous assessments, 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. 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 (Rugolo and Turnock, 2012b; Stockhausen 2015). The selectivity and retention curves used to calculate the OFL are shown in Fig.s 91-92.

To calculate the Forl 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. However, this calculation also depends on the assumed bycatch F's on Tanner crab in the snow crab, BBRKC and groundfish fisheries. For the latter two fisheries, the average F over the last 5 years is used in the calculations. Because the snow crab fishery typically accounts for the largest bycatch mortality in the bycatch fisheries, and because the FOFL for snow crab is frequently a good predictor of the actual F in the upcoming year, a different approach is used to determine the snow crab fishery F for Tanner crab bycatch. For the snow crab fishery, the ratio of the $\mathrm{F}_{\text {ofL }}$ from the snow crab assessment author's preferred model to the average F over the last 5 years is used to scale the 5 -year average bycatch F on Tanner crab. For this assessment, the snow crab FOFL is $1.24 \mathrm{yr}^{-1}$ (Szuwalski, 2016), the 5 -year average F is $0.979 \mathrm{yr}^{-1}$, the resulting ratio is 1.266 , and the fully-selected Tanner crab bycatch capture rate used in the projection model was $0.092 \mathrm{yr}^{-1}$.

OFL results from the projection model using the same approach for each of the "converged" models considered in this assessment (consequently values for Model A are missing) are listed for illustrative purposes only in Table 35. The change from the "old" (2015AMR) to the "new" (2015AMN) survey biomass cv's resulted in higher values for average recruitment ( 176.78 vs. 193.44 million crab), projected MMB-at-mating $(B)$ for $2015 / 16$ ( 51.41 vs. 63.85 thousand t ), B BSY ( 25.68 vs. 29.42 thousand t ), and OFL for 2015/16 ( 25.68 vs. 30.96 thousand $t$ ), although $\mathrm{F}_{\text {MSY }}$ was similar ( 0.58 vs. 0.56 ). Adding the 2015/16 fishery data and 2016 survey data (2015AM) reduced estimates of average recruitment (183.46 million crab), projected MMB-at-mating for 2016/17 ( 48.07 thousand $t$ ), and $\mathrm{B}_{\text {MSY }}$ ( 26.68 thousand t ), while $\mathrm{F}_{\text {MSY }}$ was similar (0.59). The OFL for 2016/17 using the 2015 assessment model configuration would be substantially smaller ( 23.79 thousand $t$ ) than that for 2015/16 from the converged model (2015AMR). Moving to the base 2016 model (Model B) involved a host of changes to the model configuration reviewed during the May 2016 CPT meeting. Compared with the 2015 model configuration run with the 2016 data (2015AM), the results from Models B and C (the author's preferred model) are really fairly similar except that $\mathrm{F}_{\text {MSY }}$ is 0.79 for the latter models and 0.59 for 2015AM. The value of $\mathrm{F}_{\text {MSY }}$ from Model $\mathrm{D}(0.09)$ does not appear to be valid, and calls into question results from the succeeding models (E through G) which build on it, although they seem more plausible. Model D, as discussed previously, was the first model to estimate the conversion from effort to fishery capture rates in the absence of bycatch data as parameters for the snow crab and BBRKC fisheries-resulting in anomalously small conversion factors.

The estimate of $B$ from Model C , the author's preferred model, is 45.34 thousand $t$ (Table 35). Male spawning biomass per recruit in an unfished stock was calculated using the TCSAM population dynamics equations (Stockhausen, 2014) with total recruitment set to 1 and fishing mortality from all sources (directed fishery and all bycatch fisheries) set to 0 , resulting in $\phi(0)=0.402 \mathrm{~kg} /$ recruit. $\mathrm{F}_{35 \%}$ was calculated for this model as $0.79 \mathrm{yr}^{-1}$, which is quite a bit larger than that calculated last year $\left(0.58 \mathrm{yr}^{-1}\right)$ but this is primarily an effect of the change to the Gmacs fishing mortality model. For the 2015 assessment, the size dependence of fishing mortality rates on males in the directed fishery followed a logistic curve. For the Gmacs fishing mortality model, the size dependence of the fishery capture rates
follows a logistic curve, but the resulting size dependence for fishing mortality is no longer a logistic shape.

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 $(\bar{R})$. After much discussion in 2012 and 2013, the SSC endorsed an averaging period of 1982+. Starting the average recruitment period in 1982 is consistent with a 5-6 year recruitment lag from 1976/77, when a well-known 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 the author's preferred model is 182.27 million. The estimates of average recruitment are reasonably similar between the 2015 assessment model and the author's preferred model (Table 33, Fig. 45). The value of $\mathrm{B}_{\text {MSY }}=\mathrm{B}_{35 \%}$ for $\bar{R}$ is 25.65 thousand t . Thus, the stock is "not overfished" because $B / \mathrm{B}_{35 \%}>0.5$ (i.e., $B>$ MSST).

Once Fofl is determined using the control rule (Fig. 90), the (total catch) OFL can be calculated based on projecting the population forward one year assuming that $F=$ FofL. In the absence of uncertainty, the OFL would then be the predicted total catch taken when fishing at $F=\mathrm{F}_{\text {ofl }}$. 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 $F=$ Fofl.

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 $\operatorname{size} \operatorname{bin} z, w_{x, z}$ 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, 2016 as estimated by the assessment model.

Assessment uncertainty was included in the calculation of OFL using the same approach as that used for previous assessments (Stockhausen, 2014, 2015). Basically, initial numbers at size on July 1, 2016 were randomized based on an assumed lognormal assessment error distribution and the cv of estimated MMB for $2015 / 16$ from the assessment model, the control rule was applied to obtain $\mathrm{F}_{\mathrm{OFL}}$, and the population projected forward to next year assuming that fishing occurred consistent with $\mathrm{F}_{\text {ofl }}$. This was repeated 10,000 times to generate a distribution of total catch OFLs. The value of OFL for 2016/17 from the author's preferred model (Model C) is $\mathbf{2 5 . 6 1}$ thousand $\mathfrak{t}$ (Table 35, Fig. 93).

Model C is the author's preferred model for calculating the $\mathrm{B}_{\text {MSY }}$ proxy as $\mathrm{B}_{35 \%}$, so MSST $=0.5 \mathrm{~B}_{\text {MSY }}=$ 12.82 thousand t . Because current $B=45.34$ thousand $\mathrm{t}>$ MSST, the stock is not overfished. The population state (directed F vs. MMB) is plotted for each year from 1965-2014 in Fig. 94 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 ACL=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 ABC control rule are: 1) a constant buffer where the ABC 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 $\left(\mathrm{P}^{*}\right)$ 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 ACL=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.

ABCs based on the $\mathrm{P}^{*}=0.49$ approach were calculated from quantiles of the associated OFL distributions such that probability that the selected ABC was greater than the true OFL was 0.49 . The resulting ABC for each scenario was almost identical to the associated OFL (Table 35). ABCs were also calculated using the SSC's 20\% OFL buffer (Table 35).

For the author's preferred model, Model C, the P* ABC ( $\mathrm{ABC}_{\max }$ ) is 25.57 thousand t while the $20 \%$ Buffer ABC is 20.49 thousand t . The author remains concerned that the projection model, based on $\mathrm{F}_{35 \%}$ as a proxy for $\mathrm{F}_{\text {MSY }}$, is overly optimistic regarding the actual productivity of the stock. Fishery-related mortality similar to these ABC levels 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. Given this uncertainty concerning the stock, the author recommends using the $\mathbf{2 0 \%}$ buffer adopted by the SSC last yearfor this stock to calculate ABC. Consequently, the author's recommended ABC is 20.49 thousand $t$.

## 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 finally been collected in the EBS on Tanner crab (molt increments observed on 100+ individuals collected in 2015 and 2016; R. Foy, AFSC, pers. comm.). More data regarding temperature-dependent effects on molting frequency would be helpful to assess potential impacts of the EBS cold pool on the stock. Information on temperature-dependent changes in crab movement and survey catchability would also e of value. In addition, it would be extremely worthwhile to develop a "better" index of reproductive potential than MMB that can be calculated in the assessment model and 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 East $166^{\circ} \mathrm{W}$ and West $166^{\circ} \mathrm{W}$ directed fisheries should be explicitly represented in the assessment model should be addressed. In addition, how, and whether or not, bycatch in the groundfish fisheries should be split into pot- and trawl-related components should be addressed.

Transition to the new model code (TCSAM2015) will occur this fall in preparation for the Modeling Workshop. Substantial progress was made this summer to allow detailed comparison of model results from the current model code (TCSAM2013) and the new code (TCSAM2015). With the implementation of TCSAM2015, several research avenues can be explored: 1) time-varying growth; 2 ) fitting molt increment data directly in the model, 3) alternative time periods for defining retention/selectivity functions, and 4) decomposing the currently "lumped" directed fishery into its eastern and western components. Development of a fully Gmacs version of the Tanner crab model will also begin.

## 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, perhaps an ideal 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 on a decadal time scale (Rugolo and Turnock, 2012), suggesting a potential 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 | unlikely to have substantial effects | minimal to none |
| HAPC biota | crab pots have a very small footprint on the bottom crab pots are unlikely to | unlikely to be having substantial effects postrationalization | minimal to none |
| Marine mammals and birds | 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 substantially reduced in | unlikely to have substantial effects | minimal to none |
| Fishery concentration in space and time | 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 | possible concern |

fishery
Fishery effects on age-atmaturity and fecundity
unknown
possible concern

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Table 1. Retained catch (males) in directed Tanner crab fisheries.

| Eastern Bering Sea Chionoecetes bairdi Retained Catch (1,000's t) |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Year | US Pot | Japan | Russia | Total |
| 1965/66 |  | 1.17 | 0.75 | 1.92 |
| 1966/67 |  | 1.69 | 0.75 | 2.44 |
| 1967/68 |  | 9.75 | 3.84 | 13.60 |
| 1968/69 | 0.46 | 13.59 | 3.96 | 18.00 |
| 1969/70 | 0.46 | 19.95 | 7.08 | 27.49 |
| 1970/71 | 0.08 | 18.93 | 6.49 | 25.49 |
| 1971/72 | 0.05 | 15.90 | 4.77 | 20.71 |
| 1972/73 | 0.10 | 16.80 |  | 16.90 |
| 1973/74 | 2.29 | 10.74 |  | 13.03 |
| 1974/75 | 3.30 | 12.06 |  | 15.24 |
| 1975/76 | 10.12 | 7.54 |  | 17.65 |
| 1976/77 | 23.36 | 6.66 |  | 30.02 |
| 1977/78 | 30.21 | 5.32 |  | 35.52 |
| 1978/79 | 19.28 | 1.81 |  | 21.09 |
| 1979/80 | 16.60 | 2.40 |  | 19.01 |
| 1980/81 | 13.47 |  |  | 13.43 |
| 1981/82 | 4.99 |  |  | 4.99 |
| 1982/83 | 2.39 |  |  | 2.39 |
| 1983/84 | 0.55 |  |  | 0.55 |
| 1984/85 | 1.43 |  |  | 1.43 |
| 1985/86 | 0.00 |  |  | 0.00 |
| 1986/87 | 0.00 |  |  | 0.00 |
| 1987/88 | 1.00 |  |  | 1.00 |
| 1988/89 | 3.15 |  |  | 3.18 |
| 1989/90 | 11.11 |  |  | 11.11 |
| 1990/91 | 18.19 |  |  | 18.19 |
| 1991/92 | 14.42 |  |  | 14.42 |
| 1992/93 | 15.92 |  |  | 15.92 |
| 1993/94 | 7.67 |  |  | 7.67 |
| 1994/95 | 3.54 |  |  | 3.54 |
| 1995/96 | 1.92 |  |  | 1.92 |
| 1996/97 | 0.82 |  |  | 0.82 |
| 1997/98 | 0.00 |  |  | 0.00 |
| 1998/99 | 0.00 |  |  | 0.00 |
| 1999/00 | 0.00 |  |  | 0.00 |
| 2000/01 | 0.00 |  |  | 0.00 |
| 2001/02 | 0.00 |  |  | 0.00 |
| 2002/03 | 0.00 |  |  | 0.00 |
| 2003/04 | 0.00 |  |  | 0.00 |
| 2004/05 | 0.00 |  |  | 0.00 |
| 2005/06 | 0.43 |  |  | 0.43 |
| 2006/07 | 0.96 |  |  | 0.96 |
| 2007/08 | 0.96 |  |  | 0.96 |
| 2008/09 | 0.88 |  |  | 0.88 |
| 2009/10 | 0.60 |  |  | 0.60 |
| 2010/11 | 0.00 |  |  | 0.00 |
| 2011/12 | 0.00 |  |  | 0.00 |
| 2012/13 | 0.00 |  |  | 0.00 |
| 2013/14 | 1.25 |  |  | 1.25 |
| 2014/15 | 6.16 |  |  | 6.16 |
| 2015/16 | 8.91 |  |  | 8.91 |

Table 2. Retained catch (males) in the US domestic pot fishery. Information from the Communnity 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 ADF\&G year (in parentheses, if different from the "Fishery Year") indicates the year ADF\&G assigned to the fishery season in compiled reports.

| year (ADF\&G year) | Total <br> Crab (no.) | Total Harvest (lbs) | $\begin{gathered} \text { GHL/TAC } \\ \text { (millions lbs) } \end{gathered}$ | 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) | closed | closed | closed | closed | closed |
| 1986/87 (1987) | closed | closed | closed | closed | closed |
| 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 |
| 2015/16 | 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 | closed | closed | closed | closed | closed |
| 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 | closed | closed | closed | closed | closed |
| 2011/12 | closed | closed | closed | closed | closed |
| 2012/13 | closed | closed | closed | closed | closed |
| 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 |

Table 3. Total bycatch (discards, 1000's t) of Tanner crab in various fisheries.

| Discards (1,000's t) of Tanner Crab by Fishery |  |  |  |  |  |  |  | Total <br> Discards <br> $(1,000$ 's t) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Tanner Crab |  | Snow Crab |  | Red King Crab |  | Groundfish |  |
| Year | Male | Female | Male | Female | Male | Female | All |  |
| 1973/74 |  |  |  |  |  |  | 17.735 | 17.735 |
| 1974/75 |  |  |  |  |  |  | 24.449 | 24.449 |
| 1975/76 |  |  |  |  |  |  | 9.408 | 9.408 |
| 1976/77 |  |  |  |  |  |  | 4.699 | 4.699 |
| 1977/78 |  |  |  |  |  |  | 2.776 | 2.776 |
| 1978/79 |  |  |  |  |  |  | 1.869 | 1.869 |
| 1979/80 |  |  |  |  |  |  | 3.397 | 3.397 |
| 1980/81 |  |  |  |  |  |  | 2.114 | 2.114 |
| 1981/82 |  |  |  |  |  |  | 1.474 | 1.474 |
| 1982/83 |  |  |  |  |  |  | 0.449 | 0.449 |
| 1983/84 |  |  |  |  |  |  | 0.671 | 0.671 |
| 1984/85 |  |  |  |  |  |  | 0.644 | 0.644 |
| 1985/86 |  |  |  |  |  |  | 0.399 | 0.399 |
| 1986/87 |  |  |  |  |  |  | 0.649 | 0.649 |
| 1987/88 |  |  |  |  |  |  | 0.640 | 0.640 |
| 1988/89 |  |  |  |  |  |  | 0.463 | 0.463 |
| 1989/90 |  |  |  |  |  |  | 0.671 | 0.671 |
| 1990/91 |  |  |  |  |  |  | 0.943 | 0.943 |
| 1991/92 |  |  |  |  |  |  | 2.545 | 2.545 |
| 1992/93 | 6.175 | 1.005 | 25.759 | 1.787 | 1.188 | 0.029 | 2.758 | 38.700 |
| 1993/94 | 3.870 | 1.028 | 14.530 | 1.814 | 2.967 | 0.198 | 1.760 | 26.167 |
| 1994/95 | 3.130 | 1.270 | 7.124 | 1.271 | 0.000 | 0.000 | 2.096 | 14.891 |
| 1995/96 | 2.762 | 1.760 | 4.797 | 1.759 | 0.000 | 0.000 | 1.524 | 12.603 |
| 1996/97 | 0.116 | 0.045 | 0.833 | 0.229 | 0.027 | 0.004 | 1.597 | 2.851 |
| 1997/98 | 0.000 | 0.000 | 1.750 | 0.226 | 0.165 | 0.003 | 1.179 | 3.323 |
| 1998/99 | 0.000 | 0.000 | 1.989 | 0.175 | 0.119 | 0.003 | 0.934 | 3.220 |
| 1999/00 | 0.000 | 0.000 | 0.695 | 0.145 | 0.076 | 0.004 | 0.630 | 1.551 |
| 2000/01 | 0.000 | 0.000 | 0.146 | 0.022 | 0.067 | 0.002 | 0.739 | 0.976 |
| 2001/02 | 0.000 | 0.000 | 0.323 | 0.011 | 0.043 | 0.002 | 1.184 | 1.563 |
| 2002/03 | 0.000 | 0.000 | 0.557 | 0.037 | 0.062 | 0.003 | 0.721 | 1.379 |
| 2003/04 | 0.000 | 0.000 | 0.193 | 0.026 | 0.056 | 0.003 | 0.422 | 0.700 |
| 2004/05 | 0.000 | 0.000 | 0.078 | 0.014 | 0.048 | 0.003 | 0.676 | 0.819 |
| 2005/06 | 0.462 | 0.044 | 0.968 | 0.043 | 0.042 | 0.002 | 0.621 | 2.182 |
| 2006/07 | 1.370 | 0.355 | 1.462 | 0.169 | 0.026 | 0.003 | 0.717 | 4.102 |
| 2007/08 | 2.041 | 0.097 | 1.872 | 0.102 | 0.056 | 0.009 | 0.694 | 4.871 |
| 2008/09 | 0.431 | 0.014 | 1.119 | 0.050 | 0.269 | 0.004 | 0.531 | 2.417 |
| 2009/10 | 0.071 | 0.002 | 1.324 | 0.014 | 0.150 | 0.001 | 0.374 | 1.937 |
| 2010/11 | 0.000 | 0.000 | 1.344 | 0.016 | 0.033 | 0.001 | 0.231 | 1.625 |
| 2011/12 | 0.000 | 0.000 | 2.119 | 0.014 | 0.017 | 0.000 | 0.203 | 2.352 |
| 2012/13 | 0.000 | 0.000 | 1.187 | 0.009 | 0.042 | 0.001 | 0.153 | 1.392 |
| 2013/14 | 0.387 | 0.023 | 1.832 | 0.015 | 0.113 | 0.001 | 0.348 | 2.720 |
| 2014/15 | 2.515 | 0.039 | 5.383 | 0.050 | 0.296 | 0.001 | 0.423 | 8.706 |
| 2015/16 | 3.045 | 0.059 | 3.519 | 0.017 | 0.174 | 0.006 | 0.352 | 7.172 |

Table 4. Bycatch (discard) mortality ( 1000 's t ) of Tanner crab in various fisheries. Discard mortality was calculated assuming mortality rates of 0.321 in the crab fisheries and 0.80 in the groundfish fisheries.

| Discard Mortality (1,000's t) of Tanner Crab by Fishery |  |  |  |  |  |  |  | Total Discard Mortality (1,000's t) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Tanner Crab |  | Snow Crab |  | Red King Crab |  | Groundfish |  |
| Year | Male | Female | Male | Female | Male | Female | All |  |
| 1973/74 |  |  |  |  |  |  | 14.188 | 14.188 |
| 1974/75 |  |  |  |  |  |  | 19.559 | 19.559 |
| 1975/76 |  |  |  |  |  |  | 7.526 | 7.526 |
| 1976/77 |  |  |  |  |  |  | 3.759 | 3.759 |
| 1977/78 |  |  |  |  |  |  | 2.221 | 2.221 |
| 1978/79 |  |  |  |  |  |  | 1.495 | 1.495 |
| 1979/80 |  |  |  |  |  |  | 2.718 | 2.718 |
| 1980/81 |  |  |  |  |  |  | 1.691 | 1.691 |
| 1981/82 |  |  |  |  |  |  | 1.179 | 1.179 |
| 1982/83 |  |  |  |  |  |  | 0.359 | 0.359 |
| 1983/84 |  |  |  |  |  |  | 0.537 | 0.537 |
| 1984/85 |  |  |  |  |  |  | 0.515 | 0.515 |
| 1985/86 |  |  |  |  |  |  | 0.319 | 0.319 |
| 1986/87 |  |  |  |  |  |  | 0.519 | 0.519 |
| 1987/88 |  |  |  |  |  |  | 0.512 | 0.512 |
| 1988/89 |  |  |  |  |  |  | 0.370 | 0.370 |
| 1989/90 |  |  |  |  |  |  | 0.537 | 0.537 |
| 1990/91 |  |  |  |  |  |  | 0.755 | 0.755 |
| 1991/92 |  |  |  |  |  |  | 2.036 | 2.036 |
| 1992/93 | 1.982 | 0.322 | 8.269 | 0.574 | 0.381 | 0.009 | 2.206 | 13.744 |
| 1993/94 | 1.242 | 0.330 | 4.664 | 0.582 | 0.952 | 0.063 | 1.408 | 9.243 |
| 1994/95 | 1.005 | 0.408 | 2.287 | 0.408 | 0.000 | 0.000 | 1.676 | 5.784 |
| 1995/96 | 0.887 | 0.565 | 1.540 | 0.565 | 0.000 | 0.000 | 1.219 | 4.776 |
| 1996/97 | 0.037 | 0.014 | 0.267 | 0.074 | 0.009 | 0.001 | 1.277 | 1.680 |
| 1997/98 | 0.000 | 0.000 | 0.562 | 0.073 | 0.053 | 0.001 | 0.943 | 1.632 |
| 1998/99 | 0.000 | 0.000 | 0.638 | 0.056 | 0.038 | 0.001 | 0.748 | 1.481 |
| 1999/00 | 0.000 | 0.000 | 0.223 | 0.047 | 0.025 | 0.001 | 0.504 | 0.800 |
| 2000/01 | 0.000 | 0.000 | 0.047 | 0.007 | 0.021 | 0.001 | 0.591 | 0.667 |
| 2001/02 | 0.000 | 0.000 | 0.104 | 0.004 | 0.014 | 0.001 | 0.947 | 1.069 |
| 2002/03 | 0.000 | 0.000 | 0.179 | 0.012 | 0.020 | 0.001 | 0.577 | 0.788 |
| 2003/04 | 0.000 | 0.000 | 0.062 | 0.008 | 0.018 | 0.001 | 0.337 | 0.427 |
| 2004/05 | 0.000 | 0.000 | 0.025 | 0.004 | 0.015 | 0.001 | 0.541 | 0.587 |
| 2005/06 | 0.148 | 0.014 | 0.311 | 0.014 | 0.014 | 0.001 | 0.497 | 0.998 |
| 2006/07 | 0.440 | 0.114 | 0.469 | 0.054 | 0.008 | 0.001 | 0.573 | 1.660 |
| 2007/08 | 0.655 | 0.031 | 0.601 | 0.033 | 0.018 | 0.003 | 0.555 | 1.896 |
| 2008/09 | 0.138 | 0.004 | 0.359 | 0.016 | 0.086 | 0.001 | 0.425 | 1.030 |
| 2009/10 | 0.023 | 0.001 | 0.425 | 0.005 | 0.048 | 0.000 | 0.299 | 0.801 |
| 2010/11 | 0.000 | 0.000 | 0.431 | 0.005 | 0.011 | 0.000 | 0.185 | 0.632 |
| 2011/12 | 0.000 | 0.000 | 0.680 | 0.004 | 0.006 | 0.000 | 0.162 | 0.852 |
| 2012/13 | 0.000 | 0.000 | 0.381 | 0.003 | 0.013 | 0.000 | 0.123 | 0.520 |
| 2013/14 | 0.124 | 0.007 | 0.588 | 0.005 | 0.036 | 0.000 | 0.278 | 1.040 |
| 2014/15 | 0.807 | 0.012 | 1.728 | 0.016 | 0.095 | 0.000 | 0.339 | 2.998 |
| 2015/16 | 0.977 | 0.019 | 1.130 | 0.005 | 0.056 | 0.002 | 0.282 | 2.471 |

Table 5. Sample sizes for retained catch-at-size in the directed fishery. $\mathrm{N}=$ number of individuals. $\mathrm{N}^{`}=$ scaled sample size used in assessment.

| year | new + old shell |  |
| :---: | ---: | ---: |
|  | N | $\mathrm{N}^{\prime}$ |
| $1980 / 81$ | 13,310 | 97.8 |
| $1981 / 82$ | 11,311 | 83.1 |
| $1982 / 83$ | 13,519 | 99.3 |
| $1983 / 84$ | 1,675 | 12.3 |
| $1984 / 85$ | 2,542 | 18.7 |
| $1988 / 89$ | 12,380 | 91.0 |
| $1989 / 90$ | 4,123 | 30.3 |
| $1990 / 91$ | 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 | 11.2 |
| $1996 / 97$ | 4,430 | 32.6 |
| $2005 / 06$ | 705 | 5.2 |
| $2006 / 07$ | 2,940 | 21.6 |
| $2007 / 08$ | 6,935 | 51.0 |
| $2008 / 09$ | 3,490 | 25.6 |
| $2009 / 10$ | 2,417 | 17.8 |
| $2013 / 14$ | 4,760 | 35.0 |
| $2014 / 15$ | 14,055 | 103.3 |
| $2015 / 16$ | 24,420 | 200.0 |

Table 6. 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 |  | $\mathrm{N}^{\prime}$ |  |
| :---: | ---: | ---: | ---: | ---: |
| year | males | females | males | females |
| $1991 / 92$ | 31,252 | 5,605 | 200.0 | 40.2 |
| $1992 / 93$ | 54,836 | 8,755 | 200.0 | 62.8 |
| $1993 / 94$ | 40,388 | 10,471 | 200.0 | 75.1 |
| $1994 / 95$ | 5,792 | 2,132 | 42.6 | 15.3 |
| $1995 / 96$ | 5,589 | 3,119 | 41.1 | 22.4 |
| $1996 / 97$ | 352 | 168 | 2.6 | 1.2 |
| $2005 / 06$ | 19,715 | 1,107 | 144.9 | 7.9 |
| $2006 / 07$ | 24,226 | 4,432 | 178.0 | 31.8 |
| $2007 / 08$ | 61,546 | 3,318 | 200.0 | 23.8 |
| $2008 / 09$ | 29,166 | 646 | 200.0 | 4.6 |
| $2009 / 10$ | 17,289 | 147 | 127.0 | 1.1 |
| $2013 / 14$ | 17,287 | 710 | 127.0 | 5.2 |
| $2014 / 15$ | 85,114 | 1,191 | 200.0 | 8.8 |
| $2015 / 16$ | 119,846 | 1,622 | 200.0 | 11.9 |

Table 7. Sample sizes for total bycatch-at-size in the snow crab fishery, from crab observer sampling. $\mathrm{N}=$ number of individuals. $\mathrm{N}^{`}=$ scaled sample size used in assessment.

| year | males | females | $\mathrm{N}^{\prime}$ |  |
| :---: | :---: | :---: | :---: | :---: |
| 1992/93 | 6,280 | 859 | 46.1 | 6.3 |
| 1993/94 | 6,969 | 1,542 | 51.2 | 11.3 |
| 1994/95 | 2,982 | 1,523 | 21.9 | 11.2 |
| 1995/96 | 1,898 | 428 | 13.9 | 3.1 |
| 1996/97 | 3,265 | 662 | 24.0 | 4.9 |
| 1997/98 | 3,970 | 657 | 29.2 | 4.8 |
| 1998/99 | 1,911 | 324 | 14.0 | 2.4 |
| 1999/00 | 976 | 82 | 7.2 | 0.6 |
| 2000/01 | 1,237 | 74 | 9.1 | 0.5 |
| 2001/02 | 3,113 | 160 | 22.9 | 1.2 |
| 2002/03 | 982 | 118 | 7.2 | 0.9 |
| 2003/04 | 688 | 152 | 5.1 | 1.1 |
| 2004/05 | 848 | 707 | 6.2 | 5.2 |
| 2005/06 | 9,792 | 368 | 72.0 | 2.7 |
| 2006/07 | 10,391 | 1,256 | 76.4 | 9.2 |
| 2007/08 | 13,797 | 728 | 101.4 | 5.3 |
| 2008/09 | 8,455 | 722 | 62.1 | 5.3 |
| 2009/10 | 11,057 | 474 | 81.2 | 3.5 |
| 2010/11 | 12,073 | 250 | 88.7 | 1.8 |
| 2011/12 | 9,453 | 189 | 69.5 | 1.4 |
| 2012/13 | 7,336 | 190 | 53.9 | 1.4 |
| 2013/14 | 12,932 | 356 | 95.0 | 2.6 |
| 2014/15 | 24,877 | 804 | 182.8 | 5.9 |
| 2015/16 | 19,838 | 230 | 145.8 | 1.7 |

Table 8. 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 |
| $1992 / 93$ | 2,056 | 105 | 15.1 | 0.8 |
| $1993 / 94$ | 7,359 | 1,196 | 54.1 | 8.8 |
| $1996 / 97$ | 114 | 5 | 0.8 | 0.0 |
| $1997 / 98$ | 1,030 | 41 | 7.6 | 0.3 |
| $1998 / 99$ | 457 | 20 | 3.4 | 0.1 |
| $1999 / 00$ | 207 | 14 | 1.5 | 0.1 |
| $2000 / 01$ | 845 | 44 | 6.2 | 0.3 |
| $2001 / 02$ | 456 | 39 | 3.4 | 0.3 |
| $2002 / 03$ | 750 | 50 | 5.5 | 0.4 |
| $2003 / 04$ | 555 | 46 | 4.1 | 0.3 |
| $2004 / 05$ | 487 | 44 | 3.6 | 0.3 |
| $2005 / 06$ | 983 | 70 | 7.2 | 0.5 |
| $2006 / 07$ | 798 | 76 | 5.9 | 0.6 |
| $2007 / 08$ | 1,399 | 91 | 10.3 | 0.7 |
| $2008 / 09$ | 3,797 | 121 | 27.9 | 0.9 |
| $2009 / 10$ | 3,395 | 72 | 24.9 | 0.5 |
| $2010 / 11$ | 595 | 30 | 4.4 | 0.2 |
| $2011 / 12$ | 344 | 4 | 2.5 | 0.0 |
| $2012 / 13$ | 618 | 48 | 4.5 | 0.4 |
| $2013 / 14$ | 2,110 | 60 | 15.5 | 0.4 |
| $2014 / 15$ | 3,110 | 32 | 22.9 | 0.2 |
| $2015 / 16$ | 2,176 | 182 | 22.9 | 0.2 |

Table 9. 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 |  | N' |  |
| :---: | :---: | :---: | :---: | :---: |
|  | males | females | males | females |
| 1973/74 | 3,155 | 2,277 | 23.2 | 16.7 |
| 1974/75 | 2,492 | 1,600 | 18.3 | 11.8 |
| 1975/76 | 1,251 | 839 | 9.2 | 6.2 |
| 1976/77 | 6,950 | 6,683 | 51.1 | 49.1 |
| 1977/78 | 10,685 | 8,386 | 78.5 | 61.6 |
| 1978/79 | 18,596 | 13,665 | 136.6 | 100.4 |
| 1979/80 | 19,060 | 11,349 | 140.1 | 83.4 |
| 1980/81 | 12,806 | 5,917 | 94.1 | 43.5 |
| 1981/82 | 6,098 | 4,065 | 44.8 | 29.9 |
| 1982/83 | 13,439 | 8,006 | 98.8 | 58.8 |
| 1983/84 | 18,363 | 8,305 | 134.9 | 61.0 |
| 1984/85 | 27,403 | 13,771 | 200.0 | 101.2 |
| 1985/86 | 23,128 | 12,728 | 170.0 | 93.5 |
| 1986/87 | 14,860 | 7,626 | 109.2 | 56.0 |
| 1987/88 | 23,508 | 15,857 | 172.7 | 116.5 |
| 1988/89 | 10,586 | 7,126 | 77.8 | 52.4 |
| 1989/90 | 59,943 | 41,234 | 200.0 | 200.0 |
| 1990/91 | 23,545 | 11,212 | 173.0 | 82.4 |
| 1991/92 | 6,817 | 3,479 | 50.1 | 25.6 |
| 1992/93 | 3,128 | 1,175 | 23.0 | 8.6 |
| 1993/94 | 1,217 | 358 | 8.9 | 2.6 |
| 1994/95 | 3,628 | 1,820 | 26.7 | 13.4 |
| 1995/96 | 3,904 | 2,669 | 28.7 | 19.6 |
| 1996/97 | 8,306 | 3,400 | 61.0 | 25.0 |
| 1997/98 | 9,949 | 3,900 | 73.1 | 28.7 |
| 1998/99 | 12,105 | 4,440 | 89.0 | 32.6 |
| 1999/00 | 11,053 | 4,522 | 81.2 | 33.2 |
| 2000/01 | 12,895 | 3,087 | 94.8 | 22.7 |
| 2001/02 | 15,788 | 3,083 | 116.0 | 22.7 |
| 2002/03 | 15,401 | 3,249 | 113.2 | 23.9 |
| 2003/04 | 9,572 | 2,733 | 70.3 | 20.1 |
| 2004/05 | 13,844 | 4,460 | 101.7 | 32.8 |
| 2005/06 | 17,785 | 3,709 | 130.7 | 27.3 |
| 2006/07 | 15,903 | 3,047 | 116.9 | 22.4 |
| 2007/08 | 16,031 | 3,788 | 117.8 | 27.8 |
| 2008/09 | 25,976 | 4,164 | 190.9 | 30.6 |
| 2009/10 | 18,852 | 2,650 | 138.5 | 19.5 |
| 2010/11 | 15,044 | 2,247 | 110.5 | 16.5 |
| 2011/12 | 16,115 | 4,237 | 118.4 | 31.1 |
| 2012/13 | 12,983 | 3,080 | 95.4 | 22.6 |
| 2013/14 | 28,781 | 6,064 | 200.0 | 44.6 |
| 2014/15 | 39,119 | 4,212 | 200.0 | 31.0 |
| 2015/16 | 26,656 | 5,705 | 195.9 | 41.9 |

Table 10. Trends in mature and total Tanner crab biomass (1000's t) in the NMFS summer bottom trawl survey.

| Year | Mature Biomass (1000 t) |  |  | Legalmales$\left(10^{6} \mathrm{crab}\right)$ |
| :---: | :---: | :---: | :---: | :---: |
|  | Male | Female | Total |  |
| 1974 | -- | -- | -- | -- |
| 1975 | 252.38 | 28.28 | 280.66 | 278.67 |
| 1976 | 127.66 | 27.02 | 154.67 | 144.48 |
| 1977 | 110.46 | 31.51 | 141.97 | 119.76 |
| 1978 | 75.30 | 20.43 | 95.73 | 83.39 |
| 1979 | 31.30 | 11.93 | 43.22 | 38.51 |
| 1980 | 79.58 | 33.79 | 113.37 | 92.05 |
| 1981 | 45.50 | 21.74 | 67.24 | 53.33 |
| 1982 | 45.60 | 29.82 | 75.42 | 58.70 |
| 1983 | 26.99 | 13.25 | 40.24 | 36.15 |
| 1984 | 22.12 | 11.10 | 33.23 | 29.07 |
| 1985 | 10.64 | 4.40 | 15.04 | 13.07 |
| 1986 | 10.80 | 3.36 | 14.16 | 11.53 |
| 1987 | 19.69 | 7.87 | 27.56 | 24.65 |
| 1988 | 53.48 | 22.89 | 76.37 | 58.41 |
| 1989 | 89.26 | 15.96 | 105.22 | 104.71 |
| 1990 | 92.45 | 28.18 | 120.63 | 110.05 |
| 1991 | 101.95 | 31.74 | 133.70 | 125.66 |
| 1992 | 100.79 | 19.22 | 120.01 | 123.66 |
| 1993 | 57.99 | 8.21 | 66.20 | 72.61 |
| 1994 | 40.05 | 7.09 | 47.13 | 49.92 |
| 1995 | 29.44 | 8.71 | 38.16 | 39.23 |
| 1996 | 24.41 | 6.76 | 31.17 | 31.43 |
| 1997 | 9.36 | 2.38 | 11.74 | 11.55 |
| 1998 | 8.79 | 1.68 | 10.47 | 10.45 |
| 1999 | 8.68 | 2.81 | 11.49 | 9.30 |
| 2000 | 13.92 | 3.14 | 17.05 | 15.85 |
| 2001 | 15.37 | 3.29 | 18.66 | 18.53 |
| 2002 | 14.36 | 2.63 | 16.99 | 16.45 |
| 2003 | 19.02 | 4.18 | 23.19 | 22.84 |
| 2004 | 22.42 | 2.86 | 25.27 | 28.63 |
| 2005 | 39.47 | 7.21 | 46.67 | 52.70 |
| 2006 | 52.55 | 10.22 | 62.77 | 69.40 |
| 2007 | 56.34 | 9.47 | 65.81 | 71.33 |
| 2008 | 58.78 | 7.91 | 66.69 | 74.83 |
| 2009 | 33.92 | 5.64 | 39.55 | 45.56 |
| 2010 | 37.05 | 4.02 | 41.07 | 49.39 |
| 2011 | 37.65 | 4.37 | 42.02 | 47.16 |
| 2012 | 29.51 | 6.75 | 36.26 | 34.34 |
| 2013 | 59.58 | 10.93 | 70.51 | 63.99 |
| 2014 | 73.33 | 9.04 | 82.37 | 85.74 |
| 2015 | 58.36 | 6.13 | 64.49 | 76.70 |
| 2016 | 53.64 | 4.24 | 57.88 | 71.58 |

Table 11. Sample sizes for NMFS survey size composition data. In the assessment model, an effective sample size of 200 is used for all survey-related compositional data.

| year | number of hauls | females |  |  |  |  |  | males |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | immature new shell |  | mature |  |  |  | immature |  | mature |  |  |  |
|  |  | number of | number of | number of | number of | number of | number of | number of | number of | number of | number of | number of | number of |
| 1975 | 136 | 73 | 1,040 | 91 | 1,861 | 39 | 706 | 127 | 2,895 | 127 | 3,993 | 80 | 399 |
| 1976 | 214 | 87 | 1,095 | 91 | 1,304 | 39 | 311 | 130 | 2,023 | 130 | 2,469 | 47 | 242 |
| 1977 | 155 | 66 | 765 | 76 | 1,183 | 60 | 738 | 114 | 1,778 | 114 | 1,971 | 79 | 485 |
| 1978 | 230 | 87 | 1,932 | 82 | 638 | 65 | 1,307 | 147 | 2,957 | 147 | 1,570 | 104 | 700 |
| 1979 | 307 | 71 | 725 | 62 | 735 | 42 | 341 | 138 | 1,805 | 138 | 808 | 68 | 306 |
| 1980 | 320 | 101 | 1,476 | 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 | 85 | 814 | 72 | 457 | 103 | 2,384 | 181 | 1,751 | 181 | 1,371 | 147 | 2,082 |
| 1983 | 353 | 102 | 2,108 | 56 | 201 | 102 | 2,154 | 166 | 2,484 | 166 | 983 | 132 | 1,181 |
| 1984 | 355 | 135 | 1,867 | 53 | 284 | 94 | 1,531 | 171 | 1,965 | 171 | 490 | 126 | 1,399 |
| 1985 | 353 | 140 | 846 | 52 | 228 | 65 | 601 | 179 | 1,060 | 179 | 381 | 86 | 459 |
| 1986 | 353 | 162 | 1,581 | 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 | 3,733 | 149 | 1,753 | 100 | 530 | 252 | 5,627 | 252 | 2,210 | 101 | 475 |
| 1989 | 373 | 204 | 3,264 | 144 | 1,241 | 108 | 882 | 237 | 4,977 | 237 | 3,201 | 135 | 1,067 |
| 1990 | 370 | 197 | 3,105 | 155 | 1,502 | 126 | 1,511 | 247 | 5,107 | 247 | 3,149 | 151 | 1,342 |
| 1991 | 371 | 159 | 2,227 | 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 | 865 | 96 | 545 | 122 | 1,337 | 207 | 2,051 | 207 | 1,677 | 180 | 1,865 |
| 1994 | 374 | 97 | 909 | 52 | 148 | 104 | 1,293 | 175 | 1,281 | 175 | 724 | 174 | 1,827 |
| 1995 | 375 | 113 | 830 | 35 | 140 | 107 | 1,057 | 153 | 958 | 153 | 220 | 137 | 1,611 |
| 1996 | 374 | 114 | 869 | 57 | 109 | 98 | 963 | 148 | 1,069 | 148 | 222 | 134 | 1,414 |
| 1997 | 375 | 116 | 1,325 | 62 | 168 | 83 | 504 | 161 | 1,336 | 161 | 289 | 125 | 582 |
| 1998 | 374 | 146 | 1,704 | 53 | 160 | 73 | 344 | 176 | 2,032 | 176 | 396 | 128 | 624 |
| 1999 | 372 | 137 | 2,608 | 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 |
| 2001 | 374 | 164 | 3,675 | 83 | 364 | 72 | 644 | 211 | 4,036 | 211 | 629 | 145 | 817 |
| 2002 | 374 | 154 | 3,583 | 81 | 350 | 70 | 500 | 186 | 3,912 | 186 | 458 | 154 | 1,089 |
| 2003 | 375 | 153 | 2,830 | 111 | 923 | 83 | 752 | 203 | 4,754 | 203 | 900 | 153 | 1,349 |
| 2004 | 374 | 173 | 3,563 | 90 | 427 | 80 | 656 | 236 | 4,568 | 236 | 1,027 | 179 | 1,873 |
| 2005 | 372 | 201 | 3,349 | 103 | 634 | 74 | 928 | 254 | 4,496 | 254 | 1,280 | 185 | 1,753 |
| 2006 | 375 | 210 | 4,355 | 143 | 1,332 | 125 | 1,327 | 254 | 6,224 | 254 | 1,757 | 211 | 4,054 |
| 2007 | 375 | 185 | 2,420 | 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 | 3,171 | 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 | 3,577 | 100 | 942 | 97 | 720 | 215 | 5,526 | 215 | 1,564 | 139 | 1,296 |
| 2013 | 375 | 163 | 2,900 | 116 | 1,417 | 101 | 1,002 | 207 | 5,592 | 207 | 2,675 | 137 | 1,344 |
| 2014 | 375 | 165 | 2,207 | 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 | 1,372 | 56 | 370 | 82 | 1,248 | 222 | 2,235 | 222 | 1,170 | 218 | 3,668 |

Table 12. Effort data (1000's potlifts) in the snow crab and BBRKC fisheries.

| Effort (1000's Potlifts) |  |  | Effort (1000's Potlifts) |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Year | BBRKC Fishery | Snow Crab Fishery | Year | BBRKC Fishery | Snow Crab Fishery |
| 1951/52 |  |  | 1986/87 | 175.753 | 616.113 |
| 1952/53 |  |  | 1987/88 | 220.971 | 747.395 |
| 1953/54 | 30.083 | -- | 1988/89 | 146.179 | 665.242 |
| 1954/55 | 17.122 | -- | 1989/90 | 205.528 | 912.718 |
| 1955/56 | 28.045 | -- | 1990/91 | 262.761 | 1382.908 |
| 1956/57 | 41.629 | -- | 1991/92 | 227.555 | 1278.502 |
| 1957/58 | 23.659 | -- | 1992/93 | 206.815 | 969.209 |
| 1958/59 | 27.932 | -- | 1993/94 | 254.389 | 716.524 |
| 1959/60 | 22.187 | -- | 1994/95 | 0.697 | 507.603 |
| 1960/61 | 26.347 | -- | 1995/96 | 0.547 | 520.685 |
| 1961/62 | 72.646 | -- | 1996/97 | 77.081 | 754.14 |
| 1962/63 | 123.643 | -- | 1997/98 | 91.085 | 930.794 |
| 1963/64 | 181.799 | -- | 1998/99 | 145.689 | 945.533 |
| 1964/65 | 180.809 | -- | 1999/00 | 151.212 | 182.634 |
| 1965/66 | 127.973 | -- | 2000/01 | 104.056 | 191.2 |
| 1966/67 | 129.306 | -- | 2001/02 | 66.947 | 326.977 |
| 1967/68 | 135.283 | -- | 2002/03 | 72.514 | 153.862 |
| 1968/69 | 184.666 | -- | 2003/04 | 134.515 | 123.709 |
| 1969/70 | 175.374 | -- | 2004/05 | 97.621 | 75.095 |
| 1970/71 | 168.059 | -- | 2005/06 | 116.32 | 117.375 |
| 1971/72 | 126.305 | -- | 2006/07 | 72.404 | 86.288 |
| 1972/73 | 208.469 | -- | 2007/08 | 113.948 | 140.857 |
| 1973/74 | 194.095 | -- | 2008/09 | 139.937 | 163.537 |
| 1974/75 | 212.915 | -- | 2009/10 | 118.521 | 136.477 |
| 1975/76 | 205.096 | -- | 2010/11 | 131.627 | 147.244 |
| 1976/77 | 321.01 | -- | 2011/12 | 45.166 | 270.602 |
| 1977/78 | 451.273 | -- | 2012/13 | 38.159 | 225.489 |
| 1978/79 | 406.165 | 190.746 | 2013/14 | 45.927 | 225.245 |
| 1979/80 | 315.226 | 255.102 | 2014/15 | 57.725 | 279.183 |
| 1980/81 | 567.292 | 435.742 | 2015/16 | 48.665 | 201.65 |
| 1981/82 | 536.646 | 469.091 |  |  |  |
| 1982/83 | 140.492 | 287.127 |  |  |  |
| 1983/84 | 0 | 173.591 |  |  |  |
| 1984/85 | 107.406 | 370.082 |  |  |  |
| 1985/86 | 84.443 | 542.346 |  |  |  |

Table 13. Effective sample sizes used for NMFS EBS trawl survey size composition data for the 2015 assessment model (2015AMO) and the author's preferred model (Model C). Effective sample sizes were estimated using the McAllister-Ianelli approach.

| year | 2015AMO input effective |  | $\begin{aligned} & \text { Mod } \\ & \text { input } \end{aligned}$ | el C effective |
| :---: | :---: | :---: | :---: | :---: |
| 1975 | 200 | 104 | 200 | 106 |
| 1976 | 200 | 167 | 200 | 175 |
| 1977 | 200 | 138 | 200 | 149 |
| 1978 | 200 | 175 | 200 | 167 |
| 1979 | 200 | 244 | 200 | 236 |
| 1980 | 200 | 132 | 200 | 142 |
| 1981 | 200 | 102 | 200 | 101 |
| 1982 | 200 | 30 | 200 | 26 |
| 1983 | 200 | 266 | 200 | 231 |
| 1984 | 200 | 134 | 200 | 162 |
| 1985 | 200 | 46 | 200 | 90 |
| 1986 | 200 | 106 | 200 | 175 |
| 1987 | 200 | 84 | 200 | 89 |
| 1988 | 200 | 214 | 200 | 220 |
| 1989 | 200 | 234 | 200 | 279 |
| 1990 | 200 | 518 | 200 | 548 |
| 1991 | 200 | 422 | 200 | 437 |
| 1992 | 200 | 491 | 200 | 629 |
| 1993 | 200 | 187 | 200 | 252 |
| 1994 | 200 | 161 | 200 | 208 |
| 1995 | 200 | 554 | 200 | 404 |
| 1996 | 200 | 521 | 200 | 448 |
| 1997 | 200 | 184 | 200 | 217 |
| 1998 | 200 | 212 | 200 | 251 |
| 1999 | 200 | 149 | 200 | 156 |
| 2000 | 200 | 247 | 200 | 251 |
| 2001 | 200 | 305 | 200 | 283 |
| 2002 | 200 | 179 | 200 | 169 |
| 2003 | 200 | 421 | 200 | 403 |
| 2004 | 200 | 269 | 200 | 304 |
| 2005 | 200 | 377 | 200 | 411 |
| 2006 | 200 | 278 | 200 | 300 |
| 2007 | 200 | 222 | 200 | 245 |
| 2008 | 200 | 346 | 200 | 406 |
| 2009 | 200 | 171 | 200 | 149 |
| 2010 | 200 | 279 | 200 | 224 |
| 2011 | 200 | 345 | 200 | 330 |
| 2012 | 200 | 279 | 200 | 280 |
| 2013 | 200 | 484 | 200 | 529 |
| 2014 | 200 | 296 | 200 | 300 |
| 2015 | 200 | 440 | 200 | 543 |
| 2016 |  |  | 200 | 268 |

Table 14. Effective sample sizes used for retained catch size composition data from the directed fishery for the 2015 assessment model (2015AMO) and the author's preferred model (Model C). Effective sample sizes were estimated using the McAllister-Ianelli approach.

| year | 2015AMO <br> input |  | effective | Model C <br> input <br> effective |  |
| ---: | ---: | ---: | ---: | ---: | :---: |
| 1980 | 97.8 | 22.8 | 97.8 | 20.2 |  |
| 1981 | 83.1 | 548.4 | 83.1 | 805.1 |  |
| 1982 | 99.3 | 1143.2 | 99.3 | 1622.3 |  |
| 1983 | 12.3 | 43.4 | 12.3 | 50.3 |  |
| 1984 | 18.7 | 560.6 | 18.7 | 342.1 |  |
| 1988 | 91.0 | 111.7 | 91.0 | 141.1 |  |
| 1989 | 30.3 | 1078.7 | 30.3 | 1042.2 |  |
| 1990 | 200.0 | 415.6 | 200.0 | 263.6 |  |
| 1991 | 200.0 | 47.1 | 200.0 | 20.7 |  |
| 1992 | 200.0 | 37.8 | 200.0 | 17.8 |  |
| 1993 | 200.0 | 48.2 | 200.0 | 23.2 |  |
| 1994 | 200.0 | 82.9 | 200.0 | 47.8 |  |
| 1995 | 11.2 | 32.4 | 11.2 | 15.5 |  |
| 1996 | 32.6 | 16.1 | 32.6 | 12.6 |  |
| 2005 | 5.2 | 7.3 | 5.2 | 6.6 |  |
| 2006 | 21.6 | 18.6 | 21.6 | 15.0 |  |
| 2007 | 51.0 | 21.5 | 51.0 | 17.0 |  |
| 2008 | 25.6 | 38.8 | 25.6 | 19.3 |  |
| 2009 | 17.8 | 158.4 | 17.8 | 70.6 |  |
| 2013 | 35.0 | 50.7 | 35.0 | 141.1 |  |
| 2014 | 103.3 | 19.5 | 103.3 | 34.5 |  |
| 2015 |  |  | 200.0 | 39.3 |  |
|  |  |  |  |  |  |

Table 15. Effective sample sizes used for total catch size composition data from the directed fishery for the 2015 assessment model (2015AMO) and the author's preferred model (Model C). Effective sample sizes were estimated using the McAllister-Ianelli approach.

| year | 2015AMO |  |  |  | Model C |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | female input effective |  | male input effective |  | female input effective |  | male input effective |  |
| 1991 | 41.2 | 218.3 | 200.0 | 11.4 | 41.2 | 322.9 | 200.0 | 12.0 |
| 1992 | 64.3 | 264.9 | 200.0 | 11.2 | 64.3 | 940.8 | 200.0 | 13.3 |
| 1993 | 76.9 | 904.9 | 200.0 | 12.3 | 76.9 | 296.2 | 200.0 | 12.9 |
| 1994 | 15.7 | 73.3 | 42.6 | 12.1 | 15.7 | 78.7 | 42.6 | 10.9 |
| 1995 | 22.9 | 71.5 | 41.1 | 60.8 | 22.9 | 152.1 | 41.1 | 80.8 |
| 1996 | 2.5 | 111.7 | 5.0 | 29.4 | 2.5 | 149.0 | 5.0 | 37.2 |
| 2005 | 8.1 | 18.6 | 144.9 | 8.0 | 8.1 | 34.3 | 144.9 | 7.8 |
| 2006 | 32.6 | 101.0 | 178.0 | 92.9 | 32.6 | 279.0 | 178.0 | 65.0 |
| 2007 | 24.4 | 61.2 | 200.0 | 13.2 | 24.4 | 310.7 | 200.0 | 10.2 |
| 2008 | 4.7 | 19.9 | 200.0 | 13.4 | 4.7 | 41.7 | 200.0 | 13.8 |
| 2009 | 1.1 | 51.7 | 127.0 | 11.0 | 1.1 | 28.2 | 127.0 | 10.9 |
| 2013 | 5.2 | 94.8 | 127.0 | 16.8 | 5.2 | 82.1 | 127.0 | 15.7 |
| 2014 | 8.8 | 121.1 | 200.0 | 8.8 | 8.8 | 208.1 | 200.0 | 7.6 |
| 2015 |  |  |  |  | 11.9 | 69.6 | 200.0 | 6.1 |

Table 16. Effective sample sizes used for bycatch size composition data from the snow crab fishery for the 2015 assessment model (2015AMO) and the author's preferred model (Model C). Effective sample sizes were estimated using the McAllister-Ianelli approach.

| year | 2015AMO |  |  |  | Model C |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | female input effective |  | male input effective |  | female input effective |  | male input effective |  |
| 1992 | 6.3 | 25.7 | 46.1 | 229.2 | 6.3 | 16.5 | 46.1 | 185.3 |
| 1993 | 11.3 | 32.5 | 51.2 | 168.9 | 11.3 | 27.4 | 51.2 | 170.8 |
| 1994 | 11.2 | 26.4 | 21.9 | 49.6 | 11.2 | 49.6 | 21.9 | 42.6 |
| 1995 | 3.1 | 29.9 | 13.9 | 128.7 | 3.1 | 38.1 | 13.9 | 122.2 |
| 1996 | 4.9 | 54.7 | 24.0 | 236.8 | 4.9 | 36.2 | 24.0 | 290.7 |
| 1997 | 4.8 | 178.6 | 29.2 | 347.3 | 4.8 | 134.6 | 29.2 | 345.9 |
| 1998 | 2.4 | 21.9 | 14.0 | 475.7 | 2.4 | 19.5 | 14.0 | 617.1 |
| 1999 | 0.6 | 30.2 | 7.2 | 118.9 | 0.6 | 27.6 | 7.2 | 134.1 |
| 2000 | 0.5 | 31.7 | 9.1 | 205.0 | 0.5 | 29.9 | 9.1 | 224.8 |
| 2001 | 1.2 | 147.4 | 22.9 | 1089.6 | 1.2 | 139.0 | 22.9 | 1123.1 |
| 2002 | 0.9 | 51.3 | 7.2 | 66.0 | 0.9 | 45.2 | 7.2 | 61.9 |
| 2003 | 1.1 | 47.6 | 5.1 | 112.1 | 1.1 | 43.8 | 5.1 | 102.8 |
| 2004 | 5.2 | 34.0 | 6.2 | 25.9 | 5.2 | 30.1 | 6.2 | 24.5 |
| 2005 | 2.7 | 167.9 | 72.0 | 145.8 | 2.7 | 95.1 | 72.0 | 127.4 |
| 2006 | 9.2 | 57.9 | 76.4 | 94.4 | 9.2 | 33.6 | 76.4 | 86.8 |
| 2007 | 5.3 | 49.7 | 101.4 | 645.0 | 5.3 | 28.8 | 101.4 | 455.6 |
| 2008 | 5.3 | 13.7 | 62.1 | 99.6 | 5.3 | 18.4 | 62.1 | 92.9 |
| 2009 | 3.5 | 19.4 | 81.2 | 404.4 | 3.5 | 31.0 | 81.2 | 430.0 |
| 2010 | 1.8 | 72.9 | 88.7 | 260.6 | 1.8 | 87.0 | 88.7 | 339.6 |
| 2011 | 1.4 | 58.2 | 69.5 | 156.6 | 1.4 | 53.7 | 69.5 | 186.9 |
| 2012 | 1.4 | 45.3 | 53.9 | 120.5 | 1.4 | 49.1 | 53.9 | 139.7 |
| 2013 | 2.6 | 274.0 | 95.0 | 192.8 | 2.6 | 128.8 | 95.0 | 222.5 |
| 2014 | 5.9 | 52.3 | 182.8 | 477.6 | 5.9 | 118.9 | 182.8 | 525.0 |
| 2015 |  |  |  |  | 1.7 | 61.8 | 145.8 | 475.2 |

Table 17. Effective sample sizes used for bycatch size composition data from the BBRKC fishery for the 2015 assessment model (2015AMO) and the author's preferred model (Model C). Effective sample sizes were estimated using the McAllister-Ianelli approach.

| year | 2015AMO |  |  |  | Model C |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | female input effective |  | male input effective |  | female input effective |  | male input effective |  |
|  |  |  |  |  |  |  |  |  |
| 1992 | 0.8 | 37.7 | 15.1 | 181.6 | 0.8 | 47.2 | 15.1 | 154.7 |
| 1993 | 8.8 | 123.4 | 54.1 | 405.8 | 8.8 | 326.2 | 54.1 | 432.7 |
| 1996 | 0.0 | 4.0 | 0.8 | 66.0 | 0.0 | 3.8 | 0.8 | 60.8 |
| 1997 | 0.3 | 16.3 | 7.6 | 26.5 | 0.3 | 17.3 | 7.6 | 24.7 |
| 1998 | 0.1 | 18.4 | 3.4 | 70.2 | 0.1 | 19.3 | 3.4 | 67.2 |
| 1999 | 0.1 | 16.1 | 1.5 | 64.1 | 0.1 | 16.6 | 1.5 | 63.0 |
| 2000 | 0.3 | 38.9 | 6.2 | 212.0 | 0.3 | 37.0 | 6.2 | 190.0 |
| 2001 | 0.3 | 53.2 | 3.4 | 139.3 | 0.3 | 46.9 | 3.4 | 131.0 |
| 2002 | 0.4 | 36.0 | 5.5 | 130.5 | 0.4 | 45.9 | 5.5 | 110.4 |
| 2003 | 0.3 | 53.1 | 4.1 | 88.2 | 0.3 | 49.0 | 4.1 | 76.5 |
| 2004 | 0.3 | 20.1 | 3.6 | 49.9 | 0.3 | 22.2 | 3.6 | 41.5 |
| 2005 | 0.5 | 7.3 | 7.2 | 36.9 | 0.5 | 8.2 | 7.2 | 38.4 |
| 2006 | 0.6 | 17.7 | 5.9 | 19.3 | 0.6 | 19.7 | 5.9 | 20.1 |
| 2007 | 0.7 | 53.7 | 10.3 | 68.7 | 0.7 | 64.9 | 10.3 | 79.0 |
| 2008 | 0.9 | 48.7 | 27.9 | 100.2 | 0.9 | 55.9 | 27.9 | 79.8 |
| 2009 | 0.5 | 110.7 | 24.9 | 23.7 | 0.5 | 119.6 | 24.9 | 21.6 |
| 2010 | 0.2 | 28.9 | 4.4 | 48.9 | 0.2 | 29.0 | 4.4 | 49.8 |
| 2011 | 0.0 | 6.7 | 2.5 | 62.2 | 0.0 | 6.4 | 2.5 | 63.8 |
| 2012 | 0.4 | 9.9 | 4.5 | 61.4 | 0.4 | 9.3 | 4.5 | 65.1 |
| 2013 | 0.4 | 16.0 | 15.5 | 84.2 | 0.4 | 14.3 | 15.5 | 83.7 |
| 2014 | 0.2 | 22.1 | 22.9 | 126.3 | 0.2 | 23.2 | 22.9 | 139.6 |
| 2015 |  |  |  |  | 0.2 | 66.4 | 22.9 | 163.2 |

Table 18. Effective sample sizes used for bycatch size composition data from the groundfish fisheries for the 2015 assessment model (2015AMO) and the author's preferred model (Model C). Effective sample sizes were estimated using the McAllister-Ianelli approach.

| year | 2015AMO |  | Model C |  |
| :---: | :---: | :---: | :---: | :---: |
| 1973 | 39.9 | 95.5 | 39.9 | 284.9 |
| 1974 | 30.1 | 172.4 | 30.1 | 396.0 |
| 1975 | 15.4 | 119.2 | 15.4 | 250.0 |
| 1976 | 100.2 | 63.9 | 100.2 | 133.6 |
| 1977 | 140.1 | 96.6 | 140.1 | 229.7 |
| 1978 | 237.1 | 100.5 | 237.1 | 208.7 |
| 1979 | 223.5 | 143.2 | 223.5 | 567.2 |
| 1980 | 137.6 | 249.3 | 137.6 | 621.7 |
| 1981 | 74.7 | 112.1 | 74.7 | 135.8 |
| 1982 | 157.6 | 102.0 | 157.6 | 128.5 |
| 1983 | 196.0 | 199.3 | 196.0 | 219.3 |
| 1984 | 301.2 | 202.2 | 301.2 | 311.2 |
| 1985 | 263.5 | 117.1 | 263.5 | 224.6 |
| 1986 | 165.2 | 105.1 | 165.2 | 224.0 |
| 1987 | 289.3 | 158.0 | 289.3 | 437.4 |
| 1988 | 130.2 | 171.4 | 130.2 | 295.9 |
| 1989 | 400.0 | 272.5 | 400.0 | 910.5 |
| 1990 | 255.4 | 413.1 | 255.4 | 625.1 |
| 1991 | 75.7 | 364.3 | 75.7 | 629.3 |
| 1992 | 31.6 | 148.3 | 31.6 | 113.2 |
| 1993 | 11.6 | 75.4 | 11.6 | 54.7 |
| 1994 | 40.0 | 82.0 | 40.0 | 69.9 |
| 1995 | 48.3 | 51.8 | 48.3 | 60.4 |
| 1996 | 86.0 | 399.0 | 86.0 | 288.0 |
| 1997 | 101.8 | 44.8 | 101.8 | 74.1 |
| 1998 | 121.6 | 95.5 | 121.6 | 246.1 |
| 1999 | 114.4 | 115.0 | 114.4 | 599.4 |
| 2000 | 117.4 | 179.0 | 117.4 | 392.0 |
| 2001 | 138.7 | 174.8 | 138.7 | 230.4 |
| 2002 | 137.0 | 88.0 | 137.0 | 122.2 |
| 2003 | 90.4 | 155.0 | 90.4 | 505.7 |
| 2004 | 134.5 | 140.6 | 134.5 | 369.3 |
| 2005 | 157.9 | 395.8 | 157.9 | 1101.6 |
| 2006 | 139.2 | 172.7 | 139.2 | 212.4 |
| 2007 | 145.6 | 223.1 | 145.6 | 596.1 |
| 2008 | 221.5 | 350.2 | 221.5 | 437.0 |
| 2009 | 156.9 | 143.0 | 158.0 | 400.9 |
| 2010 | 127.5 | 230.0 | 127.1 | 965.0 |
| 2011 | 150.1 | 79.2 | 149.6 | 60.9 |
| 2012 | 118.6 | 75.4 | 118.0 | 192.3 |
| 2013 | 244.7 | 101.0 | 244.6 | 373.6 |
| 2014 | 230.1 | 151.2 | 231.0 | 2083.9 |
| 2015 |  |  | 237.8 | 291.7 |

Table 19. Objective function components and associated applied weighting factors for the 2015 assessment model and the author's preferred model (Model C). TCF: directed Tanner crab fishery; SCF: snow crab fishery; RKF: BBRKC fishery; GTF: groundfish fisheries.

| category | description | weight | 2015AMO | Model C |
| :---: | :---: | :---: | :---: | :---: |
| likelihood: catch biomass | fishery: GTF total catch biomass | 10.0 | 2.52 | 2.43 |
| likelihood: catch biomass | fishery: RKF total catch biomass | 10.0 | 9.59 | 12.81 |
| likelihood: catch biomass | fishery: SCF total catch biomass | 10.0 | 10.52 | 6.21 |
| likelihood: catch biomass | fishery: TCF female catch biomass | 10.0 | 6.64 | 5.11 |
| likelihood: catch biomass | fishery: TCF male total catch biomass | 10.0 | 18.21 | 11.54 |
| likelihood: catch biomass | fishery: TCF retained males | 10.0 | 31.87 | 18.47 |
| likelihood: catch biomass | survey: mature crab | 1.0 | 311.35 | 199.10 |
| likelihood: size comps | fishery: GTF males+females | 1.0 | 135.17 | 463.33 |
| likelihood: size comps | fishery: RKC females | 1.0 | 2.68 | 2.25 |
| likelihood: size comps | fishery: RKC males | 1.0 | 24.21 | 26.69 |
| likelihood: size comps | fishery: SCF females | 1.0 | 13.95 | 12.49 |
| likelihood: size comps | fishery: SCF males | 1.0 | 49.26 | 52.63 |
| likelihood: size comps | fishery: TCF discarded females | 1.0 | 14.32 | 9.70 |
| likelihood: size comps | fishery: TCF retained males | 1.0 | 194.52 | 308.98 |
| likelihood: size comps | fishery: TCF total males | 1.0 | 115.60 | 184.30 |
| likelihood: size comps | survey: immature females | 1.0 | 307.31 | 281.23 |
| likelihood: size comps | survey: immature males | 1.0 | 280.47 | 269.49 |
| likelihood: size comps | survey: mature females | 1.0 | 99.13 | 128.52 |
| likelihood: size comps | survey: mature males | 1.0 | 272.48 | 250.07 |
| penalty | maturity curve smoothness (females) | 1.0 | 1.41 | 2.33 |
| penalty | maturity curve smoothness (males) | 0.5 | 0.16 | 0.79 |
| penalty | natural mortality penalty (immature females) | 1.0 | 51.27 | 36.42 |
| penalty | natural mortality penalty (immatures) | 1.0 | 0.64 | 0.59 |
| penalty | natural mortality penalty (mature males) | 1.0 | 4.21 | 5.62 |
| penalty | penalty on F-devs in BBRKC fishery | 3.0 | 0.00 | 0.13 |
| penalty | penalty on F-devs in directed fishery | 1.0 | 49.39 | 56.77 |
| penalty | penalty on F-devs in groundfish fishery | 0.5 | 11.69 | 12.98 |
| penalty | penalty on F-devs in snow crab fishery | 0.5 | 7.70 | 7.47 |
| penalty | recruitment penalty | 1.0 | 2.30 | 2.44 |
| penalty | sex ratio penalty | 0.0 | 0.00 | 0.00 |
| penalty | z50 devs for male selectivity in TCF (AR1) | 0.0 | 0.00 | 0.00 |
| penalty | z50 devs for male selectivity in TCF (norm2) | 0.0 | 0.00 | 0.00 |
| priors | female growth parameter a | 1.0 | 0.90 | 0.90 |
| priors | female growth parameter b | 1.0 | 0.68 | 0.64 |
| priors | female survey q penalty | 1.0 | 16.35 | 29.11 |
| priors | male growth parameter a | 1.0 | 0.57 | 0.23 |
| priors | male growth parameter b | 1.0 | 0.04 | 0.03 |
| priors | survey q penalty | 1.0 | 1.97 | 4.97 |

Table 20. Comparison of parameter estimates from the 2015 assessment model and the author's preferred model (Model C).

| process | description | param | index | 2015AMO | Model C |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | estimate | estimate | std. dev. |
| growth | female mean growth a parameter | pGrAF1 |  | 0.7 | 0.7 | 6.98E-05 |
|  | female mean growth b parameter | pGrBF1 |  | 0.884217 | 0.885004 | 0.0011352 |
|  | male mean growth a parameter | pGrAM1 |  | 0.411176 | 0.420826 | 0.021848 |
|  | male mean growth b parameter | pGrBM1 |  | 0.976754 | 0.972702 | 0.0051716 |
|  | size transition beta parameter | pGrBeta_x | female | 0.750005 | 0.750005 | 0 |
|  | size transition beta parameter | pGrBeta_x | male | 0.750005 | 0.750005 | 0 |
| natural mortality multipliers | multiplier for 1980-1984 | pMfac_Big | female | 1.4936 | 1.32933 | 0.10943 |
|  | multiplier for 1980-1984 | pMfac_Big | male | 3.50292 | 2.82341 | 0.33557 |
|  | multiplier for immature crab | pMfac_Imm |  | 1.05671 | 1.05437 | 0.049567 |
|  | multiplier for mature female crab | pMfac_MatF |  | 1.50633 | 1.4267 | 0.036859 |
|  | multiplier for mature male crab | pMfac_MatM |  | 1.14505 | 1.1676 | 0.041043 |
| recruitment | initial log-scale mean | pMnLnReclnit |  | 5.58529 | 5.52749 | 0.49162 |
|  | log-scale mean | pMnLnRec |  | 4.92158 | 5.00006 | 0.066058 |
|  | size distribution alpha parameter | pRecAlpha |  | 11.5 | 11.5 | 0 |
|  | size distribution beta parameter | pRecBeta |  | 4 | 4 | 0 |
| survey selectivity | male offset to 95\%-selected [-1981] | pSrv1M_dz5095 |  | 21.5698 | 22.1348 | 3.2621 |
|  | male offset to 95\%-selected [1982+] | pSrv2M_dz5095 |  | 55.6208 | 62.917 | 8.2923 |
|  | male size at 50\%-selected [-1981] | pSrv1M_z50 |  | 49.0101 | 50.2176 | 1.9188 |
|  | male size at 50\%-selected [1982+] | pSrv2M_z50 |  | 32.4911 | 32.0113 | 3.2009 |
|  | female offset to 95\%-selected [-1981] | pSrv1F_dz5095 |  | 40.8236 | 38.3361 | 6.1379 |
|  | female offset to 95\%-selected [1982+] | pSrv2F_dz5095 |  | 100 | 100 | 0.0011952 |
|  | female size at 50\%-selected [-1981] | pSrv1F_z50 |  | 53.6264 | 54.1952 | 2.7904 |
|  | female size at 50\%-selected [1982+] | pSrv2F_z50 |  | 7.10091 | -9.24299 | 15.073 |
| survey Q | females [-1981] | pSrv1_QF |  | 0.5 | 0.5 | 4.94E-05 |
|  | females [1982+] | pSrv2_QF |  | 0.594041 | 0.498521 | 0.032247 |
|  | males [-1981] | pSrv1_QM |  | 0.5 | 0.5 | 1.95E-05 |
|  | males [1982+] | pSrv2_QM |  | 0.780778 | 0.722284 | 0.036416 |

Table 21. Comparison of molt-to-maturity parameter estimates from the 2015 assessment model (ln-scale) and the author's preferred model (Model C; logit-scale).

| process | sex | index | 2015AMO <br> estimate | Model C |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | estimate | std. dev. |
| molt-to-maturity | female | 1 | -15 | -15 | 0.001669 |
|  |  | 2 | -13.7474 | -13.7599 | 0.78396 |
|  |  | 3 | -12.4437 | -12.4653 | 1.1857 |
|  |  | 4 | -11.0381 | -11.0616 | 1.288 |
|  |  | 5 | -9.47992 | -9.49471 | 1.1517 |
|  |  | 6 | -7.72241 | -7.71458 | 0.86232 |
|  |  | 7 | -5.74099 | -5.69543 | 0.52458 |
|  |  | 8 | -3.60849 | -3.5189 | 0.24124 |
|  |  | 9 | -1.84318 | -1.68486 | 0.11369 |
|  |  | 10 | -0.816855 | -0.323703 | 0.092391 |
|  |  | 11 | -0.49044 | 0.351804 | 0.097912 |
|  |  | 12 | -0.364766 | 0.624612 | 0.11199 |
|  |  | 13 | -0.116204 | 1.56765 | 0.20163 |
|  |  | 14 | -1.62E-09 | 3.35975 | 0.43493 |
|  |  | 15 | -0.004397 | 5.29665 | 0.91207 |
|  |  | 16 | -7.31E-09 | 7.25082 | 1.6735 |
| molt-to-maturity | male | 1 | -12.5966 | -12.574 | 7.6581 |
|  |  | 2 | -11.3868 | -11.3492 | 5.804 |
|  |  | 3 | -10.1769 | -10.1244 | 4.1786 |
|  |  | 4 | -8.96725 | -8.89994 | 2.8214 |
|  |  | 5 | -7.76337 | -7.68183 | 1.7702 |
|  |  | 6 | -6.58653 | -6.49274 | 1.0552 |
|  |  | 7 | -5.50199 | -5.41539 | 0.65571 |
|  |  | 8 | -4.75364 | -4.73182 | 0.42447 |
|  |  | 9 | -4.28405 | -4.29816 | 0.32128 |
|  |  | 10 | -3.73777 | -3.66934 | 0.24836 |
|  |  | 11 | -3.22015 | -3.07813 | 0.18999 |
|  |  | 12 | -2.72516 | -2.61618 | 0.15466 |
|  |  | 13 | -2.21933 | -2.15688 | 0.13134 |
|  |  | 14 | -1.69388 | -1.57984 | 0.11092 |
|  |  | 15 | -1.34277 | -1.04442 | 0.10084 |
|  |  | 16 | -1.15377 | -0.682264 | 0.095451 |
|  |  | 17 | -1.03171 | -0.491641 | 0.091504 |
|  |  | 18 | -0.744137 | -0.0111597 | 0.10251 |
|  |  | 19 | -0.457181 | 0.614424 | 0.12613 |
|  |  | 20 | -0.197996 | 1.46862 | 0.18207 |
|  |  | 21 | -0.057145 | 2.80554 | 0.32536 |
|  |  | 22 | -3.53E-09 | 4.83562 | 0.58774 |
|  |  | 23 | -1.20E-09 | 6.83313 | 1.0416 |
|  |  | 24 | -5.72E-10 | 8.57423 | 1.6365 |
|  |  | 25 | -8.69E-10 | 10.0308 | 2.258 |
|  |  | 26 | -1.11E-09 | 11.2281 | 2.7858 |
|  |  | 27 | -1.69E-09 | 12.201 | 3.1259 |
|  |  | 28 | -2.68E-09 | 12.9862 | 3.2073 |
|  |  | 29 | -6.06E-09 | 13.6211 | 2.9765 |
|  |  | 30 | -2.54E-08 | 14.1434 | 2.3927 |
|  |  | 31 | -0.02458 | 14.5905 | 1.425 |
|  |  | 32 | -0.046673 | 15 | 0.004866 |

Table 22. Comparison of recruitment dev parameter estimates from the 2015 assessment model and the author's preferred model (Model C).

| process | description | index | 2015AMO <br> estimate | Model C |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | estimate | std. dev. |
| recruitment devs | In-scale deviations | 1974 | 0.781402 | -- | -- |
|  |  | 1975 | 1.00935 | 1.40735 | 0.19124 |
|  |  | 1976 | 2.09407 | 1.99712 | 0.12382 |
|  |  | 1977 | 1.7989 | 1.76148 | 0.13002 |
|  |  | 1978 | 1.02156 | 1.09033 | 0.18136 |
|  |  | 1979 | -0.084761 | 0.165901 | 0.28812 |
|  |  | 1980 | -0.863678 | -0.465899 | 0.37249 |
|  |  | 1981 | -0.583826 | -0.0998744 | 0.21578 |
|  |  | 1982 | -1.25 | -0.492159 | 0.257 |
|  |  | 1983 | 0.697598 | 0.844003 | 0.10129 |
|  |  | 1984 | 0.664298 | 0.773732 | 0.12865 |
|  |  | 1985 | 1.59035 | 1.22589 | 0.10923 |
|  |  | 1986 | 1.32829 | 1.14466 | 0.11947 |
|  |  | 1987 | 1.26382 | 1.11144 | 0.12015 |
|  |  | 1988 | 1.17427 | 1.08617 | 0.10976 |
|  |  | 1989 | 0.206281 | 0.251569 | 0.15225 |
|  |  | 1990 | -0.659541 | -0.700321 | 0.24908 |
|  |  | 1991 | -1.21385 | -1.24123 | 0.28364 |
|  |  | 1992 | -1.49599 | -1.51533 | 0.26874 |
|  |  | 1993 | -1.59883 | -1.58988 | 0.24782 |
|  |  | 1994 | -1.4773 | -1.36351 | 0.20511 |
|  |  | 1995 | -1.19304 | -1.07756 | 0.17332 |
|  |  | 1996 | -1.08994 | -1.0552 | 0.18889 |
|  |  | 1997 | -0.187066 | -0.150971 | 0.10073 |
|  |  | 1998 | -1.09187 | -1.04219 | 0.18016 |
|  |  | 1999 | 0.0239972 | 0.0283579 | 0.10104 |
|  |  | 2000 | -0.479089 | -0.491797 | 0.1734 |
|  |  | 2001 | 0.71017 | 0.622348 | 0.091225 |
|  |  | 2002 | -0.232096 | -0.34659 | 0.19167 |
|  |  | 2003 | 0.298983 | 0.343703 | 0.12506 |
|  |  | 2004 | 0.803452 | 0.774672 | 0.088924 |
|  |  | 2005 | -0.452713 | -0.457059 | 0.19478 |
|  |  | 2006 | -0.660771 | -0.716854 | 0.21518 |
|  |  | 2007 | -0.952789 | -1.11789 | 0.27647 |
|  |  | 2008 | -0.81074 | -0.897263 | 0.25379 |
|  |  | 2009 | 0.949498 | 0.979229 | 0.099073 |
|  |  | 2010 | 1.12564 | 1.19858 | 0.093302 |
|  |  | 2011 | 0.604113 | 0.658634 | 0.12958 |
|  |  | 2012 | -0.966442 | -1.09582 | 0.38298 |
|  |  | 2013 | -0.169695 | -0.178842 | 0.17489 |
|  |  | 2014 | -0.101268 | -0.400162 | 0.19932 |
|  |  | 2015 | -0.530748 | -0.756357 | 0.26304 |
|  |  | 2016 | -- | -0.212413 | 0.24664 |

Table 23. Comparison of initial recruitment dev parameter estimates from the 2015 assessment model and the author's preferred model (Model C).

| process | description | index | 2015AMO <br> estimate | Model C |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | estimate | std. dev. |
| initial recruitment devs | In-scale deviations | 1949 | -1.49633 | -1.51108 | 1.6339 |
|  |  | 1950 | -1.49394 | -1.50848 | 1.4913 |
|  |  | 1951 | -1.48822 | -1.50227 | 1.3541 |
|  |  | 1952 | -1.47783 | -1.49106 | 1.224 |
|  |  | 1953 | -1.46091 | -1.47287 | 1.1033 |
|  |  | 1954 | -1.43472 | -1.44486 | 0.99453 |
|  |  | 1955 | -1.39531 | -1.4029 | 0.9007 |
|  |  | 1956 | -1.33677 | -1.34086 | 0.82451 |
|  |  | 1957 | -1.24998 | -1.24927 | 0.76768 |
|  |  | 1958 | -1.12031 | -1.1129 | 0.73004 |
|  |  | 1959 | -0.922636 | -0.905456 | 0.70936 |
|  |  | 1960 | -0.609611 | -0.576943 | 0.7035 |
|  |  | 1961 | -0.089749 | -0.0349116 | 0.71159 |
|  |  | 1962 | 0.696762 | 0.760147 | 0.71249 |
|  |  | 1963 | 1.54121 | 1.54366 | 0.69657 |
|  |  | 1964 | 1.98044 | 1.85947 | 0.66979 |
|  |  | 1965 | 1.9796 | 1.7515 | 0.66744 |
|  |  | 1966 | 1.75795 | 1.49285 | 0.67554 |
|  |  | 1967 | 1.51683 | 1.29124 | 0.67351 |
|  |  | 1968 | 1.3381 | 1.23276 | 0.6577 |
|  |  | 1969 | 1.24572 | 1.32514 | 0.6379 |
|  |  | 1970 | 1.19425 | 1.424 | 0.61001 |
|  |  | 1971 | 1.01783 | 1.26129 | 0.56459 |
|  |  | 1972 | 0.76483 | 0.955299 | 0.54235 |
|  |  | 1973 | 0.542804 | 0.470023 | 0.5477 |
|  |  | 1974 | -- | 0.186495 | 0.57714 |

Table 24. Comparison of fishery mortality/capture rate parameter estimates from the 2015 assessment model and the author's preferred model (Model C). GTF: groundfish fisheries; RKF: BBRKC fishery; SCF: snow crab fishery; TCF: directed Tanner crab fishery.

| process | description | param | 2015AMO | Model C |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | estimate | estimate | std. dev. |
| fishery mortality/capture rates | GTF effort extrapolation | pLnEffXtr_GTF | 1 | 1 | 0 |
|  | GTF In-scale female offset | pAvgLnF_GTFF | 0 | -1.02364 | 0.066812 |
|  | GTF In-scale mean [1973+] | pAvgLnF_GTF | -4.16128 | -4.11576 | 0.072179 |
|  | RKF effort extrapolation | pLnEffXtr_RKF | 1 | 1 | 0 |
|  | RKF In-scale female offset | pAvgLnF_RKFF | 0 | 2.43851 | 1.3139 |
|  | RKF In-scale mean [1992+] | pAvgLnF_RKF | -5.25 | -4.29718 | 0.92 |
|  | SCF effort extrapolation | pLnEffXtr_SCF | 1 | 1 | 0 |
|  | SCF In-scale female offset | pAvgLnF_SCFF | 0 | -1.48444 | 0.21286 |
|  | SCF In-scale mean [1992+] | pAvgLnF_SCF | -3.71005 | -2.55969 | 0.12387 |
|  | TCF effort extrapolation | pLnEffXtr_TCF | 1 | 1 | 0 |
|  | TCF In-scale female offset | pAvgLnF_TCFF | 0 | -1.6111 | 0.34153 |
|  | TCF In-scale mean [1965+] | pAvgLnF_TCF | -1.49637 | -1.32647 | 0.08658 |

Table 25. Comparison of fishery retention and selectivity curve parameter estimates from the 2015 assessment model and the author's preferred model (Model C). GTF: groundfish fisheries; RKF: BBRKC fishery; SCF: snow crab fishery; TCF: directed Tanner crab fishery.

| type | description | param | index | 2015AMO estimate | Model C |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  | estimate | std. dev. |
| TCF retention | size at 50\%-selected [-1990] | pRetTCFM_z50A1 |  | 137.669 | 138.347 | 0.46329 |
|  | size at 50\%-selected [1991+] | pRetTCFM_z50A2 |  | 133.078 | 133.013 | 0.5927 |
|  | slope [-1990] | pRetTCFM_slpA1 |  | 0.790725 | 0.68447 | 0.12092 |
|  | slope [1991+] | pRetTCFM_slpA2 |  | 0.366973 | 0.254571 | 0.018647 |
| TCF <br> selectivity | female size at 50\%-selected [all years] | pSelTCFF_z50 |  | 117.466 | 94.5043 | 2.1571 |
|  | female slope [all years] | pSelTCFF_slp |  | 0.140497 | 0.196036 | 0.020346 |
|  | male In-scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 1991 | 0.0832307 | 0.160928 | 0.030713 |
|  | male In-scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 1992 | 0.130107 | 0.167735 | 0.022307 |
|  | male In -scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 1993 | 0.100172 | 0.152329 | 0.026045 |
|  | male In -scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 1994 | 0.136988 | 0.245468 | 0.028421 |
|  | male In -scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 1995 | -0.00932885 | -0.116733 | 0.091221 |
|  | male In -scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 1996 | -0.431057 | -0.500471 | 0.013172 |
|  | male $\ln$-scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 2005 | -0.0562356 | -0.0691252 | 0.024499 |
|  | male In-scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 2006 | -0.0640353 | -0.0855568 | 0.023566 |
|  | male In-scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 2007 | -0.0943149 | -0.0977496 | 0.02153 |
|  | male In -scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 2008 | 0.0460822 | 0.0331269 | 0.02221 |
|  | male In -scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 2009 | 0.219118 | 0.264636 | 0.020202 |
|  | male In -scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 2013 | -0.0185012 | -0.0165809 | 0.021704 |
|  | male In -scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 2014 | -0.0422246 | -0.047993 | 0.019172 |
|  | male In -scale devs in size at 50\%-selected [1991+] | pSelTCFM_devsZ50 | 2015 | -- | -0.090013 | 0.021611 |
|  | male In-scale mean size at $50 \%$-selected | pSelTCFM_mnLnZ50A2 |  | 4.83157 | 4.75673 | 0.011685 |
|  | male slope [-1996] | pSelTCFM_slpA1 |  | 0.114058 | 0.0898399 | 0.006701 |
|  | male slope [1997+] | pSelTCFM_slpA2 |  | 0.144611 | 0.179297 | 0.014102 |
| GTF selectivity | female size at 50\%-selected [-1987] | pSelGTFF_z50A1 |  | 125.01 | 40.0799 | 1.4501 |
|  | female size at 50\%-selected [1988-1996] | pSelGTFF_z50A2 |  | 159.214 | 40 | 0.000155 |
|  | female size at 50\%-selected [1997+] | pSelGTFF_z50A3 |  | 143.991 | 79.148 | 2.4561 |
|  | female slope [-1987] | pSelGTFF_slpA1 |  | 0.0286752 | 0.152178 | 0.02319 |
|  | female slope [1988-1996] | pSelGTFF_slpA2 |  | 0.0158887 | 0.183165 | 0.037518 |
|  | female slope [1997+] | pSelGTFF_slpA3 |  | 0.052039 | 0.0768591 | 0.005855 |
|  | male size at 50\%-selected [-1987] | pSelGTFM_z50A1 |  | 57.0742 | 54.7273 | 1.8329 |
|  | male size at 50\%-selected [1988-1996] | pSelGTFM_z50A2 |  | 72.6065 | 66.3956 | 4.993 |
|  | male size at 50\%-selected [1997+] | pSelGTFM_z50A3 |  | 83.1856 | 84.6716 | 2.0078 |
|  | male slope [-1987] | pSelGTFM_slpA1 |  | 0.10874 | 0.103462 | 0.009792 |
|  | male slope [1988-1996] | pSelGTFM_slpA2 |  | 0.0427268 | 0.0483958 | 0.007576 |
|  | male slope [1997+] | pSelGTFM_slpA3 |  | 0.0777645 | 0.075398 | 0.003877 |
| ```RKF selectivity``` | female size at 50\%-selected [-1996] | pSelRKFF_z50A1 |  | 98.3537 | 97.2472 | 11.723 |
|  | female size at 50\%-selected [1997-2004] | pSelRKFF_z50A2 |  | 103.261 | 97.0295 | 10.201 |
|  | female size at 50\%-selected [2005+] | pSelRKFF_z50A3 |  | 157.074 | 114.727 | 17.968 |
|  | female slope [-1996] | pSelRKFF_slpA1 |  | 0.238438 | 0.210067 | 0.11678 |
|  | female slope [1997-2004] | pSelRKFF_slpA2 |  | 0.179464 | 0.203964 | 0.13997 |
|  | female slope [2005+] | pSelRKFF_slpA3 |  | 0.183223 | 0.164415 | 0.060323 |
|  | male size at 50\%-selected [-1996] | pSelRKFM_z50A1 |  | 150 | 150 | 0.000611 |
|  | male size at 50\%-selected [1997-2004] | pSelRKFM_z50A2 |  | 133.217 | 138.978 | 14.126 |
|  | male size at 50\%-selected [2005+] | pSelRKFM_z50A3 |  | 150 | 150 | 0.001334 |
|  | male slope [-1996] | pSelRKFM_slpA1 |  | 0.101212 | 0.113097 | 0.011114 |
|  | male slope [1997-2004] | pSelRKFM_slpA2 |  | 0.0915078 | 0.0863304 | 0.022917 |
|  | male slope [2005+] | pSelRKFM_slpA3 |  | 0.082357 | 0.0851915 | 0.006282 |
| SCF <br> selectivity | female size at 50\%-selected [-1996] | pSelSCFF_250A1 |  | 110.423 | 67.4884 | 7.1383 |
|  | female size at 50\%-selected [1997-2004] | pSelSCFF_250A2 |  | 76.1912 | 75.3363 | 4.7225 |
|  | female size at 50\%-selected [2005+] | pSelSCFF_250A3 |  | 88.6981 | 78.9834 | 3.9168 |
|  | female slope [-1996] | pSelSCFF_slpA1 |  | 0.05 | 0.206465 | 0.17212 |
|  | female slope [1997-2004] | pSelSCFF_slpA2 |  | 0.254036 | 0.271067 | 0.14346 |
|  | female slope [2005+] | pSelSCFF_slpA3 |  | 0.134828 | 0.206033 | 0.068651 |
|  | male ascending size at 50\%-selected [-1996] | pSelSCFM_z50A1 |  | 86.8038 | 87.6083 | 1.4676 |
|  | male ascending size at 50\%-selected [1997-2004] | pSelSCFM_z50A2 |  | 93.9094 | 94.1945 | 3.3921 |
|  | male ascending size at 50\%-selected [2005+] | pSelSCFM_z50A3 |  | 103.632 | 104.944 | 1.6099 |
|  | male ascending slope [-1996] | pSelSCFM_slpA1 |  | 0.404304 | 0.401603 | 0.13411 |
|  | male ascending slope [1997-2004] | pSelSCFM_slpA2 |  | 0.231803 | 0.226234 | 0.07431 |
|  | male ascending slope [2005+] | pSelSCFM_slpA3 |  | 0.178644 | 0.171992 | 0.01611 |
|  | male descending In -scale offset to size at $50 \%$-selected [-1996] | pSelSCFM_InZ50D1 |  | 3.97235 | 3.95657 | 0.036866 |
|  | male descending $\ln$-scale offset to size at $50 \%$-selected [1997-2004] | pSelSCFM_InZ50D2 |  | 3.80135 | 3.79291 | 0.16484 |
|  | male descending In-scale offset to size at $50 \%$-selected [2005+] | pSelSCFM_InZ50D3 |  | 3.53118 | 3.48534 | 0.091741 |
|  | male descending slope [-1996] | pSelSCFM_slpD1 |  | 0.499994 | 0.499999 | 0.000334 |
|  | male descending slope [1997-2004] | pSelSCFM_slpD2 |  | 0.17705 | 0.154555 | 0.090084 |
|  | male descending slope [2005+] | pSelSCFM_slpD3 |  | 0.183485 | 0.176146 | 0.027094 |

Table 26. Comparison of fishery mortality/capture rate dev parameter estimates from the 2015 assessment model and the author's preferred model (Model C). TCF: directed Tanner crab fishery.

| type | description | index | 2015AMO <br> estimate | Model C |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | estimate | std. dev. |
| TCF mortality/capture rate devs | In-scale devs [1965+] | 1965 | -0.518187 | -0.512072 | 0.49992 |
|  |  | 1966 | -0.773462 | -0.753569 | 0.38716 |
|  |  | 1967 | 0.359217 | 0.431136 | 0.34912 |
|  |  | 1968 | 0.121306 | 0.253429 | 0.32494 |
|  |  | 1969 | 0.220923 | 0.433976 | 0.31293 |
|  |  | 1970 | 0.0220202 | 0.314614 | 0.31273 |
|  |  | 1971 | -0.200343 | 0.144671 | 0.30767 |
|  |  | 1972 | -0.365518 | -0.0134198 | 0.27973 |
|  |  | 1973 | -0.570184 | -0.273418 | 0.21589 |
|  |  | 1974 | -0.323904 | -0.126451 | 0.14351 |
|  |  | 1975 | -0.040857 | 0.0557562 | 0.10496 |
|  |  | 1976 | 0.761268 | 0.81054 | 0.095966 |
|  |  | 1977 | 1.49067 | 1.60134 | 0.10925 |
|  |  | 1978 | 1.688 | 1.98097 | 0.15051 |
|  |  | 1979 | 2.38683 | 2.80725 | 0.1968 |
|  |  | 1980 | 2.44285 | 2.34269 | 0.27763 |
|  |  | 1981 | 0.596186 | 0.304394 | 0.14568 |
|  |  | 1982 | -0.350215 | -0.709751 | 0.12706 |
|  |  | 1983 | -1.2767 | -1.69005 | 0.24792 |
|  |  | 1984 | 0.0970324 | -0.611706 | 0.182 |
|  |  | 1987 | -0.866666 | -1.30304 | 0.21134 |
|  |  | 1988 | -0.113462 | -0.47743 | 0.10694 |
|  |  | 1989 | 0.879841 | 0.73493 | 0.083425 |
|  |  | 1990 | 1.37173 | 1.45872 | 0.09428 |
|  |  | 1991 | 1.28887 | 1.41528 | 0.15539 |
|  |  | 1992 | 1.66753 | 1.63773 | 0.14433 |
|  |  | 1993 | 0.961286 | 0.995718 | 0.13994 |
|  |  | 1994 | 0.761891 | 0.982647 | 0.19767 |
|  |  | 1995 | -0.070297 | -0.168372 | 0.13396 |
|  |  | 1996 | -1.2281 | -0.959074 | 0.17763 |
|  |  | 2005 | -2.14795 | -2.12915 | 0.20981 |
|  |  | 2006 | -1.65181 | -1.64818 | 0.143 |
|  |  | 2007 | -1.68988 | -1.64767 | 0.13607 |
|  |  | 2008 | -1.75263 | -1.96315 | 0.15983 |
|  |  | 2009 | -1.04851 | -1.32018 | 0.25734 |
|  |  | 2013 | -1.68639 | -1.70897 | 0.13862 |
|  |  | 2014 | -0.442409 | -0.491133 | 0.092358 |
|  |  | 2015 | -- | -0.199011 | 0.09397 |

Table 27. Comparison of fishery mortality/capture rate dev parameter estimates from the 2015 assessment model and the author's preferred model (Model C). RKF: BBRKC fishery; SCF: snow crab fishery.

| type | description | index | 2015AMO estimate | Model C |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | estimate | std. dev. |
| RKF mortality/capture rate devs |  | 1992 | 0 | -0.141197 | 0.35612 |
|  |  | 1993 | 0 | -0.0285905 | 0.37414 |
|  |  | 1994 | 0 | -0.0710423 | 0.36889 |
|  |  | 1995 | 0 | 0.0118673 | 0.38532 |
|  |  | 1996 | 0 | 0.080407 | 0.40387 |
|  |  | 1997 | 0 | 0.0817798 | 0.40921 |
|  |  | 1998 | 0 | 0.0129244 | 0.39762 |
|  |  | 1999 | 0 | -0.00110857 | 0.39589 |
|  |  | 2000 | 0 | 0.0012108 | 0.39612 |
|  |  | 2001 | 0 | -0.00950446 | 0.3933 |
|  |  | 2002 | 0 | -0.0200168 | 0.39105 |
|  | In-scale devs | 2003 | 0 | -0.00521674 | 0.39159 |
|  | [1992+] | 2004 | 0 | -0.0290172 | 0.38766 |
|  |  | 2005 | 0 | 0.00917559 | 0.39966 |
|  |  | 2006 | 0 | 0.00985092 | 0.39917 |
|  |  | 2007 | 0 | 0.0119242 | 0.39923 |
|  |  | 2008 | 0 | 0.0267412 | 0.40101 |
|  |  | 2009 | 0 | 0.0171997 | 0.39891 |
|  |  | 2010 | 0 | 0.00829416 | 0.3981 |
|  |  | 2011 | 0 | 0.00289747 | 0.39786 |
|  |  | 2012 | 0 | 0.0030385 | 0.39824 |
|  |  | 2013 | 0 | 0.0101265 | 0.39829 |
|  |  | 2014 | 0 | 0.0251161 | 0.39837 |
|  |  | 2015 | -- | -0.00686042 | 0.39308 |
| SCF mortality/capture rate devs | In-scale devs [1992+] | 1992 | 1.84979 | 1.82084 | 0.11859 |
|  |  | 1993 | 1.62748 | 1.57903 | 0.12573 |
|  |  | 1994 | 1.2734 | 1.21802 | 0.14901 |
|  |  | 1995 | 1.27571 | 1.20648 | 0.17512 |
|  |  | 1996 | 0.19664 | 0.14783 | 0.45612 |
|  |  | 1997 | 0.733603 | 0.750337 | 0.38909 |
|  |  | 1998 | 0.494163 | 0.672925 | 0.43946 |
|  |  | 1999 | $-0.381905$ | -0.326133 | 0.6841 |
|  |  | 2000 | -0.621997 | -0.654371 | 0.66115 |
|  |  | 2001 | -0.580084 | -0.618835 | 0.62982 |
|  |  | 2002 | -0.568142 | -0.547399 | 0.59508 |
|  |  | 2003 | -0.811723 | -0.853073 | 0.58876 |
|  |  | 2004 | -1.14597 | -1.08342 | 0.5689 |
|  |  | 2005 | -0.649415 | -0.609679 | 0.50401 |
|  |  | 2006 | -0.339788 | -0.33246 | 0.41964 |
|  |  | 2007 | -0.20635 | -0.224263 | 0.34989 |
|  |  | 2008 | $-0.609894$ | -0.662066 | 0.42994 |
|  |  | 2009 | -0.486074 | -0.521409 | 0.42481 |
|  |  | 2010 | -0.419701 | -0.379555 | 0.43452 |
|  |  | 2011 | 0.0130669 | 0.0832503 | 0.35008 |
|  |  | 2012 | -0.577714 | -0.525958 | 0.46695 |
|  |  | 2013 | -0.479325 | -0.494068 | 0.3501 |
|  |  | 2014 | 0.414236 | 0.353441 | 0.17733 |
|  |  | 2015 | -- | 0.000536055 | 0.23227 |

Table 28. Comparison of fishery mortality/capture rate dev parameter estimates from the 2015 assessment model and the author's preferred model (Model C). GTF: groundfish fisheries.

| type | description | index | 2015AMO <br> estimate | Model C |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | estimate | std. dev. |
| GTF mortality/capture rate devs | In-scale devs[1973+] | 1973 | 0.84482 | 1.10031 | 0.10447 |
|  |  | 1974 | 1.27268 | 1.46916 | 0.081611 |
|  |  | 1975 | 0.460622 | 0.609631 | 0.078217 |
|  |  | 1976 | -0.028137 | 0.0774622 | 0.090286 |
|  |  | 1977 | -0.248686 | -0.209844 | 0.11808 |
|  |  | 1978 | -0.419782 | -0.440285 | 0.15604 |
|  |  | 1979 | 0.218235 | 0.233132 | 0.11269 |
|  |  | 1980 | 0.0456019 | -0.0216788 | 0.15222 |
|  |  | 1981 | -0.07109 | -0.206465 | 0.19247 |
|  |  | 1982 | -0.726093 | -0.916129 | 0.39423 |
|  |  | 1983 | -0.150186 | -0.413008 | 0.35909 |
|  |  | 1984 | 0.251739 | -0.20437 | 0.39205 |
|  |  | 1985 | -0.285296 | -0.629289 | 0.47766 |
|  |  | 1986 | -0.367893 | -0.548176 | 0.38022 |
|  |  | 1987 | -0.649807 | -0.719865 | 0.37764 |
|  |  | 1988 | -1.11646 | -1.10449 | 0.40795 |
|  |  | 1989 | -1.03265 | -0.951716 | 0.34438 |
|  |  | 1990 | -0.716481 | -0.605589 | 0.27986 |
|  |  | 1991 | 0.392271 | 0.49366 | 0.12766 |
|  |  | 1992 | 0.686347 | 0.783903 | 0.11916 |
|  |  | 1993 | 0.555778 | 0.635226 | 0.16501 |
|  |  | 1994 | 1.06755 | 1.12753 | 0.1428 |
|  |  | 1995 | 1.11494 | 1.15185 | 0.18109 |
|  |  | 1996 | 1.47253 | 1.48679 | 0.17172 |
|  |  | 1997 | 1.37406 | 1.44223 | 0.23212 |
|  |  | 1998 | 1.06557 | 1.11859 | 0.33244 |
|  |  | 1999 | 0.531428 | 0.573452 | 0.50148 |
|  |  | 2000 | 0.657746 | 0.648246 | 0.4107 |
|  |  | 2001 | 1.00301 | 1.01488 | 0.25273 |
|  |  | 2002 | 0.366648 | 0.396099 | 0.37669 |
|  |  | 2003 | -0.216728 | -0.151861 | 0.48062 |
|  |  | 2004 | -0.125303 | -0.00093073 | 0.36869 |
|  |  | 2005 | $-0.353084$ | -0.222611 | 0.37665 |
|  |  | 2006 | -0.289489 | -0.174462 | 0.33252 |
|  |  | 2007 | -0.367112 | -0.280821 | 0.33126 |
|  |  | 2008 | -0.583965 | -0.517741 | 0.3744 |
|  |  | 2009 | -0.769095 | -0.672724 | 0.4316 |
|  |  | 2010 | -0.880976 | -0.74587 | 0.48448 |
|  |  | 2011 | -0.879599 | -0.7536 | 0.50303 |
|  |  | 2012 | -1.05669 | -0.946181 | 0.50307 |
|  |  | 2013 | -1.01702 | -0.932219 | 0.42678 |
|  |  | 2014 | -1.02995 | -0.963513 | 0.3941 |
|  |  | 2015 | -- | -1.02871 | 0.42894 |

Table 29. Comparison of fits to mature survey biomass by sex (in 1000's t) from the 2015 assessment model and the author's preferred model (Model C).

| year | mature female biomass (Kt) |  |  | mature male biomass (Kt) |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | observed | 2015AMO | Model C |  |  |  |
| 1975 | 31.7 | 46.4 | 47.8 | 246.0 | 155.1 | 148.1 |
| 1976 | 31.4 | 40.4 | 42.0 | 126.2 | 133.7 | 133.6 |
| 1977 | 38.8 | 34.5 | 35.8 | 110.6 | 102.2 | 105.5 |
| 1978 | 26.2 | 30.9 | 32.7 | 77.6 | 68.3 | 75.1 |
| 1979 | 19.7 | 32.2 | 34.7 | 32.2 | 59.0 | 67.0 |
| 1980 | 64.2 | 34.2 | 36.5 | 86.2 | 61.5 | 63.0 |
| 1981 | 43.1 | 28.2 | 31.5 | 49.4 | 46.4 | 53.8 |
| 1982 | 64.4 | 25.2 | 25.7 | 49.0 | 58.9 | 68.1 |
| 1983 | 20.6 | 17.2 | 19.2 | 28.5 | 37.3 | 49.1 |
| 1984 | 15.0 | 11.6 | 14.5 | 24.2 | 21.5 | 32.6 |
| 1985 | 5.6 | 8.5 | 11.7 | 11.4 | 13.0 | 23.0 |
| 1986 | 3.5 | 9.3 | 12.3 | 12.8 | 18.3 | 28.8 |
| 1987 | 5.2 | 12.3 | 14.3 | 24.1 | 31.6 | 40.7 |
| 1988 | 25.5 | 17.2 | 17.0 | 60.4 | 51.1 | 55.2 |
| 1989 | 19.5 | 22.2 | 19.8 | 91.9 | 77.0 | 70.2 |
| 1990 | 37.8 | 24.8 | 21.4 | 96.3 | 85.7 | 74.4 |
| 1991 | 45.0 | 24.6 | 21.2 | 109.7 | 74.5 | 64.8 |
| 1992 | 26.5 | 21.8 | 19.1 | 103.2 | 68.4 | 60.1 |
| 1993 | 11.7 | 16.9 | 15.3 | 60.1 | 50.4 | 45.1 |
| 1994 | 10.0 | 12.6 | 11.6 | 42.1 | 36.0 | 32.9 |
| 1995 | 12.7 | 9.2 | 8.6 | 31.1 | 25.9 | 23.9 |
| 1996 | 9.8 | 6.9 | 6.5 | 26.3 | 18.6 | 17.3 |
| 1997 | 3.5 | 5.3 | 5.1 | 10.7 | 14.6 | 13.9 |
| 1998 | 2.3 | 4.3 | 4.3 | 10.3 | 12.9 | 12.5 |
| 1999 | 3.9 | 3.9 | 4.0 | 12.5 | 12.6 | 12.4 |
| 2000 | 4.2 | 4.2 | 4.3 | 16.1 | 14.3 | 14.1 |
| 2001 | 4.6 | 4.5 | 4.7 | 17.9 | 17.6 | 17.4 |
| 2002 | 4.5 | 5.1 | 5.2 | 17.8 | 20.2 | 20.0 |
| 2003 | 8.4 | 6.0 | 6.0 | 23.3 | 24.4 | 23.7 |
| 2004 | 4.9 | 7.5 | 7.2 | 26.3 | 30.6 | 29.0 |
| 2005 | 11.6 | 8.8 | 8.3 | 43.1 | 39.6 | 36.3 |
| 2006 | 15.0 | 9.7 | 9.3 | 64.2 | 44.9 | 41.0 |
| 2007 | 13.5 | 10.8 | 10.6 | 66.4 | 49.3 | 45.4 |
| 2008 | 11.7 | 11.0 | 10.8 | 62.7 | 55.3 | 51.3 |
| 2009 | 8.6 | 9.6 | 9.6 | 36.3 | 53.9 | 50.7 |
| 2010 | 5.5 | 8.1 | 8.1 | 37.6 | 47.2 | 44.3 |
| 2011 | 5.5 | 7.8 | 7.7 | 41.5 | 41.9 | 38.8 |
| 2012 | 12.5 | 9.8 | 9.8 | 41.2 | 42.9 | 39.4 |
| 2013 | 18.0 | 13.2 | 13.5 | 65.7 | 57.4 | 53.4 |
| 2014 | 14.9 | 15.0 | 15.6 | 79.5 | 73.8 | 71.1 |
| 2015 | 11.3 | 13.8 | 14.6 | 60.2 | 72.6 | 72.2 |
| 2016 | 7.6 | -- | 12.4 | 57.6 | -- | 59.1 |

Table 30. Comparison of estimates of mature biomass-at-mating by sex (in 1000's t) from the 2015 assessment model and the author's preferred model (Model C).

| year | MMB (1000's t) |  | MFB (1000's t) |  |
| :---: | :---: | :---: | :---: | :---: |
|  | 2015AMO | Model C | 2015AMO | Model C |
| 1949 | 0.0 | 0.0 | 0.0 | 0.0 |
| 1950 | 0.0 | 0.0 | 0.0 | 0.0 |
| 1951 | 0.2 | 0.1 | 0.3 | 0.3 |
| 1952 | 1.4 | 1.2 | 1.1 | 1.1 |
| 1953 | 4.8 | 4.1 | 2.3 | 2.2 |
| 1954 | 8.7 | 7.8 | 3.3 | 3.2 |
| 1955 | 11.6 | 10.6 | 4.1 | 4.0 |
| 1956 | 13.8 | 12.7 | 4.6 | 4.5 |
| 1957 | 15.5 | 14.4 | 5.0 | 5.0 |
| 1958 | 16.9 | 15.8 | 5.4 | 5.3 |
| 1959 | 18.2 | 17.0 | 5.7 | 5.7 |
| 1960 | 19.4 | 18.2 | 6.2 | 6.2 |
| 1961 | 21.0 | 19.7 | 6.7 | 6.7 |
| 1962 | 23.1 | 21.8 | 7.7 | 7.7 |
| 1963 | 26.8 | 25.4 | 9.5 | 9.5 |
| 1964 | 34.2 | 32.5 | 13.9 | 13.9 |
| 1965 | 49.9 | 47.5 | 24.3 | 24.3 |
| 1966 | 90.2 | 84.2 | 45.3 | 43.7 |
| 1967 | 150.6 | 136.5 | 74.9 | 68.6 |
| 1968 | 233.5 | 200.1 | 103.0 | 89.0 |
| 1969 | 291.4 | 235.6 | 118.9 | 98.4 |
| 1970 | 317.0 | 244.9 | 121.9 | 98.9 |
| 1971 | 317.5 | 240.8 | 117.2 | 96.4 |
| 1972 | 305.4 | 236.2 | 109.7 | 93.9 |
| 1973 | 287.6 | 235.9 | 101.5 | 92.7 |
| 1974 | 257.2 | 229.8 | 92.2 | 89.4 |
| 1975 | 226.4 | 219.6 | 82.3 | 83.0 |
| 1976 | 171.8 | 179.3 | 71.1 | 71.8 |
| 1977 | 106.2 | 119.0 | 60.0 | 60.0 |
| 1978 | 70.3 | 81.1 | 53.8 | 55.3 |
| 1979 | 48.2 | 54.7 | 55.1 | 57.4 |
| 1980 | 31.2 | 44.9 | 52.1 | 56.0 |


| year | MMB (1000's t) |  | MFB (1000's t) |  |
| ---: | ---: | ---: | ---: | ---: |
|  | 2015AMO | Model C | 2015AMO | Model C |
| 1981 | 40.7 | 56.6 | 44.4 | 49.7 |
| 1982 | 37.9 | 54.9 | 33.3 | 40.5 |
| 1983 | 25.3 | 41.0 | 22.8 | 30.8 |
| 1984 | 12.8 | 25.7 | 15.2 | 23.1 |
| 1985 | 13.6 | 26.2 | 12.5 | 20.0 |
| 1986 | 19.1 | 32.6 | 13.7 | 20.6 |
| 1987 | 31.2 | 44.4 | 18.0 | 23.8 |
| 1988 | 48.3 | 58.5 | 25.3 | 28.5 |
| 1989 | 60.3 | 63.3 | 32.2 | 32.6 |
| 1990 | 55.1 | 54.3 | 35.1 | 34.3 |
| 1991 | 55.1 | 52.5 | 34.7 | 34.0 |
| 1992 | 48.2 | 45.2 | 30.2 | 30.6 |
| 1993 | 40.8 | 39.5 | 24.0 | 25.0 |
| 1994 | 31.5 | 31.4 | 18.0 | 19.0 |
| 1995 | 22.8 | 23.1 | 13.3 | 14.2 |
| 1996 | 17.7 | 18.1 | 10.0 | 10.8 |
| 1997 | 14.7 | 15.2 | 7.6 | 8.5 |
| 1998 | 13.2 | 13.9 | 6.3 | 7.3 |
| 1999 | 13.4 | 14.3 | 5.8 | 6.9 |
| 2000 | 15.2 | 16.3 | 6.2 | 7.3 |
| 2001 | 18.4 | 19.8 | 6.7 | 7.9 |
| 2002 | 21.5 | 23.1 | 7.5 | 8.8 |
| 2003 | 26.2 | 27.7 | 8.9 | 10.2 |
| 2004 | 32.9 | 33.8 | 11.2 | 12.4 |
| 2005 | 41.9 | 41.6 | 13.1 | 14.4 |
| 2006 | 46.8 | 46.3 | 14.4 | 16.0 |
| 2007 | 51.3 | 51.3 | 16.1 | 18.2 |
| 2008 | 58.4 | 58.9 | 16.3 | 18.5 |
| 2009 | 57.4 | 58.5 | 14.3 | 16.4 |
| 2010 | 51.0 | 51.7 | 12.1 | 13.9 |
| 2011 | 45.1 | 45.2 | 11.5 | 13.3 |
| 2012 | 46.5 | 46.2 | 14.6 | 17.0 |
| 2013 | 60.6 | 61.2 | 19.7 | 23.4 |
| 2014 | 71.6 | 75.4 | 22.0 | 26.7 |
| 2015 | -- | 73.9 | -- | 24.9 |
|  |  |  |  |  |

Table 31. Estimated population size (thousands) for females on July 1 of year. from the author's preferred model, Model C.

| year | 27.5 |  |  |  |  |  |  |  |  |  |  | 825 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 199 |  | 0.00 er | 0.00 | Doetoo | 0.008 | $0.000^{\text {a }}$ | 0.00 | 0.008 | 0.00 Ftoo | 0.00 Ftoo | 0.00 F +00 | 0.00 | 0.00 F 50 | 0.006 | $0.008+00$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }_{\substack{1950 \\ 1951}}^{1951}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }_{1951}^{1951}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }_{1} 195$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1954 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1956 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1957 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1958 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1959 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 196 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1962 |  |  |  |  |  |  |  |  |  | 1.59 |  |  |  | ${ }_{6}^{5636}$ | ${ }_{4,12}^{312}$ |  |  |  |  |  |  |  |  |  |  |  | ${ }_{5}^{4.88}$ |  |  |  |  |  |
| 1963 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1964 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1965 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1966 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\underset{1}{1967}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | (e) |
| ${ }_{1969}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | ${ }_{1.29509}^{14209}$ |
| 1970 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1971 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1972 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }_{1973}^{1974}$ | 308E |  |  |  |  |  |  |  |  | ${ }_{\text {l }}^{1.5885+02}$ | ${ }_{1.51}^{1.6}$ |  |  | ${ }_{8.0}^{8.0}$ | ${ }_{5}^{522}$ |  |  |  |  |  |  |  |  |  |  |  |  | ¢9.97e08 |  |  |  |  |
| 1975 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1976 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1977 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1980 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1981 |  | 2.40 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | 1.23 |  |  |  | 1.52E-10 |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1984 |  | ${ }_{5.856}^{150.0}$ |  |  |  | ${ }_{3} 18$ | 22 |  |  | ${ }_{3}^{4.28}$ | ${ }_{3}^{4} 7$ |  |  | 2.1 | ${ }_{1}^{194}$ |  |  |  | ${ }_{1.4}^{18}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1985 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }^{1986}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\underset{\substack{1988 \\ 1988}}{ }$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1989 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1992 |  |  |  |  |  |  |  |  |  |  |  |  |  | ${ }_{2} 2.2$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }_{1999}^{1999}$ | ${ }_{388}^{291}$ | cisfer | ${ }_{83}^{636}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 2000 |  |  | ${ }_{1}^{2}$ |  |  |  |  |  |  |  |  |  | 8.912 | 6.00¢ +00 | cose |  | 7.15 |  | ${ }_{2}^{3.066}$ |  |  | 3,7) | ${ }_{4.48}^{3688}$ |  |  |  |  |  |  |  |  |  |
|  | 2.12 | 4.848 F01 |  |  | 2.11 | 1.74 |  |  |  | 1.4 |  |  |  |  |  |  |  |  | 322 |  |  |  |  |  |  |  |  | $1.11 \mathrm{E}=08$ |  |  |  | 1.37-10 |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }_{2004}^{2003}$ |  | ${ }_{5}^{3}, 730$ | ${ }_{5}^{3217}$ | ${ }_{4}^{2} 36$ |  | ${ }_{23}^{238}$ | ${ }_{1.00}^{260}$ | ${ }_{1.72}^{21,}$ |  | ${ }_{2,60}^{214}$ | ${ }_{2}^{2}$ | ${ }_{2}^{2}$ | ${ }_{15}^{12}$ | ${ }_{1}^{8} 100$ | ${ }_{6}^{5}$ |  |  |  |  |  |  | ${ }_{6}$ | ${ }_{7}^{6}, 51$ |  |  |  |  |  |  |  |  | ${ }^{2} 2.352 .10$ |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }_{2000}^{2000}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }^{2008}$ |  | 1.08 | 1.02 | 9.4 |  |  | ${ }^{\text {l }}$ 7.83E+00 |  | 1.42F+0] | 2.4 | ${ }^{2} 888800$ | 2.6 | 22 | ${ }^{1.65}$ | 1.12 |  | 23 | 5.5 | 9 | 1.188002 | ${ }^{1200003}$ |  | 7.8 | 1.3 |  | ${ }^{1.311507}$ | ${ }^{4.212088}$ | 1. | 4,24:099 |  |  | ${ }^{1.666510}$ |
| 2009 2001 |  |  |  |  |  |  |  |  |  |  |  |  |  |  | ${ }_{8}^{10.585}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  | ${ }^{2} .2075$ |  |  |  | 4.646 |  |  |  |  |  |  | ${ }_{\text {l }}^{1.268501}$ |  |  |  |  | ${ }^{6}$ |  | 9,366-04 |  | ${ }_{1}^{12}$ | 3. |  |  |  |  |  |  |  |  |
| 2014 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 2015 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |

Table 32. Estimated population size (thousands) for males on July 1 of year. from the author's preferred mode, Model C.

|  | 27.5 | 32. | 37.5 | 42.5 | 475 | 52.5 | 57.5 | 62.5 | 67.5 | 2.5 | 77.5 | 82.5 | 87.5 | 92.5 | 975 | 1025 | 1075 | 12.5 | 7.5 | 122.5 | 127.5 | 1325 | 137.5 | 1425 | 147.5 | 2.5 | 157.5 | 1625 | 1675 | 2.5 | 7.5 |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1099 | 0.006+ | coictiol | coiole | come | coictiol | coicle | (0,0) | coicotro | (0.06too | O.OEF50 |  | - 0 OREFOO |  |  |  |  |  |  | coiole |  | coiole | (0.0Eta0 | O.0eteo | come |  |  | - |  |  |  |  |  |
| 1950 1951 1951 | $4.266+10$ | 997 | 9.56 | 8.0EE | 6.54F | 4.53E+00 | 279E | 1.58E | 8.52 E | 4.43E | ${ }^{2.22 E}$ | 10 | 5.20:02 | 245 | 1.13 | 519 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1951 |  |  |  |  |  |  |  |  |  |  |  | 1.196+00 | 7.65E.0 | 4.8.ETE1 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1953 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1954 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1955 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 195 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1957 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1958 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 196 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1962 |  | 9.43E |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1963 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1964 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1965 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1968 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1969 | 7.25 | 11.68 | ${ }_{1}^{1.59}$ |  |  |  |  |  |  |  | 6.57 | ${ }_{6}^{643}$ |  | ${ }^{6.57}$ | 691 | 6.98 |  |  |  |  |  |  | 4.00 |  | ${ }_{2}^{288}$ |  |  |  |  |  |  |  |
| 1970 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1971 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }_{1972}^{1972}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | ${ }_{26}^{2,6}$ |  |  |  |  |  |  |  | cismen |
| 1974 | ${ }_{232}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1975 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1978 |  | ${ }_{8,211}^{1+51}$ |  |  |  |  |  | ${ }^{8.346}$ | ${ }_{7}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1979 | 1.34 | ${ }^{3} 322$ | 4.18 |  | 5.54 |  |  |  |  |  |  |  |  |  | 5.04 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1982 | 6.95 | 1.66 | 1.75 | 1.8 |  | 1.32 | 1.14 |  |  |  |  | 1.08 |  | 1.3 | 15 | 1.6 |  |  | 1.78 |  | 1.61 |  |  | 9.59 | 7.95 |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }_{1}^{1984} 1$ | $\underbrace{2.46}_{3.888}$ | ${ }_{8}^{5938}$ | ${ }_{\substack{5 \\ 7,900}}^{1}$ | ${ }_{6}^{5} 5.36$ | ${ }_{4.81}^{4}$ |  |  |  |  |  | ${ }_{1}^{8.018}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  | 7.65 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1987 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1990 | 5.651 | 1.90 | 1.78 | 2.28 |  |  | 3.05 |  |  | 283 | 276 | 2.64 |  |  | 2488 | 24 E | 228 |  |  |  | 1.80 |  |  |  | 7.99 |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }_{1993}^{1992}$ | ${ }_{\text {2 }}^{2}$ | 5.4 | ${ }_{5}^{6}$ 5, 31 | ${ }_{5.1}^{6.15}$ | ${ }_{4}^{5} /{ }^{\text {5,7e }}$ |  | ${ }_{4.3}^{623}$ | ${ }_{4}^{6}$ |  |  |  | ${ }_{6.59}^{1118}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | ${ }_{5}^{6}$ |  |  | (e.03 |
| 1994 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }_{1995}^{1995}$ |  |  | ${ }_{8}^{8.14 E}$ | ${ }_{\text {coser }}^{\substack{6,822}}$ |  | ${ }_{5}^{4} 272$ | 4.46 | ${ }_{3}^{3}$ |  |  |  | 3,24 |  |  | ${ }_{3.84}^{4.98}$ | ${ }_{4}^{5} / 22$ |  | ${ }_{4.63}^{6,46}$ |  | $\underbrace{6.99}_{4.97}$ | 4.41 |  |  | ${ }_{1.95}^{2,700}$ | ${ }_{1.52}^{206}$ | ${ }_{1}^{1.45}$ |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }^{1990}$ | ${ }_{6.966}^{1176}$ | ${ }_{1.6}^{2,6}$ | ${ }_{1.83}^{2,19}$ | ${ }_{1.95}^{1,5}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| ${ }_{2002}^{2002}$ |  | 2.008 | ${ }_{3,15}^{2.55}$ |  |  |  |  |  |  |  | ${ }_{1.15}^{8.24}$ | 9,065 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  | 2.6 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  | -1, | ${ }_{88,66}^{1315}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 2011 |  | 5.28 | 5.82 C |  |  |  |  |  |  |  |  | 1.192 |  |  |  | 7.59 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |

Table 33. Comparison of estimates of recruitment (in millions) from the 2015 assessment model and the author's preferred model (Model C).

| year | 2015AMO | Model C | year | 2015AMO | Model C |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1949 | 59.6776 | 55.50094 | 1981 | 76.5356 | 134.3166 |
| 1950 | 59.8205 | 55.64543 | 1982 | 39.3139 | 90.73108 |
| 1951 | 60.1639 | 55.99151 | 1983 | 275.663 | 345.1917 |
| 1952 | 60.7919 | 56.62214 | 1984 | 266.635 | 321.7581 |
| 1953 | 61.8298 | 57.66209 | 1985 | 673.123 | 505.7285 |
| 1954 | 63.4703 | 59.29945 | 1986 | 517.949 | 466.2398 |
| 1955 | 66.0213 | 61.84307 | 1987 | 485.609 | 451.0147 |
| 1956 | 70.0018 | 65.79869 | 1988 | 444.015 | 439.7472 |
| 1957 | 76.3484 | 72.11052 | 1989 | 168.656 | 190.8714 |
| 1958 | 86.9194 | 82.64877 | 1990 | 70.9547 | 73.67769 |
| 1959 | 105.917 | 101.6972 | 1991 | 40.7613 | 42.89692 |
| 1960 | 144.847 | 141.2456 | 1992 | 30.7408 | 32.61264 |
| 1961 | 243.604 | 242.8879 | 1993 | 27.7367 | 30.2713 |
| 1962 | 534.886 | 537.8609 | 1994 | 31.3207 | 37.95875 |
| 1963 | 1244.52 | 1177.443 | 1995 | 41.6183 | 50.5266 |
| 1964 | 1930.88 | 1614.854 | 1996 | 46.1383 | 51.67117 |
| 1965 | 1929.26 | 1449.538 | 1997 | 113.808 | 127.6255 |
| 1966 | 1545.71 | 1119.122 | 1998 | 46.0495 | 52.34728 |
| 1967 | 1214.54 | 914.795 | 1999 | 140.552 | 152.6885 |
| 1968 | 1015.76 | 862.8147 | 2000 | 84.9866 | 90.76738 |
| 1969 | 926.124 | 946.3382 | 2001 | 279.151 | 276.5523 |
| 1970 | 879.663 | 1044.716 | 2002 | 108.797 | 104.9517 |
| 1971 | 737.391 | 887.8475 | 2003 | 185.039 | 209.3066 |
| 1972 | 572.562 | 653.799 | 2004 | 306.444 | 322.0478 |
| 1973 | 458.562 | 402.4215 | 2005 | 87.258 | 93.97229 |
| 1974 | 299.761 | 303.081 | 2006 | 70.8674 | 72.47198 |
| 1975 | 376.505 | 606.3152 | 2007 | 52.9206 | 48.53087 |
| 1976 | 1113.94 | 1093.567 | 2008 | 60.9981 | 60.50948 |
| 1977 | 829.217 | 863.9371 | 2009 | 354.632 | 395.1637 |
| 1978 | 381.131 | 441.598 | 2010 | 422.936 | 492.0597 |
| 1979 | 126.068 | 175.2126 | 2011 | 251.061 | 286.7756 |
| 1980 | 57.8529 | 93.14897 | 2012 | 52.203 | 49.61038 |
|  |  |  | 2013 | 115.803 | 124.1139 |
|  |  |  | 2014 | 124.004 | 99.47437 |
|  |  |  | 2015 | 80.7077 | 69.66514 |
|  |  |  | 2016 | -- | 120.013 |

Table 34. Comparison of exploitation rates (i.e., catch divided by biomass) from the 2015 assessment model and the author's preferred model (Model C).

| year | 2015AMO | Model C | year | 2015AMO | Model C |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1949 | 0.002 | 0.003 | 1981 | 0.075 | 0.070 |
| 1950 | 0.005 | 0.005 | 1982 | 0.041 | 0.035 |
| 1951 | 0.009 | 0.009 | 1983 | 0.023 | 0.017 |
| 1952 | 0.015 | 0.013 | 1984 | 0.050 | 0.033 |
| 1953 | 0.023 | 0.016 | 1985 | 0.018 | 0.019 |
| 1954 | 0.027 | 0.020 | 1986 | 0.022 | 0.027 |
| 1955 | 0.029 | 0.022 | 1987 | 0.040 | 0.042 |
| 1956 | 0.030 | 0.023 | 1988 | 0.058 | 0.052 |
| 1957 | 0.031 | 0.023 | 1989 | 0.134 | 0.117 |
| 1958 | 0.031 | 0.023 | 1990 | 0.211 | 0.197 |
| 1959 | 0.031 | 0.023 | 1991 | 0.175 | 0.171 |
| 1960 | 0.030 | 0.022 | 1992 | 0.208 | 0.208 |
| 1961 | 0.029 | 0.022 | 1993 | 0.155 | 0.153 |
| 1962 | 0.026 | 0.021 | 1994 | 0.121 | 0.118 |
| 1963 | 0.021 | 0.018 | 1995 | 0.114 | 0.110 |
| 1964 | 0.018 | 0.016 | 1996 | 0.077 | 0.073 |
| 1965 | 0.027 | 0.024 | 1997 | 0.052 | 0.047 |
| 1966 | 0.027 | 0.024 | 1998 | 0.039 | 0.037 |
| 1967 | 0.064 | 0.059 | 1999 | 0.020 | 0.019 |
| 1968 | 0.066 | 0.064 | 2000 | 0.020 | 0.018 |
| 1969 | 0.082 | 0.082 | 2001 | 0.026 | 0.023 |
| 1970 | 0.076 | 0.077 | 2002 | 0.017 | 0.016 |
| 1971 | 0.067 | 0.066 | 2003 | 0.011 | 0.011 |
| 1972 | 0.061 | 0.060 | 2004 | 0.011 | 0.011 |
| 1973 | 0.063 | 0.065 | 2005 | 0.019 | 0.018 |
| 1974 | 0.086 | 0.084 | 2006 | 0.027 | 0.025 |
| 1975 | 0.082 | 0.074 | 2007 | 0.030 | 0.027 |
| 1976 | 0.135 | 0.118 | 2008 | 0.022 | 0.020 |
| 1977 | 0.196 | 0.172 | 2009 | 0.018 | 0.017 |
| 1978 | 0.163 | 0.159 | 2010 | 0.009 | 0.009 |
| 1979 | 0.210 | 0.227 | 2011 | 0.010 | 0.010 |
| 1980 | 0.180 | 0.160 | 2012 | 0.007 | 0.006 |
|  |  |  | 2013 | 0.020 | 0.018 |
|  |  |  | 2014 | 0.069 | 0.060 |
|  |  |  | 2015 | -- | 0.081772 |

Table 35. OFL and ABC values for the models considered here. These values are presented only to illustrate the effect of incremental changes in the data used for the assessment on the OFL and ABC. The models highlighted in blue are based on data through 2014/15 (including the 2015 NMFS EBS trawl survey), while the others are based on data through 2015/16 (including the 2016 survey). Results from the author's preferred model (Model C) are highlighted in yellow.

| Model | Snow Crab <br> Fofl | Efffective <br> Snow Crab <br> F | Average <br> Recruitment | B | Fmsy | Bmsy | B/Bmsy | OFL | ABC <br> P-star | ABC <br> $(20 \%$ buffer) |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2015 Model | 1.32 | 0.049 | 179.37 | 53.70 | 0.58 | 26.79 | 2.00 | 27.19 | 27.15 | 21.75 |
| 2015AMR | 1.32 | 0.051 | 176.78 | 51.41 | 0.64 | 25.68 | 2.00 | 27.27 | 27.23 | 21.82 |
| 2015AMN | 1.32 | 0.044 | 193.44 | 63.85 | 0.56 | 29.42 | 2.17 | 30.96 | 30.91 | 24.77 |
| 2015AM | 1.24 | 0.030 | 183.46 | 48.07 | 0.59 | 26.68 | 1.80 | 23.79 | 23.75 | 19.03 |
| Model A | - | -- | - | - | - | - | - | - | -- | -- |
| Model B | 1.24 | 0.092 | 182.17 | 45.32 | 0.79 | 25.64 | 1.77 | 25.60 | 25.56 | 20.48 |
| Model C | 1.24 | 0.092 | 182.27 | 45.34 | 0.79 | 25.65 | 1.77 | 25.61 | 25.57 | 20.49 |
| Model D | 1.24 | 0.111 | 168.84 | 39.06 | 0.09 | 22.85 | 1.71 | 25.79 | 25.75 | 20.63 |
| Model E | 1.24 | 0.097 | 174.24 | 42.19 | 0.44 | 23.06 | 1.83 | 27.36 | 27.31 | 21.89 |
| Model F | 1.24 | 0.070 | 163.57 | 39.52 | 0.96 | 22.41 | 1.76 | 21.83 | 21.79 | 17.46 |
| Model G | 1.24 | 0.061 | 171.74 | 43.26 | 1.02 | 23.70 | 1.83 | 24.55 | 24.51 | 19.64 |

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case

- 2015AMO
- Model C








size (mm CW)

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## fits to mature survey biomass



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## Legal male biomass



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Figure 65. Fits to bycatch male size compositions from the snow crab fishery for the 2015 assessment (2015AMO) and the author's preferred model (Model C). Bars: observed; lines: predicted.


Figure 66. Fits to bycatch female size compositions from the snow crab fishery for the 2015 assessment (2015AMO) and the author's preferred model (Model C). Bars: observed; lines: predicted.


Figure 67. Pearson's residuals for fits to bycatch size compositions from the snow crab fishery for the 2015 assessment (2015AMO) and the author's preferred model (Model C).


Figure 68. Fits to bycatch male size compositions from the BBRKC fishery for the 2015 assessment (2015AMO) and the author's preferred model (Model C). Bars: observed; lines: predicted.


Figure 69. Fits to bycatch female size compositions from the BBRKC fishery for the 2015 assessment (2015AMO) and the author's preferred model (Model C). Bars: observed; lines: predicted.


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Figure 78. Marginal distributions for total catch (at-sea) size compositions from the directed Tanner crab fishery for the 2015 assessment (2015AMO) and the author's preferred model (Model C). Dotted lines: observed; solid lines: predicted.


Figure 79. Marginal distributions for bycatch (at-sea) size compositions from the snow crab fishery for the 2015 assessment (2015AMO) and the author's preferred model (Model C). Dotted lines: observed; solid lines: predicted.


Figure 80. Marginal distributions for bycatch (at-sea) size compositions from the BBRKC fishery for the 2015 assessment (2015AMO) and the author's preferred model (Model C). Dotted lines: observed; solid lines: predicted.


Figure 81. Marginal distributions for bycatch (at-sea) size compositions from the groundfish fisheries for the 2015 assessment (2015AMO) and the author's preferred model (Model C). Dotted lines: observed; solid lines: predicted.


Figure 82. Marginal distributions for size compositions from the NMFS EBS trawl survey for the 2015 assessment (2015AMO) and the author's preferred model (Model C). Dotted lines: observed; solid lines: predicted. Distributions are shown: top) by sex; bottom) by sex and maturity state.


Figure 83. Input and effective (McAllister-Ianelli) sample sizes for retained (upper) and total catch (lower) size compositions from the directed Tanner crab fishery for the 2015 assessment (2015AMO) and the author's preferred model (Model C). dotted lines: input; solid lines: effective.


Figure 84. Input and effective (McAllister-Ianelli) sample sizes for bycatch size compositions from the snow crab fishery (upper), BBRKC (middle), and groundfish fisheries (lower) for the 2015 assessment (2015AMO) and the author's preferred model (Model C). dotted lines: input; solid lines: effective.


Figure 85. Input and effective (McAllister-Ianelli) sample sizes for size compositions from the NMFS EBS trawl survey for the 2015 assessment (2015AMO) and the author's preferred model (Model C). dotted lines: input; solid lines: effective.

## Mature Biomass



Mature Biomass


Figure 86. Retrospective analysis for estimated mature biomass-at-mating from the author's preferred model (Model C). Model C was run for each case as though the assessment were conducted in the year indicated by the case name. Upper plot: full model time series; lower plot: recent time period.


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Figure 88. Retrospective analysis for fits to mature survey biomass from the author's preferred model (Model C). Observed: symbols and error bars; lines: predicted. Model C was run for each case as though the assessment were conducted in the year indicated by the case name. Upper plot: full model time series; lower plot: recent time period.


Figure 89. Retrospective analysis for fits to retained catch from the author's preferred model (Model C). Observed: symbols and error bars; lines: predicted. Model C was run for each case as though the assessment were conducted in the year indicated by the case name. Upper plot: full model time series; lower plot: recent time period.


Figure 90. The Fofs harvest control rule. For Tier 3 stocks such as EBS Tanner crab, F F $_{\text {mSY }}$ and $\mathrm{B}_{\text {MSY }}$ are based on spawning biomass per recruit proxies, where $\mathrm{F}_{\text {MSY }}=\mathrm{F}_{35 \%}, \mathrm{~B}_{\text {MSY }}=\mathrm{B} 35 \%$, and MMB at mating time is used as a surrogate for egg production/spawning biomass.


Figure 91 . The selectivity and retention curves for males in the directed fishery used to calculate the OFL.


Figure 92. Bycatch fishery selectivity curves used to calculate the OFL.


Figure 93. Distribution of OFL, illustrating the estimated p* ABC and 20\%-buffer ABC, for Model C.


Figure 94. Tier 3 quad plot for the author's preferred model, Model A (Dataset D). Colors indicate different time periods. Black: 1965-1979; blue: 1980-1989; cyan: 1990-1999; green: 2000-2009; red: 2010-2015.

# Appendix A: <br> Comparison of Models 2015AMO, 2015AMR, 2015AMN, 2015AM 

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## Introduction

This appendix summarizes the comparison of models 2015AMO, 2015AMR, 2015AMN, and 2015AM to document changes in progressing from the 2015 assessment model (2015AMO here) to the base model for the 2016 assessment (Model B). 2015AMR is a better-converged version of 2015AMO, with convergence evaluated using 200 runs with jittered initial parameter values. 2015AMN uses the 2015 data, but with the "new" cv's for mature survey biomass. 2015AM uses the 2016 data. Models 2015AMN and 2015AM were also evaluated for convergence using 200 runs with jittered initial parameter values.

## Evaluation

## Objective function values

Direct comparison among the four models on the basis of objective function value is not valid for drawing inferences because 2015AMO was not converged to the global minimum, uncertainties for mature survey biomass differ between 2015AMR and 2015AMN, and the 2016 data is added to 2015AM.

## Population processes

One effect of the "new" cv's was to lower estimates of natural mortality on mature crab during the "enhanced mortality" period (1980-1984). Estimated natural mortality rates were similar among the models outside the "enhanced mortality" time period, but differed for mature crab among models during this period (Fig. 1), with 2015AMO and 2015AMR exhibiting the highest rates for both mature males and females. The estimated rates on mature males during this period also increased slightly with the addition of the 2016 data. Otherwise, functions governing population processes (molt-to-maturity, growth) for all four models (Fig.s 2, 3).


Figure 1. Comparison of estimates of natural mortality from the four models.


Figure 2. Comparison of estimates of the size-specific probability of undergoing terminal molt-tomaturity from the four models.


Figure 3. Estimates of the mean post-molt size as a function of pre-molt size from the four models.

## Population quantities

Estimated trends in recruitment were quite similar for the four models (Fig.s 4, 5). The model estimates differed slightly when recruitment high for short periods, but oscillations were in-phase across models and all peaks occurred in the same year. At peaks in recruitment, the models with the "new" cv's for mature survey biomass (2015AMN, 2015AM) yielded slightly higher estimated recruitment compared with the models with the "old" cv's. Trends in population abundance were also similar for the four models, although some differences between models were discernible when the population reached its maximum abundance in the early 1970s, and again during the "enhanced mortality" period, 1980-1984. During the last 15 years, 2015AMN estimated abundance at somewhat higher levels than the other models, while 2015AMO and 2015AMR estimated abundance at the lowest levels (Fig. 6, 7). One effect of the "new" cv's was obviously to increase recruitment and population abundance estimates, while adding the 2016 data (2015AM) led to slightly decreased estimates of recruitment and abundance vis-àvis 2015AMN after 2008 (Fig.s 5, 7). Similar conclusions hold for mature biomass-at-mating (Fig.s 8, 9).


Figure 4. Estimated time series of recruitment from the four models.


Figure 5. Estimated time series of recruitment from the four models.


Figure 6. Estimated time series of population abundance from the four models.


Figure 7. Estimated time series of population abundance from the four models.


Figure 8. Estimated time series of mature biomass-at-mating from the four models.


Figure 9. Estimated time series of mature biomass-at-mating from the four models.

## Survey selectivity functions

The four models estimated almost identical survey selectivity curves and survey q's for both sexes during selectivity time period 1 (pre-1982), while in time period two the selectivity curves were similar across models but survey q's differed (with higher q's for the models using the "old" mature survey biomass cv's).


Figure 10. Comparison of estimated survey selectivity functions for the four models.

## Fishery selectivity functions

Estimated fishery retention functions were identical for the four models during the pre-1991 time period, as were those post-1990 for the 3 models using 2015 data (2015AMO, 2015AMR, 2015AMN; Fig. 11). The retention function estimated by 2015AM, using 2016 data, was left-shifted 5 mm toward smaller sizes. This may reflect accumulating evidence for shift to retention of somewhat smaller (but still legalsized) crab by industry since the fishery re-opened in 2013/14.

Estimated female selectivity in the directed fishery was essentially identical across the four models (Fig. 12). Estimated male selectivity curves before 1991 fell into two categories: those from 2015AMO and 2015AMN were left-shifted to smaller sizes by $\sim 10 \mathrm{~mm}$ relative to those from 2015AMR and 2015AM (Fig. 12). This result is rather curious, because it does not track with the change in calculated mature survey biomass cv's.

The estimated annual male selectivity curves in the directed fishery post-1990 (Fig. 13) are rather illuminating. For the years in which the directed fishery was prosecuted during this time period (1991/921996/97, 2005/06-2009/10, 2013/14-present), except 1996/97, the curves are very for all four models (only 2015AM estimates the 2015/16 curve, of course). In fact, they are practically identical in 2005/062009/10 and 2013/14-2014/15. However, they differ substantially for 1996/97, with curves from 2015AMO and 2015AMN substantially left-shifted relative to 2015AMR and 2015AM. This results in the pattern across models for the male selectivity curves pre-1991 (Fig. 12), or more likely the pattern for 1996/97 is a result of the pre-1991 pattern, because the size at $50 \%$-selected (z50) parameter in the logistic function used to describe pre-1991 male selectivity in the directed fishery is the average of the annual z50's for 1991/2-1996/97. It would be worthwhile to see how the model responds when 1996/97 is removed from the averaging time period.


Figure 11. Comparison of estimated retention functions in the directed fishery for the four models.


Figure 12. Comparison of estimated female selectivity functions and pre-1991 male total catch mortality selectivity functions in the directed fishery for the four models.


Figure 13. Comparison of estimated annual (post-1990) male total catch mortality selectivity functions in the directed fishery for the four models. The directed fishery was closed during 1997/98-2004/05 and 2010/11-2012/13. The mean selectivity function for 1991-present from
which annual deviations are taken is shown during the closures.

# Appendix B: <br> Comparison of Models 2015AM and Model B (the CPT's Base Model) 

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## Introduction

This appendix summarizes the comparison of models 2015AM and Model B to finish documenting changes in progressing from the 2015 assessment model (2015AMO) to the base model for the 2016 assessment (Model B). The progression for 2015AMO to 2015AM is discussed in Appendix A. The rationale for Model B, the CPT's base model, was discussed at the May 2016 CPT meeting. It includes a suite of changes that were evaluated in an incremental fashion by the author as part of that meeting. Model B embodies the following changes relative to the 2015AM model (which incorporates the "new" cv's for mature survey biomass and the 2016 data):

| Change | Description |
| :---: | :--- |
| A | start "current" recruitment estimation in 1975, instead of 1974 |
| B | normalize groundfish fishery size comps using original sample sizes, not input sample sizes |
| C | estimate log-scale fishing mortality/capture rate offsets for female crab |
| E | turn on fishing mortality/capture rate estimation for BBRKC |
| G | estimate probability of molt-to-maturity using logit-scale parameterization |
| I | enforce logistic selectivity =1 in largest size bin |
| J | use GMACS fishing mortality model |

The letter designations above refer to the suite of potential changes reviewed at the May meeting.

## Evaluation

## Objective function values

Direct comparison between the two models on the basis of objective function value is not valid for drawing inferences in a likelihood framework because model change B above essentially changes the bycatch size composition data for the groundfish fisheries. However, comparison of individual components of the objective function can give a sense of the size of relative fits to data, as well as the impact of penalty functions and assumed priors. In this sense, the objective function components are interpreted more as indicators of mean-squared error, in some sense.

In this regard, the size of the penalties applied in the objective function (Fig. 1) are quite similar for the two models, with perhaps the exception that the penalty on the estimate of natural mortality on mature males is larger for Model B than for 2015AM.Similarly, the size of the prior probabilities in the objective function are also similar (Fig. 2), although the prior for female catchability ( $q$ ) in the NMFS trawl survey is somewhat larger in Model B than in 2015AM.

Comparing the multinomial component values to the objective function from size fishery and survey compositions (Fig. 3), three components stand out with much larger values for Model B: the groundfish fisheries bycatch size compositions, the retained catch size compositions, and the total-catch size compositions in the directed fishery. The first of these is a non-starter, because the extended size compositions in the two models differ substantially in a number of years. It is a bit disappointing, however, that Model B does not fit the retained catch and ale total-catch size compositions better than 2015AM. This suggests there is room for improvement in the specification of selectivity and retention functions for the directed fishery, possibly in terms of allowing retention curves to vary annually as the selectivity curves are allowed to do (post-1991).

However, Model B fits the retained biomass and male total-catch biomass somewhat better than 2015AM (Fig. 4). Fitting catch biomass data at the expense of size composition data is generally considered a reasonable tradeoff, so the poorer fits to the retained catch and total-catch size composition data by Model B relative to 2015AM can be discounted in terms of overall model suitability.


Figure 1. Comparison of penalty components to the model objective function for the two models.


Figure 2. Comparison of prior probability components to the model objective function for the two models.


Figure 3. Comparison of multinomial components to the model objective function for the two models.
likelihood: catch biomass


Figure 4. Comparison of biomass components to the model objective function for the two models.

## Population processes

One effect of introducing the "new" cv's in 2015AMN was to lower estimates of natural mortality on mature crab during the "enhanced mortality" period (1980-1984; Appendix A). 2015AM, with the 2016 data, had slightly higher estimated rates than 2015AMN with only the 2015 data. Model B estimates very slightly larger rates, relative to 2015AM, for mature males outside the "enhanced mortality" period and slightly higher rates for mature males and females during the "enhanced mortality" period (Fig. 5).

The size-specific probability of undergoing the terminal molt to maturity is parameterized differently in the two models considered here: parameters (one for each size bin) are estimated on a ln-scale (with max 0 ) in 2015AM while they are estimated on a logit scale (no need to impose a maximum) in Model B. The resulting estimates, however, are remarkably similar (Fig. 6), except for the slight dip at large size for males in 2015AM (which does not seem credible, in any case).

Estimated patterns of mean growth-per-molt are almost identical for both models (Fig. 7). However, growth parameters in both models essentially hit their imposed upper bounds (as is also true of every other model considered in this assessment).


Figure 5. Comparison of estimates of natural mortality from the two models.


Figure 6. Comparison of estimates of the size-specific probability of undergoing terminal molt-tomaturity for from the four models.


Figure 7. Estimates of the mean post-molt size as a function of pre-molt size from the four models.

## Population quantities

While estimated recruitment differs somewhat in the mid-1960s between the two models (Fig. 8), the estimates are almost identical after 1980 and certainly after 2000 (Fig. 9). Similarly, the two models differ somewhat in estimated mature biomass-at-mating during the late 1960s and early 1970s (following the maturation of the recruits in the mid-1960s; Fig. 10), the estimated time series after 1980 are again very similar. During 2005-2012 (Fig. 11), estimates from 2015AM are slightly higher for males relative to Model B, but they are almost identical in 2014 and 2015. In contrast, estimates from 2015AM are slightly smaller relative to Model B during the past two years. Population abundance trends from the two models also converge to very similar values, after differing somewhat in before 1980 (Fig. 12).


Figure 8. Estimated time series of recruitment from the four models.


Figure 9. Estimated time series of recruitment from the four models.


Figure 10. Estimated time series of mature biomass-at-mating from the four models.


Figure 11. Estimated time series of mature biomass-at-mating from the two models.


Figure 12. Estimated population abundance time series from the two models.

## Survey selectivity functions

Estimated survey selectivity functions were nearly identical for the two models.


Figure 13. Comparison of estimated survey selectivity functions for the two models.

## Fishery selectivity functions

The estimated retention curves from the two models are nearly identical for the period before 1991, while the curve for 2015AM is shifted to slightly smaller sizes, relative to Model B, for the period after 1990 (Fig. 14). The estimated (bycatch) selectivity function for females in the directed fishery is substantially left-shifted to smaller sizes in Model B, relative to 2015AM (Fig. 15). This is a result of estimating a female-specific offset to male fishing mortality in the directed fishery (the size-specific fishing mortality rates are comparable). The estimated selectivity curves from the two models for males in the directed fishery should not be directly compared (despite doing so here) because they are different "beasts". The selectivity curve in 2015AM represents size-specific fishing mortality rates (retained + discard mortality: i.e., bycatch after handling mortality has been applied) while that in Model B represents size-specific capture rates (retained + bycatch before handling mortality is applied). Including handling mortality in the selectivity curve from Model B would right-shift it back toward larger sizes. Similar considerations hold for the annually-varying (1991-present) selectivity curves shown in Fig. 16, although it does not account for the really large difference between the curves in 1996. The left-shifted curve for 1996 from Model B is the result of: 1) a very small sample size for the male total-catch size composition in 1996 (with the consequence that mis-fitting this size composition has little impact on the overall objective function) and 2) the size at $50 \%$-selected ( $z 50$ ) parameter for the pre-1991 selectivity curve is the average of the $z_{50}$ 's for the 1991-1996 annually-varying selectivity functions. The small weight on fitting the 1996 size composition implies the $1996 z_{50}$ is essentially a free parameter driven by determining the $z_{50}$ for the pre-1991 selectivity curve that best minimizes the overall objective function, rather than by the size composition in 1996. The value of $z_{50}$ for the 1996 male total-catch appears to be extremely sensitive to other details of the model.

The estimated bycatch selectivity curves for males in the snow crab (Fig. 17), BBRKC (Fig. 18) and groundfish (Fig. 19) fisheries are very similar for the two models. The selectivity curves for females are substantially left-shifted to smaller sizes in Model B relative to 2015AM for two reasons: 1) female offsets to fully-selected male fishing mortality rates are estimated in Model B, but not in 2015AM; and 2) the selectivity curves are forced to equal 1 in the maximum model size bin in Model B but not in 2015AM (particularly important for the groundfish fisheries female bycatch selectivity curves).

The impact of estimating female offsets to fully-selected male fishing mortality rates in Model B vis-à-vis 2015AM is illustrated in Fig. 20, where fully-selected rates on females are identical to those estimated for males in the directed fishery in 2015AM (reaching a maximum value of $>4$ ) whereas the rates are much smaller for Model B.


Figure 14. Comparison of estimated retention functions in the directed fishery for the two models.


Figure 15. Comparison of estimated female bycatch selectivity and male selectivity prior to 1990 in the directed fishery for the two models.


Figure 16. Comparison of estimated annual selectivity functions in the directed fishery for the two models.


Figure 17. Comparison of estimated bycatch selectivity functions in the snow crab fishery for the two models.


Figure 18. Comparison of estimated bycatch selectivity functions in the BBRKC fishery for the two models.


Figure 19. Comparison of estimated bycatch selectivity functions in the groundfish fisheries for the two models.


Figure 20. Comparison of estimated mean selectivity functions in the directed fishery for the two models.

# Appendix C: <br> Comparison of Model B and Model C 

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## Introduction

This appendix summarizes the comparison of Models B and C from the 2016 Tanner crab assessment. Model C builds on Model B by eliminating the constraint imposed on bycatch F rates in the BBRKC fishery that required estimated F's to be above a minimum threshold value. Any F's that fell below this threshold were replaced by the minimum. This constraint was non-differentiable and may have complicated model convergence.

## Evaluation

Because Model C eliminated a non-differentiable constraint in the model, it would in almost any case have been preferred to Model B as a better model in terms of being consistent with AD Model Builder's minimization algorithms.

However, results for Model C were also almost identical to Model B, as indicated by very small differences in all objective function components (see below), so the constraint did not interfere with model minimization. The only "substantial" differences between the models were in some of the estimated bycatch capture rates in the BBRKC fishery:


Figure 1. Fully-selected fishery capture/mortality rates in the BBRKC fishery for Models B and C.
Consequently, there was no issue to adopting Model C as the preferred model over B.

## Objective function values



Figure 2. Differences for Model C vis-à-vis Model B (C-B) in penalty components to the model objective function.


Figure 3. Differences for Model C vis-à-vis Model B (C-B) in prior probability components to the model objective function.


Figure 4. Differences for Model C vis-à-vis Model B (C-B) in prior probability components to the model objective function.


Figure 5. Differences for Model C vis-à-vis Model B (C-B) in prior probability components to the model objective function.

## Population processes



Figure 6. Estimates of natural mortality for Models B and C.


Figure 7. Estimates of the size-specific probability of undergoing terminal molt-to-maturity for Models B and C.


Figure 8. Estimates of the mean post-molt size as a function of pre-molt size for Models B and C.

## Population quantities



Figure 9. Estimated time series of recruitment from Models B and C.


Figure 10. Estimated time series of mature biomass-at-mating from Models B and C.

# Appendix D: <br> Comparison of Models C, D, E, F, G 

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## Introduction

This appendix summarizes the comparison of Models C, D, E, F, and G from the 2016 Tanner crab assessment. Model D builds on Model C by adding two parameters, one for the snow crab fishery and one for the BBRKC fishery, to estimate fishery q's for these fisheries to convert effort (potlifts) to fishery capture rates. Model E builds on D by reducing penalties on F -devs with each estimation phase in the model convergence algorithm, then eliminating the penalties completely in the final estimation phase. Model F builds on Model D by incorporating lognormal likelihoods for catch data in all fisheries, and Model G does the same with Model E as its base (rather than Model D).

## Evaluation

Unfortunately, the (ln-scale) estimates for the fishery q parameters introduced in Model D were unreasonably small:

|  | Model D | Model E | Model F | Model G |
| :--- | :--- | :--- | :--- | :--- |
| BBRKC | -18.46 | -19.78 | -19.28 | -19.77 |
| snow crab fishery | -17.82 | -19.83 | -19.83 | -19.82 |

Table 36. Ln-scale estimates of fishery q's ( $\mathrm{F}=\mathrm{qE}$ ) for bycatch in the BBRKC and snow crab fisheries from Models D-G.
which resulted in essentially bycatch rates of 0 in the snow crab and BBRKC fisheries prior to 1992, when at-sea crab fishery observers first provided usable estimates of Tanner crab bycatch in those fisheries (Fig.s 1 and 2):


Figure 1. Fully-selected fishery capture/mortality rates in the BBRKC fishery for Models C-G.


Figure 2. Fully-selected fishery capture/mortality rates in the snow crab fishery for Models C-G.
The fishery q's in Model C are not estimated parameters, but instead are based on the ratio of mean(fishing capture rate)/mean(effort) over the period 1992-present in the two respective fisheries. This approach at least appears to give reasonable estimates of historical (pre-1992) max capture rates (see Appendix C). Thus, Model C was selected over Models D-G as the preferred model for this assessment.


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

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

