# An assessment for eastern Bering Sea snow crab 

Cody Szuwalski

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1. Stock: Eastern Bering Sea snow crab, Chionoecetes opilio.
2. Catches: trends and current levels

Retained catches increased from relatively low levels in the early 1980s (e.g. 11.85 kt during 1982) to historical highs in 1990s (retained catch during 1991, 1992, and 1998 were $143.02,104.68$, and 88.09 kt , respectively). The stock was declared overfished in 1999 at which time retained catches dropped to levels similar to the early 1980s (e.g. 11.46 kt ). Retained catches slowly increased after 1999 before dropping again in 2016. Total allowable catches were slashed with the collapse of the population in 2021 and the fishery was closed for the first time in 2022.

Discard mortality from the directed fishery is the next largest source of mortality after retained catch and approximately tracks the retained catch. The highest estimated discard mortality occurred during 1992 at 17.06 kt which was $16 \%$ of the retained catch during that year. There was no discard mortality in 2022 because there was no directed fishery.

## 3. Stock Biomass:

Observed mature male biomass (MMB) at the time of the survey increased from low levels in the early to mid-1980s to historical highs in the 1990s (observed MMB during 1990, 1991, and 1997 were 443.79, 466.61, and 326.75 kt , respectively). The stock was declared overfished in 1999 in response to the total mature biomass dropping below the 1999 minimum stock size threshold. MMB in that year decreased to 95.85 kt . Observed MMB slowly increased after 1999, and the stock was declared rebuilt in 2011 when estimated MMB at mating was above $\mathrm{B}_{35 \%}$. However, recently the observed MMB has declined to historical lows and the stock was declared overfished again in 2021. MMB at the time of the survey was 24.21 kt in 2023, the lowest value on record.

## 4. Recruitment

Estimated recruitment shifted from a period of high recruitment to a period of low recruitment in the mid1990s (corresponding with a late 1980s fertilization). A large year class recruited to the survey gear in the mid 2010s and was tracked until 2018 and 2019, but disappeared from the eastern Bering Sea shelf before reaching commercial size. Recent estimated recruitments were the lowest on record.
5. Management

Table 1: Historical status and catch specifications for snow crab (1,000t).

| Year | MSST | Biomass <br> $(M M B)$ | TAC | Retained <br> catch | Total <br> catch | OFL | ABC |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2015 / 2016$ | 75.8 | 91.6 | 18.4 | 18.4 | 21.4 | 83.1 | 62.3 |
| $2016 / 2017$ | 69.7 | 96.1 | 9.7 | 9.7 | 11 | 23.7 | 21.3 |
| $2017 / 2018$ | 71.4 | 99.6 | 8.6 | 8.6 | 10.5 | 28.4 | 22.7 |
| $2018 / 2019$ | 63 | 123.1 | 12.5 | 12.5 | 15.4 | 29.7 | 23.8 |
| $2019 / 2020$ | 56.8 | 167.3 | 15.4 | 15.4 | 20.8 | 54.9 | 43.9 |
| $2020 / 2021$ | 76.7 | 26.74 | 20.4 | 20.4 | 26.2 | 95.4 | 71.55 |
| $2021 / 2022$ | 91.6 | 41.3 | 2.5 | 2.5 | 3.6 | 7.5 | 5.6 |
| $2022 / 2023$ | 136.9 | 92.4 | 0.0 | 0.0 | 0.05 | 10.3 | 7.7 |
| $2023 / 2024$ |  | 69.2 |  |  |  | 0.31 | 0.25 |

Table 2: Historical status and catch specifications for snow crab (millions of lbs).

| Year | MSST | Biomass <br> $(M M B)$ | TAC | Retained <br> catch | Total <br> catch | OFL | ABC |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2015 / 2016$ | 167.11 | 201.94 | 40.57 | 40.57 | 47.18 | 183.2 | 137.35 |
| $2016 / 2017$ | 153.66 | 211.86 | 21.38 | 21.38 | 24.25 | 52.25 | 46.96 |
| $2017 / 2018$ | 157.41 | 219.58 | 18.96 | 18.96 | 23.15 | 62.61 | 50.04 |
| $2018 / 2019$ | 138.89 | 271.39 | 27.56 | 27.56 | 33.95 | 65.48 | 52.47 |
| $2019 / 2020$ | 125.22 | 368.83 | 33.95 | 33.95 | 45.86 | 121.03 | 96.78 |
| $2020 / 2021$ | 169.09 | 58.95 | 44.97 | 44.97 | 57.76 | 210.32 | 157.74 |
| $2021 / 2022$ | 201.94 | 91.05 | 5.51 | 5.51 | 7.94 | 16.53 | 12.35 |
| $2022 / 2023$ | 301.81 | 203.71 | 0 | 0 | 0.11 | 22.71 | 16.98 |
| $2023 / 2024$ |  | 152.56 |  |  |  | 0.68 | 0.55 |

6. Basis for the OFL

The OFL for 2023 from the author-preferred model (23.3a) was 0.31 kt fishing at $\mathrm{F}_{\text {OFL }}=0.05$ ( $17 \%$ of the calculated $\mathrm{F}_{M S Y}$ proxy, 0.29). The projected ratio of MMB at the time of mating in 2023 to the $\mathrm{B}_{M S Y}$ proxy is 0.26 under no directed fishing and 0.25 fishing at the $\mathrm{F}_{\text {OFL }}$. This OFL was calculated from the model-derived estimates of MMB, used the estimated natural mortality as a proxy for $\mathrm{F}_{M S Y}$, and the average MMB from 1982-2022 as the target biomass. This is a departure from previous assessments that have used SPR-based reference points. Rationale for this shift is included within.

Table 3: Metrics used in designation of status and OFL (1,000 t). Status represents the status of the population after the completed fishing year and is used for overfished declarations. Proj_Status represents the projected fishery status after the coming fishery removes the OFL and is used in the harvest control rule. 'Years' indicates the year range used in the calculation of the proxy for BMSY. ' M ' is the natural mortality for mature male crab. (continued below)

| Year | Tier | BMSY | MMB | Status | Proj__MMB | Proj_Status | FOFL |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2023 / 2024$ | 3 b | 273.8 | 92.4 | 0.34 | 69.18 | 0.25 | 0.05 |


| Years | M |
| :---: | :---: |
| $1982-2022$ | 0.29 |

7. Basis for ABC

The ABC for the chosen model was 0.25 kt , calculated by subtracting a $20 \%$ buffer from the OFL as recommended by the SSC.

## A. Summary of Major Changes

1. Management:

The eastern Bering Sea snow crab population was declared overfished in October 2021 and the directed fishery was closed for the 2022 season.
2. Input data:

Data added to the assessment included: 2023 eastern Bering Sea survey biomass and length composition data and non-directed discard length frequency and discard biomass from 2022.

## 3. Assessment methodology:

Management quantities were derived from maximum likelihood estimates of model parameters in a size-based, integrated assessment method using GMACS. Retrospective analyses and jittering analyses were performed for a selection of models. An application of both tier 3 and tier 4 methodologies for calculating the OFL using both assessment model output and observed survey data are included here. Assessment changes explored in this document include specifying the probability of having undergone terminal molt and parameterizing survey selectivity as a non-parametric curve (rather than logistic) informed by the inferred selectivity from the BSFRF data as priors.

## 4. Assessment results

The updated estimate from the author-preferred model of MMB on February 15, 2022 was 92.39 kt which placed the stock at $34 \%$ of the $\mathrm{B}_{M S Y}$ proxy. Projected MMB on February 15,2023 from this year's author preferred model is 69.18 kt after fishing at the OFL, which would place the stock at $25 \%$ of the $\mathrm{B}_{M S Y}$ proxy.

## B. Comments, responses and assessment summary

## SSC and CPT comments + author responses

SSC comment: The SSC agrees with the CPT to bring forward the status quo model and a Tier 4 random effects model. The SSC recommends that some variant of the simpler model be brought forward at the assessment author's discretion but does not want to be prescriptive about the configuration of the model. Any model that is brought forward should show adequate convergence properties, and the incremental effect of each change from the status quo model should be evaluated.

Convergence issues with the simpler model were solved by adjusting the way the priors on selectivity were specified. An intermediate model in which the BSFRF data are still treated as an additional survey, but the probability of having undergone terminal molt is specified based on survey observations is presented. If survey selectivity and the probability of undergoing terminal molt are not concurrently updated to reflect the best available information on the biology of snow crab (i.e. changing survey selectivity to non-parametric and using the observed probability of having undergone terminal molt), large changes in the scale of the stock occur.

SSC comment: Ideally it would be preferable to directly incorporate an environmental covariate in the assessment to inform temporal changes in natural mortality, but it must be acknowledged that such covariates are seldom available. The SSC recommends that a conservative approach be used for incorporating time-varying M. For example, it may be reasonable to fit an initial model with time-varying $M$ in all years, and use that model to identify a smaller set of years where there is a strong signal to model with time-varying M.

No additional models incorporating time-varying natural mortality are presented in this document, but will be included in upcoming January and May documents.

SSC comment: When the external estimation approach is taken, estimates of uncertainty should be included in the assessment in addition to the point estimates, so that the uncertainty in external estimation is propagated through to assessment results. The SSC supports exploration of models with pre-specified growth parameters, as well as using BSFRF survey data as a prior for survey selectivity/catchability
The growth parameters associated with the molt increment model are estimated in all models presented here, but the variance in molt increment is specified because of convergence issues. This is potentially an important point for future exploration given the larger number of large males estimated in the fishery than observed in some years.

SSC comment: Field biologists and crab life history experts should be consulted to understand the reliability of (the molt probability) data for stock assessment. Rather than adding the raw survey estimates to the model, an initial analysis in a GLM modeling framework, which treats years as random effects, should be considered. This approach could provide smoother estimates, accommodate differing sample sizes by year and length, and deal appropriately with years in which data are missing.
The data were smoothed using a GAM and the average over time was input for years in which data do not exist. This accomplishes roughly what would occur if a GLM were applied to the data without temporal autocorrelation. Temporal autocorrelation may change the estimates for years without data and this is a potential avenue for future exploration.

SSC comment: The SSC supports the CPT recommendation for a model that includes males and females, does not model groundfish bycatch, but estimates selectivity and recruitment by sex, and includes a penalty of how much recruitment can vary between the sexes. A more strategic approach to data weighting could also be considered, such that female data would receive less weight than male data in model fitting. The primary consideration should be that estimation of female parameters should not have a large effect on male parameter estimation.
The models presented include both females and males, but female selectivities are estimated separately. Recruitments are linked insofar as a single recruitment is estimated each year and an additional parameter is estimated in each year to divide that recruitment between the sexes. However, the penalty on this parameter
forcing it towards a $50 / 50$ sex ratio is small, so recruitment can vary nearly independently between the sexes. Data weighting is an on-going issue and will hopefully be addressed after a model structure is settled upon.
SSC comment: F35\% fishing mortality rate no longer results in a meaningful conservation constraint on the fishery for snow crab. To evaluate a potential alternative to the status quo, the SSC recommends that OFL and ABC estimates be provided for a modified Tier 3 approach for each model carried forward. This approach has the following characteristics: the OFL is calculated by replacing $F 35 \%$ in the Tier 3 harvest control rule by the model estimate of natural mortality. Biomass reference levels and status determination would be calculated using MMB as usual for Tier 3. The SSC requests evaluation of this approach by the assessment author and the CPT.

A tier 3 and tier 4 treatment are presented as written in the crab specs. That is, the tier 3 uses $\mathrm{B}_{35 \%}$ and $\mathrm{F}_{35 \%}$ as reference points and tier 4 uses the estimated M and the average of MMB from 1982-2022 as reference points.

## Assessment summary

Six assessment models are presented here:

- 22.1 - Last year's accepted model
- 23.1 - Last year's model fit to this year's data
- $23.2-23.1+$ specifying the probability of having undergone terminal molt based on survey data
- $23.3-23.2+$ specifying survey selectivity based on the BSFRF data
- $23.3 \mathrm{a}-23.3+$ estimating survey selectivity with the BSFRF data as priors
- $23.3 \mathrm{~b}-23.3 \mathrm{a}+$ loosening the prior on natural mortality

An overview of the model assumptions can be seen in Table 7. All models converged and produced reasonable jittering and retrospective analyses. Including the probability of having undergone terminal molt as data is recommended because these data reflect the best available science on the biology of snow crab. Incorporating the BSFRF data as priors seems like a more direct and interpretable way of using the BSFRF data to inform selectivity.
High estimated fishing mortality has been flagged as a concern in previous assessments. A more thorough exploration of the interplay between fishing mortality, selectivity, and size composition data in GMACS showed that, even when fully-selected fishing mortalities are very high, the realized exploitation rate of large males is not close to the $100 \%$ implied by the fishing mortality.
Given these observations, the author-preferred model is 23.3a. However, using this model coupled with the status quo tier 3 reference points and harvest control rule would apply an $96 \%$ exploitation rate to crab 102-105 mm carapace width under the $\mathrm{F}_{\text {OFL }}$, even with the $>101 \mathrm{~mm}$ carapace width male abundances at all time lows. Such a high exploitation rate associated with the OFL may require additional consideration in a management context.

Application of a tier 4 harvest control rule results in a much smaller $\mathrm{F}_{\text {OFL }}(0.05)$ and OFL $(0.31 \mathrm{t})$, which seem more reasonable in the current circumstances. However, the forward looking implications of adopting a tier 4 harvest control rule that still uses morphometrically mature male biomass as the currency of management and a model that estimates such low fishery selectivity in the size bins where the majority of the exploitable biomass exists is unclear. Consequently, the author recommends tier 4 HCRs be used for the 2023 assessment and suggests considering a currency of management that better reflects the impact of the fishery on the exploited fraction of the population in the future. Guidance on analyses that would be satisfactory rationale for changing the currency of management would be useful.
Analysis was also included on CPUE dynamics to provide an alternate way of quantifying the impact of the fishery on the stock. It showed that the average decline in CPUE was $11(\mathrm{sd}=11)$ crab per pot per week and declines were significantly related to the amount of catch removed from an area.

## C. Introduction

Studies and data relevant to key population and fishery processes are discussed below to provide background for the modeling choices made in this assessment. A model description is available on the github repository for GMACS and the files needed to reproduce these assessments also have a github repo, both of which are linked at the end of this document.

## Distribution

Snow crab (Chionoecetes opilio) are distributed on the continental shelf of the Bering Sea, Chukchi Sea, and in the western Atlantic Ocean as far south as Maine. In the Bering Sea, snow crab are distributed widely over the shelf and are common at depths less than $\sim 200$ meters (Figure 1 for distribution over time and Figure 2 for 2023 distribution of all males). Smaller crabs tend to occupy more inshore northern regions (Figure 3 \& Figure 4) and mature crabs occupy deeper areas to the south of the juveniles (Figure 5 \& Figure 6; Zheng et al. 2001). The eastern Bering Sea population within U.S. waters is managed as a single stock; however, the distribution of the population may extend into Russian waters to an unknown degree.

## Natural Mortality

Relatively few targeted studies exist to determine natural mortality for snow crab in the Bering Sea. Nevissi, et al. (1995) used radiometric techniques to estimate shell age from last molt (Figure 7). The total sample size was 21 male crabs (a combination of Tanner and snow crab) from a collection of 105 male crabs from various hauls in the 1992 National Marine Fishery Service (NMFS) Bering Sea survey. Representative samples for the 5 shell condition categories were collected from the available crab. Shell condition 5 crab (SC5 = very, very old shell) had a maximum age of 6.85 years (s.d. $0.58,95 \%$ CI approximately 5.69 to 8.01 years; carapace width of 110 mm ). The average age of 6 crabs with SC 4 (very old shell) and SC5, was 4.95 years (range: 2.70 to 6.85 years). Given the small sample size, this maximum age may not represent the $1.5 \%$ percentile of the population that is approximately equivalent to Hoenig's method (1983). Tag recovery evidence from eastern Canada revealed observed maximum ages in exploited populations of 17-19 years (Nevissi, et al. 1995, Sainte-Marie 2002). A maximum time at large of 11 years for tag returns of terminally molted mature male snow crab in the North Atlantic has been recorded since tagging started about 1993 (Fonseca, et al. 2008). Fonseca, et al. (2008) estimated a maximum age of 7.8 years post terminal molt using data on dactal wear.

In recent years, the mean for the prior for natural mortality used in the eastern Bering Sea snow crab assessment was based on the assumption that longevity would be at least 20 years in an unfished population of snow crab, informed by the studies above. Under negative exponential depletion, the 99th percentile corresponding to age 20 of an unexploited population corresponds to a natural mortality rate of 0.23 . Using Hoenig's (1983) method a natural mortality equal to 0.23 corresponds to a maximum age of 18 years.

In contrast to the implied natural mortalities from the methodology used above, Murphy et al. (2018) estimated time-varying natural mortality for eastern Bering Sea snow crab with a mean of 0.49 for females and 0.36 for males (based on the output of state-space models fit to NMFS survey data). Further, natural mortality estimates produced from empirical analyses by Then et al. (2015) and Hamel (2015) using similar assumed maximum ages as the methodology above produced natural mortalities larger than 0.23 (Table 5). Then et al. (2015) compared several major empirical estimation methods for M (including Hoenig's method) with an updated data set and found that maximum age was the best available predictor. A maximum age of 20 years corresponded to an M of $\sim 0.315$ in Then et al.'s analysis. Hamel (2015) developed priors in a similar manner to Then et al., but forced the regression of observed natural mortality onto maximum age through the intercept, which resulted in an M of $\sim 0.27$ for an assumed maximum age of 20 years.

Table 5: Empirical estimates of natural mortality for a range of methods over a range of assumed maximum ages (column header).

|  | 23 | 20 | 17 |
| :---: | :---: | :---: | :---: |
| Then | 0.277 | 0.315 | 0.365 |
| Hoenig (1983) | 0.19 | 0.212 | 0.257 |
| Hoenig (2013) | 0.194 | 0.223 | 0.261 |
| Hamel | 0.235 | 0.271 | 0.318 |

In addition to the results of empirical estimates of $M$ from updated methodologies and state-space modeling by Murphy et al. (2018), inspection of the survey data suggests that natural mortality for mature individuals is higher than assumed. A fraction of the mature population (which are assumed not to grow, given evidence for a terminal molt) are not selected in the fishery (e.g. sizes $50-80 \mathrm{~mm}$; Figure 8). Consequently, all mortality observed is 'natural'. The collapse in recruitment in the 1990s can be used as an instrument to understand natural mortality for mature individuals. The last large recruitment enters these size classes in the mid- to late-1990s and numbers of crab in these size classes return to low levels in less than 5 years.

The median value of the priors used in this assessment are set equal to values resulting from assuming a maximum age of 20 years and applying Hamel's methodology (0.271). A standard error of 0.0054 was used for initial priors and was estimated using the $95 \%$ CI of +-1.7 years on maximum age estimates from dactal wear and tag return analysis in Fonseca, et al. (2008). Mortality events in 2018 and 2019 are estimated as additional mortality parameters applied by sex and maturity state to allow the model to fit recent population trends.

## Maturity

Maturity of females collected during the NMFS summer survey was determined by the shape of the abdomen, by the presence of brooded eggs, or egg remnants. Maturity for males was determined by chela height measurements, which were available most years starting from the 1989 survey (Otto 1998; Figure 9). Mature male biomass referenced throughout this document refers to a morphometrically mature male (i.e. largeclawed). A maturity curve for males was estimated using the average fraction mature based on chela height data and applied to years of survey data to estimate mature survey numbers that do not have chela height data available. The separation of mature and immature males by chela height may not be adequately refined given the current measurement to the nearest millimeter. Chela height measured to the nearest tenth of a millimeter by Canadian researchers on North Atlantic snow crab showed a clear break in chela height at small and large widths and fewer mature animals at small widths than the Bering Sea data measured to the nearest millimeter. Measurements taken in 2004-2005 on Bering Sea snow crab chela to the nearest tenth of a millimeter show a similar break in chela height to the Canadian data (Rugolo et al. 2005).

Bering Sea male snow crab appear to have a terminal molt to maturity based on hormone level data and findings from molt stage analysis via setagenesis (Tamone et al. 2005). The models presented here assume a terminal molt for both males and females, which is supported by research on populations in the Bering Sea and the Atlantic Ocean (e.g. Dawe, et al. 1991). Mature male snow crab that do not molt may be important in reproduction. Paul et al. (1995) found that old shell mature male Tanner crab out-competed new shell crab of the same size in breeding in a laboratory study. Recently molted males did not breed even with no competition and may not breed until after $\sim 100$ days from molting (Paul et al. 1995). Sainte-Marie et al. (2002) stated that only old shell males take part in mating for North Atlantic snow crab.

## Mating ratio and reproductive success

Bering Sea snow crabs are managed using morphometrically mature male biomass (MMB) as a proxy for reproductive potential. MMB is used as the currency for management because the fishery only retains large
male crabs, which are nearly $100 \%$ mature. Male snow crabs are sperm conservers, using less than $4 \%$ of their sperm at each mating and females also can mate with more than one male. The amount of stored sperm and clutch fullness varies with sex ratio (Sainte-Marie 2002). If mating with only one male is inadequate to fertilize a full clutch, then females would need to mate with more than one male, necessitating a sex ratio closer to 1:1 in the mature population, than if one male is assumed to be able to adequately fertilize multiple females. Although mature male biomass is currently the currency of management, some aspect of female reproduction is likely also an important indicator of reproductive potential of the stock.

Clutch fullness is recorded for the females measured in the survey (Figure 10). However, quantifying the reproductive potential of the female population from survey data can be difficult. For example, full clutches of unfertilized eggs may be extruded and appear normal to visual examination, and may be retained for several weeks or months by snow crab. Resorption of eggs may occur if not all eggs are extruded resulting in less than a full clutch. Female snow crab at the time of the survey may have a full clutch of eggs that are unfertilized, resulting in overestimation of reproductive potential. Barren females may be a more obvious indication of low reproductive potential and increased in the early 1990s, decreased in the mid-1990s, then increased again in the late 1990s. The highest levels of barren females coincided with periods of high fishing mortality, but even then the proportion of barren females was low (Figure 11). Biennial spawning is another confounding factor in determining the reproductive potential of snow crab. Laboratory analyses showed that female snow crab collected in waters colder than 1.5 degrees C from the Bering Sea spawn only every two years.

Further complicating the process of quantifying reproductive capacity, clutch fullness and fraction of unmated females may not account for the fraction of females that may have unfertilized eggs, since these cannot be detected by eye at the time of the survey. The fraction of barren females observed in the survey may not be an accurate measure of fertilization success because females may retain unfertilized eggs for months after extrusion. To examine this hypothesis, NMFS personnel sampled mature females from the Bering Sea in winter and held them in tanks until their eggs hatched in March of the same year (Rugolo et al. 2005). All females then extruded a new clutch of eggs in the absence of males. All eggs were retained until the crabs were euthanized near the end of August. Approximately $20 \%$ of the females had full clutches of unfertilized eggs. The unfertilized eggs could not be distinguished from fertilized eggs by visual inspection at the time they were euthanized. Indices of fertilized females based on the visual inspection method of assessing clutch fullness and percent unmated females may overestimate fertilized females.

## Growth

Several studies are available to estimate the growth per molt of male and female snow crab in the Bering Sea (Table 9). These studies include:

1. Transit study (2003); 14 crab
2. Cooperative seasonality study; 6 crab
3. Dutch harbor holding study; 9 crab
4. NMFS Kodiak holding study held less than 30 days; 6 crab
5. NMFS Kodiak holding study 2016; 5 crab
6. NMFS Kodiak holding study 2017; 70 crab.
7. BSFRF/NMFS holding study 2018; 4 crab.

In the "Transit study", pre- and post-molt measurements of 14 male crabs that molted soon after being captured were collected. The crabs were measured when shells were still soft because all died after molting, so measurements may be underestimates of post-molt width (L. Rugolo, pers. com.). The holding studies include only data for crab held less than 30 days because growth of crabs held until the next spring's molting was much lower. Crab missing more than two limbs were excluded due to other studies showing lower growth. Crab from the seasonal study were excluded that were measured less than 3 days after molting due to difficulty in measuring soft crab accurately (L. Rugolo, pers. comm.). In general, growth of snow crab in the Bering Sea appears to be greater than growth of some North Atlantic snow crab stocks (Sainte-Marie
1995). Crabs in their first few years of life may molt more than once per year, however, the smallest crabs included in the model are approximately 4 years old and would be expected to molt annually.

## Management history

## ADFG harvest strategy

Before the year 2000, the Guideline Harvest Level (GHL) for retained crab only was a $58 \%$ harvest rate of the number of male crab over 101 mm CW estimated from the survey. The minimum legal size limit for snow crab is 78 mm , however, the snow crab market generally only accepts crab greater than 101 mm . In 2000, due to the decline in abundance and the declaration of the stock as overfished, the harvest rate for calculation of the GHL was reduced to $20 \%$ of male crab over 101 mm . After 2000, a rebuilding strategy was developed based on simulations by Zheng et al. (2002) using survey biomass estimates. The realized retained catch typically exceeded the GHL historically, resulting in exploitation rates for the retained catch on males $>101 \mathrm{~mm}$ ranging from about $10 \%$ to $80 \%$.

The Alaska Department of Fish and Game (ADFG) harvest strategy since 2000 sets harvest rate based on estimated mature biomass. The harvest rate scales with the status of the population relative to a proxy for $\mathrm{B}_{M S Y}$, which is calculated as the average total mature biomass at the time of the survey from 1983 to 1997 and MSST is one half the $\mathrm{B}_{M S Y}$ proxy. The harvest rate begins at 0.10 when total mature biomass exceeds $50 \%$ MSST ( 230 million lbs) and increases linearly to 0.225 when biomass is equal to or greater than the $\mathrm{B}_{M S Y}$ proxy (Zheng et al. 2002).

$$
u= \begin{cases}\text { Bycatch } & \text { if } \frac{T M B}{T M B_{M S Y}} \leq 0.25  \tag{1}\\ \frac{0.225\left(\frac{T M B}{T M B_{M S Y}}-\alpha\right)}{1-\alpha} & \text { if } 0.25<\frac{T M B}{T M B_{M S Y}}<1 \\ 0.225 & \text { ifTMB }>T M B_{M S Y}\end{cases}
$$

Where $T M B$ is the total mature biomass and $\mathrm{TMB}_{B M S Y}$ is the $T M B$ associated with maximum sustainable yield. The maximum retained catch is set as the product of the exploitation rate, $u$, calculated from the above control rule and survey mature male biomass. If the retained catch in numbers is greater than $58 \%$ of the estimated number of new shell crabs greater than 101 mm plus $25 \%$ of the old shell crab greater than 101 mm , the catch is capped at $58 \%$.

## History of BMSY

Prior to adoption of Amendment 24, $\mathrm{B}_{M S Y}$ was defined as the average total mature biomass (males and females) estimated from the survey for the years 1983 to 1997 ( 921.6 million lbs; NPFMC 1998) and MSST was defined as $50 \%$ of $\mathrm{B}_{M S Y}$. Currently, the biological reference point for biomass is calculated using a spawning biomass per recruit proxy, $\mathrm{B}_{35 \%}$ (Clark, 1993). $\mathrm{B}_{35 \%}$ is the biomass at which spawning biomass per recruit is $35 \%$ of unfished levels and has been shown to provide close to maximum sustainable yield for a range of stock productivities (Clark, 1993). Consequently, it is an often used target when a stock recruit relationship is unknown or unreliable, as is the case for snow crab. The range of years of recruitment used to calculate biomass reference points is from 1982 to the present assessment year, minus 1. However, recent analyses suggest SPR-based reference points do not provide a meaningful constraint on the snow crab fishery when the probability of having undergone terminal molt is specified to reflect observations in the survey. This is because a large fraction of the population matures (and ceases growing) at a size smaller than is harvested by the fishery.

## Fishery history

Snow crab were harvested in the Bering Sea by the Japanese from the 1960s until 1980 when the Magnuson Act prohibited foreign fishing. After the closure to foreign fleets, retained catches increased from relatively low levels in the early 1980s (e.g. retained catch of 11.85 kt during 1982) to historical highs in the early and mid-1990s (retained catches during 1991, 1992, and 1998 were $143.02,104.68$, and 88.09 kt , respectively; Table 10). The stock was declared overfished in 1999 at which time retained catches dropped to levels similar to the early 1980s (e.g. retained catch during 2000 was 11.46 kt ). Retained catches slowly increased after 1999 as the stock rebuilt. However, the fishery was closed for the first time in 2022 following the collapse observed in 2021.

Discard mortality from the directed fishery is the next largest source of mortality after retained catch and approximately tracks the retained catch. The highest estimated discard mortality occurred during 1992 at 17.06 kt which was $16 \%$ of the retained catch during that year. There was no discard mortality in 2022 because there was no directed fishery.
Discard from the directed pot fishery has been estimated from observer data since 1992 and has ranged from $11-100 \%$ of the magnitude of retained catch by numbers. In recent years, discards have reached $50-100 \%$ of the magnitude of retained catch because of the large year class entering the population. Female discard catch has been very low compared to male discard catch and has not been a significant source of mortality. Discard mortality rates for the directed fishery are assumed to be $30 \%$. Discard of snow crab in groundfish fisheries has been highest in the yellowfin sole trawl fishery, and decreases down through the flathead sole trawl fishery, Pacific cod bottom trawl fishery, rock sole trawl fishery, and the Pacific cod hook-and-line and pot fisheries, respectively (Figure 12). Bycatch in fisheries other than the groundfish trawl fishery has historically been relatively low. Discard mortality rates from non-directed fisheries are assumed to be $80 \%$. Size frequency data and catch per pot have been collected by observers on snow crab fishery vessels since 1992. Observer coverage has been $10 \%$ on catcher vessels larger than 125 ft (since 2001), and $100 \%$ coverage on catcher processors (since 1992).
Several modifications to pot gear have been introduced to reduce bycatch mortality. In the 1978/79 season, escape panels were required on pots used in the snow crab fishery to prevent ghost fishing. Escape panels consist of an opening with one-half the perimeter of the tunnel eye laced with untreated cotton twine. The size of the cotton laced panel was increased in 1991 to at least 18 inches in length. No escape mechanisms for undersized crab were required until the 1997 season when at least one-third of one vertical surface of pots had to contain not less than 5 inches stretched mesh webbing or have no less than four circular rings of no less than $33 / 4$ inches inside diameter. In the 2001 season the escapement provisions for undersized crab was increased to at least eight escape rings of no less than 4 inches placed within one mesh measurement from the bottom of the pot, with four escape rings on each side of the two sides of a four-sided pot, or one-half of one side of the pot must have a side panel composed of not less than $51 / 4$ inch stretched mesh webbing.

## D. Data

Updated time series of survey indices and size compositions were calculated from data downloaded from the AKFIN database. Bycatch data (biomass and size composition) were updated for the most recent year from the AKFIN database. Retained, total, and discarded catch (in numbers and biomass) and size composition data for each of these data sources were updated for the most recent year based on files provided by the State of Alaska.

## Catch data

Catch data and size composition of retained crab from the directed snow crab pot fishery from survey year 1982 to 2022 were used in this analysis (Table 10). Discard size composition data from 1992 to 2017 were estimated from observer data and then combined with retained catch size compositions to become the 'total
catch' size composition data, which are fit in the assessment. In 2018, observer data collection changed and only total catch size composition data and retained size composition data were produced. This is a sensible step in data collection, but the current formulation of the snow crab model accepts discarded size composition data as an input. So, from 2018 onward the discarded size compositions were calculated by subtracting the retained size compositions from the total size compositions. This mismatch of input data types will be addressed in an upcoming data overhaul for the assessment.

The discard male catch was estimated for survey years 1982 to 1991 in the model using the estimated fishery selectivities based on the observer data for the period of survey year 1992 to 2022 . The discard catch estimate was multiplied by the assumed mortality of discards from the pot fishery. The assumed mortality of discarded crab was $30 \%$ for all model scenarios. This estimate differs from the strategy used since 2001 to the present by ADFG to set the TAC, which assumes a discard mortality of $25 \%$ (Zheng, et al. 2002). The discards prior to 1992 may be underestimated due to the lack of escape mechanisms for undersized crab in the pots before 1997. See Table 6 for a summary of catch data.

Table 6: Data included in the assessment. Dates indicate survey year. The 2020 survey was cancelled due to the pandemic.

| Data component | Years |
| :--- | :---: |
| Retained male crab pot fishery size frequency by shell condition | $1982-2022$ |
| Discarded Males and female crab pot fishery size frequencey | $1992-2022$ |
| Trawl fishery bycatch size frequencies by sex | $1991-2022$ |
| Survey size frequencies by, maturity, sex and shell condition | $1982-2019,2021-2023$ |
| Retained catch estimates | $1982-2022$ |
| Discard catch estimates from crab pot fishery | $1992-2022$ |
| Trawl bycatch estimates | $1993-2022$ |
| Total survey abundance estimates and coefficients of variation | $1982-2019,2021-2023$ |
| 2009 study area biomass estimates, CVs, and size frequencey for | 2009 |
| BSFRF and NMFS tows | 2010 |
| 2010 study area biomass estimates, CVs, and size frequencey for |  |
| BSFRF and NMFS tows |  |

## Survey biomass and size composition data

Estimates of of the numbers of crab by sex and size from the annual eastern Bering Sea (EBS) bottom trawl survey conducted by NMFS (e.g. Figure $13 \&$ Figure 14; see Lang et al., 2018) are used to calculate the primary indices of abundance used in this assessment. Additional survey stations were added in 1989, which could alter the interpretation of catchability coefficient for the survey. Consequently, survey selectivity has been historically modeled in two 'eras' in the assessment (1982-1988, 1989-present). All survey data in this assessment used measured net widths instead of the fixed 50 ft net width based on Chilton et al.'s (2009) survey estimates. Carapace width and shell conditions were measured and reported for snow crab caught in the survey. Biomass and abundance of crab in several size groups are currently at or near all-time lows (Figure 15 \& Figure 16).

Mature male size composition data were calculated by multiplying the total numbers at length for new shell male crab by a vector of observed proportion of mature males at length. All old shell crab of both sexes were assumed to be mature. New shell crab were demarcated as any crab with shell condition index $<=2$. The biomass of new and old shell mature individuals was calculated by multiplying the vector of numbers at length by weight at length. These vectors were then summed by sex to provide the input for assessment (Table 11).

## Spatial distribution of survey abundance and catch

Snow crab are distributed widely over the eastern Bering Sea shelf, but their density and the extent of their distribution has changed over time (Figure $1 \&$ Figure 2). Spatial gradients exist in the survey data by maturity and size for both sexes. For example, larger males have been more prevalent on the southwest portion of the shelf (Figure 5 \& Figure 6) while smaller males have been more prevalent on the northern portion of the shelf (Figure 3 \& Figure 4). The centroids of abundance for male crab sized $45-85 \mathrm{~mm}$ carapace width have moved over time (Figure 17). Centroids of mature female abundance early in the history of the survey were farther south, but moved north during the 1990s. Since the late 1990s and early 2000s, the centroids moved south again, but not to the extent seen in the early 1980s. This phenomenon was mirrored in centroids of abundance for large males (Figure 18).
Fishing effort has generally been south of 58.5 N , even when ice cover did not restrict the fishery moving farther north (Figure 19 \& Figure 20). This is possibly due to the proximity to port and practical constraints of meeting delivery schedules. CPUE in the fishery has varied over time and an increase in average CPUE occurred after rationalization (Figure $21 \&$ Figure 22). The change in CPUE in a given spatial area within a season can reflect the impact of the fishery on the population in that area. Declines in CPUE can be seen by spatial area over time within a season (Figure 23), and the mean weekly change in CPUE is -11.6 (Figure 24). Total catch in an area is negatively correlated with the change in CPUE-that is, higher catches in an area are related to larger declines in CPUE (Figure 24).

The observed distribution of large males during the summer survey and the fishery catch have historically differed, and the origin of this difference is unknown. It is possible that crab move between the fishery and the survey, but it is also possible that fishers do not target all portions of the distribution of large male crab equally. The underlying explanation of this phenomenon could hold implications for relative exploitation rates spatially and it has been suggested that high exploitation rates in the southern portion of the snow crab range may have resulted in a northward shift in snow crab distribution (Orensanz, 2004). Snow crab larvae likely drift north and east after hatching in spring. Snow crab appear to move south and west as they age (Parada et al., 2010); however, little tagging data exists to fully characterize the ontogenetic or annual migration patterns of this stock (Murphy et al. 2010).

## Experimental study of survey selectivity

The Bering Sea Fisheries Research Foundation (BSFRF) has conducted supplementary surveys in the Bering Sea in which snow crab were caught during 2009, 2010, 2016, 2017, and 2018. The location and extent of these surveys varied over the years as the survey goals changed. In 2009, the survey consisted of 108 tows around 27 survey stations and the goal was to improve understanding snow crab densities and the selectivity of NMFS survey gear (Figure 25). In 2010, the survey area was larger and still focused on snow crab. The mature biomass and size composition data gleaned from each of these experiments (and their complimentary NMFS survey observations) are incorporated into the status quo model by fitting them as an extra survey that is linked to the NMFS survey through a shared selectivity (see GMACS documentation on the github repo to see how a survey can be 'embedded' within another and the repo holding the files for this years snow crab assessment for implementation; both linked at the end of this document). The status quo assessment model estimates a vector that represents the 'availability' of crab to the BSFRF experiments. Availability in this case means how much of the population was in the area surveyed in the BSFRF experiments. Abundances estimated by the industry surveys were generally higher than the NMFS estimates, which suggests that the catchability of the NMFS survey gear is less than 1 .
In 2016, 2017, and 2018, snow crab were not the focus of the BSFRF surveys, yet were still caught in the BSFRF gear. Comparing the ratio of the number of crab caught at length in the BSFRF gear (which is assumed to have a catchability/selectivity of 1 over all size classes) to the number of crab caught at length within the same area in the NMFS survey gear (which is assumed to have a catchability/selectivity $<=$ to 1 for at least some of the size classes) can provide an empirical estimate of catchability/selectivity (Figure 26). Empirical estimates of catchability/selectivity vary by year and size class across the different BSFRF data sets (Figure 27 \& Figure 28). The number of snow crab used to develop estimates of numbers
at length likely contribute to these differences among years (Figure 29), but other factors may also influence catchability/selectivity at size of the NMFS survey gear (e.g. Somerton et al. 2013 show substrate type can influence selectivity).

## E. Analytic approach

## History of modeling approaches for the stock

Historically, survey estimates of large males ( $>101 \mathrm{~mm}$ ) were the basis for calculating the Guideline Harvest Level (GHL) for retained catch. A harvest strategy was developed using a simulation model that pre-dated the current stock assessment model (Zheng et al. 2002). This model has been used to set the GHL (renamed total allowable catch, 'TAC', since 2009) by ADFG since the 2000/2001 fishery. Currently, NMFS uses an integrated size-structured assessment to calculate the overfishing level (OFL), which is used to set an acceptable biological catch (ABC), which in turn provides a ceiling to the TAC set by the state process.

## Model description

Recently, the Generalized Model for Assessing Crustacean Stocks (GMACS) was adopted as the assessment platform for snow crab after a demonstration that GMACS could effectively reproduce the dynamics of the status quo model and offered structural improvements. GMACS is an integrated, size-structured model developed using automatic differentiation software developed as a set of libraries under C++ (ADModel Builder). ADModel Builder can estimate a large number of parameters in a non-linear model using automatic differentiation software extended from Greiwank and Corliss (1991) and developed into C++ class libraries.

The snow crab population dynamics model tracks the number of crab of sex $s$, maturity state $m$, during year $y$ at length $l, \mathrm{~N}_{s, m, y, l}$. A terminal molt was modeled in which crab move from an immature to a mature state, after which no further molting occurred. The mid-points of the size bins tracked in the model spanned from 27.5 to 132.5 mm carapace width, with 5 mm size classes. For the author-preferred model, 407 parameters were estimated. Parameters estimated within the assessment included those associated with the population processes recruitment, growth, natural mortality (subject to an informative prior and two years of additional 'mortality events' estimated in 2018 and 2019), fishing mortality, selectivity (fishery, survey, and BSFRF experiments), catchability, and maturity. Weight at length, discard mortality, bycatch mortality, variance in growth increment, and parameters associated with proportion of recruitment allocated to size bin were estimated outside of the model or specified. See the GMACS repo linked at the end of this document for a more complete description of the population dynamics.
A 'jittering' approach has been historically used to explore the impact of different starting values on the assessment output (Turnock, 2016). Jittering was implemented for a selected number of models here. Retrospective analyses were also performed here in which the terminal year of data was removed sequentially from the model fitting process. Then time series of estimated MMB were compared between the most recent model and successive 'peels' of the data to identify retrospective patterns. A retrospective pattern is a consistent directional change in assessment estimates of management quantities (e.g. MMB) in a given year when additional years of data are added to an assessment.
Model explorations presented here include changing the way that the BSFRF data are treated in the model and modifying the way the probability of having undergone terminal molt is modeled. In the status quo assessment, the BSFRF data are fit as an additional survey and act on estimated survey selectivity through shared selectivity parameters between the full NMFS bottom trawl survey data set and the portion of it that corresponds with the area that was surveyed by BSFRF. An alternate way to incorporate these data is to calculate the inferred selectivity from the BSFRF data and use these point estimates and associated variance as direct priors on selectivity at size in the assessment (see above for description). The observed probability of having undergone terminal molt by year is based on chela height measurements and is available for the years 1989 -to present, excluding the years $1994,2008,2012,2014,2016$, and 2020 . The mean probability of having undergone terminal molt at size was used for years with no data.

## Model selection and evaluation

Models were evaluated based on their fit to the data, evidence of non-convergence, the credibility of the estimated population processes, and the strength of the influence of the assumptions of the model on the outcomes of the assessment.

## Results

All models converged with updated data and minor bug fixes to the assessment code. Retrospective patterns were relatively small compared to historical patterns (Figure 30). Jittering analyses produce bimodal management quantities in the OFL for model 23.1 and for $\mathrm{B}_{35 \%}$ in model 23.3 (Figure 31). Below, the fits to the data and estimated population processes are described for all considered models that include the most recent data. Contribution of likelihood components to the objective function are in Table 12 and parameter estimates and standard deviations are in Table 13 \& Table 14. The total objective functions of 23.1 and 23.2 are comparable to one another, but not to $23.3,23.3$ a, or 23.3 b because those models include the BSFRF data included as a direct prior on survey selectivity rather than as an additional index of abundance.

## Fits to data

## Survey biomass data

Fits to the survey mature male biomass were similar for all models for the majority of years in the the time series (Figure 32 \& Figure 33) with the same data. However, differences in the estimated survey MMB in the first survey era and the final year of data existed, with models $23.3,23.3 \mathrm{a}$, and 23.3 b producing estimates that were higher than the observation. Even with this over-prediction, model 23.3 b fit the MMB from the recent survey era (1989-present) best among the models (Table 12).

## Growth data

Small differences existed in the estimates of the relationship between pre- and post-molt increment existed among the models (Figure 34). The resulting size-transition matrix for males from author-preferred model appears to be broadly consistent with studies on crab growth (e.g. Herbert et al., 2001; Figure 35).

## Catch data

All models fit the catch data well, with few visually discernible differences among models (Figure 36). The largest differences in fit among models occurred during the early 1990s for male discards. Existing differences in fit were amplified in the objective function by the small CVs placed on the different sources of catch data, with model 23.1 and 23.3b fitting the data best (Table 12).

## Size composition data

Most years of retained and total catch size composition data were visually well fit by all models (Figure 37 \& Figure 38). In some years, the 23.3 model series estimated more crab in the largest size bin than models 23.1 and 23.2 for both retained and total catch size composition data (e.g. 1992, 2005, 2009). Model 23.1 fit the directed fishery size composition data best based on the contributions to the objective function (Table 12). Predictions of female discards in the directed fishery were right skewed for some years, potentially reflecting unmodeled time-variation in the availability of females to the directed fishery (Figure 39). Estimated size composition of the catch in non-directed fisheries was the least well fit of the catch sources, but the models were fairly consistent in their fits (Figure 40 \& Figure 41).

Size composition data for the NMFS survey were generally acceptably fit and fits were visually similar for most data sources in most models in most years (Figure 42, Figure 43, Figure 44, Figure 45, Figure 46, Figure 47, Figure 48 \& Figure 49). Poor fits often occurred at the smallest size bins, which is likely related to the interplay of poor and variable selectivity at small sizes with pseudocohorts (i.e. groups of similarly sized crab used in place of 'cohort' because we cannot age crab) that were first observed (and subsequently persisted) at larger sizes. Predicted mature male size compositions from models that estimated a single ogive for the probability of having undergone terminal molt (model 23.1 and 23.2 ) were frequently bimodal. In some years this bimodality was reflected in the data (e.g. 2012 or 2023), but in other years it was not (e.g. 2002, 2005, 2021). Model 23.1 fit 4 out of 8 survey size composition data sources at least marginally better than the other models. Fits to size composition data for the BSFRF survey selectivity experiments were similar across models (Figure 50).

## Estimated population processes and derived quantities

Estimated population processes and derived quantities varied among models. MMB estimates from 23.1 and the 23.3 model series produced similar trends in estimated MMB, except for in the early 1980s and the last five years (Figure 51). The 23.3 series estimated increases in the MMB during 2018 and 2019 that the status quo model did not track. This may be related to the higher probability of having undergone terminal molt at smaller sizes included in the 23.3 model series. Changing the way the probability of having undergone terminal molt was modeled, but not changing the way survey selectivity was modeled resulted in large increases in estimated MMB (model 23.2 in Figure 51).

The number and biomass of crab that are commercially preferred ( $>101 \mathrm{~mm}$ carapace width) are two of the most important figures to come out of the assessment because they are directly related to the OFL. The raw time series of commercially preferred males biomass is one of the time series considered in the state strategy and comparing the survey estimates to the assessment model estimates can provide context for the impact of selecting one model vs. another. Models 23.1 and 23.2 estimated much higher biomass of the commercially preferred males than was observed in the survey (Figure 52 ). However, the 23.3 model series estimates were much closer to the observed survey estimates.
Some of the differences in the estimated commercial biomass are related to estimates of survey selectivity. The scale and shape of the survey selectivity curves changed markedly among models, responding to changes in the way the BSFRF data were incorporated into the model (Figure 53). Focusing on the male selectivity in the most recent era, incorporating only the new data on the probability of having undergone terminal molt into model 23.2 resulted in much lower estimated catchability, which was reflected in the higher estimates of spawning biomass (Figure 51). Incorporating the BSFRF data as priors on survey selectivity parameters in the 23.3 series changed the shape of the selectivity curve (Figure 53). The increases in selectivity at larger carapace widths contributed to lower estimates of MMB compared to 23.2. Estimating survey selectivity with the BSFRF priors resulted in lower estimates of selectivity than inferred from the BSFRF data for individuals less than $\sim 100 \mathrm{~mm}$ carapace width, but higher estimates of selectivity above that size until $>125$ mm carapace width. Over all, estimates of survey selectivity for males mostly stayed within the implied uncertainty of the CVs associated with the BSFRF priors for model 23.3a and 23.3b (Figure 54).

The estimates of availability and selectivity from the models in which the BSFRF data were treated as an additional survey varied among models (22.1, 23.1, 23.2; Figure 55). Model 22.1 and 23.1 produced changes in the estimates of availablity in 2010 in spite of being structurally identical.
Retained fishery selectivity estimates for males were nearly identical for all models, but capture selectivity in the directed fishery varied among models (Figure 56). Selectivity associated with non-directed bycatch also varied among models ("Trawl_bycatch" in Figure 56). Estimated fully-selected fishing mortality in the directed and non-directed fleets were higher in the 23 model series than in 23.1 and 23.2 (Figure 57). High estimates of fully-selected fishing mortality in the directed fishery have been an issue of concern in previous stock assessments. Some of the models presented here still estimate seemingly unreasonably high fishing mortalities in some years (e.g. 1991 and 2020). However, the high fishing mortalities are only acting on the fully-selected portion of the stock, which, given the estimated selectivities, is comprised of only the largest crab. The proportion of exploitable male biomass that is in these largest sizes is quite low (Figure 58).

A realized exploitation rate can be calculated by dividing the retained catch by the exploitable biomass (i.e. crab $>101 \mathrm{~mm}$ carapace width; Figure 58, middle panel). Although the realized exploitation rate is high in 2020 and the early 1990s, it is no where near the estimated fully-selected fishing mortality rates. For example, in 2020 the estimated fully-selected fishing mortality rate translates to an exploitation rate of $\sim 100 \%$, but the realized exploitation rate was closer to $40 \%$.

The estimated probability of having undergone terminal molt in models 23.1 and 23.2 were very different from the specified probabilities derived from the survey data (Figure 59). The specified probabilities are calculated bas the proportion of new shell crab by size that are mature based on chela height. These proportions are used to divide the survey data into 'mature' and 'immature' data to calculate size compositions that are input into the assessment. Higher probabilities of terminally molting at smaller sizes results in much more of the population ceasing to grow beneath the size at which they are harvested in the directed fishery. This has large impacts on estimated SPR-based reference points, which will be discussed below.

Patterns and scale in recruitment by sex varied somewhat among models, particularly with respect to the size and timing of the recent large pseudocohort (Figure 60). Generally, the models estimated a period of high average recruitment during the 1980s. Following that, a period of low average recruitment persisted from the early 1990s to 2014. A large recruitment was estimated to enter the modeled fraction of the population around 2014-2016. Recruitment entering the model was distributed primarily in the first three size bins for all models (Figure 60).

Estimated natural mortality ranged from 0.27 to 0.55 for immature and mature crab (Figure 61 ). Estimated mortality events in 2018 and 2019 were most intense for immature females and males, but even the lower mortalities for mature females and immature males resulted in $>80 \%$ of crab dying.

## F. Calculation of the OFL

## Methodology for OFL

## Tier 3

The tier 3 OFL was calculated using proxies for biomass and fishing mortality reference points and a sloped control rule. Proxies for biomass and fishing mortality reference points were calculated using spawner-per-recruit methods (e.g. Clark, 1991). After fitting the assessment model to the data and estimating population parameters, the model was projected forward 100 years using the estimated parameters under no exploitation and constant recruitment to determine 'unfished' mature male biomass-per-recruit. Projections were repeated in which the bisection method was used to identify a fishing mortality that reduced the mature male biomass-per-recruit to $35 \%$ of the unfished level (i.e. $\mathrm{F}_{35 \%}$ and $\mathrm{B}_{35 \%}$ ). Calculations of $\mathrm{F}_{35 \%}$ were made under the assumption that bycatch fishing mortality was equal to the estimated average value over the last 8 years.

Calculated values of $\mathrm{F}_{35 \%}$ and $\mathrm{B}_{35 \%}$ were used in conjunction with a Tier 3 control rule to adjust the proportion of $\mathrm{F}_{35 \%}$ that is applied based on the status of the population relative to $\mathrm{B}_{35 \%}$ (Amendment 24, NMFS). To determine the $\mathrm{F}_{\mathrm{OFL}}$, the population is projected to the time of fishing for the upcoming fishery under no fishing. If the MMB at that time exceeds $25 \%$ of $\mathrm{B}_{35 \%}$, a fishery can occur and the $\mathrm{F}_{\text {ofl }}$ is calculated as:

$$
F_{O F L}= \begin{cases}\text { Bycatch } & \text { if } \frac{M M B}{B_{35}} \leq 0.25  \tag{2}\\ \frac{F_{35}\left(\frac{M M B}{B_{35}}-\alpha\right)}{1-\alpha} & \text { if } 0.25<\frac{M M B}{B_{35}}<1 \\ F_{35} & \text { if } M M B>B_{35}\end{cases}
$$

Where MMB is the projected mature male biomass in the current survey year after fishing at the $\mathrm{F}_{\text {OFL }}$, $\mathrm{B}_{35 \%}$ is the mature male biomass at the time of mating resulting from fishing at $\mathrm{F}_{35 \%}, \mathrm{~F}_{35 \%}$ is the fishing mortality that reduces the mature male biomass per recruit to $35 \%$ of unfished levels, and $\alpha$ determines the slope of the descending limb of the harvest control rule (set to 0.1 here).
Calculated tier 3 OFLs ranged from 8.58 to 37.10 kt (Table 15). Differences in OFLs were a result of differences in estimated MMB, calculated $\mathrm{B}_{35 \%}$ (which ranged from $110.01-189.24 \mathrm{kt}$ ), $\mathrm{F}_{35 \%}$ (which ranged from 1.50-205.67 $\mathrm{yr}^{-1}$ ), and $\mathrm{F}_{\text {OFL }}$ (which ranged from $0.30-37.49 \mathrm{yr}^{-1}$; Table 15).

## Tier 4

Tier 4 OFLs were calculated within GMACS using the estimated natural mortality as the proxy for $F_{M S Y}$ and the average morphometrically mature male biomass from 1982-2022 as the target biomass. A tier 4 OFL was also calculated using raw survey estimates of commercially size males ( $>101 \mathrm{~mm}$ carapace width).

Calculated tier 4 OFLs within GMACS ranged from 0.03 to 0.31 kt (Table 16). Differences in OFLs were a result of differences in estimated MMB , calculated $\mathrm{B}_{M S Y}$ proxy (which ranged from $232.32-519.67 \mathrm{kt}$ ), $\mathrm{F}_{M S Y}$ proxy (which ranged from $0.28-0.55 \mathrm{yr}^{-1}$ ), and $\mathrm{F}_{\text {OFL }}$ (which ranged from $0.00-0.05 \mathrm{yr}^{-1}$; Table 16).
The tier 4 OFL calculated from the survey data was 0 and the status was $\sim 17 \%$ of the $\mathrm{B}_{M S Y}$ proxy (Table 17).

## G. Calculation of the ABC

The acceptable biological catch (ABC) was set by subtracting a $20 \%$ buffer from the OFL to account for scientific uncertainty, as recommended by the SSC.

## Author recommendations

Two decisions need to be made to provide management advice for snow crab based on this assessment: 1) which model to choose and 2) which tier to use for the harvest control rule. All models converged and performed reasonably in jittering and retrospective analyses. The status quo model fit many of the data sources best, but it did not incorporate the best available science on the biology of snow crab. Given the importance of correctly modeling the biology, all models that include the data on terminal molt should be considered improvements over the status quo. Modeling selectivity as a non-parametric curve also is supported over logistic selectivity, given two separate data sources that suggest survey selectivity is not logistic (BSFRF and Somerton and Otto, 1998; Figure 28). Consequently, the 23.3 series have the most desirable characteristics of the models presented.
Estimating non-parametric selectivity rather than specifying it allows uncertainty in the estimate to be propagated, so 23.3 a or 23.3 b is preferable over 23.3 . A potential downside of estimating the selectivity is that its weighting relative to other data sets is poorly understood and survey selectivity is an influential parameter in determining the OFL. The prior on natural mortality is quite tight, but it is highly confounded with other parameters (e.g. selectivity, catchability) which can make its estimation difficult. Loosening the prior results in a much larger estimated natural mortality, which is in conflict with the assumption that snow crab have a maximum age of approximately 20 years. Given the influence on management quantities that changing M can have, loosening the prior should likely only be done after more extensive exploration of its impacts. Based on all of these considerations, the author-preferred model is 23.3a.
The second question for consideration is which tier to use in specifying the quantities in the harvest control rule. Tier 3 uses spawner-per-recruit proxies for fishing mortality and biomass targets; tier 4 uses natural mortality and an average mature male biomass over time for fishing mortality and biomass targets. Tier 3 rules produce fishing mortality reference points that allow for $\sim 100 \%$ exploitation rates on commerciallypreferred males in model 23.3a. This occurs because the updated information on the probability of having undergone terminal molt allows crab to mature (and stop growing) before reaching the size harvested in the fishery. This results in a large fraction of the mature biomass being protected from harvest, which then requires high exploitation rates on commercially-preferred males to reduce the MMB to $35 \%$ of unfished levels. The tier 4 rule applied within GMACS results in much lower fishing mortalities applied to the stock. The tier 4 rule applied to the survey data resulted in an even lower status than in GMACS and a closure of the fishery.

Based on these considerations, a tier 4 approach within GMACS is the author-preferred method for specifying the harvest control rule. The stock is at unprecedented lows and this warrants caution in management. Following the status quo tier 3 approach does not seem defensible given the current status of the stock. There is little practical difference between adopting the tier 4 rule within GMACS or within a survey-based rule, and retaining the model based assessment incorporates as much of the available information as possible into the analysis.

## H. Data gaps and research priorities

Although the author-preferred HCR is based on tier 4, there are some inconsistencies within its specification for snow crab. Although natural mortality is often reported to be similar to $\mathrm{F}_{M S Y}$, the shape of estimated fishing selectivity for snow crab results in a non-uniform application of the target fishing mortality to the exploitable biomass. This results in a smaller fishing mortality applied to the exploitable biomass than natural mortality. The use of morphometrically mature male biomass as the currency of management further obfuscates the impact of a chosen target fishing mortality on the population as the fishing mortality is modified based on crab that are not subject to fishing mortality. The interchangability of a 70 mm carapace width male and a 120 mm carapace male in reproductive dynamics is a key, but difficult to corroborate assumption.

A potential approach that could address these inconsistencies is modifying the currency of management to reflect the exploitable biomass more closely and then choosing some fraction of unfished levels as a
biomass target. This could provide a more interpretable analog to the status quo management given the issues translating the fully-selected fishing mortality to impacts to the stock as a result of the shape of fishery selectivity and morphometric maturity used as a currency of management. This would require more thorough testing and deliberation on what an appropriate fraction of unfished biomass levels should be to serve as a target, particularly under the potential for changing productivity of the stock.

Data weighting continues to be a topic that is acknowledged as important to modeling outcomes, but secondary to finding an appropriate model configuration. A thorough examination of the data streams in the assessment including reconstructing historical time series (rather than appending a year of data to the existing data file) and reevaluating the data sets to which the assessment is fit (e.g. should immature crab or very large crab also be fit) should be undertaken.

Considerable effort has been expended since the May CPT meeting exploring male only models that are not presented here. These models may be useful in understanding the impacts of modeling both sexes in the same model and will be explored further in future documents.

## I. Ecosystem considerations

Key questions related to ecosystem change include more thoroughly understanding the mortality event in 2018 and 2019, anticipating the potential for impacts of warming on mortality, recruitment, and other population processes, and understanding the potential for shifts in distribution and their impacts on the population and fishery. Several on-going projects are being conducted to inform these questions and results will be shared as they become available.

See the ESP for snow crab specific indices of environmental variation that may be relevant to stock dynamics.
Input and output for the models described here can be found at https://github.com/szuwalski/snow_2023 9.

GMACS code and (some) documentation can be found at: https://github.com/GMACS-project.

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Table 7: Key differences in presented models. (continued below)

| Process | 23.1 | 23.2 | 23.3 | 23.3a |
| :---: | :---: | :---: | :---: | :---: |
| Sex | Both | Both | Both | Both |
| Maturity | Single estimated ogive | Input | Input | Input |
| BSFRF | Survey | Survey | Prior | Prior |
| Survey | Estimated logistic by sex and era | Estimated logistic by sex and era | Specified non-parametric | Estimated non-parametric |
| Growth | Linear estimated | Linear estimated | Linear estimated | Linear estimated |
| Natural.M | By sex and maturity + 2018/19 | By sex and maturity + 2018/19 | By sex and maturity $+2018 / 19$ | By sex and maturity $+2018 / 19$ |
| Fishery | Logistic | Logistic | Logistic | Logistic |


| 23.3 b |
| :--- |
| Both |
| Input |
| Prior |
| Estimated |
| non-parametric |
| Linear estimated |
| By sex and maturity + |
| $2018 / 19+$ looser prior |
| Logistic |

Table 9: Observed growth increment data by sex

| Male premolt length (mm) | Male growth increment (mm) | Female premolt length (mm) | Female growth increment (mm) |
| :---: | :---: | :---: | :---: |
| 16.1 | 6.9 | 93.8 | 23.8 |
| 19.2 | 7.4 | 18.6 | 6.6 |
| 19.8 | 6.7 | 19.3 | 5.9 |
| 20 | 6.3 | 19.37 | 4.87 |
| 20 | 6.3 | 19.8 | 7.1 |
| 20.1 | 7.9 | 20.2 | 4.7 |
| 20.3 | 6.1 | 20.3 | 5.9 |
| 20.6 | 8.3 | 20.4 | 6 |
| 20.7 | 7 | 20.4 | 6.3 |
| 20.7 | 8.5 | 20.6 | 4.5 |
| 21 | 6.8 | 20.7 | 6.3 |
| 21.23 | 5.18 | 20.7 | 6.7 |
| 21.9 | 6.5 | 20.8 | 6.5 |
| 22.2 | 5.9 | 20.8 | 6.5 |
| 23.48 | 4.79 | 20.8 | 6.8 |
| 24 | 8.3 | 21.25 | 7.48 |
| 25.2 | 7.6 | 21.4 | 6.6 |
| 25.6 | 5.8 | 21.6 | 6.1 |
| 25.9 | 5.2 | 21.94 | 6.77 |
| 26 | 6.2 | 22 | 6.2 |
| 29.9 | 10 | 22.2 | 7.5 |
| 30.3 | 10 | 22.3 | 7.1 |
| 30.7 | 9.8 | 22.8 | 6.8 |
| 44.2 | 14.5 | 22.8 | 7.4 |
| 44.7 | 12.6 | 22.9 | 5.7 |
| 56.5 | 13.5 | 23 | 8.2 |
| 57 | 13 | 23.09 | 6.17 |
| 57.63 | 10.97 | 24.2 | 6.7 |
| 58.7 | 13.8 | 24.2 | 7.2 |
| 59.3 | 15.8 | 24.4 | 6.3 |
| 60.3 | 14.8 | 25.2 | 6.8 |
| 60.8 | 17.6 | 25.4 | 6.3 |
| 62.3 | 19.5 | 25.5 | 9.1 |
| 64 | 20.7 | 25.5 | 7.4 |
| 64.7 | 18 | 25.7 | 6.8 |
| 67.6 | 18.4 | 25.9 | 6.8 |
| 67.9 | 17.4 | 26 | 7.1 |
| 74.5 | 19.4 | 26.2 | 6.4 |
| 79.9 | 17.9 | 26.4 | 5.4 |
| 89.8 | 20.2 | 26.5 | 7.4 |
| 89.9 | 22.2 | 26.9 | 7.5 |
| 89.9 | 22.4 | 26.9 | 7.6 |
| 93.8 | 23.8 | 27.4 | 7.7 |
|  |  | 27.5 | 7.3 |
|  |  | 28.1 | 6.4 |
|  |  | 28.2 | 8.02 |
|  |  | 28.2 | 7.6 |
|  |  | 28.7 | 8.4 |
|  |  | 28.7 | 7.3 |
|  |  | 29 | 7.7 |

$\left.\begin{array}{ccc}\hline \begin{array}{c}\text { Male premolt length } \\ (\mathrm{mm})\end{array} & \begin{array}{c}\text { Male growth } \\ \text { increment }(\mathrm{mm})\end{array} & \begin{array}{c}\text { Female premolt } \\ \text { length }(\mathrm{mm})\end{array}\end{array} \begin{array}{c}\text { Female growth } \\ \text { increment (mm) }\end{array}\right]$

Table 10: Observed retained catches, discarded catch, and bycatch.
Discards and bycatch have assumed mortalities applied.

| Survey Year | Retained catch (kt) | Discarded females (kt) | Discarded males (kt) | Non-directed bycatch (kt) |
| :---: | :---: | :---: | :---: | :---: |
| 1982 | 11.85 | 1.27 | 0.02 | 0.37 |
| 1983 | 12.16 | 1.24 | 0.01 | 0.47 |
| 1984 | 29.94 | 2.76 | 0.01 | 0.5 |
| 1985 | 44.45 | 4.01 | 0.01 | 0.43 |
| 1986 | 46.22 | 4.25 | 0.02 | 0 |
| 1987 | 61.4 | 5.52 | 0.03 | 0 |
| 1988 | 67.79 | 5.82 | 0.04 | 0 |
| 1989 | 73.4 | 6.68 | 0.05 | 0.1 |
| 1990 | 149.1 | 15.21 | 0.05 | 0.71 |
| 1991 | 143 | 12 | 0.06 | 1.5 |
| 1992 | 104.7 | 17.06 | 0.12 | 2.28 |
| 1993 | 67.94 | 5.32 | 0.08 | 1.57 |
| 1994 | 34.13 | 4.03 | 0.06 | 2.67 |
| 1995 | 29.81 | 5.75 | 0.02 | 1.01 |
| 1996 | 54.22 | 7.44 | 0.07 | 0.66 |
| 1997 | 114.4 | 5.73 | 0.01 | 0.82 |
| 1998 | 88.09 | 4.67 | 0.01 | 0.54 |
| 1999 | 15.1 | 0.52 | 0 | 0.47 |
| 2000 | 11.46 | 0.62 | 0 | 0.41 |
| 2001 | 14.8 | 1.89 | 0 | 0.31 |
| 2002 | 12.84 | 1.47 | 0 | 0.17 |
| 2003 | 10.86 | 0.57 | 0 | 0.46 |
| 2004 | 11.29 | 0.51 | 0 | 0.63 |
| 2005 | 16.77 | 1.36 | 0 | 0.2 |
| 2006 | 16.49 | 1.78 | 0 | 0.42 |
| 2007 | 28.59 | 2.53 | 0.01 | 0.18 |
| 2008 | 26.56 | 2.06 | 0.01 | 0.18 |
| 2009 | 21.78 | 1.23 | 0.01 | 0.47 |
| 2010 | 24.61 | 0.62 | 0.01 | 0.14 |
| 2011 | 40.29 | 1.69 | 0.18 | 0.15 |
| 2012 | 30.05 | 2.32 | 0.03 | 0.22 |
| 2013 | 24.49 | 3.27 | 0.07 | 0.11 |
| 2014 | 30.82 | 3.52 | 0.17 | 0.13 |
| 2015 | 18.42 | 2.96 | 0.07 | 0.13 |
| 2016 | 9.67 | 1.31 | 0.02 | 0.06 |
| 2017 | 8.6 | 1.93 | 0.02 | 0.04 |
| 2018 | 12.51 | 2.86 | 0.02 | 0.23 |
| 2019 | 15.43 | 5.07 | 0.02 | 0.24 |
| 2020 | 20.41 | 5.8 | 0 | 0.07 |
| 2021 | 2.48 | 1.16 | 0 | 0.06 |
| 2022 | 0 | 0 | 0 | 0.05 |

Table 11: Observed mature male and female biomass (1000 t) at the time of the survey and coefficients of variation.

| Survey year | Female mature biomass | Female CV | Mature male biomass | Male CV | $\begin{gathered} \text { Males } \\ >101 \mathrm{~mm} \\ (\mathrm{kt}) \end{gathered}$ | $\begin{gathered} \text { Males } \\ >101 \mathrm{~mm} \\ \text { (million) } \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1982 | 144.4 | 0.15 | 176.8 | 0.14 | 34.82 | 65.04 |
| 1983 | 90.13 | 0.2 | 161.6 | 0.13 | 35.09 | 65.57 |
| 1984 | 42.32 | 0.19 | 177.7 | 0.12 | 85.1 | 148.3 |
| 1985 | 6.12 | 0.2 | 71.84 | 0.11 | 43.1 | 73.82 |
| 1986 | 15.74 | 0.18 | 89.81 | 0.11 | 45.97 | 78.15 |
| 1987 | 122.6 | 0.16 | 194.6 | 0.11 | 74.29 | 130.8 |
| 1988 | 169.9 | 0.17 | 259.4 | 0.15 | 105.7 | 178.4 |
| 1989 | 264.2 | 0.25 | 299.2 | 0.11 | 92.42 | 162 |
| 1990 | 182.9 | 0.19 | 443.8 | 0.14 | 225.1 | 395.1 |
| 1991 | 214.9 | 0.19 | 466.6 | 0.15 | 278.7 | 439.7 |
| 1992 | 131.4 | 0.18 | 235.5 | 0.09 | 139 | 223.3 |
| 1993 | 132.1 | 0.16 | 183.9 | 0.1 | 77.23 | 127.6 |
| 1994 | 126.2 | 0.15 | 171.3 | 0.08 | 44.64 | 73.79 |
| 1995 | 168.7 | 0.14 | 220.5 | 0.13 | 38.18 | 67.3 |
| 1996 | 107.3 | 0.14 | 288.4 | 0.12 | 89.02 | 161.4 |
| 1997 | 103.8 | 0.2 | 326.8 | 0.1 | 171.5 | 290.8 |
| 1998 | 72.73 | 0.25 | 206.4 | 0.09 | 127.5 | 214.9 |
| 1999 | 30.89 | 0.21 | 95.85 | 0.09 | 52.04 | 85.72 |
| 2000 | 96.46 | 0.52 | 96.39 | 0.14 | 41.13 | 69.78 |
| 2001 | 77.24 | 0.28 | 136.5 | 0.12 | 39.99 | 69.26 |
| 2002 | 30.22 | 0.28 | 93.17 | 0.23 | 37.17 | 66.58 |
| 2003 | 41.71 | 0.31 | 79.07 | 0.12 | 31.53 | 54.97 |
| 2004 | 50.16 | 0.26 | 79.57 | 0.14 | 35.58 | 58 |
| 2005 | 64.85 | 0.17 | 123.5 | 0.11 | 39.85 | 62.96 |
| 2006 | 51.93 | 0.17 | 139.3 | 0.26 | 72.34 | 126.4 |
| 2007 | 55.89 | 0.22 | 153.1 | 0.15 | 74.72 | 132.5 |
| 2008 | 57.15 | 0.19 | 142 | 0.1 | 60.33 | 105.1 |
| 2009 | 52.16 | 0.21 | 148.2 | 0.13 | 77.51 | 129.9 |
| 2010 | 98.01 | 0.17 | 162.8 | 0.12 | 87.1 | 138.2 |
| 2011 | 175.8 | 0.18 | 167.1 | 0.11 | 94.38 | 150.1 |
| 2012 | 149.4 | 0.2 | 122.2 | 0.12 | 53.15 | 87 |
| 2013 | 131.4 | 0.17 | 97.46 | 0.12 | 43.13 | 73.64 |
| 2014 | 119.7 | 0.19 | 163.5 | 0.16 | 79.51 | 138.5 |
| 2015 | 85.13 | 0.17 | 80.04 | 0.12 | 35.84 | 57.19 |
| 2016 | 55.39 | 0.21 | 63.21 | 0.11 | 22 | 37.43 |
| 2017 | 106.8 | 0.21 | 83.96 | 0.13 | 20.74 | 36 |
| 2018 | 165.9 | 0.18 | 198.4 | 0.17 | 27.02 | 49.41 |
| 2019 | 110.4 | 0.2 | 169.1 | 0.17 | 28.95 | 53.7 |
| 2021 | 31.66 | 0.43 | 62.25 | 0.13 | 12.44 | 23.53 |
| 2022 | 22.44 | 0.41 | 37.5 | 0.15 | 13.49 | 24.59 |
| 2023 | 14.96 | 0.24 | 24.21 | 0.13 | 11.44 | 20.03 |

Table 12: Contribution to the objective function by individual likelihood component by model. Total likelihoods from models 23.1 and 23.2 are not comparable to the other models because they still fit the BSFRF data as an extra survey. Models 23.3a and 23.3b estimate parametric survey selectivity with a prior; 23.3 specifies survey selectivity.

| Component | Fishery | 23.1 | 23.2 | 23.3 | 23.3a | 23.3b |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| catch | Retained | -7.08 | -4.15 | 5.03 | 2.92 | -7.27 |
| catch | Discard (male) | 140.7 | 130.44 | 79.02 | 88.95 | 69.18 |
| catch | Discard (female) | -69.66 | -69.66 | -69.66 | -69.66 | -69.66 |
| catch | Trawl | -52.03 | -52.03 | -52.02 | -52.02 | -52.02 |
| cpue | NMFS survey (era 1; females) | 43.44 | 54.65 | 71.7 | 53.59 | 36.06 |
| cpue | NMFS survey (era 2, females) | -28.82 | -13.3 | -8.25 | -2.38 | -16.41 |
| cpue | NMFS survey (era 1, males) | 32.15 | 35.28 | 49.98 | 46.62 | 43.13 |
| cpue | NMFS survey (era 2, males) | 21.33 | -0.98 | 28.32 | 31.23 | -5.25 |
| growth_inc | 1 | 1020.3 | 1061.75 | 1049.82 | 1038.86 | 1033.27 |
| growth_inc | 2 | 0 | 0 | 0 | 0 | 0 |
| rec_dev | 1 | 0.78 | 0.78 | 0.78 | 0.78 | 0.78 |
| rec_dev | 2 | 0 | 0 | 0 | 0 | 0 |
| rec_dev | 3 | 93.91 | 79.3 | 96.65 | 80.87 | 78.4 |
| size_comp | Retained males | -3702.08 | -3608.48 | -3632.29 | -3641.29 | -3665.38 |
| size_comp | Survey mature females (1982-1988) | -688.42 | -685.48 | -650.21 | -678.99 | -681.7 |
| size_comp | Survey mature females (1989-present) | -3168.88 | -3056.79 | -2981.02 | -3198.41 | -3196.99 |
| size_comp | Survey mature males (1982-1988) | -595.71 | -581.17 | -576.56 | -582.62 | -585.6 |
| size_comp | Survey mature males (1989-present) | -2828.43 | -2694.15 | -2725.15 | -2800.92 | -2829.29 |
| size_comp | Total males | -2708.96 | -2630.54 | -2629.61 | -2638.41 | -2662.37 |
| size_comp | Discard females | -2283.93 | -2275.2 | -2272.08 | -2270.88 | -2275.2 |
| size_comp | Non-directed bycatch (females) | -2539.63 | -2481.05 | -2486.45 | -2460.89 | -2436.49 |
| size_comp | Non-directed bycatch (male) | -2435.53 | -2426.65 | -2349.29 | -2358.69 | -2369.79 |
| size_comp | Survey immature females (1982-1988) | -624.89 | -624.2 | -588.64 | -613.95 | -628.55 |
| size_comp | Survey immature females (1989-present) | -2966.22 | -2999.83 | -2840.13 | -3045.46 | -3034.88 |
| size_comp | Survey immature males (1982-1988) | -577.99 | -569.18 | -521.78 | -541.84 | -544.7 |
| size_comp | Survey immature males (1989-present) | -2828.2 | -2815.83 | -2711.59 | -2791.83 | -2807.43 |
| Total | Total | -23714.45 | -25638.68 | $-24378.36$ | -25155.73 | -25297.88 |

Table 13: Parameter estimates and standard deviations. See .CTL files for names on github repo. A fix to display the names of the parameters is on the to do list.

| Parameter | 23.1 | SD | 23.2 | SD |
| :---: | :---: | :---: | :---: | :---: |
| theta[1] | 0.29 | 0 | 0.29 | 0 |
| theta[2] | 0.28 | 0 | 0.28 | 0 |
| theta[5] | 18.95 | 375.26 | 6.93 | 407.52 |
| theta[13] | 10.54 | 0.7 | 11.76 | 0.69 |
| theta[14] | 10.56 | 0.56 | 11.78 | 0.55 |
| theta[15] | 10.64 | 0.44 | 11.87 | 0.44 |
| theta[16] | 11.1 | 0.4 | 12.32 | 0.41 |
| theta[17] | 11.76 | 0.37 | 12.93 | 0.38 |
| theta[18] | 12.32 | 0.31 | 13.39 | 0.32 |
| theta[19] | 12.69 | 0.27 | 13.68 | 0.28 |
| theta[20] | 12.69 | 0.25 | 13.62 | 0.27 |
| theta[21] | 12.52 | 0.25 | 13.3 | 0.28 |
| theta[22] | 12.52 | 0.24 | 13.21 | 0.28 |
| theta[23] | 12.54 | 0.24 | 13.24 | 0.27 |
| theta[24] | 12.37 | 0.24 | 13.08 | 0.27 |
| theta[25] | 12.24 | 0.25 | 13 | 0.26 |
| theta[26] | 12.18 | 0.25 | 13.19 | 0.25 |
| theta[27] | 12.26 | 0.24 | 13.48 | 0.24 |
| theta[28] | 12.24 | 0.2 | 13.1 | 0.22 |
| theta[29] | 11.96 | 0.21 | 12.54 | 0.23 |
| theta[30] | 11.45 | 0.24 | 11.92 | 0.27 |
| theta[31] | 10.6 | 0.28 | 11 | 0.31 |
| theta[32] | 9.57 | 0.33 | 9.89 | 0.36 |
| theta[33] | 8.62 | 0.36 | 8.85 | 0.38 |
| theta[34] | 8.07 | 0.41 | 8.27 | 0.43 |
| theta[35] | 11.88 | 0.46 | 12.57 | 0.45 |
| theta[36] | 12.03 | 0.31 | 12.66 | 0.29 |
| theta[37] | 12.77 | 0.29 | 13.33 | 0.28 |
| theta[38] | 13.32 | 0.23 | 13.72 | 0.25 |
| theta[39] | 12.72 | 0.21 | 13.27 | 0.21 |
| theta[40] | 12.79 | 0.2 | 13.38 | 0.21 |
| theta[41] | 12.69 | 0.2 | 13.33 | 0.21 |
| theta[42] | 12.43 | 0.22 | 13.08 | 0.23 |
| theta[43] | 12.24 | 0.24 | 12.88 | 0.24 |
| theta[44] | 12.04 | 0.24 | 12.7 | 0.26 |
| theta[45] | 11.65 | 0.28 | 12.22 | 0.31 |
| theta[46] | 11.32 | 0.31 | 11.64 | 0.35 |
| theta[47] | 11.14 | 0.32 | 11.23 | 0.4 |
| theta[48] | 10.62 | 0.34 | 10.64 | 0.45 |
| theta[49] | 9.67 | 0.39 | 9.66 | 0.46 |
| theta[50] | 8.71 | 0.42 | 8.68 | 0.46 |
| theta[51] | 7.9 | 0.44 | 7.89 | 0.47 |
| theta[52] | 7.29 | 0.47 | 7.31 | 0.5 |
| theta[53] | 6.86 | 0.51 | 6.89 | 0.54 |
| theta [54] | 6.56 | 0.56 | 6.61 | 0.59 |
| theta[55] | 6.37 | 0.62 | 6.44 | 0.65 |
| theta[56] | 6.28 | 0.7 | 6.35 | 0.73 |
| theta[57] | 13.55 | 0.79 | 12.8 | 0.78 |


| Parameter | 23.1 | SD | 23.2 | SD |
| :---: | :---: | :---: | :---: | :---: |
| theta[58] | 13.54 | 0.64 | 12.8 | 0.64 |
| theta[59] | 13.55 | 0.48 | 12.82 | 0.48 |
| theta[60] | 13.76 | 0.36 | 13.06 | 0.37 |
| theta[61] | 14.53 | 0.29 | 13.83 | 0.34 |
| theta[62] | 14.69 | 0.27 | 14.13 | 0.33 |
| theta[63] | 13.79 | 0.26 | 13.4 | 0.34 |
| theta[64] | 12.58 | 0.28 | 12.24 | 0.35 |
| theta[65] | 11.42 | 0.33 | 11.1 | 0.39 |
| theta[66] | 10.19 | 0.37 | 9.88 | 0.42 |
| theta[67] | 9.44 | 0.41 | 9.12 | 0.46 |
| theta[68] | 9 | 0.46 | 8.62 | 0.5 |
| theta[69] | 8.76 | 0.52 | 8.32 | 0.56 |
| theta[70] | 8.57 | 0.56 | 8.08 | 0.61 |
| theta[71] | 8.38 | 0.59 | 7.86 | 0.64 |
| theta[72] | 8.2 | 0.6 | 7.65 | 0.65 |
| theta[73] | 8.05 | 0.62 | 7.47 | 0.66 |
| theta[74] | 7.94 | 0.64 | 7.32 | 0.67 |
| theta[75] | 7.86 | 0.68 | 7.2 | 0.69 |
| theta[76] | 7.8 | 0.72 | 7.1 | 0.71 |
| theta[77] | 7.76 | 0.78 | 7.04 | 0.76 |
| theta[78] | 7.73 | 0.86 | 7 | 0.83 |
| theta[79] | 7.44 | 1.25 | 6.79 | 1.18 |
| theta[80] | 7.4 | 1.14 | 6.77 | 1.1 |
| theta[81] | 7.35 | 1.03 | 6.64 | 1.05 |
| theta[82] | 7.45 | 0.99 | 6.55 | 0.96 |
| theta[83] | 7.55 | 0.99 | 6.64 | 0.95 |
| theta[84] | 7.68 | 1.01 | 6.9 | 1 |
| theta[85] | 6.47 | 1.03 | 5.79 | 1.02 |
| theta[86] | 5.15 | 1.05 | 4.52 | 1.04 |
| theta[87] | 4.11 | 1.07 | 3.5 | 1.06 |
| theta[88] | 3.35 | 1.08 | 2.75 | 1.07 |
| theta[89] | 2.86 | 1.1 | 2.27 | 1.09 |
| theta[90] | 2.54 | 1.13 | 1.95 | 1.12 |
| theta[91] | 2.31 | 1.15 | 1.71 | 1.14 |
| theta[92] | 2.12 | 1.17 | 1.53 | 1.16 |
| theta[93] | 1.98 | 1.2 | 1.39 | 1.19 |
| theta[94] | 1.87 | 1.22 | 1.28 | 1.21 |
| theta[95] | 1.78 | 1.25 | 1.19 | 1.24 |
| theta[96] | 1.71 | 1.27 | 1.12 | 1.26 |
| theta[97] | 1.65 | 1.3 | 1.06 | 1.29 |
| theta[98] | 1.61 | 1.34 | 1.02 | 1.33 |
| theta[99] | 1.59 | 1.38 | 1 | 1.37 |
| theta[100] | 1.58 | 1.43 | 0.99 | 1.42 |
| Grwth[1] | 2.23 | 0.08 | 2.35 | 0.08 |
| Grwth[2] | -0.22 | 0 | -0.21 | 0 |
| Grwth[4] | -0.18 | 0.11 | 0.48 | 0.1 |
| Grwth[5] | -0.3 | 0 | -0.27 | 0 |
| Grwth[10] | 0.04 | 0 | NA | NA |
| Grwth[11] | 0.06 | 0.01 | NA | NA |
| Grwth[12] | 0.1 | 0.01 | NA | NA |
| Grwth[13] | 0.14 | 0.01 | NA | NA |
| Grwth[14] | 0.2 | 0.01 | NA | NA |


| Parameter | 23.1 | SD | 23.2 | SD |
| :---: | :---: | :---: | :---: | :---: |
| Grwth[15] | 0.22 | 0.01 | NA | NA |
| Grwth[16] | 0.24 | 0.01 | NA | NA |
| Grwth[17] | 0.28 | 0.02 | NA | NA |
| Grwth[18] | 0.28 | 0.02 | NA | NA |
| Grwth[19] | 0.25 | 0.02 | NA | NA |
| Grwth[20] | 0.26 | 0.02 | NA | NA |
| Grwth[21] | 0.42 | 0.02 | NA | NA |
| Grwth[22] | 0.82 | 0.02 | NA | NA |
| Grwth[30] | 0.05 | 0.04 | NA | NA |
| Grwth[31] | 0.06 | 0.02 | NA | NA |
| Grwth[32] | 0.43 | 0.04 | NA | NA |
| Grwth[33] | 0.77 | 0.02 | NA | NA |
| Grwth[34] | 0.92 | 0.01 | NA | NA |
| Grwth[35] | 0.96 | 0 | NA | NA |
| log_slx_pars[1] | 4.63 | 0.01 | 4.66 | 0 |
| log_slx_pars[2] | 1.59 | 0.03 | 1.46 | 0.03 |
| log_slx_pars[3] | 4.26 | 0.01 | 4.24 | 0.01 |
| log_slx_pars[4] | 0.98 | 0.03 | 1.04 | 0.03 |
| log_slx_pars[5] | 4.54 | 0.01 | 4.7 | 0.01 |
| log_slx_pars[6] | 2.19 | 0.02 | 2.38 | 0.02 |
| log_slx_pars[7] | 3.76 | 0.05 | 3.81 | 0.06 |
| log_slx_pars[8] | 2.05 | 0.14 | 2.06 | 0.15 |
| log_slx_pars[9] | 3.94 | 0.02 | 3.91 | 0.02 |
| log_slx_pars[10] | 1.27 | 0.05 | 1.22 | 0.06 |
| log_slx_pars[11] | 3.72 | 0.02 | 3.69 | 0.01 |
| log_slx_pars[12] | 1.64 | 0.07 | 1.38 | 0.06 |
| log_slx_pars[13] | 3.86 | 0.01 | 3.78 | 0.01 |
| log_slx_pars[14] | 1.27 | 0.03 | 1.2 | 0.03 |
| log_slx_pars[15] | -3.94 | 0.94 | -4.39 | 0.96 |
| log_slx_pars[16] | -3.94 | 0.85 | -4.39 | 0.87 |
| log_slx_pars[17] | -3.94 | 0.75 | -4.39 | 0.77 |
| log_slx_pars[18] | -3.93 | 0.63 | -4.39 | 0.66 |
| log_slx_pars[19] | -3.91 | 0.52 | -4.38 | 0.53 |
| log_slx_pars[20] | -3.76 | 0.44 | -4.25 | 0.46 |
| log_slx_pars[21] | -3.42 | 0.41 | -3.95 | 0.42 |
| log_slx_pars[22] | -2.95 | 0.39 | -3.55 | 0.39 |
| log_slx_pars[23] | -2.52 | 0.36 | -3.22 | 0.36 |
| log_slx_pars[24] | -2.08 | 0.35 | -2.8 | 0.34 |
| log_slx_pars[25] | -1.89 | 0.33 | -2.55 | 0.33 |
| log_slx_pars[26] | -1.7 | 0.32 | -2.34 | 0.32 |
| log_slx_pars[27] | -1.27 | 0.32 | -1.93 | 0.31 |
| log_slx_pars[28] | -0.91 | 0.32 | -1.46 | 0.31 |
| log_slx_pars[29] | -1.01 | 0.31 | -1.22 | 0.3 |
| log_slx_pars[30] | -1.46 | 0.3 | -1.22 | 0.3 |
| log_slx_pars[31] | -1.7 | 0.31 | -1.25 | 0.31 |
| log_slx_pars[32] | -1.81 | 0.32 | -1.29 | 0.33 |
| log_slx_pars[33] | -1.77 | 0.33 | -1.26 | 0.34 |
| log_slx_pars[34] | -1.58 | 0.37 | -1.17 | 0.38 |
| log_slx_pars[35] | -1.48 | 0.45 | -1.23 | 0.41 |
| log_slx_pars[36] | -1.4 | 0.6 | -1.32 | 0.46 |
| log_slx_pars[37] | -3.36 | 1.19 | -2.72 | 1.17 |
| log_slx_pars[38] | -3.36 | 1.12 | -2.72 | 1.1 |


| Parameter | 23.1 | SD | 23.2 | SD |
| :---: | :---: | :---: | :---: | :---: |
| log_slx_pars[39] | -3.42 | 1.01 | -2.77 | 1 |
| log_slx_pars[40] | -3.65 | 0.88 | -3.1 | 0.88 |
| log_slx_pars[41] | -3.77 | 0.85 | -3.41 | 0.85 |
| log_slx_pars[42] | -2.84 | 0.84 | -2.64 | 0.84 |
| log_slx_pars[43] | -2.31 | 0.84 | -2.16 | 0.84 |
| log_slx_pars[44] | -2.07 | 0.84 | -2.26 | 0.84 |
| log_slx_pars[45] | -2.11 | 0.86 | -2.53 | 0.85 |
| log_slx_pars[46] | -2.29 | 0.88 | -2.93 | 0.87 |
| log_slx_pars[47] | -2.34 | 0.94 | -3.07 | 0.91 |
| log_slx_pars[48] | -2.34 | 1.02 | -3.07 | 1 |
| log_slx_pars[49] | -2.33 | 1.1 | -3.07 | 1.08 |
| log_slx_pars[50] | -2.33 | 1.17 | -3.07 | 1.15 |
| log_slx_pars[51] | -2.33 | 1.24 | -3.07 | 1.22 |
| log_slx_pars[52] | -2.33 | 1.31 | -3.07 | 1.29 |
| log_slx_pars[53] | -2.33 | 1.37 | -3.07 | 1.35 |
| log_slx_pars[54] | -2.33 | 1.43 | -3.07 | 1.41 |
| log_slx_pars[55] | -2.33 | 1.49 | -3.07 | 1.47 |
| log_slx_pars[56] | -2.33 | 1.54 | -3.07 | 1.52 |
| log_slx_pars[57] | -2.33 | 1.59 | -3.07 | 1.58 |
| log_slx_pars[58] | -2.33 | 1.65 | -3.07 | 1.63 |
| log_slx_pars[62] | -0.35 | 0.25 | -0.27 | 0.25 |
| log_slx_pars[63] | -0.66 | 0.22 | -0.58 | 0.22 |
| log_slx_pars[64] | -0.34 | 0.19 | -0.29 | 0.19 |
| log_slx_pars[69] | -0.4 | 0.22 | -0.66 | 0.21 |
| log_slx_pars[70] | -0.49 | 0.26 | -0.81 | 0.26 |
| log_slx_pars[71] | -0.56 | 0.27 | -0.89 | 0.29 |
| log_slx_pars[72] | -0.68 | 0.27 | -0.97 | 0.29 |
| log_slx_pars[73] | -0.6 | 0.3 | -0.9 | 0.31 |
| log_slx_pars[74] | -0.58 | 0.31 | -0.83 | 0.32 |
| log_slx_pars[75] | -0.63 | 0.29 | -0.78 | 0.33 |
| log_slx_pars[76] | -0.75 | 0.25 | -0.77 | 0.34 |
| log_slx_pars[77] | -0.83 | 0.25 | -0.81 | 0.35 |
| log_slx_pars[78] | -0.8 | 0.26 | -0.86 | 0.39 |
| log_slx_pars[79] | -0.74 | 0.28 | -0.92 | 0.45 |
| log_slx_pars[80] | -0.67 | 0.35 | -0.98 | 0.53 |
| log_slx_pars[81] | -0.6 | 0.48 | -1.02 | 0.61 |
| log_slx_pars[82] | -0.58 | 0.63 | -1.05 | 0.7 |
| log_slx_pars[83] | -0.38 | 0.54 | 0 | 0 |
| log_slx_pars[84] | -1 | 0.32 | -0.39 | 0.23 |
| log_slx_pars[85] | -1.29 | 0.29 | -0.23 | 0.27 |
| log_slx_pars[86] | -0.68 | 0.28 | -0.12 | 0.16 |
| log_slx_pars[87] | -0.39 | 0.16 | 0 | 0 |
| log_slx_pars[88] | -0.56 | 0.16 | 0 | 0 |
| log_slx_pars[89] | -0.62 | 0.18 | 0 | 0 |
| log_slx_pars [90] | -0.79 | 0.23 | -0.31 | 0.23 |
| log_slx_pars[91] | -0.73 | 0.35 | -0.43 | 0.33 |
| log_slx_pars[92] | -0.75 | 0.48 | -0.58 | 0.41 |
| log_slx_pars[93] | -0.75 | 0.62 | -0.61 | 0.54 |
| log_slx_pars[94] | -0.74 | 0.75 | -0.61 | 0.67 |
| log_slx_pars[95] | -0.74 | 0.85 | -0.61 | 0.79 |
| log_slx_pars[96] | -0.74 | 0.94 | -0.61 | 0.89 |
| log_slx_pars [97] | -0.74 | 1.03 | -0.61 | 0.98 |


| Parameter | 23.1 | SD | 23.2 | SD |
| :---: | :---: | :---: | :---: | :---: |
| log_slx_pars[98] | -0.74 | 1.11 | -0.61 | 1.06 |
| log_slx_pars[99] | -0.74 | 1.18 | -0.61 | 1.14 |
| log_slx_pars[100] | -0.74 | 1.25 | -0.61 | 1.21 |
| log_slx_pars[101] | -0.74 | 1.31 | -0.61 | 1.27 |
| log_slx_pars[102] | -0.74 | 1.38 | -0.61 | 1.34 |
| log_slx_pars[103] | -0.74 | 1.43 | -0.61 | 1.4 |
| log_slx_pars[104] | -0.74 | 1.49 | -0.61 | 1.46 |
| log_slx_pars[107] | 4.58 | 0 | 4.58 | 0 |
| log_slx_pars[108] | 0.31 | 0.21 | 0.49 | 0.14 |
| log_fbar [1] | -1.06 | 0.08 | -0.97 | 0.1 |
| log_fbar [2] | -6.6 | 0.09 | -6.29 | 0.11 |
| log_fdev[1] | NA | NA | NA | NA |
| log_fdev[2] | NA | NA | NA | NA |
| log_foff[1] | -5.83 | 0.16 | -6.47 | 0.14 |
| log_fdov[1] | NA | NA | NA | NA |
| rec_dev_est | NA | NA | NA | NA |
| logit_rec_prop_est | NA | NA | NA | NA |
| m__dev_est[1] | 1.77 | 0.14 | 1.98 | 0.1 |
| m_dev_est[2] | 2.4 | 0.09 | 0.92 | 0.18 |
| m_dev_est[4] | 0.62 | 0.39 | 0 | 0 |
| m__dev_est[5] | 2.23 | 0.08 | 2.26 | 0.08 |
| m_dev_est[7] | 0.8 | 0.4 | 0.91 | 0.33 |
| m__dev_est[8] | 1.88 | 0.17 | 1.63 | 0.2 |
| m_dev__est[10] | 2.5 | 0.22 | 2.48 | 0.23 |
| m_dev_est[11] | 2.51 | 0.71 | 2.19 | 0.88 |
| m_mat_mult[1] | 0 | 0.05 | -0.07 | 0.04 |
| m_mat_mult[2] | 0.06 | 0.05 | 0.28 | 0.05 |
| survey_q[1] | 0.34 | 0.08 | 0.52 | 0.15 |
| survey_q[2] | 0.39 | 0.04 | 0.46 | 0.04 |
| survey_q[3] | 0.22 | 0.03 | 0.11 | 0.02 |
| survey_q[4] | 0.49 | 0.03 | 0.27 | 0.01 |
| sd_log_recruits | NA | NA | NA | NA |
| ParsOut | NA | NA | NA | NA |
| sd__log_ssb | NA | NA | NA | NA |
| sd_last_ssb | 38.36 | 2.71 | 63.24 | 4.44 |
| log_slx_pars[61] | NA | NA | 0 | 0 |

Table 14: Parameter estimates and standard deviations from considered models.

| Parameter | 23.3b | SD | 23.3 | SD | 23.3 | SD |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| theta[1] | 0.29 | 0 | 0.29 | 0 | 0.55 | 0.02 |
| theta[2] | 0.27 | 0 | 0.27 | 0 | 0.48 | 0.03 |
| theta[5] | 5.27 | 172.79 | 6.23 | 1308.1 | 6.79 | 166.31 |
| theta[13] | 8.84 | 0.69 | 9.57 | 0.73 | 10.34 | 0.76 |
| theta[14] | 8.86 | 0.55 | 9.58 | 0.6 | 10.35 | 0.64 |
| theta[15] | 8.92 | 0.43 | 9.63 | 0.47 | 10.39 | 0.51 |
| theta[16] | 9.3 | 0.38 | 9.87 | 0.38 | 10.56 | 0.41 |
| theta[17] | 9.98 | 0.36 | 10.57 | 0.39 | 11.19 | 0.41 |
| theta[18] | 10.76 | 0.32 | 11.25 | 0.35 | 11.83 | 0.38 |
| theta[19] | 11.45 | 0.26 | 11.68 | 0.27 | 12.23 | 0.3 |
| theta[20] | 11.69 | 0.22 | 11.86 | 0.25 | 12.38 | 0.27 |
| theta[21] | 11.61 | 0.22 | 11.9 | 0.25 | 12.39 | 0.27 |
| theta[22] | 11.65 | 0.22 | 11.93 | 0.25 | 12.37 | 0.26 |
| theta[23] | 11.73 | 0.21 | 11.92 | 0.24 | 12.29 | 0.25 |
| theta[24] | 11.56 | 0.21 | 11.71 | 0.24 | 12.02 | 0.25 |
| theta[25] | 11.38 | 0.21 | 11.49 | 0.24 | 11.74 | 0.26 |
| theta[26] | 11.3 | 0.2 | 11.33 | 0.22 | 11.46 | 0.24 |
| theta[27] | 11.34 | 0.18 | 11.37 | 0.21 | 11.41 | 0.23 |
| theta[28] | 11.12 | 0.15 | 11.16 | 0.16 | 11.31 | 0.18 |
| theta[29] | 10.43 | 0.15 | 10.46 | 0.17 | 10.59 | 0.18 |
| theta[30] | 9.55 | 0.19 | 9.57 | 0.2 | 9.71 | 0.21 |
| theta[31] | 8.53 | 0.25 | 8.56 | 0.26 | 8.72 | 0.26 |
| theta[32] | 7.54 | 0.31 | 7.56 | 0.31 | 7.76 | 0.32 |
| theta[33] | 6.74 | 0.34 | 6.77 | 0.34 | 7 | 0.35 |
| theta[34] | 6.31 | 0.4 | 6.34 | 0.4 | 6.58 | 0.41 |
| theta [35] | 13.74 | 0.51 | 14.04 | 0.41 | 14.86 | 0.43 |
| theta[36] | 13.45 | 0.29 | 13.75 | 0.21 | 14.58 | 0.22 |
| theta[37] | 13.02 | 0.23 | 13.3 | 0.24 | 14.13 | 0.23 |
| theta[38] | 13.11 | 0.2 | 13.34 | 0.2 | 14.15 | 0.19 |
| theta[39] | 12.39 | 0.19 | 12.67 | 0.19 | 13.41 | 0.19 |
| theta[40] | 12.46 | 0.21 | 12.71 | 0.2 | 13.38 | 0.2 |
| theta[41] | 12.3 | 0.2 | 12.49 | 0.19 | 13.1 | 0.18 |
| theta[42] | 11.92 | 0.24 | 12.13 | 0.22 | 12.71 | 0.22 |
| theta[43] | 11.62 | 0.25 | 11.81 | 0.25 | 12.36 | 0.24 |
| theta[44] | 11.18 | 0.24 | 11.37 | 0.24 | 11.87 | 0.23 |
| theta [45] | 10.51 | 0.28 | 10.69 | 0.28 | 11.16 | 0.28 |
| theta[46] | 10.1 | 0.3 | 10.26 | 0.3 | 10.71 | 0.29 |
| theta[47] | 9.97 | 0.27 | 10.09 | 0.27 | 10.52 | 0.25 |
| theta[48] | 9.31 | 0.28 | 9.44 | 0.28 | 9.8 | 0.27 |
| theta[49] | 8.16 | 0.33 | 8.29 | 0.33 | 8.63 | 0.33 |
| theta[50] | 7.13 | 0.37 | 7.25 | 0.37 | 7.6 | 0.37 |
| theta[51] | 6.35 | 0.42 | 6.46 | 0.42 | 6.82 | 0.42 |
| theta[52] | 5.79 | 0.46 | 5.89 | 0.46 | 6.24 | 0.46 |
| theta[53] | 5.38 | 0.51 | 5.47 | 0.51 | 5.81 | 0.51 |
| theta[54] | 5.08 | 0.56 | 5.17 | 0.55 | 5.49 | 0.55 |
| theta[55] | 4.88 | 0.61 | 4.96 | 0.61 | 5.27 | 0.6 |
| theta[56] | 4.77 | 0.69 | 4.85 | 0.69 | 5.15 | 0.67 |
| theta[57] | 9.63 | 0.56 | 11.23 | 0.57 | 12.55 | 0.68 |
| theta[58] | 9.69 | 0.41 | 11.41 | 0.5 | 12.61 | 0.58 |
| theta[59] | 9.99 | 0.3 | 11.71 | 0.42 | 12.7 | 0.46 |


| Parameter | 23.3b | SD | 23.3 | SD | 23.3 | SD |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| theta[60] | 11.03 | 0.24 | 12.19 | 0.29 | 12.86 | 0.29 |
| theta[61] | 12.78 | 0.15 | 13.24 | 0.2 | 13.75 | 0.21 |
| theta[62] | 13.77 | 0.11 | 13.57 | 0.15 | 13.93 | 0.16 |
| theta[63] | 13.32 | 0.12 | 13.12 | 0.15 | 13.37 | 0.15 |
| theta[64] | 12.35 | 0.16 | 12.29 | 0.19 | 12.56 | 0.2 |
| theta[65] | 11.29 | 0.25 | 11.24 | 0.25 | 11.56 | 0.26 |
| theta[66] | 10.11 | 0.3 | 10.09 | 0.31 | 10.46 | 0.31 |
| theta[67] | 9.3 | 0.36 | 9.28 | 0.37 | 9.72 | 0.37 |
| theta[68] | 8.76 | 0.43 | 8.71 | 0.42 | 9.22 | 0.43 |
| theta[69] | 8.41 | 0.5 | 8.38 | 0.5 | 8.87 | 0.49 |
| theta[70] | 8.11 | 0.56 | 8.1 | 0.55 | 8.59 | 0.53 |
| theta[71] | 7.83 | 0.59 | 7.83 | 0.58 | 8.35 | 0.57 |
| theta[72] | 7.54 | 0.6 | 7.55 | 0.59 | 8.14 | 0.6 |
| theta[73] | 7.29 | 0.6 | 7.3 | 0.59 | 7.97 | 0.62 |
| theta[74] | 7.07 | 0.61 | 7.09 | 0.59 | 7.82 | 0.65 |
| theta[75] | 6.89 | 0.61 | 6.91 | 0.6 | 7.7 | 0.67 |
| theta[76] | 6.74 | 0.63 | 6.77 | 0.61 | 7.6 | 0.7 |
| theta[77] | 6.64 | 0.66 | 6.66 | 0.64 | 7.53 | 0.74 |
| theta[78] | 6.58 | 0.73 | 6.61 | 0.71 | 7.5 | 0.82 |
| theta[79] | -16.35 | 1.44 | -13.47 | 3757.8 | -14.18 | 2.58 |
| theta[80] | -16.4 | 1.39 | -13.45 | 3757.8 | -14.26 | 2.54 |
| theta[81] | -16.66 | 1.33 | -13.6 | 3757.8 | -14.42 | 2.49 |
| theta[82] | -16.11 | 1.32 | -13.59 | 3757.8 | -14.56 | 2.45 |
| theta[83] | -15.14 | 1.32 | -13.25 | 3757.8 | -14.37 | 2.44 |
| theta[84] | -14.17 | 1.31 | -12.87 | 3757.8 | -14.19 | 2.44 |
| theta[85] | -14.99 | 1.31 | -13.73 | 3757.8 | -15.17 | 2.44 |
| theta[86] | -16.12 | 1.33 | -14.8 | 3757.8 | -16.23 | 2.45 |
| theta[87] | -17.08 | 1.34 | -15.72 | 3757.8 | -17.13 | 2.45 |
| theta[88] | -17.8 | 1.34 | -16.41 | 3757.8 | -17.82 | 2.46 |
| theta[89] | -18.3 | 1.35 | -16.9 | 3757.8 | -18.3 | 2.46 |
| theta[90] | -18.64 | 1.36 | -17.24 | 3757.8 | -18.64 | 2.47 |
| theta[91] | -18.91 | 1.36 | -17.51 | 3757.8 | -18.91 | 2.47 |
| theta[92] | -19.13 | 1.35 | -17.74 | 3757.8 | -19.13 | 2.46 |
| theta[93] | -19.32 | 1.34 | -17.93 | 3757.8 | -19.32 | 2.45 |
| theta[94] | -19.48 | 1.32 | -18.09 | 3757.8 | -19.48 | 2.44 |
| theta[95] | -19.62 | 1.28 | -18.23 | 3757.8 | -19.62 | 2.43 |
| theta [96] | -19.74 | 1.24 | -18.35 | 3757.8 | -19.74 | 2.4 |
| theta [97] | -19.84 | 1.19 | -18.45 | 3757.8 | -19.84 | 2.38 |
| theta[98] | -19.92 | 1.13 | -18.53 | 3757.8 | -19.92 | 2.35 |
| theta[99] | -19.97 | 1.05 | -18.58 | 3757.8 | -19.97 | 2.31 |
| theta[100] | -20 | 0.95 | -18.61 | 3757.8 | -20 | 2.27 |
| Grwth[1] | 2.19 | 0.08 | 2.26 | 0.08 | 2.19 | 0.08 |
| Grwth[2] | -0.22 | 0 | -0.22 | 0 | -0.22 | 0 |
| Grwth[4] | 0.45 | 0.1 | 0.24 | 0.11 | 0.2 | 0.11 |
| Grwth[5] | -0.27 | 0 | -0.28 | 0 | -0.28 | 0 |
| log_slx_pars[1] | 4.68 | 0.01 | 4.68 | 0 | 4.67 | 0 |
| log_slx_pars [2] | 1.46 | 0.03 | 1.45 | 0.03 | 1.4 | 0.03 |
| log_slx_pars[3] | 4.23 | 0.01 | 4.23 | 0.01 | 4.24 | 0 |
| log_slx_pars[4] | 1.05 | 0.03 | 1.05 | 0.03 | 1.01 | 0.03 |
| log_slx_pars[5] | 4.8 | 0.02 | 4.79 | 0.02 | 4.73 | 0.01 |
| log_slx_pars[6] | 2.44 | 0.02 | 2.41 | 0.02 | 2.28 | 0.02 |
| log_slx_pars[95] | 4.58 | 0 | 4.58 | 0 | 4.58 | 0 |


| Parameter | 23.3b | SD | 23.3 | SD | 23.3 | SD |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| log_slx_pars [96] | 0.55 | 0.11 | 0.55 | 0.11 | 0.61 | 0.1 |
| log_fbar [1] | 0.17 | 0.07 | 0.09 | 0.07 | 0.15 | 0.06 |
| log_fbar[2] | -5.08 | 0.09 | -5.15 | 0.09 | -5.15 | 0.08 |
| log_fdev[1] | NA | NA | NA | NA | NA | NA |
| log_fdev[2] | NA | NA | NA | NA | NA | NA |
| log _foff[1] | -7.61 | 0.1 | -7.41 | 0.1 | -7.23 | 0.1 |
| log_fdov[1] | NA | NA | NA | NA | NA | NA |
| rec_dev_est | NA | NA | NA | NA | NA | NA |
| logit_rec_prop_est | NA | NA | NA | NA | NA | NA |
| m__dev_est[1] | 1.51 | 0.13 | 1.78 | 0.11 | 1.15 | 0.12 |
| m__dev_est[2] | 0.39 | 0.28 | 0.9 | 0.17 | 0 | 0 |
| m_dev_est[4] | 0 | 0 | 0 | 0 | 0 | 0 |
| m_dev_est[5] | 2.52 | 0.08 | 2.47 | 0.07 | 2.08 | 0.06 |
| m__dev_est [7] | 0.56 | 0.48 | 0.89 | 0.35 | 0.13 | 0.43 |
| m__dev_est [8] | 1.99 | 0.15 | 1.82 | 0.17 | 0.98 | 0.25 |
| $m \ldots d e v \_$est [10] | 2.67 | 0.19 | 2.73 | 0.29 | 2.08 | 0.23 |
| $m \_$dev__est[11] | 2.48 | 0.81 | 1.74 | 1.3 | 1.32 | 0.96 |
| m_mat_mult[1] | -0.47 | 0.04 | -0.21 | 0.04 | -0.38 | 0.04 |
| m_mat_mult[2] | -0.01 | 0.05 | 0.08 | 0.05 | 0.09 | 0.05 |
| sd_log_recruits | NA | NA | NA | NA | NA | NA |
| ParsOut | NA | NA | NA | NA | NA | NA |
| sd__log_ssb | NA | NA | NA | NA | NA | NA |
| sd_last_ssb | 50.04 | 3.14 | 55.01 | 3.5 | 29.05 | 2.14 |
| log_slx_pars[7] | NA | NA | -2.73 | 0.24 | -3.39 | 0.25 |
| log_slx_pars[8] | NA | NA | -2.31 | 0.16 | -2.94 | 0.19 |
| log_slx_pars [9] | NA | NA | -1.95 | 0.15 | -2.55 | 0.18 |
| log_slx_pars[10] | NA | NA | -1.2 | 0.12 | -1.67 | 0.16 |
| log_slx_pars[11] | NA | NA | -1.3 | 0.12 | -1.74 | 0.15 |
| log_slx_pars[12] | NA | NA | -1.18 | 0.11 | -1.55 | 0.13 |
| log_slx_pars[13] | NA | NA | -0.97 | 0.1 | -1.27 | 0.12 |
| log_slx_pars[14] | NA | NA | -0.99 | 0.1 | -1.24 | 0.12 |
| log_slx_pars[15] | NA | NA | -1.14 | 0.11 | -1.35 | 0.12 |
| log_slx_pars[16] | NA | NA | -1.13 | 0.11 | -1.28 | 0.12 |
| log_slx_pars[17] | NA | NA | -1.05 | 0.11 | -1.15 | 0.12 |
| log_slx_pars[18] | NA | NA | -1.01 | 0.12 | -1.08 | 0.12 |
| log_slx_pars[19] | NA | NA | -0.97 | 0.12 | -1 | 0.12 |
| log_slx_pars [20] | NA | NA | -0.84 | 0.11 | -0.84 | 0.11 |
| log_slx_pars[21] | NA | NA | -0.78 | 0.11 | -0.76 | 0.11 |
| log_slx_pars[22] | NA | NA | -0.68 | 0.11 | -0.68 | 0.11 |
| log_slx_pars[23] | NA | NA | -0.54 | 0.1 | -0.56 | 0.1 |
| log_slx_pars [24] | NA | NA | -0.43 | 0.1 | -0.44 | 0.1 |
| log_slx_pars[25] | NA | NA | -0.33 | 0.09 | -0.34 | 0.09 |
| log_slx_pars[26] | NA | NA | -0.21 | 0.08 | -0.22 | 0.08 |
| log_slx_pars[27] | NA | NA | -0.1 | 0.08 | -0.11 | 0.08 |
| log_slx_pars[28] | NA | NA | -0.04 | 0.11 | -0.04 | 0.11 |
| log_slx_pars[29] | NA | NA | -3.14 | 0.31 | -4.62 | 0.25 |
| log_slx_pars[30] | NA | NA | -3.5 | 0.32 | -4.97 | 0.22 |
| log_slx_pars[31] | NA | NA | -3.39 | 0.23 | -4.41 | 0.2 |
| log_slx_pars[32] | NA | NA | -2.02 | 0.16 | -2.45 | 0.15 |
| log_slx_pars[33] | NA | NA | -1.24 | 0.13 | -1.53 | 0.13 |
| log_slx_pars[34] | NA | NA | -0.6 | 0.09 | -0.66 | 0.1 |
| log_slx_pars[35] | NA | NA | -0.58 | 0.09 | -0.51 | 0.09 |


| Parameter | 23.3b | SD | 23.3 | SD | 23.3 | SD |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| log_slx_pars[36] | NA | NA | -0.76 | 0.11 | -0.72 | 0.11 |
| log_slx_pars [37] | NA | NA | -0.81 | 0.12 | -0.8 | 0.12 |
| log_slx_pars [38] | NA | NA | -0.91 | 0.14 | -0.92 | 0.14 |
| log_slx_pars [39] | NA | NA | -0.91 | 0.14 | -0.91 | 0.15 |
| log_slx_pars[40] | NA | NA | -0.87 | 0.14 | -0.87 | 0.14 |
| log_slx_pars[41] | NA | NA | -0.83 | 0.14 | -0.84 | 0.14 |
| log_slx_pars[42] | NA | NA | -0.78 | 0.13 | -0.79 | 0.13 |
| log_slx_pars[43] | NA | NA | -0.72 | 0.13 | -0.72 | 0.13 |
| log_slx_pars[44] | NA | NA | -0.63 | 0.12 | -0.63 | 0.12 |
| log_slx_pars[45] | NA | NA | -0.53 | 0.11 | -0.53 | 0.11 |
| log_slx_pars[46] | NA | NA | -0.42 | 0.1 | -0.42 | 0.1 |
| log_slx_pars[47] | NA | NA | -0.31 | 0.09 | -0.31 | 0.09 |
| log_slx_pars[48] | NA | NA | -0.2 | 0.08 | -0.2 | 0.08 |
| log_slx_pars[49] | NA | NA | -0.08 | 0.08 | -0.08 | 0.08 |
| log_slx_pars [50] | NA | NA | -0.01 | 0.11 | -0.01 | 0.11 |
| log_slx_pars[51] | NA | NA | -3.93 | 0.23 | -4.31 | 0.17 |
| log_slx_pars [52] | NA | NA | -2.92 | 0.11 | -3.2 | 0.14 |
| log_slx_pars[53] | NA | NA | -2.05 | 0.1 | -2.37 | 0.11 |
| log_slx_pars [54] | NA | NA | -1.26 | 0.08 | -1.58 | 0.1 |
| log_slx_pars[55] | NA | NA | -1.18 | 0.07 | -1.39 | 0.08 |
| log_slx_pars[56] | NA | NA | -1.08 | 0.06 | -1.21 | 0.07 |
| log_slx_pars[57] | NA | NA | -1.04 | 0.06 | -1.1 | 0.06 |
| log_slx_pars[58] | NA | NA | -1.05 | 0.06 | -1.07 | 0.06 |
| log_slx_pars [59] | NA | NA | -1.21 | 0.06 | -1.19 | 0.06 |
| log_slx_pars [60] | NA | NA | -1.26 | 0.06 | -1.19 | 0.06 |
| log_slx_pars[61] | NA | NA | -1.25 | 0.06 | -1.15 | 0.06 |
| log_slx_pars [62] | NA | NA | -1.24 | 0.06 | -1.1 | 0.06 |
| log_slx_pars[63] | NA | NA | -1.17 | 0.06 | -1.03 | 0.06 |
| log_slx_pars [64] | NA | NA | -0.99 | 0.06 | -0.84 | 0.06 |
| log_slx_pars[65] | NA | NA | -0.78 | 0.07 | -0.67 | 0.06 |
| log_slx_pars [66] | NA | NA | -0.55 | 0.07 | -0.51 | 0.07 |
| log_slx_pars [67] | NA | NA | -0.38 | 0.07 | -0.38 | 0.07 |
| log_slx_pars [68] | NA | NA | -0.26 | 0.07 | -0.27 | 0.07 |
| log_slx_pars [69] | NA | NA | -0.2 | 0.07 | -0.22 | 0.07 |
| log_slx_pars[70] | NA | NA | -0.16 | 0.07 | -0.16 | 0.07 |
| log_slx_pars[71] | NA | NA | -0.12 | 0.08 | -0.13 | 0.08 |
| log_slx_pars[72] | NA | NA | -0.09 | 0.11 | -0.09 | 0.11 |
| log_slx_pars[73] | NA | NA | -4.12 | 0.16 | -4.67 | 0.21 |
| log_slx_pars[74] | NA | NA | -3.42 | 0.09 | -3.81 | 0.11 |
| log_slx_pars[75] | NA | NA | -2.85 | 0.09 | -3.08 | 0.09 |
| log_slx_pars[76] | NA | NA | -1.22 | 0.07 | -1.38 | 0.07 |
| log_slx_pars [77] | NA | NA | -0.63 | 0.06 | -0.76 | 0.06 |
| log_slx_pars[78] | NA | NA | -0.27 | 0.05 | -0.35 | 0.05 |
| log_slx_pars [79] | NA | NA | -0.39 | 0.05 | -0.41 | 0.06 |
| log_slx_pars[80] | NA | NA | -0.99 | 0.07 | -0.97 | 0.08 |
| log_slx_pars[81] | NA | NA | -1.06 | 0.1 | -1 | 0.1 |
| log_slx_pars[82] | NA | NA | -1.25 | 0.12 | -1.17 | 0.12 |
| log_slx_pars[83] | NA | NA | -1.2 | 0.15 | -1.12 | 0.14 |
| log_slx_pars[84] | NA | NA | -0.91 | 0.15 | -0.89 | 0.14 |
| log_slx_pars[85] | NA | NA | -0.83 | 0.14 | -0.83 | 0.14 |
| log_slx_pars[86] | NA | NA | -0.78 | 0.13 | -0.78 | 0.13 |
| log_slx_pars[87] | NA | NA | -0.71 | 0.12 | -0.71 | 0.12 |


| Parameter | 23.3 b | SD | 23.3 | SD | 23.3 | SD |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| log_slx_pars[88] | NA | NA | -0.62 | 0.12 | -0.62 | 0.12 |
| log_slx_pars[89] | NA | NA | -0.52 | 0.11 | -0.52 | 0.11 |
| log_slx_pars[90] | NA | NA | -0.42 | 0.1 | -0.42 | 0.1 |
| log_slx_pars[91] | NA | NA | -0.31 | 0.09 | -0.31 | 0.09 |
| log_slx_pars[92] | NA | NA | -0.2 | 0.08 | -0.2 | 0.08 |
| log_slx_pars[93] | NA | NA | -0.08 | 0.08 | -0.08 | 0.08 |
| log_slx_pars[94] | NA | NA | -0.01 | 0.11 | -0.01 | 0.11 |

Table 15: Management quantities derived from maximum likelihood estimates by model using Tier 3 reference points. Reported natural mortality is for mature males, average recruitment is for males, and status and MMB were estimates for February 15 of the completed crab year.

| Model | MMB | B35 | F35 | FOFL | OFL | M | avg_rec | Status |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 22.1 | 41.21 | 183.15 | 1.50 | 0.32 | 10.32 | 0.28 | 164.02 | 0.23 |
| 23.1 | 56.41 | 189.24 | 1.60 | 0.30 | 8.58 | 0.29 | 169.90 | 0.30 |
| 23.2 | 135.43 | 132.46 | 71.89 | 30.14 | 37.10 | 0.29 | 222.75 | 1.02 |
| 23.3 | 81.96 | 130.98 | 33.47 | 10.49 | 12.12 | 0.29 | 91.92 | 0.63 |
| 23.3 a | 92.39 | 155.91 | 53.25 | 14.96 | 15.44 | 0.29 | 141.66 | 0.59 |
| 23.3 b | 68.15 | 110.01 | 205.67 | 37.49 | 11.56 | 0.55 | 351.66 | 0.62 |

Table 16: Management quantities derived from maximum likelihood estimates by model using Tier 4 reference points. Reported natural mortality is for mature males, average recruitment is for males, and status and MMB were estimates for February 15 of the completed crab year.

| Model | MMB | BMSY | FMSY | FOFL | OFL | M | avg_rec | Status |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 23.1 | 55.85 | 264.57 | 0.28 | 0.00 | 0.09 | 0.28 | 166.64 | 0.21 |
| 23.2 | 135.43 | 519.67 | 0.29 | 0.00 | 0.05 | 0.29 | 222.75 | 0.26 |
| 23.3 | 81.96 | 236.84 | 0.29 | 0.05 | 0.29 | 0.29 | 91.92 | 0.35 |
| 23.3 a | 92.39 | 273.83 | 0.29 | 0.05 | 0.31 | 0.29 | 141.66 | 0.34 |
| 23.3 b | 68.15 | 232.32 | 0.55 | 0.00 | 0.03 | 0.55 | 351.66 | 0.29 |

Table 17: Survey-based tier 4 status and OFL (1,000 t). 'Males_com' is the observed biomass of $>101 \mathrm{~mm}$ carapace width males. Status represents the status of the population after the completed fishing year and is used for overfished declarations. Proj_Status represents the projected fishery status after the coming fishery removes the OFL and is used in the harvest control rule. 'Years' indicates the year range used to calculate reference points. ' M ' is the natural mortality for mature male crab.

| Year | Tier | BMSY | Males_com | Status | FOFL | OFL | Years | M |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2023 / 2024$ | $3 c$ | 59.64 | 9.996 | 0.1676 | 0 | 0 | $1982-2022$ | 0.27 |

Table 18: Maximum likelihood estimates of mature male biomass (MMB), mature female biomass (FMB), and males $>101 \mathrm{~mm}$ biomass (1000 t) and numbers (in millions) at the time of the survey from the author-preferred model. Columns 2-5 are subject to survey selectivity; columns 6-9 are the population values.

| Survey year | FMB | MMB | Male >101 biomass | Male >101 (millions) | FMB | MMB | Male >101 biomass | Male $>101$ (millions) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1982 | 90.34 | 122.8 | 30.91 | 55.91 | 168.8 | 290.2 | 51.75 | 95.61 |
| 1983 | 68.67 | 134 | 35.23 | 61.38 | 128.3 | 316.8 | 57.05 | 102.2 |
| 1984 | 52.49 | 159.6 | 51.92 | 86.59 | 98.07 | 365.1 | 80.85 | 139.6 |
| 1985 | 41.16 | 176.4 | 58.81 | 95.54 | 76.93 | 402.6 | 89.54 | 151.1 |
| 1986 | 51.19 | 189.6 | 54.11 | 88.3 | 94.72 | 446.9 | 82.74 | 140.2 |
| 1987 | 126.5 | 224.7 | 60.11 | 99.26 | 237.3 | 536.9 | 92.84 | 158.9 |
| 1988 | 198 | 274.6 | 75.44 | 124 | 377.9 | 654.4 | 116 | 197.6 |
| 1989 | 274.5 | 314.8 | 110.2 | 182.3 | 454.8 | 763.6 | 150.5 | 255.6 |
| 1990 | 248.4 | 345.5 | 145 | 238 | 422.1 | 780.5 | 196.9 | 331.9 |
| 1991 | 204.6 | 284.4 | 126.8 | 210.7 | 350.4 | 654.1 | 173.5 | 295.7 |
| 1992 | 164.2 | 244.3 | 112.6 | 174.4 | 281.6 | 545.2 | 147.8 | 235.8 |
| 1993 | 137.1 | 204.9 | 73.01 | 116.8 | 233.5 | 489.6 | 97.47 | 160.2 |
| 1994 | 125.8 | 207.7 | 48.27 | 78.32 | 211.8 | 549.2 | 65.26 | 108.9 |
| 1995 | 131.7 | 220.6 | 41.64 | 71.11 | 218.6 | 610.4 | 58.29 | 102 |
| 1996 | 145.3 | 286.2 | 88.33 | 148.9 | 241.8 | 720.8 | 122.1 | 210.9 |
| 1997 | 140.8 | 305.7 | 126.7 | 207.1 | 238.1 | 701.3 | 171.6 | 288.4 |
| 1998 | 121 | 250.3 | 113.6 | 181.6 | 207.2 | 557.5 | 151.7 | 249.2 |
| 1999 | 96.71 | 175.2 | 69.37 | 108.3 | 166.7 | 404.1 | 91.74 | 147.8 |
| 2000 | 79.47 | 141.9 | 53.1 | 81.92 | 135.8 | 331.2 | 69.78 | 111.2 |
| 2001 | 75.38 | 114.7 | 37.43 | 59.19 | 126.9 | 279.2 | 49.97 | 81.57 |
| 2002 | 73.35 | 100.5 | 31.27 | 52.37 | 123.9 | 247.4 | 43.23 | 74.37 |
| 2003 | 65.17 | 97.85 | 38.52 | 63.49 | 111.2 | 225.5 | 52.51 | 88.89 |
| 2004 | 58.49 | 93.6 | 42.53 | 67.97 | 99.07 | 207.8 | 56.74 | 93.23 |
| 2005 | 71.61 | 98.93 | 37.87 | 59.12 | 115.6 | 232.9 | 49.97 | 80.44 |
| 2006 | 111 | 103.7 | 30.78 | 49.96 | 180.4 | 261.1 | 41.73 | 69.77 |
| 2007 | 113.6 | 132.1 | 44.1 | 72.48 | 192.1 | 327.4 | 59.89 | 101 |
| 2008 | 98.43 | 150.9 | 55.83 | 91.56 | 168.7 | 369.1 | 75.88 | 127.9 |
| 2009 | 83.57 | 170.3 | 76.51 | 121.1 | 142.5 | 381.2 | 101.6 | 165.5 |
| 2010 | 93.91 | 160.9 | 77.55 | 121.9 | 153.6 | 352.2 | 102.5 | 165.9 |
| 2011 | 128.8 | 170.1 | 82.98 | 125.8 | 211.2 | 364.6 | 107.2 | 167.4 |
| 2012 | 129.5 | 124.2 | 42.29 | 67.95 | 219 | 301.4 | 56.85 | 94.13 |
| 2013 | 113.5 | 110.9 | 33.68 | 56.9 | 194 | 278.5 | 46.68 | 80.85 |
| 2014 | 98.1 | 110.6 | 39.13 | 65.03 | 167.2 | 265.1 | 53.57 | 91.34 |
| 2015 | 91.07 | 85.47 | 24.45 | 40.54 | 154.3 | 215.5 | 33.59 | 57.27 |
| 2016 | 88.37 | 77.73 | 17.19 | 28.61 | 147.8 | 205.6 | 23.72 | 40.59 |
| 2017 | 127.1 | 93.34 | 17.89 | 30.24 | 202.4 | 252.5 | 24.87 | 43.11 |
| 2018 | 251.5 | 160.6 | 26.52 | 44.31 | 394.1 | 444.3 | 36.52 | 62.61 |
| 2019 | 131.8 | 162.6 | 33.43 | 59.9 | 206.9 | 432.2 | 48.31 | 88.12 |
| 2020 | 24.69 | 76.12 | 11.66 | 21.33 | 38.87 | 209.5 | 17.25 | 32.06 |
| 2021 | 18.95 | 51.1 | 5.84 | 10.29 | 29.85 | 146.6 | 8.41 | 15.15 |
| 2022 | 15.48 | 41.39 | 7.02 | 11.33 | 24.21 | 114 | 9.5 | 15.79 |
| 2023 | 14.73 | 31.9 | 5.98 | 9.5 | 23.15 | 86.75 | 8 | 13.1 |

Table 19: Maximum likelihood estimates of total numbers of crab (billions), not subject to survey selectivity at the time of the survey.

| Survey year | Total numbers |
| :---: | :---: |
| 1983 | 9.346 |
| 1984 | 10.58 |
| 1985 | 19.84 |
| 1986 | 24.53 |
| 1987 | 24.15 |
| 1988 | 20.88 |
| 1989 | 16.71 |
| 1990 | 14.09 |
| 1991 | 15.76 |
| 1992 | 18.03 |
| 1993 | 16.36 |
| 1994 | 15.29 |
| 1995 | 12.83 |
| 1996 | 10.26 |
| 1997 | 8.271 |
| 1998 | 7.68 |
| 1999 | 6.685 |
| 2000 | 6.159 |
| 2001 | 5.22 |
| 2002 | 6.237 |
| 2003 | 6.782 |
| 2004 | 11.49 |
| 2005 | 11.01 |
| 2006 | 9.161 |
| 2007 | 7.392 |
| 2008 | 8.72 |
| 2009 | 11.87 |
| 2010 | 9.899 |
| 2011 | 8.579 |
| 2012 | 7.062 |
| 2013 | 7.473 |
| 2014 | 7.018 |
| 2015 | 14.32 |
| 2016 | 23.31 |
| 2017 | 32.05 |
| 2018 | 24.74 |
| 2019 | 8.068 |
| 2020 | 1.894 |
| 2021 | 1.696 |
| 2022 | 1.412 |
| 2023 |  |
|  |  |

Table 20: Maximum likelihood estimates of mature male biomass at mating, male recruitment (billions), and fully-selected total fishing mortaltiy.

| Survey year | Mature male biomass | Male recruits | Fishing mortality |
| :---: | :---: | :---: | :---: |
| 1982 | 221.1 | 2.34 | 0.54 |
| 1983 | 242.2 | 3.31 | 0.48 |
| 1984 | 263.1 | 4.55 | 0.93 |
| 1985 | 278.9 | 4.32 | 1.34 |
| 1986 | 315.7 | 0.34 | 1.45 |
| 1987 | 376.5 | 2.02 | 1.75 |
| 1988 | 462.3 | 0.21 | 1.49 |
| 1989 | 543.6 | 1.29 | 1.22 |
| 1990 | 485.7 | 4.68 | 2.56 |
| 1991 | 406.8 | 5.16 | 2.92 |
| 1992 | 321.8 | 0.91 | 3.49 |
| 1993 | 327.5 | 0.23 | 1.97 |
| 1994 | 400 | 0.08 | 1.38 |
| 1995 | 454.8 | 0.1 | 1.44 |
| 1996 | 525.2 | 0.54 | 1.15 |
| 1997 | 474.6 | 1.41 | 1.46 |
| 1998 | 370.6 | 0.06 | 1.42 |
| 1999 | 312.7 | 0.25 | 0.27 |
| 2000 | 254.6 | 0.3 | 0.31 |
| 2001 | 206.4 | 1.99 | 0.76 |
| 2002 | 185.6 | 1.17 | 0.69 |
| 2003 | 171.6 | 2.17 | 0.39 |
| 2004 | 156.6 | 1.81 | 0.38 |
| 2005 | 168.2 | 0.21 | 0.81 |
| 2006 | 192.3 | 0.32 | 0.97 |
| 2007 | 234.9 | 1.89 | 1.25 |
| 2008 | 272.3 | 1.26 | 0.8 |
| 2009 | 286.1 | 0.27 | 0.43 |
| 2010 | 261.1 | 0.47 | 0.4 |
| 2011 | 250.2 | 0.26 | 0.83 |
| 2012 | 209.5 | 1.09 | 1.38 |
| 2013 | 196.3 | 1.01 | 1.57 |
| 2014 | 179.3 | 6.55 | 1.83 |
| 2015 | 151.7 | 4.78 | 1.75 |
| 2016 | 153.3 | 0.36 | 1.04 |
| 2017 | 190.1 | 0.05 | 1 |
| 2018 | 126.2 | 0.05 | 3.25 |
| 2019 | 245 | 0.01 | 1.36 |
| 2020 | 145.5 | 0.11 | 4.96 |
| 2021 | 114.8 | 0.08 | 0.79 |
| 2022 | 92.39 | 0.05 | 0 |



Figure 1: Observed relative density of all males over time during the NMFS summer survey. Each colored square in a facet represents a survey tow. Red line is the border between the EBS and NBS.


Figure 2: Observed relative density of all males at the time of the 2022 NMFS summer survey. Each colored square in a facet represents a survey tow. Red line is the border between the EBS and NBS.


Figure 3: Observed relative density of $45-55 \mathrm{~mm}$ carapace width males over time during the NMFS summer survey. Each colored square in a facet represents a survey tow. Red line is the border between the EBS and NBS.


Figure 4: Observed relative density of males $45-55 \mathrm{~mm}$ carapace width at the time of the 2023 NMFS summer survey. Each colored square in a facet represents a survey tow. Red line is the border between the EBS and NBS.


Figure 5: Observed relative density of $>101 \mathrm{~mm}$ carapace width males over time during the NMFS summer survey. Each colored square in a facet represents a survey tow. Red line is the border between the EBS and NBS.


Figure 6: Observed relative density of males $>101 \mathrm{~mm}$ carapace width at the time of the 2023 NMFS summer survey. Each colored square in a facet represents a survey tow. Red line is the border between the EBS and NBS.

| Shell <br> condition | CW <br> $(\mathrm{mm})$ | Age <br> (years) | Error <br> (years) | Coordinates | Depth <br> $(\mathrm{m})$ | Species |
| :--- | ---: | :--- | :--- | :--- | :--- | :--- |
| $0^{+}$ | 121 | 0.05 | 0.26 | $59^{\circ} 20^{\prime} \mathrm{N}, 171^{\circ} 49^{\prime} \mathrm{W}$ | 43 | C. opilio |
| $0^{+}$ | 110 | 0.11 | 0.27 | $59^{\circ} 20^{\prime} \mathrm{N}, 171^{\circ} 49^{\prime} \mathrm{W}$ | 43 | C. opilio |
| $0^{+}$ | 132 | 0.11 | 0.19 | $59^{\circ} 20^{\prime} \mathrm{N}, 171^{\circ} 49^{\prime} \mathrm{W}$ | 43 | C. opilio |
| 1 | 118 | 0.15 | 0.26 | $59^{\circ} 20^{\prime} \mathrm{N}, 171^{\circ} 49^{\prime} \mathrm{W}$ | 43 | C. opilio |
| 1 | 130 | 0.23 | 0.27 | $59^{\circ} 20^{\prime} \mathrm{N}, 171^{\circ} 49^{\prime} \mathrm{W}$ | 43 | C. opilio |
| 1 | 116 | 0.25 | 0.24 | $59^{\circ} 20^{\prime} \mathrm{N}, 171^{\circ} 49^{\prime} \mathrm{W}$ | 43 | C. opilio |
| $2^{+}$ | 93 | 0.33 | 0.28 | $57^{\circ} 00^{\prime} \mathrm{N}, 167^{\circ} 43^{\prime} \mathrm{W}$ | 42 | C. bairdi |
| $2^{+}$ | 122 | 0.42 | 0.26 | $57^{\circ} 00^{\prime} \mathrm{N}, 167^{\circ} 43^{\prime} \mathrm{W}$ | 42 | C. bairdi |
| $2^{+}$ | 97 | 0.66 | 0.30 | $59^{\circ} 00^{\prime} \mathrm{N}, 171^{\circ} 47^{\prime} \mathrm{W}$ | 46 | C. opilio |
| $2^{+}$ | 123 | 0.78 | 0.32 | $59^{\circ} 00^{\prime} \mathrm{N}, 171^{\circ} 47^{\prime} \mathrm{W}$ | 46 | C. opilio |
| $2^{+}$ | 121 | 0.85 | 0.27 | $57^{\circ} 00^{\prime} \mathrm{N}, 167^{\circ} 43^{\prime} \mathrm{W}$ | 42 | C. opilio |
| $2^{+}$ | 66 | 1.07 | 0.29 | $59^{\circ} 00^{\prime} \mathrm{N}, 171^{\circ} 47^{\prime} \mathrm{W}$ | 46 | C. opilio |
| 3 | 117 | 0.92 | 0.34 | $59^{\circ} 00^{\prime} \mathrm{N}, 171^{\circ} 47^{\prime} \mathrm{W}$ | 46 | C. opilio |
| 3 | 69 | 1.04 | 0.28 | $59^{\circ} 00^{\prime} \mathrm{N}, 171^{\circ} 47^{\prime} \mathrm{W}$ | 46 | C. opilio |
| 3 | 78 | 1.10 | 0.30 | $59^{\circ} 00^{\prime} \mathrm{N}, 171^{\circ} 47^{\prime} \mathrm{W}$ | 46 | C. opilio |
| 4 | 100 | 4.43 | 0.33 | $57^{\circ} 21^{\mathrm{N}, 167^{\circ} 45^{\prime} \mathrm{W}} ⿻$39 <br> C. opilio <br> 4 | 93 | 4.89 |
| 0.37 | $58^{\circ} 20^{\prime} \mathrm{N}, 171^{\circ} 38^{\prime} \mathrm{W}$ | 52 | C. bairdi |  |  |  |
| 4 | 100 | 6.60 | 0.33 | $57^{\circ} 00^{\prime} \mathrm{N}, 167^{\circ} 43^{\prime} \mathrm{W}$ | 42 | C. opilio |
| 5 | 111 | 2.70 | 0.44 | $58^{\circ} 60^{\prime} \mathrm{N}, 169^{\circ} 12^{\prime} \mathrm{W}$ | 28 | C. opilio |
| 5 | 100 | 4.21 | 0.34 | $59^{\circ} 00^{\prime} \mathrm{N}, 171^{\circ} 47^{\prime} \mathrm{W}$ | 46 | C. bairdi |
| 5 | 110 | 6.85 | 0.58 | $58^{\circ} 60^{\prime} \mathrm{N}, 169^{\circ} 12^{\prime} \mathrm{W}$ | 28 | C. opilio |

Figure 7: Radiometric estimates of shell age in male snow and tanner crabs collected during the NMFS survey of 1992. Reproduced from Ernst et al. 2005's presentation of Nevissi et al. 1995.


Figure 8: Observed numbers at length of old shell mature males by size class. The presented size bins are not vulnerable to the fishery, so all mortality is 'natural'. The decline in numbers in a size class after the recruitment collapse in the early 1990s demonstrates expected natural mortality for mature male individuals.


Figure 9: Observed probability of having undergone terminal molt at size for new shell male crab based on chelae height. Blue lines occurred farther back in history; red lines are most recent.


Figure 10: Clutch fullness scores from the 1982-2023 NMFS summer survey. Scores: $0=$ immature, $1=$ mature no eggs, $2=$ trace to $0.125,3=0.25,4=0.5,5=0.75,6=$ full of eggs; $7=$ overflowing.


Figure 11: Time series of the average clutch fullness score (top) and the proportion of observed crab with full clutches (green) and empty clutches (blue) in the NMFS summer survey (bottom). Scores: $0=$ immature, $1=$ mature no eggs, $2=$ trace to $0.125,3=0.25,4=0.5,5=0.75,6=$ full of eggs; $7=$ overflowing..


Figure 12: Time series of non-directed bycatch by gear in numbers of crab.


Figure 13: Raw total numbers at size of male crab observed in the survey. Blue are all numbers at size; green are males $>101 \mathrm{~mm}$ carapace width.


Figure 14: Raw total numbers at size of female crab observed in the survey.


Figure 15: Abundance of males estimated from the NMFS summer survey over time for different size classes. GE102 means greater than or equal to 102 mm carapace width. Grey shading is 95 th percent confidence interval. Left side allows for free $y$-axis; right side retains a common $y$-axis.


Figure 16: Biomass of males estimated from the NMFS summer survey over time for commercially relevant size classes. GE102 means greater than or equal to 102 mm carapace width. Grey shading is 95 th percent confidence interval. Grey shading is 95 th percent confidence interval.

$64{ }^{\circ} \mathrm{N}-$

$$
2022
$$

$$
202316
$$

$60^{\circ} \mathrm{N}$ -


2014
$58^{\circ} \mathrm{N}-$



Figure 17: Centroids of abundance for males $45-85 \mathrm{~mm}$ carapace width. Map shows the centroid in space by year; blue colors are farther in the past. Bottom figures isolate the latidudinal and longitudinal components.


Figure 18: Centroids of abundance for males greater than 101 mm carapace width. Map shows the centroid in space by year; blue colors are farther in the past. Bottom figures isolate the latidudinal and longitudinal components.


Figure 19: Distribution of effort in terms of potlifts in the directed fishery on the Bering Sea shelf summed from 1990-present. Squares are statistical areas defined by the state. Numbers are generated to give context to the following figures. Only data in areas that had three or more fishers and processors represented were used to make this figure. That accounts for $87 \%$ of the data points available.


Figure 20: Yearly distribution of effort in terms of potlifts in the directed fishery on the Bering Sea shelf displayed from 1990-present.


Figure 21: Yearly distribution of catch per unit effort across from 1990-present


Figure 22: Catch per unit effort in the snow crab fleet (top) and total crab caught (bottom), courtesy of Ben Daly.



Figure 24: Distribution of the slopes of trends in inseason cpue by spatial area shown in previous figure. Slopes plotted against the catches removed in a given season and area.


Figure 25: Location of BSFRF survey selectivity experiments that provided data used in this assessment over time.


Figure 26: Observed numbers at length extrapolated from length composition data and estimates of total numbers within the survey selectivity experimental areas by year (left). Inferred selectivity (i.e. the ratio of crab at length in the NMFS gear to crab at length in the BSFRF gear.


Figure 27: Inferred selectivity for all available years of BSFRF data.


Figure 28: Inferred selectivity from BSFRF experiments with selectivity at size class estimated by generalized additive model (top). Inferred selectivity from BSFRF experiments with selectivity at size class estimated by sample size-weighted means and variances (middle). Somerton and Otto (1998) underbag experimental data. Point estimates and associated CVs from the GAM were used as priors in model series 23.3.


Figure 29: Number of crab collected in the BSFRF experimental areas by the NMFS survey and the BSFRF survey.


Figure 30: Retrospective patterns in estimated mature male biomass for selected models.


Figure 31: Output of 100 jittered model fittings for selected models. Top left is the maximum gradient component, top right is the overfishing level, bottom left is F35, and bottom right is B35. Each dot represent an instance of a jittered fitted model and are colored based on the OFL resulting from that run.


Figure 32: Model fits to the observed mature biomass at survey.


Figure 33: Model fits to the observed mature biomass at survey 2009-present


Figure 34: Model fits (colored lines) to the growth data (black dots).


Figure 35: Size transition matrix from the author-preferred model.


Figure 36: Model fits to catch data.


Figure 37: Model fits (lines) to the retained catch size composition data (grey bars).


Figure 38: Model fits (lines) to the total catch size composition data (grey bars).


Figure 39: Model fits (lines) to the female discard size composition data (grey bars).


Figure 40: Model fits (lines) to the male non-directed fishery size composition data (grey bars).


Figure 41: Model fits (lines) to the female non-directed size composition data (grey bars).


1986






$$
\begin{gathered}
\text { model }-{ }^{22.1}{ }^{23.1}{ }^{23.2}{ }^{23.3}{ }^{23.2}{ }^{23.3 a} \\
\end{gathered}
$$

Figure 42: Model fits to immature male survey size composition data from 1982-1988.


Figure 43: Model fits to immature female survey size composition data from 1982-1988.


Figure 44: Model fits to mature male survey size composition data from 1982-1988.


Figure 45: Model fits to mature female survey size composition data from 1982-1988.


Figure 46: Model fits to immature male survey size composition data from 1989-present.


$$
\begin{array}{r}
\text { model }-22.1-23.2-23.3 \mathrm{a} \\
\\
-23.1-23.3-23.3 \mathrm{~b}
\end{array}
$$

Figure 47: Model fits to immature female survey size composition data from 1989-present.


Figure 48: Model fits to mature male survey size composition data from 1989-present.


Figure 49: Model fits to mature female survey size composition data from 1989-present.


Figure 50: Model fits to BSFRF survey selectivity experiment size composition data. Figure labels indicate the year of the survey (top), the entity completing the survey (middle), and the sex of crab collected (bottom).


Figure 51: Model predicted mature biomass at mating time in 1,000 tonnes.


Figure 52: Estimated biomass of male crab $>101 \mathrm{~mm}$ carapace width from the survey (black line and dots with gray 95 th CI) and from each model in the assessment (colored lines).


Figure 53: Estimated selectivities by NMFS survey, sex, and time period.


Figure 54: Estimated survey selectivity (lines) with normal priors derived from BSFRF selectivity experiment data (points + intervals).


Figure 55: Estimated availability/selectivity for the BSFRF experimental data. Curves for BSFRF are availability*selectivity and curves for NMFS are only availability, which is then multiplied by the estimated selectivity in the previous figure to calculate size composition data.


Figure 56: Estimated selectivities by fishing fleet and sex for capture and retained catches.


Figure 57: Estimated fishing mortalities for the directed and non-directed fisheries.


Figure 58: Proportion of biomass by size over time (bottom), realized exploitation rate by model (middle), and proportion of total biomass for select sizes over time (top).


Figure 59: Estimated (black line) or specified (colored lines) probability(s) of maturing for male crab.


model

- 22.1
- 23.1
- 23.2
- 23.3
- 23.3a
- 23.3b

Figure 60: Estimated recruitment by sex (bottom) and proportions recruiting to length bin (top) by model.


Figure 61: Estimated natural mortality by sex and maturity state. Natural mortality in all years previous to 2018 and after 2019 are equal to the estimated M in 2017.


Figure 62: Prevalence of bitter crab syndrome over time. Top figure is the unweighted prevalence of visual evidence of BCS in crab observed in the survey. Bottom is the prevalence of BCS weighted by the sampling factor.

