# BRISTOL BAY RED KING CRAB STOCK ASSESSMENT IN FALL 2022 

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## Executive Summary

1. Stock: Red king crab (RKC), Paralithodes camtschaticus, in Bristol Bay, Alaska.
2. Catches: The domestic RKC fishery began to expand in the late 1960s and peaked in 1980 with a catch of 129.95 million $\mathrm{lb}(58,943 \mathrm{t})$. The catch declined dramatically in the early 1980 s and remained at low levels during the last three decades. After rationalization, catches were relatively high before the 2010/11 season but have been on a declining trend since 2014. The retained catch in 2020/21 was approximately 2.65 million lb ( $1,257 \mathrm{t}$ ), compared to 4.5 million $\mathrm{lb}(2,027 \mathrm{t}$ ) in 2018/19, following a reduction in total allowable catch (TAC). The magnitude of bycatch from groundfish trawl and fixed gear fisheries has been stable and small relative to stock abundance during the last 10 years. The decline of the directed pot fishery crab/pot lift (CPUE) has been much less than the retained catch decline, with the 2020/21 CPUE having about $12.5 \%$ reduction from the average CPUE during the recent 20 years.
3. Stock biomass: Estimated mature biomass increased dramatically in the mid-1970s, then decreased precipitously in the early 1980s. Estimated mature crab abundances increased during 1985-2007 with mature females being about four times more abundant in 2007 than in 1985 and mature males being about two times more abundant in 2007 than in 1985. Estimated mature abundance has steadily declined since 2007. The projected mature male biomass in 2022 is approximately $43 \%$ of the estimate mean survey biomass for the entire time series. The estimated mature female biomass has also been very low during the last four years, with the 2022 values begin approximately $42 \%$ of the mean. The estimated mature female abundance was below the state of Alaska harvest strategy threshold of 8.4 million of crab for a fishery opening in 2022.
4. Recruitment: Estimated recruitment was high during the 1970s and early 1980s and has generally been low since 1985 (1979-year class). During 1984-2020, estimated recruitment was above the historical average (1976-2019 reference years) only in 1984, 1986, 1990, 1995, 1999, 2002, and 2005. Estimated recruitment was extremely low during the last 12 years, and even lower during the recent eight years. With the low recruitment in recent
years, the projected mature biomass is expected to decline during the next few years with a below-average fishing mortality of 0.167 to $0.25 \mathrm{yr}^{-1}$.
5. Management performance:

Status and catch specifications (1,000 t) (model 21.1b):

| Year | MSST | Biomass <br> $($ MMB $)$ | TAC | Retained <br> Catch | Total <br> Catch | OFL | ABC |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2018 / 19$ | $10.62^{\mathrm{B}}$ | $16.92^{\mathrm{B}}$ | 1.95 | 2.03 | 2.65 | 5.34 | 4.27 |
| $2019 / 20$ | $12.72^{\mathrm{C}}$ | $14.24^{\mathrm{C}}$ | 1.72 | 1.78 | 2.22 | 3.40 | 2.72 |
| $2020 / 21$ | $12.12^{\mathrm{D}}$ | $13.96^{\mathrm{D}}$ | 1.20 | 1.26 | 1.57 | 2.14 | 1.61 |
| $2021 / 22$ | 12.01 | 16.64 | 0 | 0.02 | 0.10 | 2.23 | 1.78 |
| $2022 / 23$ |  | 16.95 |  |  |  | 3.04 | 2.43 |

The stock was above MSST in 2021/22 (69\% of BMSY) and hence was not overfished. Since total catch was below OFL, overfishing did not occur. The projection using the lowest recruitment periods during 2013-2021 would not likely result in "approaching an overfished condition" based on the current harvest strategy. The relatively low MSST in 2018/19 and $B_{35 \%}$ in 2019/20 below was caused by a problem of the previous GMACS version using the only sex ratio of recruitment in the terminal year for $B_{35 \%}$ computation in 2019. The lower estimated male recruitment ratio in the terminal year in 2019 resulted in a lower mean male recruitment for $B_{35 \%}$ computation. The current version of GMACS uses an average of sex ratios of recruitment during the reference period to estimate $B_{35 \%}$, which results in a stable sex ratio (about 50\%) for the reference point calculation.

The ABC buffer was increased from $10 \%$ to $20 \%$ in 2018, and an additional buffer of $5 \%$ was added in 2020 due to the lack of a 2020 survey. A $20 \%$ buffer was recommended by the CPT and SSC for ABC estimation for 2021/22. Reoccurring concerns for this stock are still present (cold pool distributional shifts, declining trends in mature biomass, lack of large recruitment pulses, retrospective patterns), in addition to low mature female biomass the last two years, all contribute to a recommended 20\% buffer for 2022/23.

Status and catch specifications (million lb, model 21.1b):

| Year | MSST | Biomass <br> (MMB) | TAC | Retained <br> Catch | Total <br> Catch | OFL | ABC |
| :---: | :---: | :---: | ---: | :---: | :---: | :---: | :---: |
| $2018 / 19$ | $23.4^{\mathrm{B}}$ | $37.3^{\mathrm{B}}$ | 4.31 | 4.31 | 5.85 | 11.76 | 9.41 |
| $2019 / 20$ | $28.0^{\mathrm{C}}$ | $31.4^{\mathrm{C}}$ | 3.80 | 3.91 | 4.89 | 7.50 | 6.00 |
| $2020 / 21$ | $26.7^{\mathrm{D}}$ | $308^{\mathrm{D}}$ | 2.77 | 2.65 | 3.47 | 4.72 | 3.54 |
| $2021 / 22$ | 26.5 | 36.7 | 0 | 0.04 | 0.22 | 4.91 | 3.92 |
| $2022 / 23$ |  | 37.4 |  |  |  | 6.70 | 5.35 |

Notes:
A - Calculated from the assessment reviewed by the Crab Plan Team in September 2018

B - Calculated from the assessment reviewed by the Crab Plan Team in September 2019
C - Calculated from the assessment reviewed by the Crab Plan Team in September 2020
D - Calculated from the assessment reviewed by the Crab Plan Team in September 2021
6. Basis for the OFL: Values are in $1,000 \mathrm{t}$ (model 21.1b):

| Year | Tier | $\mathbf{B}_{\text {MSY }}$ | Current <br> MMB | B/BMSY <br> (MMB) | FofL | Years to <br> define <br> BMSY | Natural <br> Mortality |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2018 / 19$ | 3b | 25.5 | 20.8 | 0.82 | 0.25 | $1984-2017$ | 0.18 |
| $2019 / 20$ | $3 b$ | 21.2 | 16.0 | 0.75 | 0.22 | $1984-2018$ | 0.18 |
| $2020 / 21$ | $3 b$ | 25.4 | 14.9 | 0.59 | 0.16 | $1984-2019$ | 0.18 |
| $2021 / 22$ | 3b | 24.2 | 14.9 | 0.62 | 0.17 | $1984-2020$ | 0.18 |
| $2022 / 23$ | 3b | 24.03 | 17.0 | 0.71 | 0.20 | $1984-2021$ | 0.18 |

Basis for the OFL: Values are in million lb:

| Year | Tier | BMSY | Current <br> MMB | B/BMSY <br> (MMB) | FofL | Years to <br> define <br> BMSY | Natural <br> Mortality |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2018 / 19$ | $3 b$ | 56.2 | 45.9 | 0.82 | 0.25 | $1984-2017$ | 0.18 |
| $2019 / 20$ | $3 b$ | 46.8 | 35.2 | 0.75 | 0.22 | $1984-2018$ | 0.18 |
| $2020 / 21$ | $3 b$ | 56.1 | 32.9 | 0.59 | 0.16 | $1984-2019$ | 0.18 |
| $2021 / 22$ | $3 b$ | 53.4 | 33.0 | 0.62 | 0.17 | $1984-2020$ | 0.18 |
| $2022 / 23$ | 3b | 53.0 | 37.4 | 0.71 | 0.20 | $1984-2021$ | 0.18 |

## A. Summary of Major Changes

1. Changes to management of the fishery: None.
2. Changes to the input data:
a. Updated groundfish fisheries bycatch data during 1986-2021.
b. Updated NMFS survey data for 2022 , biomass and length compositions
c. Updated length composition data for directed and non-directed fisheries.

## 3. Changes to the assessment methodology:

a. Updated version of GMACS (version 2.01.E, Feb. 6, 2022) is used.
b. The analyses of terminal years of recruitment are updated.
c. Three models are compared in this report (See Section E.3.a for details):
21.1b: model 21.1a + updated groundfish fisheries bycatch data.
22.0: model $21.1 \mathrm{~b}+$ starting in 1985.
22.0a: model $22.0+$ estimating a constant $M$ for males.

These models are designed for evaluating starting the model in 1985 and estimating M for males.

## 4. Changes to assessment results:

Three model scenarios are compared in this report. In the May 2022 draft report small updates were made to the accepted model in 2021 (21.1), which included an updated GMACS version 2.01.E (finalized on Feb. 6, 2022) and groundfish bycatch that was updated when the AKFIN database was updated in spring of 2022, resulting in a new "base" - model 21.1b. These updates are minor and considered appropriate to replace the base mofel of 21.1 (refer to CPT May minutes and SSC June minutes 2022). Model 21.1b was used to compare to the other model scenarios.

The two additional models considered starting the model in 1985, rather that the 1975 start date of the base model (21.1b). These models were used to evaluate model starting year and estimating a base $M$ for males. Model 22.0 is the reduced time series data version of model 21.1 b , and the overall results are similar. The notable differences are smaller $B_{35 \%}(22,896 \mathrm{t}$ vs $24,026 \mathrm{t}$ ) and NMFS survey catchability ( 0.94 vs 0.97 ), and higher MMB in the terminal year (2022) ( $17,158 \mathrm{t}$ vs $16,953 \mathrm{t}$ ) and higher OFL ( $3,482 \mathrm{t}$ vs $3,036 \mathrm{t}$ ) for model 22.0. These differences are likely caused by a high recruitment in 1984 (associated with the very high $M$ ) being used for $B_{35 \%}$ computation for model 21.1 b and more influence of BSFRF survey data for model 22.0. Estimating $M$ for males ( 0.23 compared to fixing at 0.18 for the base model) in model 22.0a reduces total likelihood compared to model 22.0, slightly increases annual mature male biomass, except for the terminal year projection, and results in an estimated $B_{35 \%}$ about $10 \%$ lower than model 22.0. A higher $M$ also results in higher $\mathrm{F}_{35 \%}$ and OFL for model 22.0a.
Based on the model results, it appears that the choice of preferred models depends on two factors: model starting year and estimation of M. Moving the starting year to 1985 greatly simplifies this model by removing early years of high biomass and subsequent dramatic decline in biomass in the early 80s. Additionally, a 1985 start date coincides to gear changes in the NMFS trawl survey in the early 80s. Considerations for $M$ estimation are whether to estimate a base $M$ for males for the whole time series or keep the base $M$ for males fixed at 0.18 . Estimating the base M for males does reduce the retrospective bias from model 22.0 (the base model that starts in 1985). The concern with estimating a base $M$ for males for the whole time series is potential confounding with estimating trawl survey catchability.
Based on the above considerations, we recommend model 21.b or 22.0 as the base model for overfishing definition determination in September 2022 due their simple approach: a fixed base $M$ of 0.18 for males, less confounding between estimating $M$ and survey catchability, and acceptable data fittings. Model 22.0 has an additional advantage of avoiding the dramatic abundance decline during the early 1980s and not including the recruitment associated with an extremely high $M$ being used for estimating $B_{35 \%}$. If estimating a base $M$ value for males in the model is acceptable, then model 22.0a is also a good alternative, which fits the data well and
greatly reduces retrospective bias from model 22.0. Model 21.1 b results are presented in the specification tables in the executive summary but values for management-related quantities for all models are summarized in likelihood Tables 5b.

## B. Responses to SSC and CPT Comments

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

## Response to SSC Comments (from February 2022):

"The SSC supports the CPT general recommendations that all stock assessments include results from the currently accepted model with new data (base model) so that changes in model performance can be assessed. Values for management-related quantities for all models that may be recommended by the CPT or SSC should also be available."

Response: We have followed these recommendations.
2. Responses to the most recent two sets of SSC and CPT comments specific to this assessment:

## Response to CPT Comments (from May 2021):

"The CPT was concerned that the 'information' content of the data with respect to natural mortality could be related to strong assumptions elsewhere in the model, and recommended further exploration of natural mortality after September and suggested attending the June 2021 CAPAM workshop on natural mortality, which may provide some insights into best practices. A large increase in estimated natural mortality would likely increase fishing mortality reference points, with management implications."

Response: $M$ was explored in May 2022. Estimated $M$ values in the length-based crab models tend to have higher values than the other approaches, and confounding among estimated $M$, survey selectivity/catchability, and recruitment in a length-based model makes it difficult to accurately estimate $M$ in the model. Among the three models in September 2022, one estimates a base $M$ for males (22.0a).
"The CPT recommended presenting Models 19.3d, 19.3e, and 19.3g in September with updated data."

Response: We ran these three models as well as another model suggested by the SSC for September 2021.
"The CPT was interested in more exploration of the retrospective patterns, which seem to have increased since the last assessment despite no new data being added. Reported Mohn's rhos were starting to reach concerning magnitudes in the proposed models?"

Response: The catch and bycatch updates make the retrospective patterns slightly worse than before. Higher than expected BSFRF survey biomass during 2007-2008 and 2013-2016 and NMFS survey biomass in 2014 likely caused these biases. Also, much lower than expected NMFS survey biomass during 2018-2019 and 2021 results in lower biomass estimates in 2020 and 2021. The biases for total abundance are much smaller than mature male biomass.

In May 2022, we examined the retrospective patterns further. It appears that adding a time block of $M$ during 2015-2018 and dropping BSFRF survey data reduces retrospective bias but these changes were not considered to be biologically or scientifically acceptable.
"Model 19.3c probably should have been labeled model 21.0, given the large change in inputs?"
Response: To avoid confusion, we do not change the model label this time. The year in the model label will be changed when the major model changes, such as the model suggested by the SSC in June 2021, which is named as model 21.0 in the draft SAFE report in September 2021.
"When calculating the probability of being overfished via MCMC, it is necessary to calculate $B_{35 \%}$ for each draw to compare the MMB from that draw. If this is not done, the comparison is not consistent."

Response: We have followed this recommendation.

## Response to CPT Comments (from September 2021):

"When projecting the stock to determine whether it is approaching an overfished condition, identify the uncertainties included and ignored in the projection. It is particularly important to distinguish those that are captured in the projection (i.e. those associated with the model) and the additional uncertainties that form the basis for the ABC buffer."

Response: we list uncertainties in the projection section.
"When projecting MMB, label figures with the date to which it is projected (e.g., Feb. 15, 2022), not just the year (which can lead to confusion)."

Response: We follow this recommendation.
"Consider a model with constant M, but estimated separately for males and females (i.e., similar to Model 21.0, but with sex-specific M's) for May 2022."

Response: Models 22.0, 22.0a, 22.0b, 22.0c and 22.0d with starting year of 1985 in May 2022 are constant $M$ and estimated separately for males and females.
"Consider a model in which the data starts in 1985 (as suggested by the CIE reviewers)."
Response: Models 22.0, 22.0a, 22.0b, 22.0c, 22.0d, and 22.0e in May 2022 start in 1985. Of these models 22.0 and 22.0a were recommended to bring forth in September 2022 for consideration. They are included in this report.

## Response to SSC Comments specific to this assessment (from June 2021):

"The SSC supports exploring more modern methods for estimating natural mortality, but notes that this method still relies strongly on the maximum age for BBRKC. The SSC recommends continued research to validate the ages for this stock."

Response: We agree with this suggestion. The maximum age was determined by old tagging data, and due to funding and personnel constraint, age validation for BBRKC is more like a long-term goal than a short-term project.
"The likelihood profile suggests that the values of $M$ for male and female might be similar and that the current difference may be because of the constraint of base $M$ to a low value. When $M$ is misspecified, it can be the cause of a strong positive retrospective pattern, which BBRKC has. The SSC would have liked to have seen compositional fits and a retrospective analysis for model 19.6 or some model with a higher $M$ value, particularly to see if it fits the plus group better. Despite the increase in $F 35 \%$, there was not a commensurate increase in OFL. An exploration of the underlying reasons for this outcome is needed."

Response: Based on our past modelling experience, when $M$ values for males and females are estimated separately, estimated $M$ values tended to be always higher for females than for males. The likelihood profile was created through fixing $M$ values for males and estimating $M$ values for females, and when the fixed $M$ values for males were very high, estimated $M$ values for females tended to be similar to $M$ values for males. The increase in $F 35 \%$ but not a commensurate increase in $O F L$ is due to reduction of mature male biomass caused by the high $M$.

As a reference, we copied the likelihood profile computed in May 2020 below. Model 19.6 uses male base $M$ of 0.257 estimated by Then et al. (2015), and the likelihood profile of base $M$ from 0.1 to 0.4 is as follow:


It appears that the maximum likelihood value is achieved with a base $M$ of 0.31 for males and 0.321 for females.

In May 2022, instead of continuing to examine model 19.6, we examine new models starting in 1985 with constant $M$ over time: models 22.0, 22.0a, 22.0b, 22.0c, and 22.0d. Model 22.0a with model estimated $M$ of 0.226 for males and 0.261 for females is close to model 19.6. Bubble plots are illustrated for models 22.0, 22.0a, and 22.0d in Figures 25c, 25d, 25e, 26c, 26d, and 26e. Higher base $M$ values do improve the plus group fittings somewhat for model 22.0a.
"In addition to the CPT recommended models (19.3d, 19.3e, and 19.3g), the SSC recommends a simplified version of model $19.3 d$ that estimates one natural mortality parameter across sex and time, and one shared catchability and selectivity curve for the NMFS trawl survey to help make several selectivity parameters better defined."

Response: We named this as model 21.0 and included it in the September 2021 assessment.
"The SSC requests that the current crab management zones be included in the maps of VAST model-derived spatial distributions of BBRKC."

Response: We will ask Dr. Jon Richar to add the current crab management zones to the VAST spatial plots.
"The SSC also looks forward to the summary report from the March 2021 CIE Review for this stock."

Response: The summary report of the 2021 CIE review is included in Appendix D.

## Response to SSC Comments specific to this assessment (from October 2021):

"The SSC requests that in addition to temperature effects on the timing of the molt-mate cycle, the authors explore other potential drivers (e.g., prey quality or quantity) that could underlie the incomplete molt-mate cycle observed in 2021. Based on NMFS trawl survey female biomass estimates, the State of Alaska closed the BBRKC fishery. Next year's assessment should estimate the probability that the stock is currently in the overfished condition."

Response: NMFS staff is doing evaluation of re-tow survey protocol. Probabilities in the overfished condition for some models were estimated in September 2021 and are estimated in this report for the base model (21.1b).
"The SSC recommends that authors should carefully consider assessment implications of the stock boundaries given the evidence of crabs outside of the managed area. The SSC suggests that the authors should still be able to use data from outside stock boundaries, even if not used in the input survey abundance estimates. For example, the abundance seen outside stock boundaries could be treated as covariate informing catchability within the model. This analysis seems particularly important for females that are increasingly outside of the current stock boundaries and are at low abundance, triggering the State closure. The SSC recommends that the authors formulate separate survey abundance time series inside and outside of the defined area that could prove useful in the assessment model (e.g., informing catchability). If this is not an option in the stock assessment, then it highlights the need for ESRs or ESPs to track movement of these crabs both through survey results and developing indices from local knowledge."

Response: The current version of GMACS seems not to be able to use the Northern RKC survey index to inform BBRKC survey catchability. We tried to add a model to include both BBRKC and Northern RKC data, but the groundfish fisheries bycatch is not currently available in the Northern area. We plot more proportional data of the Northern RKC in Figures 35a and 35b. Overall, the proportions of different size groups of the Northern RKC during a recent dozen years are higher than in the past and do not trend higher except for mature females in 2021. The high survey mature female abundance in the Northern area in 2021 was primarily from three tows and one of them is more than $50 \%$ of total mature females. The survey abundance of the Northern RKC will continue to be plotted in the SAFE report in the future. After migration patterns between BBRKC and the Northern RKC are fully understood, we will model them in the stock assessment.

[^0]Response: We agree and have urged tagging studies for a few years.
"It would be useful to investigate if there is a mechanism for higher natural mortality or fishing mortality for females only during that early time period while following the CPT recommendation of looking at model 21.0 with constant but separate Ms by sex. Since Model 21.0 estimates a very high level of fishing mortality, but does seem to account for the decline in large females, there may be a fishery selectivity issue in that period. If the modelers choose not to continue to use historic data prior to 1985, this suggestion may not be useful."

Response: Figuring out the exact causes of high mortality in the early 1980s is always difficult and we summarize the potential causes in Appendix A, section C-vi, "Potential Reasons for High Mortality during the Early 1980s". The directed fishery does not catch many large females and small crab, so it is difficult to remove these crab from the fishery. We prefer to start the model in 1985, which has two advantages: avoiding the early 1980s period so that a constant $M$ over time can be used, and the same NMFS survey gear throughout the whole model time period.
"The SSC supports continued exploration of the use of VAST estimates for this assessment, particularly if their use will inform mechanisms underlying shifting distributions outside of the current management area."

Response: We also support improvement of VAST estimates and are willing to provide feedback to Jon for further improvement. Hopefully, we will be able to use the VAST results in the models soon.

## C. Introduction

## 1. Species

Red king crab (RKC), Paralithodes camtschaticus, in Bristol Bay, Alaska.

## 2. General distribution

Red king crab inhabit intertidal waters to depths $>200 \mathrm{~m}$ of the North Pacific Ocean from British Columbia, Canada, to the Bering Sea, and south to Hokkaido, Japan, and are found in several areas of the Aleutian Islands, eastern Bering Sea, and the Gulf of Alaska.

## 3. Stock Structure

The State of Alaska divides the Aleutian Islands and eastern Bering Sea into three management registration areas to manage RKC fisheries: Aleutian Islands, Bristol Bay, and Bering Sea (ADF\&G 2012). The Bristol Bay area includes all waters north of the latitude of Cape Sarichef ( $54^{\circ} 36^{\prime} \mathrm{N}$ lat.), east of $168^{\circ} 00^{\prime} \mathrm{W}$ long., and south of the latitude of Cape Newenham ( $58^{\circ} 39^{\prime} \mathrm{N}$ lat.) and the fishery for RKC in this area is managed separately from fisheries for RKC outside of this area; i.e., the red king crab in the Bristol Bay area are assumed to be a separate stock from
red king crab outside of this area. This report summarizes the stock assessment results for the Bristol Bay RKC stock.

## 4. Life History

Red king crab have a complex life history. Fecundity is a function of female size, ranging from tens of thousands to hundreds of thousands (Haynes 1968; Swiney et al. 2012). The eggs are extruded by females, fertilized in the spring, and held by females for about 11 months (Powell and Nickerson 1965). Fertilized eggs are hatched in the spring, most during April-June (Weber 1967). Primiparous females are bred a few weeks earlier in the season than multiparous females.
Larval duration and juvenile crab growth depend on temperature (Stevens 1990; Stevens and Swiney 2007). Male and female RKC mature at 5-12 years old, depending on stock and temperature (Stevens 1990; Loher et al. 2001) and may live >20 years (Matsuura and Takeshita 1990). Males and females attain a maximum size of 227 mm and 195 mm carapace length (CL), respectively (Powell and Nickerson 1965). Female maturity is evaluated by the size at which females are observed to carry egg clutches. Male maturity can be defined by multiple criteria including spermataphore production and size, chelae vs. carapace allometry, and participation in mating in situ (reviewed by Webb 2014). For management purposes, females $>89 \mathrm{~mm}$ CL and males >119 mm CL are assumed to be mature for Bristol Bay RKC. Juvenile RKC molt multiple times per year until age 3 or 4 ; thereafter, molting continues annually in females for life and in males until maturity. Male molting frequency declines after attaining functional maturity.

## 5. Fishery

The RKC stock in Bristol Bay, Alaska, supports one of the most valuable fisheries in the United States. A review of the history of the Bristol Bay RKC fishery is provided in Fitch et al. (2012) and Otto (1989). The Japanese fleet started the fishery in the early 1930s, stopped fishing from 1940 to 1952, and resumed the fishery from 1953 until 1974. The Russian fleet fished for RKC from 1959 to 1971. The Japanese fleet employed primarily tanglenets with a very small proportion of catch from trawls and pots. The Russian fleet used only tanglenets. United States trawlers started fishing Bristol Bay RKC in 1947, but the effort and catch declined in the 1950s. The domestic RKC pot fishery began to expand in the late 1960s and peaked in 1980 with a catch of 129.95 million lb ( $58,943 \mathrm{t}$ ), worth an estimated $\$ 115.3$ million ex-vessel value. The catch declined dramatically in the early 1980s and has remained at low levels during the last two decades (Tables 1a and 1b). After the early 1980s stock collapse, the Bristol Bay RKC fishery took place during a short period in the fall (usually lasting about a week) with the catch quota based on the stock assessment conducted the previous summer (Zheng and Kruse 2002). Beginning with the 2005/2006 season, new regulations associated with fishery rationalization resulted in an increase in the duration of the fishing season (October 15 to January 15). With the implementation of crab rationalization, the annual guideline harvest level (GHL) was changed to a total allowable catch (TAC). Before rationalization, the implementation errors were quite high for some years and sum of actual catches from 1980 to 2007 was about $6 \%$ less than the sum of GHL/TAC over that period.

## 6. Fisheries Management

King and Tanner crab stocks in the Bering Sea and Aleutian Islands are managed by the State of Alaska through a federal king and Tanner crab fishery management plan (FMP). Under the FMP, management measures are divided into three categories: (1) fixed in the FMP, (2) frameworked in the FMP, and (3) discretion of the State of Alaska. The State of Alaska is responsible for determining and establishing the GHL/TAC under the framework in the FMP.
Harvest strategies for the Bristol Bay RKC fishery have changed over time. Two major management objectives for the fishery are to maintain a healthy stock that ensures reproductive viability and to provide for sustained levels of harvest over the long term (ADF\&G 2012). In attempting to meet these objectives, the GHL/TAC is coupled with size-sex-season restrictions. Only males $\geq 6.5$-in carapace width (equivalent to $135-\mathrm{mm} \mathrm{CL}$ ) may be harvested and no fishing is allowed during molting and mating periods (ADF\&G 2012). Specification of TAC is based on a harvest rate strategy. Before 1990, harvest rates on legal males were based on population size, abundance of prerecruits to the fishery, postrecruit abundance, and rates varied from less than $20 \%$ to $60 \%$ (Schmidt and Pengilly 1990). In 1990, the harvest strategy was modified, and a $20 \%$ mature male harvest rate was applied to the abundance of mature-sized ( $\geq 120-\mathrm{mm}$ CL) males with a maximum $60 \%$ harvest rate cap of legal ( $\geq 135-\mathrm{mm}$ CL) males (Pengilly and Schmidt 1995). In addition, a minimum threshold of 8.4 million mature-sized females ( $\geq 90-\mathrm{mm} \mathrm{CL}$ ) was added to existing management measures to avoid recruitment overfishing (Pengilly and Schmidt 1995). Based on a new assessment model and research findings (Zheng et al. 1995a, 1995b, 1997a, 1997b), the Alaska Board of Fisheries adopted a new harvest strategy in 1996. That strategy had two mature male harvest rates: $10 \%$ when effective spawning biomass (ESB) is between 14.5 and 55.0 million lb and $15 \%$ when ESB is at or above 55.0 million lb (Zheng et al. 1996). The maximum harvest rate cap of legal males was changed from $60 \%$ to $50 \%$. A threshold of 14.5 million lb of ESB was also added. In 1997, a minimum threshold of 4.0 million lb was established as the minimum GHL for opening the fishery and maintaining fishery viability and manageability when the stock abundance is low. The Board modified the current harvest strategy in 2003 by adding a mature harvest rate of $12.5 \%$ when the ESB is between 34.75 and 55.0 million lb and in 2012 eliminated the minimum GHL threshold. The current harvest strategy is illustrated in Figure 1.

## D. Data

## 1. Summary of New Information

a. Updated groundfish fisheries bycatch data during 1986-2021.
b. Updated survey data for 2022
c. Updated length-frequencies distributions for all data sets for 2021/2022

Data types and availability periods are illustrated in Figure 2.

## 2. Catch Data

Data on landings of Bristol Bay RKC by length and year and catch per unit effort from 1960 to

1973 were obtained from annual reports of the International North Pacific Fisheries Commission (Hoopes et al. 1972; Jackson 1974; Phinney 1975) and from the Alaska Department of Fish and Game from 1974 to 2020 (Tables 1a and 1b). Bycatch data are available starting from 1990 and were obtained from the ADF\&G observer database and reports (Gaeuman 2013) (Table 2). Sample sizes for catch by length and shell condition are summarized in Table 3a. Relatively large samples were taken from the retained catch each year. Sample sizes for trawl bycatch were the annual sums of length frequency samples in the National Marine Fisheries Service (NMFS) database.

## (i). Catch Biomass

Retained catch and estimated bycatch biomasses are summarized in Tables 1a and 1 b and illustrated in Figure 3. Retained catch and estimated bycatch from the directed fishery include the general, open-access fishery (prior to rationalization), or the individual fishery quota (IFQ) fishery (after rationalization), as well as the Community Development Quota (CDQ) fishery and the ADF\&G cost-recovery harvest. Starting in 1973, the fishery generally occurred during the late summer and fall. Before 1973, a small portion of retained catch in some years was caught from April to June. The years in Tables 1a and 1 b are defined as crab year from July 1 to June 30. Bycatch data for the cost-recovery fishery before 2006 were not available. In this report, pot fisheries include both the directed fishery and RKC bycatch in the Tanner crab pot fishery, and trawl fisheries and fixed gear fisheries are groundfish fisheries. Observers did not separate retained and discarded catch of legalsized crab after 2017 in the directed pot fishery, so the male discarded biomass from the directed fishery has been estimated by the subtraction method since 2018 (B. Daly, ADF\&G, personal communication).

## (ii). Catch Size Composition

Retained catches by length and shell condition and bycatches by length, shell condition, and sex were obtained for stock assessments. From 1960 to 1966, only retained catch length compositions from the Japanese fishery were available. Retained catches from the Russian and U.S. fisheries were assumed to have the same length compositions as the Japanese fishery during this period. From 1967 to 1969 , the length compositions from the Russian fishery were assumed to be the same as those from the Japanese and U.S. fisheries. After 1969, foreign catch declined sharply and only length compositions from the U.S. fishery were used to distribute catch by length.

## (iii). Catch per Unit Effort

Catch per unit effort (CPUE) is defined as the number of retained crab per tan (a unit fishing effort for tanglenets) for the Japanese and Russian tanglenet fisheries and the number of retained crab per potlift for the U.S. fishery (Table 1b). Soak time, while an important factor influencing CPUE, is difficult to standardize. Furthermore, complete historical soak time data from the U.S. fishery are not available. Based on the approach of Balsiger (1974), all fishing effort from Japan, Russia, and U.S. were standardized to the Japanese tanglenet from 1960 to 1971, and the CPUE was standardized as crab per tan. Except for the peak-to-crash years of the late 1970s and early 1980s, the correspondence between U.S. fishery CPUE and area-swept survey abundance is poor (Figure
4). Due to the difficulty in estimating commercial fishing catchability and crab availability to the NMFS annual trawl survey data, commercial CPUE data were not used in the model.

## 3. NMFS Survey Data

The NMFS has conducted annual trawl surveys of the eastern Bering Sea since 1968. Two vessels, each towing an eastern otter trawl with an 83 ft headrope and a 112 ft footrope, conducted this multispecies, crab-groundfish survey during the summer. Stations were sampled in the center of a systematic 20 X 20 nm grid overlaid in an area of $\approx 140,000 \mathrm{~nm}^{2}$. Since 1972 , the trawl survey has covered the full stock distribution except in nearshore waters. The survey in Bristol Bay occurs primarily during late May and June. Tow-by-tow trawl survey data for Bristol Bay RKC during 1975-2021 were provided by NMFS. Due to survey data quality issue, only survey data after 1974 are used in the assessment models.
Abundance estimates by sex, carapace length, and shell condition were derived from survey data using an area-swept approach (Figures 5a and 5b). Until the late 1980s, NMFS used a poststratification approach, but subsequently treated Bristol Bay as a single stratum; the estimates shown for Bristol Bay in Figures 4, 5a, and 5b were made without post-stratification. If multiple tows were made at a single station in a given year, the average of the abundances from all tows within that station was used as the estimate of abundance for that station. The new time series since 2015 discards all "hot spot" tows. We used the new area-swept estimates provided by NMFS in 2022. The VAST estimated biomasses were not considered in this year's assessment but may be considered in the future (Figure 6).
In addition to the standard surveys conducted in early June (late May to early June in 1999 and 2000), a portion of the distribution of Bristol Bay RKC was resurveyed in 1999, 2000, 2006-2012, and 2021 to better assess mature female abundance. Resurveys performed in late July, about six weeks after the standard survey, included 31 stations (1999), 23 stations (2000), 31 stations (2006, 1 bad tow and 30 valid tows), 32 stations (2007-2009), 23 stations (2010), and 20 stations (2011, 2012, and 2021) with high female densities. The resurveys were necessary because a high proportion of mature females had not yet molted or mated when sampled during the standard survey time. Differences in area-swept estimates of abundance between the standard surveys and resurveys of these same stations are attributed to survey measurement errors or to seasonal changes in distribution between survey and resurvey periods. More large females were observed in the resurveys than during the standard surveys in 1999 and 2000, presumably because most mature females had not molted prior to the standard surveys. As in 2006, area-swept estimates of males $>89 \mathrm{~mm}$ CL, mature males, and legal males within the 32 resurvey stations in 2007 were not significantly different ( $P=0.74,0.74$ and 0.95 ; paired $t$-test of sample means) between the standard survey and resurvey tows. However, similar to 2006, area-swept estimates of mature females within the 32 resurvey stations in 2007 were significantly different ( $P=0.03$; paired $t$-test) between the standard survey and resurvey tows. Resurvey stations were close to shore during 2010-2012, and mature and legal male abundance estimates were lower for the re-tow than the standard survey. Following the CPT recommendation, we used the standard survey data for male abundance estimates and only the resurvey data, plus the standard survey data outside the resurveyed stations, to assess female abundances during resurvey years.

## 4. Bering Sea Fisheries Research Foundation Survey Data

The BSFRF conducted trawl surveys for Bristol Bay RKC in 2007 and 2008 with a small-mesh trawl net and 5-minute tows (S. Goodman, BSFRF, pers. com.). The surveys occurred at similar times as the NMFS standard surveys and covered about $97 \%$ of the Bristol Bay survey area. Few Bristol Bay RKC were found outside the BSFRF survey area. Because of the small mesh size, the BSFRF surveys were expected to catch more RKC within the swept area. Crab abundances of different size groups were estimated by the kriging method. Mature male abundances were estimated to be 22.331 million $\operatorname{crab}(\mathrm{CV}=0.0634)$ in 2007 and 19.747 million crab $(\mathrm{CV}=$ 0.0765 ) in 2008. BSFRF also conducted a side-by-side survey concurrent with the NMFS trawl survey during 2013-2016 in Bristol Bay. In May 2017, survey biomass and size composition estimates from 2016 BSFRF side-by-side trawl survey data were updated. Ratios of NMFS survey abundances/total NMFS and BSFRF side-by-side trawl survey abundances are illustrated in Figure 7a, and ratios of NMFS survey abundances/BSFRF side-by-side trawl survey abundances are shown in Figures 7b and 7c.

As a comparison to the estimated NMFS survey catchability (0.896) at 162.5 mm CL by the double-bag experiment, we computed an overall ratio ( $q=0.891$ ) of NMFS survey abundances/BSFRF side-by-side trawl survey abundances for legal crab ( $\geq 135 \mathrm{~mm}$ carapace length) as follow:
$q=\sum_{y=2013, l=135 \mathrm{~mm}}^{y=2016, l=\infty} r_{y, l} n_{y, l} / \sum_{y=2013, l=135 \mathrm{~mm}}^{y=2016, l=\infty} n_{y, l}$
where $r_{y, l}$ is the ratio of NMFS survey abundance/BSFRF side-by-side trawl survey abundance in year $y$ and length group $l$, and $n_{y, l}$ is the combined survey abundance of side-by-side surveys in year $y$ and length group $l$. Due to small catch, all haul data were combined to compute the ratios for each length group and year.

## E. Analytic Approach

## 1. History of Modeling Approaches

To reduce annual measurement errors associated with abundance estimates derived from the area-swept method, ADF\&G developed a length-based analysis (LBA) in 1994 that incorporates multiple years of data and multiple data sources in the estimation procedure (Zheng et al. 1995a). Annual abundance estimates of the Bristol Bay RKC stock from the LBA have been used to manage the directed crab fishery and to set crab bycatch limits in the groundfish fisheries since 1995 (Figure 1). An alternative length-based model (research model) was developed in 2004 to include small size crab to determine federal overfishing limits. Given that the crab abundance declined sharply during the early 1980s, the LBA estimated natural mortality for different periods of years, whereas the research model estimated additional mortality beyond a base constant natural mortality during 1980-1984. In this report, we present only the research model that was fit to the data from 1975 to 2022.

## 2. Model Description

The original LBA model was described in detail by Zheng et al. (1995a, 1995b) and Zheng and Kruse (2002). The model combines multiple sources of survey, catch, and bycatch data using a maximum likelihood approach to estimate abundance, recruitment, selectivity, fishing mortality, catch, and bycatch of commercial pot fisheries and groundfish trawl fisheries. Since 2019, GMACS (General Model for Alaska Crab Stocks) has been used for this stock assessment. A full model description is provided in Appendix A.
a-f. See Appendix A.
g. Critical assumptions of the model:
i. The base natural mortality is kept constant at $0.18 \mathrm{yr}^{-1}$ for males, shell condition, and length and was estimated assuming a maximum age of 25 and applying the $1 \%$ rule (Zheng 2005).
ii. Survey and fisheries selectivities are a function of length and were constant over shell condition. Selectivities may or may not be a function of sex except for groundfish fisheries bycatch selectivities, which are the same for both sexes. Two different NMFS survey selectivities were estimated: (1) 1975-1981 and (2) 19822021, based on modifications to the trawl gear used in the assessment survey.
iii. Growth is a function of length. For females, growth-per-molt increments as a function of length are estimated for three periods (1975-1982, 1983-1993, and 19942022) based on sizes at maturity. Once mature, female red king crab have a much smaller growth increment per molt.
iv. Molting probabilities are an inverse logistic function of length for males. Females molt annually.
v. Annual fishing seasons for the directed fishery are short.
vi. The prior mean for NMFS survey catchability $(Q)$ is estimated to be 0.896 with a standard deviation of 0.025 for some models, based on a trawl experiment by Weinberg et al. (2004); $Q$ is assumed to be constant over time and is estimated in the model. The BSFRF survey catchability is assumed to be 1.0 . The prior mean of 0.896 for NMFS survey $Q$ (at 162.5 mm carapace length) is also close to the abundance-weighted average ratio of 0.891 for crab $\geq 135 \mathrm{~mm}$ CL across four years of side-by-side NMFS and BSFRF survey data (Figure 7c).
vii. Males mature at sizes $\geq 120 \mathrm{~mm}$ CL. For convenience, female abundance is summarized at sizes $\geq 90 \mathrm{~mm}$ CL as an index of mature females.
viii. Measurement errors are assumed to be normally distributed for length compositions and are log-normally distributed for biomasses.
h. Changes to the above since previous assessment: see Section A. 3 for changes to the assessment methodology.
i. Outline of methods used to validate the code used to implement the model and whether the code is available: Assessment results by GMACS have been compared to the previous assessment models, and the code is online and available from the first author.

## 3. Model Selection and Evaluation

a. Alternative model configurations (models):
21.1b:: the base model for September 2021 with accepted updates May 2022 (12,13). Basic features of this model include:
(1) An estimated constant $M$ for males during 1980-1984, a constant (base) $M$ of 0.18 for males during the other years, and an estimated constant multiplier being used to multiply male $M$ for female $M$. That is, $M$ for females is relative to $M$ for males each year.
(2) Including BSFRF survey data during 2007-2008 and 2013-2016.
(3) Estimating a constant NMFS survey catchability over time in the model and assuming BSFRF survey catchability to be 1.0.
(4) Assuming the BSFRF survey selectivities as the availability to the NMFS trawl survey because the BSFRF survey gear has very small mesh sizes and has tighter contact to the sea floor. This implies that crab occurring in nearshore areas are not available to trawl survey gears.
(5) Two levels of molting probabilities for males: one before 1980 and one after 1979, based on survey shell condition data. Each level has two parameters.
(6) Estimating effective sample size from observed sample sizes. Stage-1 effective sample sizes are estimated as $\min (0.25 * \mathrm{n}, \mathrm{N})$ for trawl surveys and $\min (0.05 * \mathrm{n}, \mathrm{N})$ for catch and bycatch, where n is the sum of observed sample sizes for two sexes, and N is the maximum sample size ( 200 for trawl surveys, 150 for retained catch and total males from the directed pot fishery and 50 for females from the pot fishery and for both males and females from the Tanner crab and groundfish fisheries). There is justification for enforcing a maximum limit to effective sample sizes because the number of length measurements is large (Fournier et al. 1998).
(7) Standard survey data for males and NMFS survey re-tow data (if available during cold years) for females.
(8) Estimating initial year length compositions.
(9) Using total observer male biomass and total observer male length composition data in the directed pot fishery to replace discarded male biomass and discarded male length composition data.
(10) Using total male selectivity and retained proportions in the directed pot fishery to replace retained selectivity and discarded male selectivity; and due to high grading
problems in some years since rationalization, estimating two logistic curves for retained proportions: one before rationalization (before 2005) and another after 2004.
(11) Equal annual effective sample sizes of male and female length compositions for all size composition data sets.
(12) using the recently updated version of GMACS (version 2.01.E).
(13) updated groundfish fisheries bycatch data
22.0: model $21.1 \mathrm{~b}+$ starting in 1985.
22.0a: model $22.0+$ estimating a constant M for males.
b. Progression of results: See the new results at the beginning of the report.
c. Evidence of search for balance between realistic and simpler models: NA.
d. Convergence status/criteria: ADMB default convergence criteria.
e. Sample sizes for length composition data: observed sample sizes are summarized in Table 3a.
f. Credible parameter estimates: All estimated parameters seem to be credible and within bounds.
g. Model selection criteria: The likelihood values are used to select among alternatives that could be legitimately compared by that criterion.
h. Residual analysis: Residual plots are illustrated in various figures.
i. Model evaluation is provided under Results, below.
j. Jittering: The Stock Synthesis Approach is used to perform jittering to find the optimum:

The Jitter factor of 0.1 is multiplied by a random normal deviation $r d e v=N(0,1)$, to a transformed parameter value based upon the predefined parameter:

$$
\begin{equation*}
\text { temp }=0.5 \mathrm{rdev} \text { Jitter } \ln \left(\frac{P_{\max }-P_{\min }+0.0000002}{P_{\text {val }}-P_{\min }+0.0000001}-1\right), \tag{6}
\end{equation*}
$$

with the final jittered starting parameter value back-transformed as:

$$
\begin{equation*}
P_{\text {new }}=P_{\min }+\frac{P_{\max }-P_{\min }}{1.0+\exp (-2.0 \text { temp })}, \tag{7}
\end{equation*}
$$

where $P_{\max }$ and $P_{\text {min }}$ are upper and lower bounds of parameters and $P_{\text {val }}$ is the estimated parameter value before the jittering. Jittering results are not updated and presented in this report.

## 4. Results

a. Effective sample sizes and weighting factors.
i. CVs are assumed to be 0.03 for retained catch biomass, 0.04 for total male biomass, 0.07 for pot bycatch biomasses, 0.10 for groundfish bycatch biomasses, and 0.23 for recruitment sex ratio. Models also estimate sigmaR for recruitment variation and have a penalty on $M$ variation and many prior-densities.
ii. Initial trawl survey catchability $(Q)$ is estimated to be 0.896 with a standard deviation of 0.025 ( CV about 0.03 ) based on the double-bag experiment results (Weinberg et al. 2004). These values are used to set a prior for estimating $Q$ in all models.
iii. Harmonic means of implied sample sizes and maximum caps of effective sample sizes for models 21.1b, 22.0, and 22.0a are summarized in Table 4.
b. Tables of estimates.
i. Negative log-likelihood values and parameter estimates are summarized in Tables $5 \mathrm{a}, 5 \mathrm{~b}, 6 \mathrm{a}, 6 \mathrm{~b}$, and 6 c for all three models.
ii. Natural mortality estimates are shown in Table 7 for three models.
iii. Area-swept estimates of mature female abundance and model estimates of effective spawning biomass (Zheng et al. 1995b) during 2011-2022 for groundfish fisheries bycatch calculation are provided in Table 8.
iv. Abundance and biomass time series are provided in Tables 9a, 9b, and 9c for models 21.1b, 22.0, and 22.0a.
v. Recruitment time series for models 21.1b, 22.0, 22.0a are provided in Tables 9a, 9 b , and 9c.
vi. Time series of catch biomass is provided in Tables 1 a and 1 b .

Length-specific fishing mortality is equal to selectivity-at-length times the full selection fishing mortality. Estimated full pot fishing mortalities for females and full fishing mortalities for groundfish fisheries bycatch are low due to low bycatch and handling mortality rates less than 1.0. Estimated recruits varied greatly among years (Tables 9a, $9 b$, and 9 c ). Estimated selectivities for female pot bycatch are close to 1.0 for all mature females, and the estimated full fishing mortalities for female pot bycatch are lower than those for male retained catch and bycatch (Tables $6 \mathrm{a}, 6 \mathrm{~b}$, and 6 c for models 21.1b, 22.0, and 22.0a).
c. Graphs of estimates.
i. Estimated selectivities by length are provided in Figures 8a, 8b, 8c, and 8d and estimated molting probabilities by length are illustrated in Figures 9a and 9b.

One of the most important results is estimated trawl survey selectivity (Figures 8a). Survey selectivity affects not only the fitting of the data but also the absolute
abundance estimates. These estimated survey selectivities are generally smaller than the capture probabilities in Figure A1 because survey selectivities include capture probabilities and crab availability. The NMFS survey catchability is estimated to be 0.896 from the trawl experiment. The reliability of estimated survey selectivities will greatly affect the application of the model to fisheries management. Under- or overestimates of survey selectivities will cause a systematic upward or downward bias of abundance estimates, respectively. Information about crab availability in the survey area at survey times will help estimate the survey selectivities. Higher estimated natural mortalities generally result in lower NMFS survey selectivities, while the estimated survey selectivities after 1981 are similar among the models.

For all models, estimated molting probabilities during 1975-2021 (Figures 9a, 9b, and 9c) are generally lower than those estimated from the 1954-1961 and 1966-1969 tagging data (Balsiger 1974). Lower molting probabilities mean more oldshell crab, possibly due to changes in molting probabilities over time or shell aging errors. Overestimates or underestimates of oldshell crab will result in lower or higher estimates of male molting probabilities.
ii. Estimated male and female survey biomasses are shown for NMFS surveys (Figures 10a and 10b) and BSFRF surveys (Figure 10c). Absolute mature male biomasses are illustrated in Figures 11a and 11b.

The survey male biomass estimates in 2022 are slightly higher than those in 2018, 2019, and 2021, but the survey female biomass estimates are lower than 2018 and 2019 , and up slightly from 2021. Estimated population biomass increased dramatically in the mid-1970s then decreased precipitously in the early 1980s. Estimated biomass had increased during 1985-2003 for males and during 1985-2007 for females, then declined, and have steadily declined since the late 2000s (Figures 10a-10b, 11a, 11b, and 11c). Absolute mature male biomasses for all models have a similar trend over time (Figures 11a and 11b). Among the three models, model estimated relative NMFS survey biomasses are similar for two models 21.1b and 22.0. Model 22.0a estimates a constant $M$ for males, resulting in slightly higher NMFS survey biomass estimates from the early 2000s and lower in recent years than the other models. All models fit the catch and bycatch biomasses very well.

The fit to BSFRF survey data and estimated survey selectivities are illustrated in Figures 10c-10d, but are all similar in their results.
iii. Estimated recruitment time series are plotted in Figure 12a and recruitment length distributions in Figure 12b for models 21.1b, 22.0, and 22.0a. Recruitment is estimated at the end of year in GMACS and is moved up one year for the beginning of next year. Estimated recruitment time series for models 21.1b, 22.0, and 22.0a are similar. Estimated recruitments among models with higher $M$ values are generally higher.

Like the results of previous models, the terminal year recruitment analysis with model 21.1b suggests the estimated recruitment in the last year should not be used for estimating $B_{35 \%}$ (Figure 28c).
iv. Estimated fishing mortality rates are plotted against mature male biomass in Figures 13a, 13.b, and 13c for models 21.1b, 22.0, and 22.0e, and estimated $M$ and directed pot fishing mortality values over time are illustrated in Figure 13d for models 21.1b, 22.0, and 22.0a.
The average of estimated male recruits from 1984 to 2021 for models starting in 1975 and from 1986 to 2021 for models starting in 1985 (Figure 12a) and mature male biomass per recruit are used to estimate $B_{35 \%}$. The full fishing mortalities for the directed pot fishery at the time of fishing are plotted against mature male biomass on Feb. 15 (Figures 13a, 13.b and 13c). Estimated fishing mortalities in most years before the current harvest strategy was adopted in 1996 were above $F_{35 \%}$ (Figures 13a and 13b). Under the current harvest strategy, estimated fishing mortalities were at or above the $F_{35 \%}$ limits in 1998-1999, 2005, 2007-2010, and 2014-2019 for models 21.1b and 22.0, but below the $F_{35 \%}$ limits in the other post1995 years.
For model 21.1b, estimated full pot fishing mortalities ranged from 0.00 to 2.27 during 1975-2020, with estimated values over 0.40 during 1975-1982, 1984-1987, 1990-1991, 1993, 1998 and 2007-2009 (Table 6a, Figure 13a). For model 22.0, estimated full pot fishing mortalities ranged from 0.00 to 0.70 during 1985-2020, with estimated values over 0.40 in the same years as model 21.1b. Estimated fishing mortalities for pot female and groundfish fisheries bycatches are generally small and less than 0.07.

For model 21.1b, estimated $M$ values are 0.88 during 1980-1984 and 0.18 for the other years for males, and 1.17 during 1980-1984 and 0.24 for the other years for females, with estimated female $M$ values equaling to 1.325 times male $M$ values (Figure 13c). For model 22.0, estimated $M$ values 0.18 for all years for males, and 0.23 for females, with estimated female $M$ values equaling to 1.327 times male $M$ values. For model 22.0 a, estimates $M$ for males is 0.23 , higher than the fixed value of 0.18 in the other models, while $M$ for females is estimated at 0.26 , only slightly higher than the base model. Biologically, females mature earlier than males and likely have higher $M$ values. $M$ values for all models are listed in Table 7.
v. Estimated mature male biomass and recruitment are plotted to illustrate their relationships with model 21.1b (Figure 14a). Annual stock productivities are illustrated in Figure 14b.
Stock productivity (recruitment/mature male biomass) is generally lower during the last 20 years (Figure 14b). However, there are high variations for the relation of stock productivity against mature male biomass.

Egg clutch data collected during summer surveys may provide information about mature female reproductive conditions (Figures 15a, b). Although egg clutch data are subject to rating errors as well as sampling errors, data trends over time may be useful. Proportions of empty clutches for newshell mature females >89 mm CL are high in some years before 1990 but have been low since 1990 (Figure 15a). The high proportion of empty clutches ( 0.2 ) was in 1986, and primarily involved soft shell females (shell condition 1). Clutch fullness fluctuated annually around average levels during two periods: before 1991 and after 1990 (Figure 15a). The average clutch fullness is similar for these two periods (Figure 15). Egg clutch fullness during 2016-2018 was relatively low, then increased in 2019, and declined again in 2021.
d. Graphic evaluation of the fit to the data.
i. Observed vs. estimated catches are plotted in Figure 16a, with bycatch mortalities from different sources shown in Figure 16b for all models.
ii. Model fits to NMFS survey biomass are shown in Figure 10 with a standardized residual plot in Figure 17 for models 21.1b, 22.0, and 22.0a.
iii. Model fits to catch and survey proportions by length are illustrated in Figures 1824 and residual bubble plots are shown in Figures 25-26.
All models fit the fishery biomass data well and the survey biomass reasonably well (Figures 10 and 16). Because the model estimates annual fishing mortality for directed pot male catch, pot female bycatch, and trawl and fixed gear bycatch, the deviations of observed and predicted (estimated) fishery biomass are mainly due to size composition differences. All models fit the NMFS area-swept biomass data almost identically (Figure 10a).

All models also fit the length composition data well (Figures 18-24). Modal progressions are tracked well in the trawl survey data, particularly beginning in mid-1990s (Figures 18 and 19). Cohorts first seen in the trawl survey data in 1975, 1986, 1990, 1995, 1999, 2002 and 2005 can be tracked over time. Some cohorts can be tracked over time in the pot bycatch as well (Figure 21), but the bycatch data did not track the cohorts as well as the survey data. Groundfish bycatch data provide little information to track modal progression (Figures 23 and 24).

Residuals of survey biomasses and proportions of length are plotted to examine their patterns. Residuals were calculated as observed minus predicted and standardized by the estimated standard deviation. Residuals of survey biomasses did not show any consistent patterns for all models (Figures 17). Generally, residuals of proportions of survey males and females appear to be random over length and year for all models (Figures 25a-25e and 26a-26e). Models with higher base $M$ values like model 22.0 a improve the plus group fittings slightly.
e. Retrospective and historic analyses.

Retrospective analyses were conducted for this report using the 2022 models. The 2022 model hindcast results are based on sequentially excluding one-year of data to evaluate the current model performance with fewer data.
i. Retrospective analysis (retrospective bias in base model or models).

The performance of the 2022 model includes sequentially excluding one-year of data. Model 21.1b produces some upward biases during 2012-2022 with higher terminal year estimates of mature male biomass in 2014-2022 (Figure 27a). Higher than expected BSFRF survey biomass during 2007-2008 and 2013-2016 and NMFS survey biomass in 2014 likely caused these biases. Also, much lower than expected NMFS survey biomass during 2018-2019 results in lower biomass estimates in 2020 and 2021. Models 22.0 and 22.0a with starting year of 1985 have similar results (Figures 27a-27c). Mohn's rho calculations for these retrospective runs were high ( 0.329 to 0.453 ) but were reduced some in model 22.0a, which estimates a base $M$ for males in the model.

Ratios of estimated retrospective recruitments to terminal estimates in 2021 as a function of number of years estimated in the model show converging to 1.0 as the number of years increases (Figure 28b). Standard deviations of the ratios drop sharply from one year estimated in the model to two years (Figure 28c), showing great uncertainty of recruitment estimates for terminal years. Based on these results, we suggest not using recruitment estimates in a terminal year for overfishing/overfished determination.
f. Uncertainty and sensitivity analyses
i. Estimated standard deviations of parameters are summarized in Tables 6a-6c for models 21.1b, 22.0, and 22.0a. Estimated standard deviations of mature male biomass are listed in Table 9.
ii. Probabilities for mature male biomass and OFL in 2023 were illustrated in Figures 30a-30b for model 21.1b using the MCMC approach.
iii. Probabilities for mature male biomass below the minimum threshold ( $0.5^{*} B_{35 \%}$ ) in 2023 were plotted in Figure 31 for model 21.1b using the MCMC approach.
iv. Sensitivity analysis for handling mortality rate was included in the SAFE report in May 2010. The baseline handling mortality rate for the directed pot fishery was set at 0.2 . A $50 \%$ reduction and $100 \%$ increase respectively resulted in 0.1 and 0.4 as alternatives. Overall, a higher handling mortality rate resulted in slightly higher estimates of mature abundance, and a lower rate resulted in a minor reduction of estimated mature abundance. Differences of estimated legal male abundance and mature male biomass were small for these handling mortality rate changes.
v. Sensitivity of weights. Sensitivity of weights was examined in the SAFE report in May 2010. Weights to biomasses (trawl survey biomass, retained catch biomass, and bycatch biomasses) were reduced to $50 \%$ or increased to $200 \%$ to examine their sensitivity to abundance estimates. Weights to the penalty terms (recruitment
variation and sex ratio) were respectively reduced or increased. Overall, estimated biomasses were similar under different weights except during the mid-1970s. The variation of estimated biomasses in the mid-1970s was mainly caused by the changes in estimates of additional mortalities in the early 1980s.
g. Comparison of alternative models

These comparisons, based on the data through 2010, were reported in the SAFE report in May 2011. Estimating length proportions in the initial year (scenario 1a) resulted in a better fit of survey length compositions at an expense of 36 more parameters than model 1.
Abundance and biomass estimates with model 1a were similar between models. Using only standard survey data (scenario 1 b) resulted in a poorer fit of survey length compositions and biomass than scenarios using both standard and re-tow data (scenarios 1, 1a, and 1c) and had the lowest likelihood value. Although the likelihood value was higher for using both standard survey and re-tow data for males (scenario 1) than using only standard survey for males (scenario 1c), estimated abundances and biomasses were almost identical. The higher likelihood value for scenario 1 over scenario 1 c was due to trawl bycatch length compositions.

In the SAFE report in September 2020, seven models were compared. The population biomass estimates in 2020 were slightly higher than those in 2019. Absolute mature male biomasses for all models had a similar trend over time. Among the seven models, model estimated relative NMFS survey biomasses and mature biomasses were similar, especially for models 19.0a and 19.0b and for models 19.3 and 19.3a. Biomass estimates for models 19.0a and 19.0b were higher during recent years than the other five model scenarios. As expected, model 19.3b estimated a higher trawl survey catchability (>1.0), thus resulting in overall lower absolute biomass estimates. Differences of biomass estimates between models 19.0a and 19.0b and models 19.3, 19.3a, 19.31, and 19.3 h could largely be explained by different structures of natural mortality. All seven models fitted the catch and bycatch biomasses very well.

In this report (September 2022), three models are compared. For negative likelihood value comparisons (Table 5b), models 22.0a has a higher negative likelihood value. High base $M$ values estimated inside the models generally result in significantly higher total likelihood values.

The first model, 21.1b, has very close results to the base model in 2021 and was considered to be the "base" for this assessment as of May 2022. The GMACS update and the groundfish fisheries bycatch update hardly affected the results. Model 21.1b was used to compare the other two model scenarios, both of which were presented in May 2022 and chosen as potential candidates for specification setting.

The other models presented here both start in 1985 and were used to evaluate model starting year and estimating a $M$ value for males in the model. Model 22.0 is the short data version of model 21.1b and the overall results are similar. The notable differences are smaller $B_{35 \%}$ ( $21,896 \mathrm{t}$ vs $24,026 \mathrm{t}$ ) and NMFS survey catchability ( 0.94 vs 0.97 ), and higher MMB in the
terminal year (2022) ( $17,158 \mathrm{t}$ vs $16,953 \mathrm{t}$ ) and higher OFL ( $3,482 \mathrm{t}$ vs $3,036 \mathrm{t}$ ) for model 22.0. These differences are probably caused by a high recruitment in 1984 (associated with the very large $M$ ) being used for $B_{35 \%}$ computation for model 21.1 b and more influence of BSFRF survey data for model 22.0. Estimating a base $M$ for males for model 22.0a reduces total likelihood compared to model 22.0, slightly increases annual mature male biomass, and results in an estimated $B_{35 \%}$ about $10 \%$ lower than model 22.0 . A high $M$ also results in higher $\mathrm{F}_{35 \%}$ and OFL for model 22.0a.

Based on the model results, it appears that the choice of preferred models depends on two factors: preferred starting year and estimation of $M$. Considerations of $M$ estimation are whether to estimate a base $M$ for males for the whole time series versus a fixed base $M$. Either of these two approaches reduces the retrospective bias considerably. Concern of estimating a base $M$ for males for the whole time series is potential confounding with estimating trawl survey catchability.

Based on the above considerations, we recommend model 21.1b (a fixed base $M$ of 0.18 for males, less confounding between estimating $M$ and survey catchability) or model 22.0 (avoiding dramatic abundance decline during the early 1980s, no recruitment associated with an extremely high $M$ being used for estimating $B_{35 \%}$, and acceptable data fittings) for specification setting for September 2022. The base model is considered a good option if moving the starting date is not a desired change this assessment cycle. Values for management-related quantities for all models are summarized in likelihood Tables 5b.

## F. Calculation of the OFL and ABC

1. Bristol Bay RKC is currently placed in Tier 3b (NPFMC 2007).
2. For Tier 3 stocks, estimated biological reference points include $B_{35 \%}$ and $F_{35 \%}$. Estimated model parameters are used to conduct mature male biomass-per-recruit analysis.
3. Specification of the OFL:

The Tier 3 control rule formula is as follows:
a) $\frac{B}{B^{*}}>1$
$F_{O F L}=F^{*}$
b) $\quad \beta<\frac{B}{B^{*}} \leq 1$
$F_{O F L}=F^{*}\left(\frac{B / B^{*}-\alpha}{1-\alpha}\right)$
c) $\frac{B}{B^{*}} \leq \beta$
directed pot fishery $F=0$ and $F_{O F L} \leq F^{*}$

Where
$B=$ a measure of the productive capacity of the stock such as spawning biomass or fertilized egg production. A proxy of $B$ is mature male biomass (MMB) estimated at the time of primiparous female mating (February 15).
$F^{*}=F_{35 \%}$, a proxy of $F_{M S Y}$, which is a full selection instantaneous $F$ that will produce MSY at the MSY producing biomass.
$B^{*}=B_{35 \%}$, a proxy of $B_{M S Y}$, which is the value of biomass at the MSY producing level.
$\beta=$ a parameter with a restriction that $0 \leq \beta<1$. A default value of 0.25 is used.
$\alpha=$ a parameter with a restriction that $0 \leq \alpha \leq \beta$. A default value of 0.1 is used.
Because trawl bycatch fishing mortality is not related to pot fishing mortality, average trawl bycatch fishing mortality during 2017 to 2021 is used for the per recruit analysis as well as for projections in the next section. Some discards of legal males occurred after the Individual Fishery Quota (IFQ) fishery started in 2005, but the discard rates were much lower during 2007-2013 than in 2005 after the fishing industry minimized discards of legal males. However, due to high proportions of large oldshell males, the discard rate increased greatly in 2014. The current models estimate two levels of retained proportions before 2005 and after 2004. The retained proportions after 2004 and total male selectivities are used to represent current trends for per recruit analysis and projections. Average molting probabilities during 2016-2021 are used for per recruit analysis and projections. For the models in 2022, the averages are the same since they are constant over time during at least the last 15 years.
Average recruitments during 1984-2021 for models starting in 1975 and during 1986-2021 for models starting in 1985 are used to estimate $B_{35 \%}$ (Figure 12a). Estimated $B_{35 \%}$ is compared with historical mature male biomass in Figure 14a. The period of 1984-2021 corresponds to the 1976/77 regime shift, and the recruitment period 1984-present has been used since 2011 to set the overfishing limits. Several factors support our recommendation. First, estimated recruitment was lower after 1983 than before 1984, which corresponded to brood years 1978 and later, after the 1976/77 regime shift. Second, high recruitments during the late 1960s and 1970s generally occurred when the spawning stock was primarily located in the southern Bristol Bay, whereas the recent spawning stock has been concentrated in the middle of Bristol Bay. Oceanic current flows favor larvae hatched in the southern Bristol Bay (see the section on Ecosystem Considerations for SAFE reports in 2008 and 2009). Finally, stock productivity (recruitment/mature male biomass) was higher before the 1976/1977 regime shift.

The control rule is used for stock status determination. If total catch exceeds OFL estimated at $B$, then "overfishing" occurs. If $B$ equals or declines below $0.5 B_{M S Y}$ (i.e., MSST), the stock is "overfished." If $B / B_{M S Y}$ or $B / B_{M S Y}$-proxy equals or declines below $\beta$, then the stock productivity is severely depleted, and the directed fishery is closed.

The estimated probability distributions of MMB in 2023 are illustrated in Figure 30 for model 21.1b. Based on SSC suggestions in 2011, $\mathrm{ABC}=0.9^{*}$ OFL and in October 2018, $\mathrm{ABC}=0.8^{*} \mathrm{OFL}$. The CPT then recommended $\mathrm{ABC}=0.8^{*} \mathrm{OFL}$ in May 2018 (accepted by the SSC ), which is used to estimate ABC in this report. Due to the stock being close to an overfished condition and the lack of a 2020 survey, the CPT recommended an additional 5\%
buffer in September 2020, resulting in $\mathrm{ABC}=0.75^{*}$ OFL for 2020. A $20 \%$ buffer was suggested by the CPT for 2021.
MCMC runs with 500,000 replicates and 500 draws with model 21.1 b are used for estimating the probability of estimated mature male biomass being below the minimum threshold ( $0.5 * B_{35 \%}$ ) (Figure 31). The probability (converted to a percentage) is estimated to be about $0 \%$ for model 21.1b (Figure 31).

Status and catch specifications (1,000 t) (model 21.1b):

| Year | MSST | Biomass <br> $(\mathbf{M M B})$ | TAC | Retained <br> Catch | Total <br> Catch | OFL | ABC |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2018 / 19$ | 10.62 B | 16.92 B | 1.95 | 2.03 | 2.65 | 5.34 | 4.27 |
| $2019 / 20$ | 12.72 C | 14.24 C | 1.72 | 1.78 | 2.22 | 3.40 | 2.72 |
| $2020 / 21$ | 12.12 D | 13.96 D | 1.20 | 1.26 | 1.57 | 2.14 | 1.61 |
| $2021 / 22$ | 12.01 | 16.64 | 0 | 0.02 | 0.10 | 2.23 | 1.78 |
| $2022 / 23$ |  | 16.95 |  |  |  | 3.04 | 2.43 |

The stock was above MSST in 2021/22 and hence was not overfished. Since total catch was below OFL, overfishing did not occur. The relatively low MSST in 2018/19 and $B_{M S Y}$ in 2019/20 below was caused by a problem of the previous GMACS version using the only sex ratio of recruitment in the terminal year for $B_{35 \%}$ computation in 2019. The lower estimated male recruitment ratio in the terminal year in 2019 resulted in a lower mean male recruitment for $B_{35 \%}$ computation. The current version of GMACS uses an average of sex ratios of recruitment during the reference period to estimate $B_{35 \%}$, which results in a much more stable sex ratio (about 50\%) for the reference point calculation.

Status and catch specifications (million lb) (model 21.1b):

| Year | MSST | Biomass <br> (MMB) | TAC | Retained <br> Catch | Total <br> Catch | OFL | ABC |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2018 / 19$ | 23.4 B | 37.3 B | 4.31 | 4.31 | 5.85 | 11.76 | 9.41 |
| $2019 / 20$ | 28.0 C | 31.4 C | 3.80 | 3.91 | 4.89 | 7.50 | 6.00 |
| $2020 / 21$ | 26.7 D | 30.8 D | 2.77 | 2.65 | 3.47 | 4.72 | 3.54 |
| $2021 / 22$ | 26.5 | 36.7 | 0 | 0.04 | 0.22 | 4.91 | 3.92 |
| $2022 / 23$ |  | 37.4 |  |  |  | 6.70 | 5.35 |

Notes:
A - Calculated from the assessment reviewed by the Crab Plan Team in September 2018
B - Calculated from the assessment reviewed by the Crab Plan Team in September 2019
C - Calculated from the assessment reviewed by the Crab Plan Team in September 2020
D - Calculated from the assessment reviewed by the Crab Plan Team in September 2021

Basis for the OFL: Values are in $1,000 \mathrm{t}$ (model 21.1b):

| Year | Tier | BMSY | Current <br> MMB | B/BMSY <br> (MMB) | FofL | Years to <br> define <br> BMSY | Natural <br> Mortality |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2018 / 19$ | 3 b | 25.5 | 20.8 | 0.82 | 0.25 | $1984-2017$ | 0.18 |
| $2019 / 20$ | 3 b | 21.2 | 16.0 | 0.75 | 0.22 | $1984-2018$ | 0.18 |
| $2020 / 21$ | 3 b | 25.4 | 14.9 | 0.59 | 0.16 | $1984-2019$ | 0.18 |
| $2021 / 22$ | 3b | 24.2 | 14.9 | 0.62 | 0.17 | $1984-2020$ | 0.18 |
| $2022 / 23$ | 3b | 24.03 | 17.0 | 0.71 | 0.20 | $1984-2021$ | 0.18 |

Basis for the OFL: Values are in million lb (model 21.1b):

| Year | Tier | BMSY | Current <br> MMB | B/BMSY <br> (MMB) | FofL | Years to <br> define <br> BMSY | Natural <br> Mortality |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2018 / 19$ | 3b | 56.2 | 45.9 | 0.82 | 0.25 | $1984-2017$ | 0.18 |
| $2019 / 20$ | 3 b | 46.8 | 35.2 | 0.75 | 0.22 | $1984-2018$ | 0.18 |
| $2020 / 21$ | 3 b | 56.1 | 32.9 | 0.59 | 0.16 | $1984-2019$ | 0.18 |
| $2021 / 22$ | 3b | 53.4 | 33.0 | 0.62 | 0.17 | $1984-2020$ | 0.18 |
| $2022 / 23$ | 3b | 53.0 | 37.4 | 0.71 | 0.20 | $1984-2021$ | 0.18 |

Based on the $B_{35 \%}$ estimated from the average male recruitment during 1984-2021, the biological reference points and OFL are illustrated in Table 5 b.

Based on the CPT/SSC recommendation of 20\% buffer rule in May 2018 and an additional buffer of $5 \%$ by the CPT for 2020 due to the lack of a 2020 survey, $\mathrm{ABC}=0.75 *$ OFL (Table 4 ). A $20 \%$ buffer was recommended by the CPT for ABC estimation for 2021/22. A $20 \%$ buffer is also recommended for 2022/23 for similar reasoning as 2021/22.

## G. Rebuilding Analyses

NA

## H. Data Gaps and Research Priorities

1. The following data gaps exist for this stock:
a. Information about changes in natural mortality in the early 1980s,
b. Un-observed trawl bycatch in the early 1980s,
c. Natural mortality,
d. Crab availability to the trawl surveys,
e. Juvenile crab abundance,
f. Female growth per molt as a function of size and maturity,
g. Changes in male molting probability over time,
h. A better understanding of larval distribution and subsequent recruit distribution.
2. Research priorities:
a. Estimating natural mortality,
b. Estimating crab availability to the trawl surveys,
c. Surveying juvenile crab abundance in nearshore,
d. Studying environmental factors that affect the survival rates from larvae to recruitment.

## I. Projections and Future Outlook

## 1. Projections

Future population projections primarily depend on future recruitment, but crab recruitment is difficult to predict. Therefore, annual recruitment for the projections is a random selection from estimated recruitments during 2013-2021, a low recruitment period. Four levels of fishing mortality for the directed pot fishery are used in the projections: $0,0.083,0.167$ and 0.25 . A fishing mortality of 0.167 is similar to the estimated $F_{\text {of }}$ of 0.173 in 2021 with model 21.1b. MCMC runs with 500,000 replicates and 500 draws are used for the projection.

As expected, projected mature male biomasses are much higher without the directed fishing mortality than under other positive mortality values. At the end of 10 years, projected mature male biomass is below $B_{35 \%}$ for all models with a fishing mortality of 0.083 or higher due to low recruitments (Figure 32). Due to the poor recruitment in recent years, the projected biomass is expected to decline during the next few years with a fishing mortality of greater than $\mathrm{F}=0.167$.

Even though the stock is not overfished in 2021/22, there is still a question whether the stock is "approaching an overfished condition", which is defined as "when it is projected that there is more than a 50 percent chance that the biomass of the stock or stock complex will decline below the MSST within two years" by the National Standards 1 (NS1). If the stock is not fished more than a fishing mortality of 0.167 for the directed pot fishery in the 2022/2023and 2023/2024 seasons, the projection using the lowest recruitment periods during 2013-2021 would not likely result in "approaching an overfished condition" for model 21.1b (Figure 33). With additional low recruitment estimate used to compute $B_{35 \%}$, the estimated MSST would decline further in 2023.

The projections are subject to many uncertainties. Constant population parameters estimated in the models used for the projections include $M$, growth, and fishery selectivities. The uncertainty of abundance and biomass estimates in the terminal year also affects the projections. Uncertainties of the projections caused by these constant parameters and abundance estimates in the terminal year would be reduced by the $20 \%$ ABC buffer. However, if an extreme event occurs, like a sharp increase of $M$ during the projection period, the ABC buffer would be inadequate, and the projections might underestimate uncertainties. The largest uncertainty is likely from recruitments used for the projections. Higher or lower assumed recruitments would cause too optimistic or too pessimistic projections. Overall, recruitments and $M$ used for projections are main factors for projection uncertainties.

## 2. Near Future Outlook

The near future outlook for the Bristol Bay RKC stock is a steady to declining trend. The three recent above-average year classes (hatching years 1990, 1994, and 1997) had entered the legal population by 2006 (Figure 34). Most individuals from the 1997-year class will continue to gain weight to offset loss of the legal biomass to fishing and natural mortalities. The above-average year class (hatching year 2000) with lengths centered around 87.5 mm CL for both males and females in 2006 and with lengths centered around $112.5-117.5 \mathrm{~mm}$ CL for males and around 107.5 mm CL for females in 2008 has largely entered the mature male population in 2009 and the legal population by 2014 (Figures 5a, b). However, no additional strong cohorts were observed in the survey data after this cohort through 2010 (Figure 5a, 5b, and 34). A huge tow of juvenile crab of size $45-55 \mathrm{~mm}$ in 2011 was not tracked during 2012-2022 surveys and is unlikely to be a strong cohort. The high survey abundances of large males and mature females in 2014 cannot be explained by the survey data during the previous years and were also inconsistent with the 2016-2022 survey results (Figure 34). Due to lack of recruitment, mature and legal crab may continue to decline next year in the presence of fishing pressure. However, this past year suggests that lack of a directed fishery and a small increase in recruitment contributed to an increase in abundance. Current crab abundance is still low relative to the late 1970s, and without favorable environmental conditions, recovery to the high levels of the late 1970s is unlikely.

Although mature crab abundance in Bristol Bay has declined in recent years, mature crab abundance and biomass north of Bristol Bay has been generally stable during last 16 years (Figures 35 a and 35 b). Overall, the proportions of different size groups of the Northern RKC during a recent dozen years are higher than in the past and do not trend higher except for mature females in 2021. The high survey mature female abundance in the Northern area in 2021 was primarily from three tows and one of them is more than $50 \%$ of total mature females. The survey abundance of the Northern RKC will continue to be provided in figures in the SAFE report in the future. After migration patterns between BBRKC and the Northern RKC are more fully understood, we will examine their relationships and model them in the stock assessment.

## J. Acknowledgements

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## Tables

Table 1a. Bristol Bay red king crab annual catch and bycatch mortality biomass (t) from July 1 to June 30. A handling mortality rate of $20 \%$ for the directed pot, $25 \%$ for the Tanner fishery, $80 \%$ for trawl, and $50 \%$ or fixed gear was assumed to estimate bycatch mortality biomass. The male bycatch biomass in the directed pot fishery is not estimated outside of a model and not included in this Table. Pot bycatch and Tanner crab fishery bycatch are estimated through expanding the mean observer bycatch per pot to total fishery pot. The pot male bycatch after 2017 is estimated through the subtraction method (B. Daly, ADF\&G, personal communication). The trawl and fixed gear fishery bycatches are obtained from the NMFS database. The directed pot bycatch before 1990 and Tanner crab fishery bycatch before 1991 are not available from the observer data and thus not included in this table. These include recently updated estimates from the pot fisheries observer data in 2021.

| Year | Retained Catch |  |  |  | Pot BycatchFemales | Trawl <br> Bycatch | Fixed <br> Bycatch | Tanner Fishery Bycatch |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | U.S. | CostRecovery | Foreign | Total |  |  |  |  |
| 1953 | 1331.3 |  | 4705.6 | 6036.9 |  |  |  |  |
| 1954 | 1149.9 |  | 3720.4 | 4870.2 |  |  |  |  |
| 1955 | 1029.2 |  | 3712.7 | 4741.9 |  |  |  |  |
| 1956 | 973.4 |  | 3572.9 | 4546.4 |  |  |  |  |
| 1957 | 339.7 |  | 3718.1 | 4057.8 |  |  |  |  |
| 1958 | 3.2 |  | 3541.6 | 3544.8 |  |  |  |  |
| 1959 | 0.0 |  | 6062.3 | 6062.3 |  |  |  |  |
| 1960 | 272.2 |  | 12200.7 | 12472.9 |  |  |  |  |
| 1961 | 193.7 |  | 20226.6 | 20420.3 |  |  |  |  |
| 1962 | 30.8 |  | 24618.7 | 24649.6 |  |  |  |  |
| 1963 | 296.2 |  | 24930.8 | 25227.0 |  |  |  |  |
| 1964 | 373.3 |  | 26385.5 | 26758.8 |  |  |  |  |
| 1965 | 648.2 |  | 18730.6 | 19378.8 |  |  |  |  |
| 1966 | 452.2 |  | 19212.4 | 19664.6 |  |  |  |  |
| 1967 | 1407.0 |  | 15257.0 | 16664.1 |  |  |  |  |
| 1968 | 3939.9 |  | 12459.7 | 16399.6 |  |  |  |  |
| 1969 | 4718.7 |  | 6524.0 | 11242.7 |  |  |  |  |
| 1970 | 3882.3 |  | 5889.4 | 9771.7 |  |  |  |  |
| 1971 | 5872.2 |  | 2782.3 | 8654.5 |  |  |  |  |
| 1972 | 9863.4 |  | 2141.0 | 12004.3 |  |  |  |  |
| 1973 | 12207.8 |  | 103.4 | 12311.2 |  |  |  |  |
| 1974 | 19171.7 |  | 215.9 | 19387.6 |  |  |  |  |
| 1975 | 23281.2 |  | 0 | 23281.2 |  |  |  |  |
| 1976 | 28993.6 |  | 0 | 28993.6 |  | 682.8 |  |  |
| 1977 | 31736.9 |  | 0 | 31736.9 |  | 1249.9 |  |  |
| 1978 | 39743.0 |  | 0 | 39743.0 |  | 1320.6 |  |  |
| 1979 | 48910.0 |  | 0 | 48910.0 |  | 1331.9 |  |  |
| 1980 | 58943.6 |  | 0 | 58943.6 |  | 1036.5 |  |  |
| 1981 | 15236.8 |  | 0 | 15236.8 |  | 219.4 |  |  |
| 1982 | 1361.3 |  | 0 | 1361.3 |  | 574.9 |  |  |
| 1983 | 0.0 |  | 0 | 0.0 |  | 420.4 |  |  |
| 1984 | 1897.1 |  | 0 | 1897.1 |  | 1094.0 |  |  |
| 1985 | 1893.8 |  | 0 | 1893.8 |  | 390.1 |  |  |
| 1986 | 5168.2 |  | 0 | 5168.2 |  | 200.6 |  |  |
| 1987 | 5574.2 |  | 0 | 5574.2 |  | 186.4 |  |  |
| 1988 | 3351.1 |  | 0 | 3351.1 |  | 598.4 |  |  |
| 1989 | 4656.0 |  | 0 | 4656.0 |  | 175.2 |  |  |
| 1990 | 9236.2 | 36.6 | 0 | 9272.8 | 639.2 | 259.9 |  |  |
| 1991 | 7791.8 | 93.4 | 0 | 7885.1 | 46.8 | 349.4 |  | 1401.8 |


| 1992 | 3648.2 | 33.6 | 0 | 3681.8 | 395.3 | 293.5 | 244.4 |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1993 | 6635.4 | 24.1 | 0 | 6659.6 | 628.3 | 401.4 | 54.6 |  |
| 1994 | 0.0 | 42.3 | 0 | 42.3 | 0.4 | 87.3 | 10.8 |  |
| 1995 | 0.0 | 36.4 | 0 | 36.4 | 0.3 | 82.1 | 0.0 |  |
| 1996 | 3812.7 | 49.0 | 0 | 3861.7 | 1.0 | 90.8 | 41.4 | 0.0 |
| 1997 | 3971.9 | 70.2 | 0 | 4042.1 | 36.5 | 57.5 | 22.5 | 0.0 |
| 1998 | 6693.8 | 85.4 | 0 | 6779.2 | 553.9 | 186.1 | 18.5 | 0.0 |
| 1999 | 5293.5 | 84.3 | 0 | 5377.9 | 5.6 | 150.5 | 50.1 | 0.0 |
| 2000 | 3698.8 | 39.1 | 0 | 3737.9 | 164.4 | 81.7 | 4.7 | 0.0 |
| 2001 | 3811.5 | 54.6 | 0 | 3866.2 | 120.8 | 192.8 | 35.3 | 0.0 |
| 2002 | 4340.9 | 43.6 | 0 | 4384.5 | 9.1 | 151.2 | 29.2 | 0.0 |
| 2003 | 7120.0 | 15.3 | 0 | 7135.3 | 356.9 | 136.9 | 12.7 | 0.0 |
| 2004 | 6915.2 | 91.4 | 0 | 7006.7 | 171.8 | 173.5 | 15.2 | 0.0 |
| 2005 | 8305.0 | 94.7 | 0 | 8399.7 | 405.4 | 124.7 | 19.9 | 0.0 |
| 2006 | 7005.3 | 137.9 | 0 | 7143.2 | 37.5 | 151.7 | 19.6 | 3.8 |
| 2007 | 9237.9 | 66.1 | 0 | 9303.9 | 159.9 | 154.1 | 32.3 | 1.8 |
| 2008 | 9216.1 | 0.0 | 0 | 9216.1 | 144.8 | 136.6 | 15.6 | 4.0 |
| 2009 | 7226.9 | 45.5 | 0 | 7272.5 | 88.3 | 94.9 | 5.8 | 1.6 |
| 2010 | 6728.5 | 33.0 | 0 | 6761.5 | 118.5 | 83.2 | 2.4 | 0.0 |
| 2011 | 3553.3 | 53.8 | 0 | 3607.1 | 25.0 | 56.2 | 10.9 | 0.0 |
| 2012 | 3560.6 | 61.1 | 0 | 3621.7 | 11.2 | 34.1 | 18.4 | 0.0 |
| 2013 | 3901.1 | 89.9 | 0 | 3991.0 | 98.1 | 66.9 | 55.1 | 28.5 |
| 2014 | 4530.0 | 8.6 | 0 | 4538.6 | 84.9 | 34.5 | 118.7 | 42.0 |
| 2015 | 4522.3 | 91.4 | 0 | 4613.7 | 239.1 | 45.1 | 77.4 | 84.2 |
| 2016 | 3840.4 | 83.4 | 0 | 3923.9 | 123.4 | 67.3 | 29.7 | 0.0 |
| 2017 | 2994.1 | 99.6 | 0 | 3093.7 | 53.4 | 91.7 | 130.0 | 0.0 |
| 2018 | 1954.1 | 72.4 | 0 | 2026.5 | 150.1 | 78.0 | 154.7 | 0.0 |
| 2019 | 1719.8 | 55.5 | 0 | 1775.3 | 43.3 | 80.7 | 45.1 | 0.0 |
| 2020 | 1200.6 | 56.4 | 0 | 1257.0 | 15.2 | 80.7 | 37.6 | 0.0 |
| 2021 | 0.0 | 17.4 | 0 | 17.4 | 5.9 | 34.4 | 40.3 | 0.0 |

Table 1b. Annual retained catch (millions of crab) and catch per unit effort of the Bristol Bay red king crab fishery.

| Year | Japanese Tanglenet |  | Russian Tanglenet |  | U.S. Pot |  | Standardized Crab/tan |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Catch | Crab/tan | Catch | Crab/tan | Catch | Crab/Potlift |  |
| 1960 | 1.949 | 15.2 | 1.995 | 10.4 | 0.088 |  | 15.8 |
| 1961 | 3.031 | 11.8 | 3.441 | 8.9 | 0.062 |  | 12.9 |
| 1962 | 4.951 | 11.3 | 3.019 | 7.2 | 0.010 |  | 11.3 |
| 1963 | 5.476 | 8.5 | 3.019 | 5.6 | 0.101 |  | 8.6 |
| 1964 | 5.895 | 9.2 | 2.800 | 4.6 | 0.123 |  | 8.5 |
| 1965 | 4.216 | 9.3 | 2.226 | 3.6 | 0.223 |  | 7.7 |
| 1966 | 4.206 | 9.4 | 2.560 | 4.1 | 0.140 | 52 | 8.1 |
| 1967 | 3.764 | 8.3 | 1.592 | 2.4 | 0.397 | 37 | 6.3 |
| 1968 | 3.853 | 7.5 | 0.549 | 2.3 | 1.278 | 27 | 7.8 |
| 1969 | 2.073 | 7.2 | 0.369 | 1.5 | 1.749 | 18 | 5.6 |
| 1970 | 2.080 | 7.3 | 0.320 | 1.4 | 1.683 | 17 | 5.6 |
| 1971 | 0.886 | 6.7 | 0.265 | 1.3 | 2.405 | 20 | 5.8 |
| 1972 | 0.874 | 6.7 |  |  | 3.994 | 19 |  |
| 1973 | 0.228 |  |  |  | 4.826 | 25 |  |
| 1974 | 0.476 |  |  |  | 7.710 | 36 |  |
| 1975 |  |  |  |  | 8.745 | 43 |  |
| 1976 |  |  |  |  | 10.603 | 33 |  |
| 1977 |  |  |  |  | 11.733 | 26 |  |
| 1978 |  |  |  |  | 14.746 | 36 |  |
| 1979 |  |  |  |  | 16.809 | 53 |  |
| 1980 |  |  |  |  | 20.845 | 37 |  |
| 1981 |  |  |  |  | 5.308 | 10 |  |
| 1982 |  |  |  |  | 0.541 | 4 |  |
| 1983 |  |  |  |  | No directed | fishery |  |
| 1984 |  |  |  |  | 0.794 | 7 |  |
| 1985 |  |  |  |  | 0.796 | 9 |  |
| 1986 |  |  |  |  | 2.100 | 12 |  |
| 1987 |  |  |  |  | 2.122 | 10 |  |
| 1988 |  |  |  |  | 1.236 | 8 |  |
| 1989 |  |  |  |  | 1.685 | 8 |  |
| 1990 |  |  |  |  | 3.130 | 12 |  |
| 1991 |  |  |  |  | 2.661 | 12 |  |
| 1992 |  |  |  |  | 1.208 | 6 |  |
| 1993 |  |  |  |  | 2.270 | 9 |  |
| 1994 |  |  |  |  | No directed | fishery |  |
| 1995 |  |  |  |  | No directed | fishery |  |
| 1996 |  |  |  |  | 1.264 | 16 |  |
| 1997 |  |  |  |  | 1.338 | 15 |  |
| 1998 |  |  |  |  | 2.238 | 15 |  |
| 1999 |  |  |  |  | 1.923 | 12 |  |
| 2000 |  |  |  |  | 1.272 | 12 |  |
| 2001 |  |  |  |  | 1.287 | 19 |  |
| 2002 |  |  |  |  | 1.484 | 20 |  |
| 2003 |  |  |  |  | 2.510 | 18 |  |
| 2004 |  |  |  |  | 2.272 | 23 |  |
| 2005 |  |  |  |  | 2.763 | 30 |  |
| 2006 |  |  |  |  | 2.477 | 31 |  |
| 2007 |  |  |  |  | 3.154 | 28 |  |
| 2008 |  |  |  |  | 3.064 | 22 |  |
| 2009 |  |  |  |  | 2.553 | 21 |  |
| 2010 |  |  |  |  | 2.410 | 18 |  |
| 2011 |  |  |  |  | 1.298 | 28 |  |
| 2012 |  |  |  |  | 1.176 | 30 |  |
| 2013 |  |  |  |  | 1.272 | 27 |  |
| 2014 |  |  |  |  | 1.501 | 26 |  |
| 2015 |  |  |  |  | 1.527 | 31 |  |
| 2016 |  |  |  |  | 1.281 | 38 |  |
| 2017 |  |  |  |  | 0.997 | 20 |  |
| 2018 |  |  |  |  | 0.630 | 20 |  |
| 2019 |  |  |  |  | 0.549 | 16 |  |
| 2020 |  |  |  |  | 0.455 | 21 |  |
| 2021 |  |  |  |  | No directed | fishery |  |

Table 2. Total observer catch and bycatch (metric ton) of Bristol Bay red king crab. No handling mortality rates are applied. These include recently updated estimates from the pot fishery observer data in 2021.

| Year | Directed Pot Total |  | Trawl Bycatch | Fixed Bycatch | Tanner Bycatch |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | Males | Females |  |  |  |
| 1975 |  |  | 0.000 |  |  |
| 1976 |  |  | 853.494 |  |  |
| 1977 |  |  | 1,562.313 |  |  |
| 1978 |  |  | 1,650.775 |  |  |
| 1979 |  |  | 1,664.925 |  |  |
| 1980 |  |  | 1,295.625 |  |  |
| 1981 |  |  | 274.229 |  |  |
| 1982 |  |  | 718.610 |  |  |
| 1983 |  |  | 525.554 |  |  |
| 1984 |  |  | 1,367.550 |  |  |
| 1985 |  |  | 487.576 |  |  |
| 1986 |  |  | 250.758 |  |  |
| 1987 |  |  | 233.045 |  |  |
| 1988 |  |  | 747.996 |  |  |
| 1989 |  |  | 219.023 |  |  |
| 1990 | 11621.800 | 3196.200 | 324.883 |  |  |
| 1991 | 9792.900 | 233.900 | 436.783 |  | 5,580.843 |
| 1992 | 5916.200 | 1976.300 | 366.816 |  | 962.846 |
| 1993 | 9516.800 | 3141.500 | 501.770 |  | 218.112 |
| 1994 | 62.300 | 1.877 | 109.129 |  | 39.395 |
| 1995 | 52.800 | 1.612 | 102.623 |  | 0.000 |
| 1996 | 3845.200 | 5.100 | 113.495 | 82.859 | 0.000 |
| 1997 | 3758.800 | 182.700 | 71.862 | 44.979 | 0.000 |
| 1998 | 15644.800 | 2769.300 | 232.580 | 36.916 | 0.000 |
| 1999 | 12112.300 | 28.000 | 188.101 | 100.242 | 0.000 |
| 2000 | 6579.700 | 821.900 | 102.161 | 9.446 | 0.000 |
| 2001 | 5711.500 | 604.000 | 241.011 | 70.553 | 0.000 |
| 2002 | 6961.400 | 45.600 | 189.018 | 58.382 | 0.000 |
| 2003 | 12166.500 | 1784.400 | 171.114 | 25.351 | 0.000 |
| 2004 | 10692.000 | 859.200 | 216.889 | 30.422 | 0.000 |
| 2005 | 13615.900 | 2027.100 | 155.924 | 39.802 | 0.000 |
| 2006 | 9254.000 | 187.400 | 189.660 | 39.134 | 15.217 |
| 2007 | 13871.900 | 799.400 | 192.571 | 64.655 | 7.142 |
| 2008 | 14894.900 | 724.200 | 170.754 | 31.158 | 16.070 |
| 2009 | 12218.800 | 441.300 | 118.672 | 11.614 | 6.499 |
| 2010 | 10095.400 | 592.600 | 104.005 | 4.944 | 0.000 |
| 2011 | 5665.300 | 124.800 | 70.286 | 21.726 | 0.000 |
| 2012 | 4495.500 | 55.900 | 42.641 | 36.897 | 0.000 |
| 2013 | 5305.900 | 490.700 | 83.613 | 110.208 | 113.063 |
| 2014 | 8113.800 | 424.300 | 43.129 | 237.374 | 137.786 |
| 2015 | 6726.800 | 1195.600 | 56.410 | 154.775 | 639.573 |
| 2016 | 5651.800 | 617.200 | 84.127 | 59.418 | 0.000 |
| 2017 | 4077.200 | 266.900 | 114.624 | 260.011 | 0.000 |
| 2018 | 3423.200 | 750.400 | 97.561 | 309.415 | 0.000 |
| 2019 | 3144.600 | 218.000 | 100.915 | 90.291 | 0.000 |
| 2020 | 2299.700 | 76.100 | 100.842 | 75.130 | 0.000 |
| 2021 | 33.800 | 29.400 | 42.99 | 80.602 | 0.000 |

Table 3a. Annual sample sizes (>64 mm CL) in numbers of crab for trawl surveys, retained catch, directed pot, Tanner crab, trawl, and fixed gear fishery bycatches of Bristol Bay red king crab. These include recently updated estimates from the pot fisheries observer data and are used for models 2022.

| Year | Trawl Survey |  | Retained Catch | Pot Total <br> Males | Pot <br> Bycatch <br> Females | Trawl <br> Bycatch <br> Combined | Fixed G. <br> BycatchCombined | Tanner <br> Fishery <br> Bycatch <br> Combined |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Males | Females |  |  |  |  |  |  |
| 1975 | 2,815 | 2,042 | 29,570 |  |  |  |  |  |
| 1976 | 2,699 | 1,466 | 26,450 |  |  | 3,003 |  |  |
| 1977 | 2,734 | 2,424 | 32,596 |  |  | 14,703 |  |  |
| 1978 | 2,735 | 2,793 | 27,529 |  |  | 10,439 |  |  |
| 1979 | 1,158 | 1,456 | 27,900 |  |  | 10,049 |  |  |
| 1980 | 1,917 | 1,301 | 34,747 |  |  | 87,152 |  |  |
| 1981 | 591 | 664 | 18,029 |  |  | 91,806 |  |  |
| 1982 | 1,911 | 1,948 | 11,466 |  |  | 131,469 |  |  |
| 1983 | 1,343 | 733 | 0 |  |  | 309,374 |  |  |
| 1984 | 1,209 | 778 | 4,404 |  |  | 505,115 |  |  |
| 1985 | 790 | 414 | 4,582 |  |  | 200,460 |  |  |
| 1986 | 959 | 341 | 5,773 |  |  | 2,126 |  |  |
| 1987 | 1,123 | 1,011 | 4,230 |  |  | 998 |  |  |
| 1988 | 708 | 478 | 9,833 |  |  | 630 |  |  |
| 1989 | 764 | 403 | 32,858 |  |  | 4,641 |  |  |
| 1990 | 729 | 535 | 7,218 | 2,544 | 696 | 908 |  |  |
| 1991 | 1,180 | 490 | 36,928 | 4,696 | 375 | 275 |  | 3,131 |
| 1992 | 509 | 357 | 25,550 | 4,775 | 2,379 | 333 |  | 965 |
| 1993 | 725 | 576 | 32,942 | 10,200 | 5,944 | 5 |  | 497 |
| 1994 | 416 | 239 | 0 | 0 | 0 | 571 |  | 17 |
| 1995 | 685 | 407 | 0 | 0 | 0 | 120 |  |  |
| 1996 | 755 | 753 | 8,896 | 642 | 11 | 1,209 | 756 |  |
| 1997 | 1,280 | 702 | 16,143 | 10,016 | 906 | 339 | 1,269 |  |
| 1998 | 1,067 | 1,123 | 17,116 | 24,537 | 9,655 | 1,430 | 1,036 |  |
| 1999 | 765 | 618 | 18,685 | 6,892 | 40 | 629 | 1,602 |  |
| 2000 | 734 | 730 | 14,143 | 32,709 | 8,470 | 729 | 591 |  |
| 2001 | 599 | 736 | 13,735 | 25,135 | 5,436 | 795 | 5,029 |  |
| 2002 | 972 | 826 | 16,837 | 32,317 | 706 | 1,139 | 3,503 |  |
| 2003 | 1,360 | 1,250 | 18,178 | 44,600 | 12,474 | 516 | 1,872 |  |
| 2004 | 1,852 | 1,271 | 22,465 | 38,772 | 6,666 | 636 | 2,184 |  |
| 2005 | 1,198 | 1,563 | 27,971 | 94,622 | 26,782 | 1,040 | 2,146 |  |
| 2006 | 1,178 | 1,432 | 18,451 | 73,315 | 3,991 | 1,168 | 1,868 | 140 |
| 2007 | 1,228 | 1,305 | 22,809 | 115,507 | 12,691 | 1,225 | 785 | 53 |
| 2008 | 1,228 | 1,183 | 24,997 | 89,771 | 8,564 | 1,596 | 1,164 | 145 |
| 2009 | 837 | 941 | 19,336 | 97,868 | 6,055 | 1,170 | 1,089 | 193 |
| 2010 | 708 | 1,004 | 20,347 | 69,276 | 6,872 | 901 | 513 |  |
| 2011 | 531 | 912 | 10,904 | 42,931 | 1,920 | 439 | 1,190 |  |
| 2012 | 585 | 707 | 9,084 | 21,404 | 563 | 281 | 2,977 |  |
| 2013 | 647 | 569 | 10,396 | 32,332 | 6,051 | 481 | 8,523 | 814 |
| 2014 | 1,107 | 1,257 | 9,718 | 31,216 | 2,663 | 261 | 4,285 | 631 |
| 2015 | 615 | 681 | 11,971 | 24,533 | 7,457 | 409 | 4,472 | 2,872 |
| 2016 | 378 | 812 | 11,003 | 30,030 | 5,832 | 617 | 4,329 |  |
| 2017 | 385 | 508 | 10,067 | 30,002 | 4,043 | 718 | 1,415 |  |
| 2018 | 285 | 359 | 7,825 | 25,635 | 9,840 | 893 | 5,382 |  |
| 2019 | 273 | 299 | 8,134 | 25,999 | 2,894 | 823 | 863 |  |
| 2020 |  |  | 3,850 | 16,650 | 961 | 764 | 246 |  |
| 2021 | 324 | 247 | 101 | 1,100 | 1433 | 503 | 120 |  |
| 2022 | 401 | 319 |  |  |  |  |  |  |

Table 3b. Comparison of area-swept and VAST-based male Bristol Bay red king crab biomass estimates from the NMFS trawl survey. Difference $=($ area-swept - VAST $) /[($ area-swept + VAST $) / 2]$. Reduction $=$ (area-swept - VAST)/area-swept. VAST estimates were not computed for 2022 or included in models in 2022.

| Year | Area-swept |  | VAST |  | Biomass <br> Differ.\% | CVReduction\% |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Biomass | CV | Biomass | CV |  |  |
| 1975 | 133.084 | 0.171 | 148.119 | 0.099 | -10.69 | 42.37 |
| 1976 | 256.362 | 0.222 | 243.853 | 0.089 | 5.00 | 59.74 |
| 1977 | 232.539 | 0.176 | 239.346 | 0.080 | -2.89 | 54.39 |
| 1978 | 199.542 | 0.200 | 196.698 | 0.090 | 1.44 | 54.94 |
| 1979 | 102.448 | 0.239 | 96.579 | 0.101 | 5.90 | 57.79 |
| 1980 | 166.524 | 0.240 | 141.622 | 0.096 | 16.16 | 59.90 |
| 1981 | 68.294 | 0.144 | 73.903 | 0.081 | -7.89 | 44.07 |
| 1982 | 72.296 | 0.263 | 60.766 | 0.096 | 17.33 | 63.40 |
| 1983 | 34.762 | 0.210 | 34.590 | 0.088 | 0.50 | 58.16 |
| 1984 | 96.418 | 0.549 | 47.590 | 0.108 | 67.81 | 80.41 |
| 1985 | 26.819 | 0.154 | 29.607 | 0.090 | -9.88 | 41.62 |
| 1986 | 40.549 | 0.481 | 27.200 | 0.098 | 39.41 | 79.62 |
| 1987 | 46.769 | 0.225 | 42.384 | 0.095 | 9.84 | 57.78 |
| 1988 | 35.374 | 0.168 | 37.874 | 0.092 | -6.83 | 45.42 |
| 1989 | 42.358 | 0.222 | 40.527 | 0.094 | 4.42 | 57.83 |
| 1990 | 38.728 | 0.227 | 37.492 | 0.099 | 3.24 | 56.50 |
| 1991 | 66.528 | 0.543 | 36.916 | 0.149 | 57.25 | 72.63 |
| 1992 | 25.096 | 0.178 | 26.546 | 0.099 | -5.62 | 44.19 |
| 1993 | 35.671 | 0.210 | 36.554 | 0.109 | -2.45 | 48.32 |
| 1994 | 23.003 | 0.173 | 25.230 | 0.105 | -9.23 | 39.35 |
| 1995 | 27.252 | 0.327 | 23.646 | 0.103 | 14.17 | 68.56 |
| 1996 | 26.816 | 0.187 | 28.476 | 0.104 | -6.01 | 44.62 |
| 1997 | 59.638 | 0.244 | 55.682 | 0.101 | 6.86 | 58.76 |
| 1998 | 46.209 | 0.162 | 50.277 | 0.092 | -8.43 | 43.25 |
| 1999 | 44.529 | 0.210 | 46.095 | 0.109 | -3.46 | 48.10 |
| 2000 | 38.391 | 0.164 | 46.505 | 0.101 | -19.12 | 38.40 |
| 2001 | 27.943 | 0.146 | 31.181 | 0.088 | -10.95 | 39.84 |
| 2002 | 45.140 | 0.195 | 48.796 | 0.101 | -7.78 | 48.09 |
| 2003 | 74.641 | 0.406 | 60.035 | 0.101 | 21.69 | 75.04 |
| 2004 | 90.354 | 0.395 | 64.126 | 0.104 | 33.96 | 73.78 |
| 2005 | 54.790 | 0.181 | 55.097 | 0.098 | -0.56 | 46.06 |
| 2006 | 51.215 | 0.197 | 54.277 | 0.088 | -5.80 | 55.27 |
| 2007 | 58.144 | 0.184 | 62.256 | 0.091 | -6.83 | 50.34 |
| 2008 | 67.214 | 0.302 | 61.024 | 0.103 | 9.65 | 65.93 |
| 2009 | 43.170 | 0.365 | 39.091 | 0.113 | 9.92 | 69.05 |
| 2010 | 39.021 | 0.237 | 40.329 | 0.101 | -3.30 | 57.57 |
| 2011 | 27.385 | 0.207 | 29.640 | 0.106 | -7.91 | 48.65 |
| 2012 | 30.655 | 0.255 | 34.232 | 0.117 | -11.02 | 54.08 |
| 2013 | 39.650 | 0.207 | 42.819 | 0.105 | -7.68 | 49.11 |
| 2014 | 60.649 | 0.192 | 64.111 | 0.097 | -5.55 | 49.56 |
| 2015 | 37.085 | 0.174 | 42.030 | 0.093 | -12.50 | 46.41 |
| 2016 | 27.185 | 0.148 | 30.230 | 0.091 | -10.61 | 38.84 |
| 2017 | 25.335 | 0.174 | 26.252 | 0.086 | -3.56 | 50.61 |
| 2018 | 16.034 | 0.138 | 18.270 | 0.091 | -13.03 | 33.75 |
| 2019 | 15.170 | 0.163 | 16.262 | 0.093 | -6.95 | 42.65 |
| 2021 | 18.235 | 0.202 | 17.185 | 0.133 | 5.93 | 34.14 |
| Mean | 61.631 | 0.234 | 58.942 | 0.099 | 2.69 | 53.24 |
| Min |  |  |  |  | -19.12 | 33.75 |
| Max |  |  |  |  | 67.81 | 80.41 |

Table 3c. Comparison of area-swept and VAST-based female Bristol Bay red king crab biomass estimates from the NMFS trawl survey. Difference $=($ area-swept - VAST $) /[($ area-swept + VAST $) / 2]$. Reduction $=$ (area-swept -VAST)/area-swept. VAST estimates were not computed for 2022 or included in models in 2022.

| Year | Area-swept |  | VAST |  | Biomass <br> Differ.\% | CVReduction\% |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Biomass | CV | Biomass | CV |  |  |
| 1975 | 66.559 | 0.301 | 58.081 | 0.127 | 13.60 | 57.79 |
| 1976 | 71.252 | 0.235 | 68.255 | 0.106 | 4.30 | 55.08 |
| 1977 | 138.684 | 0.188 | 134.450 | 0.097 | 3.10 | 48.60 |
| 1978 | 143.647 | 0.196 | 125.444 | 0.099 | 13.53 | 49.30 |
| 1979 | 63.001 | 0.179 | 53.741 | 0.091 | 15.86 | 49.34 |
| 1980 | 80.701 | 0.327 | 67.448 | 0.118 | 17.89 | 63.92 |
| 1981 | 62.850 | 0.257 | 55.937 | 0.107 | 11.64 | 58.30 |
| 1982 | 69.601 | 0.251 | 61.728 | 0.103 | 11.99 | 58.91 |
| 1983 | 13.714 | 0.247 | 11.953 | 0.106 | 13.72 | 56.95 |
| 1984 | 56.189 | 0.710 | 19.191 | 0.154 | 98.16 | 78.28 |
| 1985 | 7.319 | 0.251 | 6.680 | 0.116 | 9.12 | 53.59 |
| 1986 | 6.885 | 0.331 | 5.835 | 0.122 | 16.51 | 63.20 |
| 1987 | 22.476 | 0.320 | 17.208 | 0.125 | 26.55 | 61.01 |
| 1988 | 19.224 | 0.411 | 13.843 | 0.153 | 32.55 | 62.72 |
| 1989 | 12.778 | 0.347 | 9.644 | 0.121 | 27.95 | 65.03 |
| 1990 | 20.723 | 0.401 | 14.301 | 0.138 | 36.67 | 65.47 |
| 1991 | 17.364 | 0.415 | 11.900 | 0.124 | 37.34 | 70.14 |
| 1992 | 12.238 | 0.247 | 10.797 | 0.116 | 12.51 | 53.03 |
| 1993 | 17.235 | 0.248 | 15.702 | 0.127 | 9.31 | 48.83 |
| 1994 | 9.102 | 0.219 | 8.425 | 0.126 | 7.72 | 42.42 |
| 1995 | 10.816 | 0.247 | 9.454 | 0.117 | 13.44 | 52.54 |
| 1996 | 17.143 | 0.270 | 14.672 | 0.126 | 15.54 | 53.41 |
| 1997 | 24.392 | 0.352 | 19.315 | 0.131 | 23.23 | 62.79 |
| 1998 | 37.893 | 0.250 | 31.954 | 0.113 | 17.01 | 54.82 |
| 1999 | 20.225 | 0.339 | 19.950 | 0.138 | 1.37 | 59.28 |
| 2000 | 28.991 | 0.330 | 31.734 | 0.143 | -9.04 | 56.73 |
| 2001 | 24.513 | 0.294 | 21.338 | 0.123 | 13.85 | 58.20 |
| 2002 | 23.947 | 0.289 | 20.469 | 0.122 | 15.66 | 57.63 |
| 2003 | 41.119 | 0.221 | 37.258 | 0.114 | 9.85 | 48.58 |
| 2004 | 40.202 | 0.255 | 32.518 | 0.109 | 21.13 | 57.43 |
| 2005 | 50.937 | 0.205 | 44.651 | 0.109 | 13.15 | 46.93 |
| 2006 | 43.262 | 0.200 | 54.154 | 0.113 | -22.36 | 43.48 |
| 2007 | 45.183 | 0.223 | 53.047 | 0.105 | -16.01 | 53.10 |
| 2008 | 45.867 | 0.322 | 47.268 | 0.124 | -3.01 | 61.57 |
| 2009 | 47.377 | 0.327 | 45.385 | 0.120 | 4.29 | 63.32 |
| 2010 | 41.480 | 0.271 | 42.706 | 0.119 | -2.91 | 56.21 |
| 2011 | 39.023 | 0.256 | 41.777 | 0.121 | -6.82 | 52.62 |
| 2012 | 30.042 | 0.334 | 30.582 | 0.150 | -1.78 | 55.21 |
| 2013 | 22.567 | 0.359 | 22.856 | 0.145 | -1.27 | 59.51 |
| 2014 | 52.486 | 0.227 | 65.939 | 0.129 | -22.72 | 43.09 |
| 2015 | 27.090 | 0.295 | 30.854 | 0.133 | -12.99 | 54.81 |
| 2016 | 33.773 | 0.259 | 36.498 | 0.114 | -7.75 | 55.92 |
| 2017 | 27.599 | 0.250 | 29.231 | 0.106 | -5.74 | 57.70 |
| 2018 | 12.771 | 0.224 | 14.247 | 0.117 | -10.93 | 47.79 |
| 2019 | 13.369 | 0.185 | 15.989 | 0.100 | -17.85 | 46.11 |
| 2021 | 10.241 | 0.244 | 10.576 | 0.109 | -3.23 | 55.17 |
| Mean | 37.475 | 0.285 | 34.674 | 0.120 | 9.22 | 56.00 |
| Min |  |  |  |  | -22.72 | 42.42 |
| Max |  |  |  |  | 98.16 | 78.28 |

Table 4. Comparison of harmonic means of implied sample sizes and maximum caps ( N ) of effective sample sizes for models 21.1b, 22.0, and 22.0a.

|  |  | Models |  |  |  |  |
| :--- | ---: | ---: | ---: | ---: | :---: | :---: |
|  | 21.1 b |  |  | 22.0 |  | 22.0 a |
|  | N | Harm.S | Harm.S | Harm.S |  |  |
| Retained catch | 150 | 156.6 | 161.5 | 171.0 |  |  |
| Pot total males | 150 | 216.9 | 218.1 | 221.0 |  |  |
| Pot total females | 50 | 29.4 | 29.4 | 29.4 |  |  |
| Trawl bycatch | 50 | 58.8 | 57.4 | 61.5 |  |  |
| Tanner fishery bycatch | 50 | 25.4 | 25.2 | 25.6 |  |  |
| Fixed gear bycatch | 50 | 42.0 | 41.8 | 42.5 |  |  |
| NMFS survey | 200 | 175.1 | 201.6 | 210.2 |  |  |
| BSFRF survey | 200 | 118.8 | 115.3 | 124.4 |  |  |

Table 5a. Number of parameters for the model (Models 21.1b, 22.0, and 22.0a). Red values indicate different values among models.

| Parameter counts | $\mathbf{2 1 . 1 b}$ | $\mathbf{2 2 . 0}$ | $\mathbf{2 2 . 0 a}$ |
| :--- | :---: | :--- | :--- |
| Fixed growth parameters | 0 | 0 | 0 |
| Fixed recruitment parameters | 2 | 2 | 2 |
| Fixed length-weight relationship parameters | 6 | 6 | 6 |
| Fixed mortality parameters | 5 | 5 | 4 |
| Fixed survey catchability parameter | 1 | 1 | 1 |
| Fixed high grading parameters | 0 | 0 | 0 |
| Total number of fixed parameters | 14 | 14 | 13 |
|  |  |  |  |
| Free survey catchability parameter | 1 | 1 | 1 |
| Free growth parameters | 6 | 4 | 4 |
| Initial abundance (1975 or 1985) | 1 | 1 | 1 |
| Recruitment-distribution parameters | 2 | 2 | 2 |
| Mean recruitment parameters | 1 | 1 | 1 |
| Male recruitment deviations | 47 | 37 | 37 |
| Female recruitment deviations | 47 | 37 | 37 |
| Natural mortality parameters | 2 | 1 | 2 |
| Mean \& offset fishing mortality parameters | 6 | 6 | 6 |
| Pot male fishing mortality deviations | 47 | 37 | 37 |
| Bycatch mortality from Tanner crab fishery | 50 | 30 | 30 |
| Pot female bycatch fishing mortality devia. | 32 | 32 | 32 |
| Trawl bycatch fishing mortality deviations | 47 | 37 | 37 |
| Fixed gear bycatch fishing mortality devia. | 26 | 26 | 26 |
| Initial (1975 or 1985) length compositions | 35 | 35 | 35 |
| Survey extra CV | 1 | 1 | 1 |
| Free selectivity parameters | 22 | 20 | 20 |
| Total number of free parameters | 372 | 308 | 309 |
| Total number of fixed and free parameters | 386 | 322 | 323 |
|  |  |  |  |
|  |  |  |  |

Table 5b. Negative log likelihood components and their differences for Models 21.1b. 22.0, 22.0a, and some management quantities. Highlighted cells in yellow color show prior density values and total negative likelihood values without prior density. Biomass is in metric ton, and abundance is in millions of crab.

|  | Model |  |  |  |
| :--- | ---: | ---: | ---: | ---: |
|  | $\underline{21.1 \mathrm{~b}}$ | $\underline{22.0}$ | $\underline{22.0 \mathrm{a}}$ | $22.0 \mathrm{a}-22.0$ |
| Pot-ret-catch | -60.88 | -34.88 | -35.81 | -0.93 |
| Pot-totM-catch | 26.54 | 26.55 | 25.62 | -0.93 |
| Pot-F-discC | -55.69 | -55.70 | -55.28 | 0.42 |
| Trawl-discC | -63.74 | -51.28 | -51.28 | 0 |
| Tanner-M-discC | -43.54 | -26.12 | -26.12 | 0 |
| Tanner-F-discC | -43.48 | -26.08 | -26.10 | -0.02 |
| Fixed-discC | -36.04 | -36.04 | -36.04 | 0 |
| Trawl-suv-bio | -35.47 | -44.09 | -47.68 | -3.59 |
| BSFRF-sur-bio | -2.94 | -3.33 | -4.73 | -1.4 |
| Pot-ret-comp | -3932.20 | -3131.80 | -3134.52 | -2.72 |
| Pot-totM-comp | -2369.46 | -2370.52 | -2371.39 | -0.87 |
| Pot-discF-comp | -1449.36 | -1449.09 | -1450.22 | -1.13 |
| Trawl-disc-comp | -5836.10 | -4681.05 | -4685.50 | -4.45 |
| TC-disc-comp | -1274.28 | -1273.40 | -1276.25 | -2.85 |
| Fixed-disc-comp | -3393.50 | -3394.74 | -3392.59 | 2.15 |
| Trawl-sur-comp | -6984.67 | -5503.89 | -5516.02 | -12.13 |
| BSFRF-sur-comp | -843.53 | -842.35 | -844.98 | -2.63 |
| Recruit-dev | 70.56 | 41.28 | 41.83 | 0.55 |
| Recruit-sex-R | 76.98 | 60.67 | 60.63 | -0.04 |
| Log_fdev=0 | 0.00 | 0.00 | 0.00 | 0 |
| M-deviation | 43.83 | 0.00 | 0.00 |  |
| Sex-specific-R | 0.01 | 0.15 | 0.18 | 0.03 |
| Ini-size-struct. | 30.88 | 50.88 | 55.77 | 4.89 |
| PriorDensity | 267.30 | 233.94 | 221.50 | -12.44 |
| Tot-likelihood | -25908.79 | -22510.90 | -22549.41 | -38.51 |
| Tot-likeli-no-PD | -25641.5 | -22276.96 | -22317.91 | -40.95 |
| Tot-parameter | 372 | 308 | 309 | 1 |
| MMB $35 \%$ | 24026.11 | 21896.23 | 19512.93 | -2383.3 |
| MMB-terminal | 16952.82 | 17157.89 | 15713.76 | -1444.13 |
| F35\% | 0.298 | 0.299 | 0.395 | 0.096 |
| Fofl | 0.200 | 0.227 | 0.309 | 0.082 |
| OFL | 3035.63 | 3481.84 | 4319.04 | 837.2 |
| ABC | 2428.50 | 2785.47 | 3455.23 | 669.76 |
| NMFS Q | 0.967 | 0.940 | 0.922 | -0.018 |
| Mature females | 10.20 | 10.99 | 11.688 | 0.698 |
| Mohn's rho, 10yr | 0.373 | 0.453 | 0.329 |  |
|  |  |  |  |  |

Table 6a. Summary of estimated model parameter values and standard deviations for model 21.1 b for Bristol Bay red king crab.

| index | name | value | std.dev | index | name | value | std.dev |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | theta[2] | 0.281 | 0.014 | 47 | log_slx_pars[1] | 4.759 | 0.008 |
| 2 | theta[4] | 19.823 | 0.049 | 48 | log_slx_pars[2] | 2.264 | 0.046 |
| 3 | theta[5] | 16.213 | 0.138 | 49 | log_slx_pars[3] | 4.507 | 0.016 |
| 4 | theta[7] | 0.680 | 0.127 | 50 | log_slx_pars[4] | 2.020 | 0.113 |
| 5 | theta[9] | -0.472 | 0.230 | 51 | log_slx_pars[5] | 5.153 | 0.056 |
| 6 | theta[13] | 0.959 | 0.420 | 52 | log_slx_pars[6] | 2.852 | 0.045 |
| 7 | theta[14] | 0.651 | 0.469 | 53 | log_slx_pars[7] | 4.720 | 0.220 |
| 8 | theta[15] | 0.858 | 0.333 | 54 | log_slx_pars[8] | 2.164 | 0.306 |
| 9 | theta[16] | 0.707 | 0.305 | 55 | log_slx_pars[9] | 4.746 | 0.078 |
| 10 | theta[17] | 0.544 | 0.295 | 56 | log_slx_pars[10] | 0.900 | 0.304 |
| 11 | theta[18] | 0.499 | 0.277 | 57 | log_slx_pars[11] | 4.790 | 0.023 |
| 12 | theta[19] | 0.342 | 0.278 | 58 | log_slx_pars[12] | 2.340 | 0.086 |
| 13 | theta[20] | 0.376 | 0.264 | 59 | log_slx_pars[13] | 4.106 | 0.176 |
| 14 | theta[21] | 0.408 | 0.259 | 60 | log_slx_pars[14] | 2.234 | 0.378 |
| 15 | theta[22] | 0.181 | 0.282 | 61 | log_slx_pars[15] | 3.773 | 0.604 |
| 16 | theta[23] | 0.158 | 0.277 | 62 | log_slx_pars[16] | 3.290 | 0.409 |
| 17 | theta[24] | 0.053 | 0.287 | 63 | log_slx_pars[17] | 4.427 | 0.029 |
| 18 | theta[25] | 0.169 | 0.263 | 64 | log_slx_pars[18] | 2.435 | 0.072 |
| 19 | theta[26] | -0.010 | 0.204 | 65 | log_slx_pars[19] | 4.923 | 0.002 |
| 20 | theta[27] | -0.239 | 0.196 | 66 | log_slx_pars[20] | 0.673 | 0.053 |
| 21 | theta[28] | -0.390 | 0.198 | 67 | log_slx_pars[21] | 4.932 | 0.002 |
| 22 | theta[29] | -0.738 | 0.211 | 68 | log_slx_pars[22] | 0.727 | 0.099 |
| 23 | theta[30] | -1.198 | 0.233 | 69 | log_fbar[1] | -1.574 | 0.042 |
| 24 | theta[31] | -1.243 | 0.235 | 70 | log_fbar[2] | -4.306 | 0.075 |
| 25 | theta[52] | 1.294 | 0.679 | 71 | log_fbar[3] | -5.599 | 0.289 |
| 26 | theta[53] | 1.454 | 0.462 | 72 | log_fbar[4] | -6.514 | 0.072 |
| 27 | theta[54] | 1.395 | 0.367 | 73 | log_fdev[1] | 0.818 | 0.119 |
| 28 | theta[55] | 1.170 | 0.336 | 74 | log_fdev[1] | 0.776 | 0.090 |
| 29 | theta[56] | 1.083 | 0.295 | 75 | log_fdev[1] | 0.688 | 0.074 |
| 30 | theta[57] | 0.600 | 0.318 | 76 | log_fdev[1] | 0.781 | 0.060 |
| 31 | theta[58] | 0.214 | 0.353 | 77 | log_fdev[1] | 0.993 | 0.054 |
| 32 | theta[59] | -0.025 | 0.362 | 78 | log_fdev[1] | 1.861 | 0.056 |
| 33 | theta[60] | -0.214 | 0.355 | 79 | log_fdev[1] | 2.394 | 0.119 |
| 34 | theta[61] | -0.545 | 0.374 | 80 | log_fdev[1] | 0.816 | 0.177 |
| 35 | theta[62] | -0.932 | 0.386 | 81 | log_fdev[1] | -8.897 | 0.126 |
| 36 | theta[63] | -1.190 | 0.390 | 82 | $\log$ _fdev[1] | 1.143 | 0.112 |
| 37 | theta[64] | -1.421 | 0.389 | 83 | log_fdev[1] | 1.217 | 0.089 |
| 38 | theta[65] | -1.791 | 0.377 | 84 | log_fdev[1] | 1.386 | 0.073 |
| 39 | theta[66] | -1.897 | 0.373 | 85 | log_fdev[1] | 0.920 | 0.064 |
| 40 | theta[67] | -1.838 | 0.352 | 86 | log_fdev[1] | -0.018 | 0.053 |
| 41 | Grwth[21] | 0.972 | 0.185 | 87 | log_fdev[1] | 0.097 | 0.047 |
| 42 | Grwth[42] | 1.454 | 0.122 | 88 | log_fdev[1] | 0.745 | 0.039 |
| 43 | Grwth[85] | 142.500 | 1.727 | 89 | $\log$ _fdev[1] | 0.758 | 0.041 |
| 44 | Grwth[86] | 0.058 | 0.010 | 90 | $\mathrm{log}_{\text {_fdev [1] }}$ | 0.242 | 0.046 |
| 45 | Grwth[87] | 139.890 | 0.596 | 91 | log_fdev[1] | 0.910 | 0.051 |


| 46 | Grwth[88] | 0.071 | 0.003 | 92 | log_fdev[1] | -4.242 | 0.049 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 93 | log_fdev[1] | -4.651 | 0.042 | 143 | log_fdev[2] | -0.222 | 0.104 |
| 94 | log_fdev[1] | -0.181 | 0.041 | 144 | log_fdev[2] | -0.981 | 0.103 |
| 95 | log_fdev[1] | -0.132 | 0.041 | 145 | log_fdev[2] | -0.211 | 0.103 |
| 96 | log_fdev[1] | 0.783 | 0.044 | 146 | log_fdev[2] | -0.511 | 0.103 |
| 97 | log_fdev[1] | 0.426 | 0.043 | 147 | log_fdev[2] | -0.604 | 0.102 |
| 98 | log_fdev[1] | -0.160 | 0.041 | 148 | log_fdev[2] | -0.371 | 0.102 |
| 99 | log_fdev[1] | -0.239 | 0.041 | 149 | log_fdev[2] | -0.646 | 0.102 |
| 100 | log_fdev[1] | -0.128 | 0.040 | 150 | log_fdev[2] | -0.477 | 0.102 |
| 101 | log_fdev[1] | 0.337 | 0.038 | 151 | log_fdev[2] | -0.399 | 0.102 |
| 102 | log_fdev[1] | 0.294 | 0.038 | 152 | log_fdev[2] | -0.426 | 0.102 |
| 103 | log_fdev[1] | 0.585 | 0.039 | 153 | log_fdev[2] | -0.784 | 0.102 |
| 104 | log_fdev[1] | 0.339 | 0.038 | 154 | log_fdev[2] | -0.932 | 0.102 |
| 105 | log_fdev[1] | 0.704 | 0.038 | 155 | log_fdev[2] | -1.394 | 0.102 |
| 106 | log_fdev[1] | 0.874 | 0.040 | 156 | log_fdev[2] | -1.914 | 0.102 |
| 107 | log_fdev[1] | 0.690 | 0.041 | 157 | log_fdev[2] | -1.199 | 0.102 |
| 108 | log_fdev[1] | 0.560 | 0.040 | 158 | log_fdev[2] | -1.762 | 0.103 |
| 109 | log_fdev[1] | -0.075 | 0.039 | 159 | log_fdev[2] | -1.378 | 0.103 |
| 110 | log_fdev[1] | -0.151 | 0.038 | 160 | log_fdev[2] | -0.852 | 0.105 |
| 111 | log_fdev[1] | 0.036 | 0.038 | 161 | log_fdev[2] | -0.418 | 0.107 |
| 112 | log_fdev[1] | 0.366 | 0.038 | 162 | log_fdev[2] | -0.483 | 0.110 |
| 113 | log_fdev[1] | 0.438 | 0.041 | 163 | log_fdev[2] | -0.389 | 0.113 |
| 114 | log_fdev[1] | 0.437 | 0.046 | 164 | log_fdev[2] | -0.421 | 0.115 |
| 115 | log_fdev[1] | 0.347 | 0.054 | 165 | log_fdev[2] | -1.414 | 0.115 |
| 116 | log_fdev[1] | 0.156 | 0.064 | 166 | log_fdev[3] | -0.116 | 0.068 |
| 117 | log_fdev[1] | 0.095 | 0.072 | 167 | log_fdev[3] | 0.670 | 0.068 |
| 118 | log_fdev[1] | -0.342 | 0.076 | 168 | log_fdev[3] | 1.228 | 0.068 |
| 119 | log_fdev[1] | -4.797 | 0.075 | 169 | log_fdev[3] | 1.093 | 0.068 |
| 120 | log_fdev[2] | 0.195 | 0.125 | 170 | log_fdev[3] | 1.383 | 0.068 |
| 121 | log_fdev[2] | 0.634 | 0.116 | 171 | log_fdev[3] | 1.424 | 0.068 |
| 122 | log_fdev[2] | 0.614 | 0.111 | 172 | log_fdev[3] | 0.993 | 0.068 |
| 123 | log_fdev[2] | 0.690 | 0.109 | 173 | log_fdev[3] | 0.476 | 0.068 |
| 124 | log_fdev[2] | 1.406 | 0.112 | 174 | log_fdev[3] | -0.987 | 0.068 |
| 125 | log_fdev[2] | 1.176 | 0.131 | 175 | log_fdev[3] | -0.579 | 0.068 |
| 126 | log_fdev[2] | 2.458 | 0.132 | 176 | log_fdev[3] | -1.099 | 0.068 |
| 127 | log_fdev[2] | 2.178 | 0.119 | 177 | log_fdev[3] | -0.256 | 0.068 |
| 128 | log_fdev[2] | 3.398 | 0.116 | 178 | log_fdev[3] | 0.940 | 0.068 |
| 129 | log_fdev[2] | 2.192 | 0.111 | 179 | log_fdev[3] | 1.418 | 0.068 |
| 130 | log_fdev[2] | 1.129 | 0.111 | 180 | log_fdev[3] | 3.239 | 0.075 |
| 131 | log_fdev[2] | 0.676 | 0.109 | 181 | log_fdev[3] | 1.284 | 0.095 |
| 132 | log_fdev[2] | 1.452 | 0.104 | 182 | log_fdev[3] | 0.581 | 0.119 |
| 133 | log_fdev[2] | 0.022 | 0.104 | 183 | log_fdev[3] | -0.757 | 0.082 |
| 134 | log_fdev[2] | 0.475 | 0.104 | 184 | log_fdev[3] | -2.137 | 0.073 |
| 135 | log_fdev[2] | 0.899 | 0.105 | 185 | log_fdev[3] | -2.991 | 0.093 |
| 136 | log_fdev[2] | 0.735 | 0.105 | 186 | log_fdev[3] | -2.412 | 0.112 |
| 137 | log_fdev[2] | 1.212 | 0.108 | 187 | log_fdev[3] | -3.494 | 0.075 |
| 138 | log_fdev[2] | -0.556 | 0.105 | 188 | log_fdev[3] | -0.845 | 0.094 |
| 139 | log_fdev[2] | -0.842 | 0.103 | 189 | log_fdev[3] | -0.119 | 0.111 |
| 140 | log_fdev[2] | -0.774 | 0.104 | 190 | log_fdev[3] | 1.064 | 0.133 |


| 141 | log_fdev[2] | -1.240 | 0.103 | 191 | log_fdev[4] | 0.562 | 0.103 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 142 | log_fdev[2] | 0.058 | 0.104 | 192 | $\mathrm{log}_{-} \mathrm{fdev}[4]$ | -0.100 | 0.102 |
| 193 | log_fdev[4] | -0.314 | 0.102 | 243 | $\mathrm{log}_{-} \mathrm{fdov}[1]$ | -0.226 | 0.078 |
| 194 | log_fdev[4] | 0.606 | 0.102 | 244 | $\mathrm{log}_{-} \mathrm{fdov}[1]$ | 0.834 | 0.079 |
| 195 | log_fdev[4] | -1.822 | 0.101 | 245 | $\mathrm{log}_{\text {f }} \mathrm{fdov}[1]$ | 0.287 | 0.080 |
| 196 | log_fdev[4] | 0.133 | 0.101 | 246 | $\mathrm{log}_{\text {_f }} \mathrm{fdov}[1]$ | -0.364 | 0.083 |
| 197 | log_fdev[4] | -0.124 | 0.100 | 247 | $\mathrm{log}_{\mathrm{f}} \mathrm{fdov}[1]$ | 0.964 | 0.087 |
| 198 | log_fdev[4] | -0.957 | 0.100 | 248 | log_fdov[1] | -0.106 | 0.091 |
| 199 | log_fdev[4] | -0.783 | 0.100 | 249 | $\log _{-} \mathrm{fdov}[1]$ | -0.626 | 0.093 |
| 200 | log_fdev[4] | -0.509 | 0.100 | 250 | $\mathrm{log}_{\text {_f dov [1] }}$ | 2.972 | 0.094 |
| 201 | log_fdev[4] | -0.555 | 0.100 | 251 | log_fdov[3] | 0.000 | 0.096 |
| 202 | log_fdev[4] | -0.008 | 0.100 | 252 | log_fdov[3] | 0.000 | 0.096 |
| 203 | log_fdev[4] | -0.707 | 0.100 | 253 | log_fdov[3] | 0.000 | 0.096 |
| 204 | log_fdev[4] | -1.703 | 0.100 | 254 | log_fdov[3] | 0.000 | 0.096 |
| 205 | log_fdev[4] | -2.538 | 0.099 | 255 | log_fdov[3] | 0.000 | 0.096 |
| 206 | log_fdev[4] | -1.057 | 0.099 | 256 | $\mathrm{log}_{\text {_f }} \mathrm{fdov}[3]$ | 0.000 | 0.096 |
| 207 | log_fdev[4] | -0.501 | 0.099 | 257 | log_fdov[3] | 0.000 | 0.096 |
| 208 | log_fdev[4] | 0.639 | 0.099 | 258 | log_fdov[3] | 0.000 | 0.096 |
| 209 | log_fdev[4] | 1.491 | 0.099 | 259 | log_fdov[3] | 0.000 | 0.096 |
| 210 | log_fdev[4] | 1.175 | 0.100 | 260 | log_fdov[3] | 0.000 | 0.096 |
| 211 | log_fdev[4] | 0.345 | 0.101 | 261 | $\mathrm{log}_{-} \mathrm{fdov}[3]$ | 0.000 | 0.096 |
| 212 | log_fdev[4] | 1.949 | 0.102 | 262 | log_fdov[3] | 0.000 | 0.096 |
| 213 | log_fdev[4] | 2.207 | 0.103 | 263 | $\mathrm{log}_{\text {_f }} \mathrm{fdov}[3]$ | 0.000 | 0.096 |
| 214 | log_fdev[4] | 1.005 | 0.105 | 264 | log_fdov[3] | 0.001 | 0.096 |
| 215 | log_fdev[4] | 0.795 | 0.107 | 265 | log_fdov[3] | 1.546 | 0.169 |
| 216 | log_fdev[4] | 0.771 | 0.109 | 266 | log_fdov[3] | 1.804 | 0.120 |
| 217 | log_foff[1] | -2.781 | 0.039 | 267 | log_fdov[3] | 0.573 | 0.141 |
| 218 | log_foff[3] | -0.095 | 0.414 | 268 | log_fdov[3] | -3.440 | 0.108 |
| 219 | log_fdov[1] | 1.969 | 0.083 | 269 | log_fdov[3] | -2.134 | 0.145 |
| 220 | log_fdov[1] | -0.701 | 0.083 | 270 | $\mathrm{log}_{\text {_fdov }}$ [3] | -0.775 | 0.126 |
| 221 | log_fdov[1] | 1.973 | 0.084 | 271 | log_fdov[3] | 0.043 | 0.132 |
| 222 | log_fdov[1] | 1.810 | 0.086 | 272 | $\mathrm{log}_{-} \mathrm{fdov}[3]$ | 0.387 | 0.102 |
| 223 | log_fdov[1] | -0.421 | 0.084 | 273 | log_fdov[3] | 0.943 | 0.167 |
| 224 | log_fdov[1] | -0.191 | 0.082 | 274 | log_fdov[3] | 0.162 | 0.152 |
| 225 | log_fdov[1] | -3.696 | 0.081 | 275 | log_fdov[3] | 0.889 | 0.167 |
| 226 | log_fdov[1] | -0.328 | 0.082 | 276 | rec_dev_est | 1.066 | 0.269 |
| 227 | log_fdov[1] | 1.457 | 0.082 | 277 | rec_dev_est | 0.627 | 0.295 |
| 228 | log_fdov[1] | -2.773 | 0.081 | 278 | rec_dev_est | 1.073 | 0.240 |
| 229 | log_fdov[1] | 1.155 | 0.081 | 279 | rec_dev_est | 1.658 | 0.207 |
| 230 | log_fdov[1] | 0.881 | 0.080 | 280 | rec_dev_est | 1.921 | 0.216 |
| 231 | log_fdov[1] | -1.866 | 0.080 | 281 | rec_dev_est | 1.127 | 0.257 |
| 232 | log_fdov[1] | 1.218 | 0.080 | 282 | rec_dev_est | 2.396 | 0.165 |
| 233 | log_fdov[1] | 0.425 | 0.080 | 283 | rec_dev_est | 1.442 | 0.179 |
| 234 | log_fdov[1] | 0.958 | 0.079 | 284 | rec_dev_est | 1.072 | 0.166 |
| 235 | log_fdov[1] | -1.226 | 0.079 | 285 | rec_dev_est | -0.768 | 0.250 |
| 236 | log_fdov[1] | -0.187 | 0.079 | 286 | rec_dev_est | 0.322 | 0.163 |
| 237 | log_fdov[1] | -0.450 | 0.079 | 287 | rec_dev_est | -0.843 | 0.245 |
| 238 | log_fdov[1] | -0.714 | 0.079 | 288 | rec_dev_est | -1.269 | 0.277 |
| 239 | log_fdov[1] | -0.233 | 0.079 | 289 | rec_dev_est | -1.008 | 0.223 |


| 240 | log_fdov[1] | -1.129 | 0.078 | 290 | rec_dev_est | -0.054 | 0.164 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 241 | log_fdov[1] | -1.845 | 0.078 | 291 | rec_dev_est | -0.518 | 0.184 |
| 242 | log_fdov[1] | 0.178 | 0.078 | 292 | rec_dev_est | -1.985 | 0.357 |
| 293 | rec_dev_est | -0.886 | 0.197 | 339 | logit_rec_prop_est | 0.751 | 0.737 |
| 294 | rec_dev_est | -2.023 | 0.424 | 340 | logit_rec_prop_est | 0.231 | 0.284 |
| 295 | rec_dev_est | 0.994 | 0.147 | 341 | logit_rec_prop_est | -0.222 | 0.692 |
| 296 | rec_dev_est | -0.936 | 0.260 | 342 | logit_rec_prop_est | -0.298 | 0.087 |
| 297 | rec_dev_est | -1.599 | 0.339 | 343 | logit_rec_prop_est | 1.317 | 0.660 |
| 298 | rec_dev_est | -0.573 | 0.198 | 344 | logit_rec_prop_est | 0.431 | 0.646 |
| 299 | rec_dev_est | 0.423 | 0.155 | 345 | logit_rec_prop_est | 0.498 | 0.324 |
| 300 | rec_dev_est | -0.568 | 0.225 | 346 | logit_rec_prop_est | -0.049 | 0.141 |
| 301 | rec_dev_est | -0.534 | 0.239 | 347 | logit_rec_prop_est | 0.193 | 0.363 |
| 302 | rec_dev_est | 0.850 | 0.154 | 348 | logit_rec_prop_est | -0.531 | 0.373 |
| 303 | rec_dev_est | -0.627 | 0.265 | 349 | logit_rec_prop_est | -0.489 | 0.124 |
| 304 | rec_dev_est | -0.698 | 0.265 | 350 | logit_rec_prop_est | -0.419 | 0.426 |
| 305 | rec_dev_est | 0.586 | 0.156 | 351 | logit_rec_prop_est | 0.009 | 0.444 |
| 306 | rec_dev_est | -0.150 | 0.182 | 352 | logit_rec_prop_est | -0.393 | 0.139 |
| 307 | rec_dev_est | -0.542 | 0.189 | 353 | logit_rec_prop_est | -0.091 | 0.237 |
| 308 | rec_dev_est | -1.124 | 0.236 | 354 | logit_rec_prop_est | 0.360 | 0.280 |
| 309 | rec_dev_est | -0.992 | 0.236 | 355 | logit_rec_prop_est | -0.185 | 0.370 |
| 310 | rec_dev_est | -0.010 | 0.178 | 356 | logit_rec_prop_est | -0.429 | 0.359 |
| 311 | rec_dev_est | -0.557 | 0.227 | 357 | logit_rec_prop_est | -0.786 | 0.194 |
| 312 | rec_dev_est | -1.103 | 0.233 | 358 | logit_rec_prop_est | -0.460 | 0.317 |
| 313 | rec_dev_est | -1.421 | 0.222 | 359 | logit_rec_prop_est | -0.522 | 0.346 |
| 314 | rec_dev_est | -1.890 | 0.268 | 360 | logit_rec_prop_est | -0.228 | 0.332 |
| 315 | rec_dev_est | -1.428 | 0.225 | 361 | logit_rec_prop_est | -0.287 | 0.428 |
| 316 | rec_dev_est | -0.756 | 0.174 | 362 | logit_rec_prop_est | -0.314 | 0.327 |
| 317 | rec_dev_est | -1.619 | 0.257 | 363 | logit_rec_prop_est | 0.331 | 0.218 |
| 318 | rec_dev_est | -0.816 | 0.196 | 364 | logit_rec_prop_est | 0.537 | 0.474 |
| 319 | rec_dev_est | -1.425 | 0.277 | 365 | logit_rec_prop_est | 0.800 | 0.308 |
| 320 | rec_dev_est | -1.335 | 0.279 | 366 | logit_rec_prop_est | 0.187 | 0.467 |
| 321 | rec_dev_est | -1.203 | 0.273 | 367 | logit_rec_prop_est | 0.662 | 0.523 |
| 322 | rec_dev_est | -0.893 | 0.316 | 368 | logit_rec_prop_est | 0.194 | 0.444 |
| 323 | logit_rec_prop_est | -0.093 | 0.433 | 369 | logit_rec_prop_est | -0.755 | 0.513 |
| 324 | logit_rec_prop_est | -0.844 | 0.513 | 370 | m_dev_est[1] | 1.594 | 0.029 |
| 325 | logit_rec_prop_est | -0.235 | 0.357 | 371 | survey_q[1] | 0.967 | 0.025 |
| 326 | logit_rec_prop_est | -0.439 | 0.266 | 372 | log_add_cv[2] | -0.775 | 0.273 |
| 327 | logit_rec_prop_est | 0.078 | 0.255 |  |  |  |  |
| 328 | logit_rec_prop_est | 0.273 | 0.336 |  |  |  |  |
| 329 | logit_rec_prop_est | 0.343 | 0.140 |  |  |  |  |
| 330 | logit_rec_prop_est | 0.395 | 0.231 |  |  |  |  |
| 331 | logit_rec_prop_est | -0.095 | 0.175 |  |  |  |  |
| 332 | logit_rec_prop_est | 0.462 | 0.463 |  |  |  |  |
| 333 | logit_rec_prop_est | -0.497 | 0.165 |  |  |  |  |
| 334 | logit_rec_prop_est | 0.206 | 0.419 |  |  |  |  |
| 335 | logit_rec_prop_est | -0.106 | 0.458 |  |  |  |  |
| 336 | logit_rec_prop_est | 0.433 | 0.389 |  |  |  |  |
| 337 | logit_rec_prop_est | -0.091 | 0.167 |  |  |  |  |
| 338 | logit_rec_prop_est | 0.167 | 0.242 |  |  |  |  |

Table 6 b. Summary of estimated model parameter values and standard deviations for model 22.0 for Bristol Bay red king crab.

| index | name | value | std.dev | index | name | value | std.dev |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | theta[2] | 0.253 | 0.016 | 47 | log_slx_pars[3] | 4.501 | 0.016 |
| 2 | theta[4] | 17.851 | 0.041 | 48 | log_slx_pars[4] | 1.993 | 0.116 |
| 3 | theta [5] | 15.817 | 0.156 | 49 | log_slx_pars[5] | 5.208 | 0.093 |
| 4 | theta[7] | 0.648 | 0.123 | 50 | log_slx_pars[6] | 2.932 | 0.054 |
| 5 | theta[9] | -0.456 | 0.250 | 51 | log_slx_pars[7] | 4.731 | 0.223 |
| 6 | theta[13] | 0.759 | 0.497 | 52 | log_slx_pars[8] | 2.165 | 0.306 |
| 7 | theta[14] | 0.778 | 0.475 | 53 | log_slx_pars[9] | 4.718 | 0.091 |
| 8 | theta[15] | 1.131 | 0.350 | 54 | log_slx_pars[10] | 0.903 | 0.302 |
| 9 | theta[16] | 1.320 | 0.285 | 55 | log_slx_pars[11] | 4.787 | 0.023 |
| 10 | theta[17] | 1.255 | 0.265 | 56 | log_slx_pars[12] | 2.339 | 0.088 |
| 11 | theta[18] | 0.989 | 0.272 | 57 | log_slx_pars[13] | 3.946 | 0.380 |
| 12 | theta[19] | 0.945 | 0.259 | 58 | log_slx_pars[14] | 2.956 | 0.384 |
| 13 | theta[20] | 1.202 | 0.222 | 59 | log_slx_pars[15] | 4.435 | 0.033 |
| 14 | theta[21] | 1.194 | 0.216 | 60 | log_slx_pars[16] | 2.425 | 0.090 |
| 15 | theta[22] | 1.011 | 0.223 | 61 | log_slx_pars[17] | 4.924 | 0.002 |
| 16 | theta[23] | 0.958 | 0.215 | 62 | log_slx_pars[18] | 0.672 | 0.070 |
| 17 | theta[24] | 0.815 | 0.218 | 63 | log_slx_pars[19] | 4.932 | 0.002 |
| 18 | theta[25] | 0.486 | 0.223 | 64 | log_slx_pars[20] | 0.734 | 0.099 |
| 19 | theta[26] | 0.044 | 0.194 | 65 | $\log _{-} \mathrm{fbar}[1]$ | -1.645 | 0.048 |
| 20 | theta[27] | -0.427 | 0.197 | 66 | log_fbar[2] | -4.694 | 0.082 |
| 21 | theta[28] | -1.082 | 0.220 | 67 | $\log _{-} \mathrm{fbar}[3]$ | -5.973 | 0.307 |
| 22 | theta[29] | -1.662 | 0.253 | 68 | log_fbar[4] | -6.548 | 0.073 |
| 23 | theta[30] | -2.340 | 0.276 | 69 | log_fdev[1] | 1.030 | 0.119 |
| 24 | theta[31] | -2.004 | 0.361 | 70 | log_fdev[1] | 1.250 | 0.079 |
| 25 | theta[52] | -0.099 | 0.599 | 71 | log_fdev[1] | 0.857 | 0.063 |
| 26 | theta[53] | 0.396 | 0.656 | 72 | log_fdev[1] | 0.005 | 0.052 |
| 27 | theta[54] | 0.869 | 0.546 | 73 | log_fdev[1] | 0.153 | 0.046 |
| 28 | theta[55] | 1.077 | 0.428 | 74 | log_fdev[1] | 0.809 | 0.037 |
| 29 | theta[56] | 1.224 | 0.336 | 75 | log_fdev[1] | 0.821 | 0.039 |
| 30 | theta[57] | 1.049 | 0.314 | 76 | log_fdev[1] | 0.305 | 0.043 |
| 31 | theta[58] | 0.830 | 0.311 | 77 | log_fdev[1] | 0.969 | 0.047 |
| 32 | theta[59] | 0.368 | 0.348 | 78 | log_fdev[1] | -4.182 | 0.045 |
| 33 | theta[60] | -0.372 | 0.393 | 79 | log_fdev[1] | -4.591 | 0.039 |
| 34 | theta[61] | -0.822 | 0.386 | 80 | log_fdev[1] | -0.120 | 0.038 |
| 35 | theta[62] | -1.523 | 0.376 | 81 | log_fdev[1] | -0.074 | 0.038 |
| 36 | theta[63] | -1.616 | 0.373 | 82 | log_fdev[1] | 0.842 | 0.040 |
| 37 | theta[64] | -1.546 | 0.373 | 83 | log_fdev[1] | 0.482 | 0.039 |
| 38 | theta[65] | -1.769 | 0.363 | 84 | log_fdev[1] | -0.106 | 0.038 |
| 39 | theta[66] | -1.907 | 0.353 | 85 | log_fdev[1] | -0.185 | 0.038 |
|  | theta[67] | -1.874 | 0.344 | 86 | log_fdev[1] | -0.071 | 0.037 |
| 41 | Grwth[21] | 0.897 | 0.192 | 87 | log_fdev[1] | 0.394 | 0.036 |


| 42 | Grwth[42] | 1.488 | 0.134 | 88 | log_fdev[1] | 0.351 | 0.037 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 43 | Grwth[64] | 139.700 | 0.612 | 89 | log_fdev[1] | 0.641 | 0.037 |
| 44 | Grwth[65] | 0.071 | 0.003 | 90 | log_fdev[1] | 0.395 | 0.036 |
| 45 | log_slx_pars[1] | 4.759 | 0.008 | 91 | log_fdev[1] | 0.759 | 0.036 |
| 46 | log_slx_pars[2] | 2.266 | 0.046 | 92 | log_fdev[1] | 0.928 | 0.037 |
| 93 | log_fdev[1] | 0.743 | 0.037 | 143 | log_fdev[3] | -0.727 | 0.066 |
| 94 | log_fdev[1] | 0.612 | 0.037 | 144 | log_fdev[3] | 0.116 | 0.066 |
| 95 | log_fdev[1] | -0.024 | 0.036 | 145 | log_fdev[3] | 1.312 | 0.066 |
| 96 | log_fdev[1] | -0.100 | 0.035 | 146 | log_fdev[3] | 1.790 | 0.066 |
| 97 | log_fdev[1] | 0.088 | 0.035 | 147 | log_fdev[3] | 3.616 | 0.076 |
| 98 | log_fdev[1] | 0.416 | 0.035 | 148 | log_fdev[3] | 1.665 | 0.095 |
| 99 | log_fdev[1] | 0.486 | 0.038 | 149 | log_fdev[3] | 0.960 | 0.127 |
| 100 | log_fdev[1] | 0.482 | 0.043 | 150 | log_fdev[3] | -0.383 | 0.080 |
| 101 | log_fdev[1] | 0.387 | 0.051 | 151 | log_fdev[3] | -1.764 | 0.074 |
| 102 | log_fdev[1] | 0.192 | 0.061 | 152 | log_fdev[3] | -2.614 | 0.091 |
| 103 | log_fdev[1] | 0.129 | 0.069 | 153 | log_fdev[3] | -2.036 | 0.116 |
| 104 | log_fdev[1] | -0.309 | 0.073 | 154 | log_fdev[3] | -3.123 | 0.077 |
| 105 | $\log _{\text {_f }}$ dev[1] | -4.762 | 0.073 | 155 | log_fdev[3] | -0.479 | 0.096 |
| 106 | log_fdev[2] | 2.366 | 0.114 | 156 | log_fdev[3] | 0.243 | 0.114 |
| 107 | log_fdev[2] | 1.339 | 0.112 | 157 | log_fdev[3] | 1.422 | 0.137 |
| 108 | log_fdev[2] | 0.944 | 0.109 | 158 | log_fdev[4] | 0.578 | 0.103 |
| 109 | log_fdev[2] | 1.778 | 0.104 | 159 | log_fdev[4] | -0.086 | 0.102 |
| 110 | log_fdev[2] | 0.381 | 0.103 | 160 | log_fdev[4] | -0.305 | 0.103 |
| 111 | log_fdev[2] | 0.847 | 0.103 | 161 | log_fdev[4] | 0.614 | 0.102 |
| 112 | log_fdev[2] | 1.271 | 0.104 | 162 | log_fdev[4] | -1.814 | 0.101 |
| 113 | log_fdev[2] | 1.108 | 0.104 | 163 | log_fdev[4] | 0.142 | 0.101 |
| 114 | log_fdev[2] | 1.579 | 0.106 | 164 | log_fdev[4] | -0.115 | 0.100 |
| 115 | log_fdev[2] | -0.183 | 0.104 | 165 | log_fdev[4] | -0.949 | 0.100 |
| 116 | log_fdev[2] | -0.468 | 0.102 | 166 | log_fdev[4] | -0.775 | 0.100 |
| 117 | log_fdev[2] | -0.403 | 0.103 | 167 | log_fdev[4] | -0.501 | 0.100 |
| 118 | log_fdev[2] | -0.874 | 0.102 | 168 | log_fdev[4] | -0.547 | 0.100 |
| 119 | log_fdev[2] | 0.417 | 0.103 | 169 | log_fdev[4] | -0.001 | 0.100 |
| 120 | log_fdev[2] | 0.138 | 0.103 | 170 | log_fdev[4] | -0.702 | 0.100 |
| 121 | log_fdev[2] | -0.620 | 0.102 | 171 | log_fdev[4] | -1.700 | 0.100 |
| 122 | log_fdev[2] | 0.150 | 0.102 | 172 | log_fdev[4] | -2.536 | 0.099 |
| 123 | log_fdev[2] | -0.149 | 0.102 | 173 | log_fdev[4] | -1.055 | 0.099 |
| 124 | log_fdev[2] | -0.243 | 0.102 | 174 | log_fdev[4] | -0.500 | 0.099 |
| 125 | log_fdev[2] | -0.011 | 0.102 | 175 | log_fdev[4] | 0.639 | 0.099 |
| 126 | log_fdev[2] | -0.289 | 0.101 | 176 | log_fdev[4] | 1.488 | 0.099 |
| 127 | log_fdev[2] | -0.118 | 0.101 | 177 | log_fdev[4] | 1.168 | 0.100 |
| 128 | log_fdev[2] | -0.043 | 0.101 | 178 | log_fdev[4] | 0.334 | 0.101 |
| 129 | log_fdev[2] | -0.074 | 0.101 | 179 | log_fdev[4] | 1.932 | 0.102 |
| 130 | log_fdev[2] | -0.431 | 0.101 | 180 | log_fdev[4] | 2.187 | 0.104 |
| 131 | log_fdev[2] | -0.577 | 0.101 | 181 | log_fdev[4] | 0.981 | 0.105 |
| 132 | log_fdev[2] | -1.037 | 0.101 | 182 | log_fdev[4] | 0.771 | 0.107 |
| 133 | log_fdev[2] | -1.557 | 0.101 | 183 | log_fdev[4] | 0.751 | 0.109 |
| 134 | log_fdev[2] | -0.843 | 0.101 | 184 | log_foff[1] | -2.787 | 0.039 |
| 135 | log_fdev[2] | -1.410 | 0.102 | 185 | log_foff[3] | -0.208 | 0.425 |
| 136 | log_fdev[2] | -1.029 | 0.102 | 186 | log_fdov[1] | 2.000 | 0.084 |


| 137 | log_fdev[2] | -0.507 | 0.104 | 187 | log_fdov[1] | -0.674 | 0.083 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 138 | log_fdev[2] | -0.078 | 0.106 | 188 | log_fdov[1] | 1.995 | 0.084 |
| 139 | log_fdev[2] | -0.148 | 0.108 | 189 | log_fdov[1] | 1.830 | 0.086 |
| 140 | log_fdev[2] | -0.059 | 0.111 | 190 | log_fdov[1] | -0.408 | 0.084 |
| 141 | log_fdev[2] | -0.089 | 0.113 | 191 | log_fdov[1] | -0.183 | 0.082 |
| 142 | log_fdev[2] | -1.077 | 0.114 | 192 | log_fdov[1] | -3.692 | 0.081 |
| 193 | log_fdov[1] | -0.323 | 0.082 | 243 | rec_dev_est | -0.556 | 0.265 |
| 194 | log_fdov[1] | 1.462 | 0.082 | 244 | rec_dev_est | -1.183 | 0.334 |
| 195 | log_fdov[1] | -2.767 | 0.081 | 245 | rec_dev_est | -0.172 | 0.208 |
| 196 | log_fdov[1] | 1.161 | 0.080 | 246 | rec_dev_est | 0.807 | 0.171 |
| 197 | log_fdov[1] | 0.884 | 0.080 | 247 | rec_dev_est | -0.179 | 0.234 |
| 198 | log_fdov[1] | -1.866 | 0.080 | 248 | rec_dev_est | -0.124 | 0.246 |
| 199 | log_fdov[1] | 1.219 | 0.080 | 249 | rec_dev_est | 1.218 | 0.171 |
| 200 | log_fdov[1] | 0.428 | 0.080 | 250 | rec_dev_est | -0.225 | 0.271 |
| 201 | log_fdov[1] | 0.963 | 0.079 | 251 | rec_dev_est | -0.300 | 0.268 |
| 202 | log_fdov[1] | -1.223 | 0.079 | 252 | rec_dev_est | 0.967 | 0.172 |
| 203 | log_fdov[1] | -0.183 | 0.079 | 253 | rec_dev_est | 0.231 | 0.195 |
| 204 | log_fdov[1] | -0.447 | 0.079 | 254 | rec_dev_est | -0.131 | 0.201 |
| 205 | log_fdov[1] | -0.713 | 0.079 | 255 | rec_dev_est | -0.739 | 0.249 |
| 206 | log_fdov[1] | -0.233 | 0.079 | 256 | rec_dev_est | -0.574 | 0.244 |
| 207 | log_fdov[1] | -1.132 | 0.078 | 257 | rec_dev_est | 0.364 | 0.194 |
| 208 | log_fdov[1] | -1.852 | 0.078 | 258 | rec_dev_est | -0.131 | 0.236 |
| 209 | log_fdov[1] | 0.168 | 0.078 | 259 | rec_dev_est | -0.730 | 0.248 |
| 210 | log_fdov[1] | -0.238 | 0.078 | 260 | rec_dev_est | -1.007 | 0.233 |
| 211 | log_fdov[1] | 0.821 | 0.079 | 261 | rec_dev_est | -1.458 | 0.272 |
| 212 | log_fdov[1] | 0.272 | 0.080 | 262 | rec_dev_est | -1.020 | 0.231 |
| 213 | log_fdov[1] | -0.380 | 0.083 | 263 | rec_dev_est | -0.328 | 0.187 |
| 214 | log_fdov[1] | 0.948 | 0.088 | 264 | rec_dev_est | -1.182 | 0.262 |
| 215 | log_fdov[1] | -0.126 | 0.091 | 265 | rec_dev_est | -0.398 | 0.209 |
| 216 | log_fdov[1] | -0.650 | 0.093 | 266 | rec_dev_est | -0.986 | 0.282 |
| 217 | log_fdov[1] | 2.939 | 0.093 | 267 | rec_dev_est | -0.917 | 0.285 |
| 218 | log_fdov[3] | 0.000 | 0.093 | 268 | rec_dev_est | -0.778 | 0.286 |
| 219 | log_fdov[3] | 0.000 | 0.093 | 269 | rec_dev_est | -0.442 | 0.330 |
| 220 | log_fdov[3] | 0.000 | 0.093 | 270 | logit_rec_prop_est | -0.452 | 0.150 |
| 221 | log_fdov[3] | 0.001 | 0.093 | 271 | logit_rec_prop_est | 0.248 | 0.420 |
| 222 | log_fdov[3] | 1.546 | 0.142 | 272 | logit_rec_prop_est | -0.070 | 0.459 |
| 223 | log_fdov[3] | 1.829 | 0.119 | 273 | logit_rec_prop_est | 0.463 | 0.371 |
| 224 | log_fdov[3] | 0.598 | 0.145 | 274 | logit_rec_prop_est | -0.061 | 0.163 |
| 225 | log_fdov[3] | -3.425 | 0.108 | 275 | logit_rec_prop_est | 0.234 | 0.244 |
| 226 | log_fdov[3] | -2.181 | 0.144 | 276 | logit_rec_prop_est | 0.580 | 0.665 |
| 227 | log_fdov[3] | -0.801 | 0.117 | 277 | logit_rec_prop_est | 0.313 | 0.287 |
| 228 | log_fdov[3] | 0.026 | 0.136 | 278 | logit_rec_prop_est | -0.463 | 0.653 |
| 229 | log_fdov[3] | 0.376 | 0.104 | 279 | logit_rec_prop_est | -0.239 | 0.088 |
| 230 | log_fdov[3] | 0.961 | 0.151 | 280 | logit_rec_prop_est | 1.304 | 0.604 |
| 231 | log_fdov[3] | 0.168 | 0.145 | 281 | logit_rec_prop_est | 0.416 | 0.602 |
| 232 | log_fdov[3] | 0.901 | 0.174 | 282 | logit_rec_prop_est | 0.542 | 0.320 |
| 233 | rec_dev_est | 0.742 | 0.175 | 283 | logit_rec_prop_est | -0.018 | 0.142 |
| 234 | rec_dev_est | -0.478 | 0.253 | 284 | logit_rec_prop_est | 0.219 | 0.364 |
| 235 | rec_dev_est | -0.901 | 0.285 | 285 | logit_rec_prop_est | -0.503 | 0.369 |


| 236 | rec_dev_est | -0.583 | 0.227 | 286 | logit_rec_prop_est | -0.444 | 0.128 |
| :--- | :--- | ---: | :--- | :--- | :--- | ---: | ---: |
| 237 | rec_dev_est | 0.327 | 0.178 | 287 | logit_rec_prop_est | -0.417 | 0.422 |
| 238 | rec_dev_est | -0.132 | 0.197 | 288 | logit_rec_prop_est | 0.046 | 0.434 |
| 239 | rec_dev_est | -1.581 | 0.363 | 289 | logit_rec_prop_est | -0.350 | 0.139 |
| 240 | rec_dev_est | -0.481 | 0.209 | 290 | logit_rec_prop_est | -0.069 | 0.239 |
| 241 | rec_dev_est | -1.543 | 0.410 | 291 | logit_rec_prop_est | 0.381 | 0.279 |
| 242 | rec_dev_est | 1.377 | 0.164 | 292 | logit_rec_prop_est | -0.184 | 0.377 |
| 293 | logit_rec_prop_est | -0.451 | 0.356 |  |  |  |  |
| 294 | logit_rec_prop_est | -0.724 | 0.200 |  |  |  |  |
| 295 | logit_rec_prop_est | -0.511 | 0.313 |  |  |  |  |
| 296 | logit_rec_prop_est | -0.461 | 0.358 |  |  |  |  |
| 297 | logit_rec_prop_est | -0.237 | 0.332 |  |  |  |  |
| 298 | logit_rec_prop_est | -0.306 | 0.422 |  |  |  |  |
| 299 | logit_rec_prop_est | -0.278 | 0.321 |  |  |  |  |
| 300 | logit_rec_prop_est | 0.327 | 0.212 |  |  |  |  |
| 301 | logit_rec_prop_est | 0.516 | 0.456 |  |  |  |  |
| 302 | logit_rec_prop_est | 0.811 | 0.307 |  |  |  |  |
| 303 | logit_rec_prop_est | 0.122 | 0.454 |  |  |  |  |
| 304 | logit_rec_prop_est | 0.587 | 0.488 |  |  |  |  |
| 305 | logit_rec_prop_est | 0.096 | 0.448 |  |  |  |  |
| 306 | logit_rec_prop_est | -0.969 | 0.567 |  |  |  |  |
| 307 | survey_q[1] | 0.941 | 0.027 |  |  |  |  |
| 308 | log_add_cv[2] | -0.816 | 0.276 |  |  |  |  |

Table 6c. Summary of estimated model parameter values and standard deviations for model 22.0a for Bristol Bay red king crab.

| index | name | value | std.dev | index | name | value | std.dev |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | theta[1] | 0.228 | 0.007 | 47 | log_slx_pars[2] | 2.275 | 0.043 |
| 2 | theta[2] | 0.135 | 0.020 | 48 | log_slx_pars[3] | 4.548 | 0.019 |
| 3 | theta[4] | 18.053 | 0.053 | 49 | log_slx_pars[4] | 2.177 | 0.099 |
| 4 | theta[5] | 16.129 | 0.162 | 50 | log_slx_pars[5] | 5.176 | 0.070 |
| 5 | theta[7] | 0.684 | 0.127 | 51 | log_slx_pars[6] | 2.857 | 0.049 |
| 6 | theta[9] | -0.433 | 0.248 | 52 | log_slx_pars[7] | 4.724 | 0.239 |
| 7 | theta[13] | 0.907 | 0.505 | 53 | log_slx_pars[8] | 2.168 | 0.305 |
| 8 | theta[14] | 0.941 | 0.477 | 54 | log_slx_pars[9] | 4.701 | 0.104 |
| 9 | theta[15] | 1.263 | 0.352 | 55 | log_slx_pars[10] | 0.905 | 0.302 |
| 10 | theta[16] | 1.419 | 0.288 | 56 | log_slx_pars[11] | 4.809 | 0.023 |
| 11 | theta[17] | 1.336 | 0.266 | 57 | log_slx_pars[12] | 2.345 | 0.079 |
| 12 | theta[18] | 1.052 | 0.272 | 58 | log_slx_pars[13] | 4.008 | 0.335 |
| 13 | theta[19] | 0.981 | 0.258 | 59 | log_slx_pars[14] | 3.375 | 0.502 |
| 14 | theta[20] | 1.199 | 0.221 | 60 | log_slx_pars[15] | 4.480 | 0.033 |
| 15 | theta[21] | 1.156 | 0.216 | 61 | log_slx_pars[16] | 2.565 | 0.105 |
| 16 | theta[22] | 0.944 | 0.224 | 62 | log_slx_pars[17] | 4.924 | 0.002 |
| 17 | theta[23] | 0.863 | 0.217 | 63 | log_slx_pars[18] | 0.673 | 0.070 |
| 18 | theta[24] | 0.687 | 0.222 | 64 | log_slx_pars[19] | 4.933 | 0.002 |
| 19 | theta[25] | 0.344 | 0.227 | 65 | log_slx_pars[20] | 0.737 | 0.097 |
| 20 | theta[26] | -0.102 | 0.196 | 66 | $\log _{-} \mathrm{fbar}[1]$ | -1.678 | 0.049 |
| 21 | theta[27] | -0.582 | 0.198 | 67 | log_fbar[2] | -4.707 | 0.082 |
| 22 | theta[28] | -1.236 | 0.221 | 68 | log_fbar[3] | -6.067 | 0.344 |
| 23 | theta[29] | -1.812 | 0.252 | 69 | log_fbar[4] | -6.553 | 0.078 |
| 24 | theta[30] | -2.483 | 0.275 | 70 | log_fdev[1] | 1.080 | 0.123 |
| 25 | theta[31] | -2.110 | 0.352 | 71 | log_fdev[1] | 1.291 | 0.083 |
| 26 | theta[52] | -0.079 | 0.608 | 72 | log_fdev[1] | 0.873 | 0.066 |
| 27 | theta[53] | 0.446 | 0.691 | 73 | log_fdev[1] | 0.003 | 0.054 |
| 28 | theta[54] | 0.990 | 0.566 | 74 | log_fdev[1] | 0.148 | 0.048 |
| 29 | theta[55] | 1.213 | 0.431 | 75 | log_fdev[1] | 0.802 | 0.038 |
| 30 | theta[56] | 1.350 | 0.336 | 76 | $\mathrm{log}_{\text {_f }} \mathrm{fdev}[1]$ | 0.806 | 0.040 |
| 31 | theta[57] | 1.152 | 0.314 | 77 | log_fdev[1] | 0.291 | 0.044 |
| 32 | theta[58] | 0.910 | 0.309 | 78 | log_fdev[1] | 0.949 | 0.048 |
| 33 | theta[59] | 0.434 | 0.342 | 79 | log_fdev[1] | -4.216 | 0.046 |
| 34 | theta[60] | -0.326 | 0.387 | 80 | log_fdev[1] | -4.617 | 0.040 |
| 35 | theta[61] | -0.798 | 0.380 | 81 | log_fdev[1] | -0.131 | 0.039 |
| 36 | theta[62] | -1.534 | 0.372 | 82 | log_fdev[1] | -0.068 | 0.039 |
| 37 | theta[63] | -1.635 | 0.368 | 83 | log_fdev[1] | 0.853 | 0.041 |
| 38 | theta[64] | -1.560 | 0.365 | 84 | log_fdev[1] | 0.469 | 0.041 |
| 39 | theta[65] | -1.802 | 0.356 | 85 | log_fdev[1] | -0.124 | 0.039 |
| 40 | theta[66] | -1.957 | 0.346 | 86 | log_fdev[1] | -0.189 | 0.039 |
| 41 | theta[67] | -1.928 | 0.336 | 87 | log_fdev[1] | -0.067 | 0.038 |
| 42 | Grwth[21] | 0.888 | 0.192 | 88 | log_fdev[1] | 0.391 | 0.037 |


| 43 | Grwth[42] | 1.460 | 0.138 | 89 | log_fdev[1] | 0.348 | 0.037 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 44 | Grwth[64] | 141.050 | 0.637 | 90 | log_fdev[1] | 0.641 | 0.038 |
| 45 | Grwth[65] | 0.068 | 0.003 | 91 | log_fdev[1] | 0.387 | 0.037 |
| 46 | log_slx_pars[1] | 4.779 | 0.009 | 92 | $\mathrm{log}_{-} \mathrm{fdev}[1]$ | 0.749 | 0.037 |
| 93 | log_fdev[1] | 0.918 | 0.038 | 143 | log_fdev[2] | -0.990 | 0.114 |
| 94 | log_fdev[1] | 0.720 | 0.039 | 144 | log_fdev[3] | -0.727 | 0.066 |
| 95 | log_fdev[1] | 0.575 | 0.038 | 145 | log_fdev[3] | 0.116 | 0.066 |
| 96 | log_fdev[1] | -0.063 | 0.037 | 146 | log_fdev[3] | 1.312 | 0.066 |
| 97 | log_fdev[1] | -0.129 | 0.036 | 147 | log_fdev[3] | 1.790 | 0.066 |
| 98 | log_fdev[1] | 0.071 | 0.036 | 148 | log_fdev[3] | 3.619 | 0.076 |
| 99 | log_fdev[1] | 0.403 | 0.037 | 149 | $\log _{-}$fdev[3] | 1.661 | 0.107 |
| 100 | log_fdev[1] | 0.472 | 0.039 | 150 | log_fdev[3] | 0.932 | 0.133 |
| 101 | log_fdev[1] | 0.475 | 0.043 | 151 | log_fdev[3] | -0.394 | 0.085 |
| 102 | log_fdev[1] | 0.398 | 0.050 | 152 | log_fdev[3] | -1.751 | 0.075 |
| 103 | log_fdev[1] | 0.229 | 0.059 | 153 | log_fdev[3] | -2.609 | 0.099 |
| 104 | log_fdev[1] | 0.189 | 0.067 | 154 | log_fdev[3] | -2.043 | 0.122 |
| 105 | log_fdev[1] | -0.241 | 0.071 | 155 | log_fdev[3] | -3.136 | 0.077 |
| 106 | log_fdev[1] | -4.689 | 0.071 | 156 | log_fdev[3] | -0.470 | 0.099 |
| 107 | log_fdev[2] | 2.414 | 0.116 | 157 | log_fdev[3] | 0.256 | 0.123 |
| 108 | log_fdev[2] | 1.369 | 0.114 | 158 | log_fdev[3] | 1.443 | 0.152 |
| 109 | log_fdev[2] | 0.951 | 0.110 | 159 | log_fdev[4] | 0.556 | 0.103 |
| 110 | log_fdev[2] | 1.771 | 0.105 | 160 | log_fdev[4] | -0.094 | 0.102 |
| 111 | log_fdev[2] | 0.367 | 0.104 | 161 | log_fdev[4] | -0.314 | 0.103 |
| 112 | log_fdev[2] | 0.825 | 0.103 | 162 | log_fdev[4] | 0.592 | 0.102 |
| 113 | log_fdev[2] | 1.241 | 0.104 | 163 | log_fdev[4] | -1.836 | 0.101 |
| 114 | log_fdev[2] | 1.079 | 0.104 | 164 | log_fdev[4] | 0.126 | 0.101 |
| 115 | log_fdev[2] | 1.542 | 0.107 | 165 | log_fdev[4] | -0.128 | 0.101 |
| 116 | log_fdev[2] | -0.215 | 0.104 | 166 | log_fdev[4] | -0.965 | 0.101 |
| 117 | log_fdev[2] | -0.494 | 0.103 | 167 | log_fdev[4] | -0.789 | 0.100 |
| 118 | log_fdev[2] | -0.422 | 0.103 | 168 | log_fdev[4] | -0.516 | 0.100 |
| 119 | log_fdev[2] | -0.878 | 0.103 | 169 | log_fdev[4] | -0.565 | 0.100 |
| 120 | log_fdev[2] | 0.416 | 0.103 | 170 | log_fdev[4] | -0.018 | 0.100 |
| 121 | log_fdev[2] | 0.126 | 0.103 | 171 | log_fdev[4] | -0.720 | 0.100 |
| 122 | log_fdev[2] | -0.638 | 0.102 | 172 | log_fdev[4] | -1.724 | 0.100 |
| 123 | log_fdev[2] | 0.138 | 0.102 | 173 | log_fdev[4] | -2.565 | 0.100 |
| 124 | log_fdev[2] | -0.153 | 0.102 | 174 | log_fdev[4] | -1.082 | 0.099 |
| 125 | log_fdev[2] | -0.249 | 0.102 | 175 | log_fdev[4] | -0.517 | 0.099 |
| 126 | log_fdev[2] | -0.019 | 0.102 | 176 | log_fdev[4] | 0.631 | 0.099 |
| 127 | log_fdev[2] | -0.297 | 0.102 | 177 | log_fdev[4] | 1.485 | 0.100 |
| 128 | log_fdev[2] | -0.129 | 0.101 | 178 | log_fdev[4] | 1.173 | 0.100 |
| 129 | log_fdev[2] | -0.058 | 0.102 | 179 | log_fdev[4] | 0.350 | 0.101 |
| 130 | log_fdev[2] | -0.091 | 0.102 | 180 | log_fdev[4] | 1.965 | 0.102 |
| 131 | log_fdev[2] | -0.452 | 0.102 | 181 | log_fdev[4] | 2.238 | 0.104 |
| 132 | log_fdev[2] | -0.606 | 0.102 | 182 | log_fdev[4] | 1.046 | 0.105 |
| 133 | log_fdev[2] | -1.065 | 0.102 | 183 | log_fdev[4] | 0.844 | 0.107 |
| 134 | log_fdev[2] | -1.577 | 0.102 | 184 | log_fdev[4] | 0.828 | 0.109 |
| 135 | log_fdev[2] | -0.852 | 0.102 | 185 | log_foff[1] | -2.815 | 0.044 |
| 136 | log_fdev[2] | -1.409 | 0.102 | 186 | log_foff[3] | -0.269 | 0.480 |
| 137 | log_fdev[2] | -1.023 | 0.103 | 187 | log_fdov[1] | 1.972 | 0.084 |


| 138 | log_fdev[2] | -0.491 | 0.104 | 188 | log_fdov[1] | -0.689 | 0.083 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 139 | log_fdev[2] | -0.046 | 0.106 | 189 | log_fdov[1] | 1.976 | 0.084 |
| 140 | log_fdev[2] | -0.094 | 0.108 | 190 | log_fdov[1] | 1.812 | 0.086 |
| 141 | log_fdev[2] | 0.015 | 0.111 | 191 | log_fdov[1] | -0.410 | 0.084 |
| 142 | log_fdev[2] | -0.005 | 0.113 | 192 | log_fdov[1] | -0.193 | 0.082 |
| 193 | log_fdov[1] | -3.696 | 0.081 | 243 | rec_dev_est | 1.368 | 0.163 |
| 194 | log_fdov[1] | -0.344 | 0.082 | 244 | rec_dev_est | -0.441 | 0.255 |
| 195 | log_fdov[1] | 1.420 | 0.083 | 245 | rec_dev_est | -1.147 | 0.332 |
| 196 | log_fdov[1] | -2.793 | 0.082 | 246 | rec_dev_est | -0.171 | 0.209 |
| 197 | log_fdov[1] | 1.146 | 0.081 | 247 | rec_dev_est | 0.820 | 0.171 |
| 198 | log_fdov[1] | 0.860 | 0.081 | 248 | rec_dev_est | -0.134 | 0.231 |
| 199 | log_fdov[1] | -1.898 | 0.080 | 249 | rec_dev_est | -0.170 | 0.255 |
| 200 | log_fdov[1] | 1.199 | 0.080 | 250 | rec_dev_est | 1.239 | 0.170 |
| 201 | log_fdov[1] | 0.407 | 0.081 | 251 | rec_dev_est | -0.187 | 0.266 |
| 202 | log_fdov[1] | 0.929 | 0.080 | 252 | rec_dev_est | -0.284 | 0.268 |
| 203 | log_fdov[1] | -1.244 | 0.079 | 253 | rec_dev_est | 0.942 | 0.173 |
| 204 | log_fdov[1] | -0.199 | 0.079 | 254 | rec_dev_est | 0.291 | 0.192 |
| 205 | log_fdov[1] | -0.462 | 0.080 | 255 | rec_dev_est | -0.111 | 0.199 |
| 206 | log_fdov[1] | -0.713 | 0.080 | 256 | rec_dev_est | -0.696 | 0.243 |
| 207 | log_fdov[1] | -0.215 | 0.080 | 257 | rec_dev_est | -0.542 | 0.240 |
| 208 | log_fdov[1] | -1.096 | 0.079 | 258 | rec_dev_est | 0.331 | 0.191 |
| 209 | log_fdov[1] | -1.813 | 0.078 | 259 | rec_dev_est | -0.108 | 0.227 |
| 210 | log_fdov[1] | 0.201 | 0.078 | 260 | rec_dev_est | -0.722 | 0.243 |
| 211 | log_fdov[1] | -0.203 | 0.079 | 261 | rec_dev_est | -1.031 | 0.233 |
| 212 | log_fdov[1] | 0.864 | 0.079 | 262 | rec_dev_est | -1.506 | 0.274 |
| 213 | log_fdov[1] | 0.318 | 0.081 | 263 | rec_dev_est | -1.071 | 0.227 |
| 214 | log_fdov[1] | -0.338 | 0.083 | 264 | rec_dev_est | -0.408 | 0.186 |
| 215 | log_fdov[1] | 0.975 | 0.087 | 265 | rec_dev_est | -1.226 | 0.256 |
| 216 | log_fdov[1] | -0.109 | 0.090 | 266 | rec_dev_est | -0.471 | 0.204 |
| 217 | log_fdov[1] | -0.630 | 0.092 | 267 | rec_dev_est | -1.072 | 0.281 |
| 218 | log_fdov[1] | 2.963 | 0.093 | 268 | rec_dev_est | -0.954 | 0.282 |
| 219 | log_fdov[3] | 0.000 | 0.093 | 269 | rec_dev_est | -0.839 | 0.283 |
| 220 | log_fdov[3] | 0.000 | 0.093 | 270 | rec_dev_est | -0.510 | 0.322 |
| 221 | log_fdov[3] | 0.000 | 0.093 | 271 | logit_rec_prop_est | -0.441 | 0.148 |
| 222 | log_fdov[3] | 0.001 | 0.093 | 272 | logit_rec_prop_est | 0.216 | 0.403 |
| 223 | log_fdov[3] | 1.494 | 0.120 | 273 | logit_rec_prop_est | -0.075 | 0.450 |
| 224 | log_fdov[3] | 1.809 | 0.130 | 274 | logit_rec_prop_est | 0.450 | 0.371 |
| 225 | log_fdov[3] | 0.604 | 0.151 | 275 | logit_rec_prop_est | -0.072 | 0.166 |
| 226 | log_fdov[3] | -3.435 | 0.114 | 276 | logit_rec_prop_est | 0.207 | 0.233 |
| 227 | log_fdov[3] | -2.230 | 0.149 | 277 | logit_rec_prop_est | 0.648 | 0.658 |
| 228 | log_fdov[3] | -0.825 | 0.122 | 278 | logit_rec_prop_est | 0.308 | 0.285 |
| 229 | log_fdov[3] | 0.021 | 0.142 | 279 | logit_rec_prop_est | -0.490 | 0.671 |
| 230 | log_fdov[3] | 0.385 | 0.103 | 280 | logit_rec_prop_est | -0.299 | 0.090 |
| 231 | log_fdov[3] | 1.010 | 0.143 | 281 | logit_rec_prop_est | 1.198 | 0.569 |
| 232 | log_fdov[3] | 0.214 | 0.151 | 282 | logit_rec_prop_est | 0.451 | 0.607 |
| 233 | log_fdov[3] | 0.951 | 0.194 | 283 | logit_rec_prop_est | 0.536 | 0.324 |
| 234 | rec_dev_est | 0.753 | 0.174 | 284 | logit_rec_prop_est | -0.057 | 0.140 |
| 235 | rec_dev_est | -0.429 | 0.247 | 285 | logit_rec_prop_est | 0.217 | 0.354 |
| 236 | rec_dev_est | -0.864 | 0.281 | 286 | logit_rec_prop_est | -0.462 | 0.385 |


| 237 | rec_dev_est | -0.574 | 0.227 | 287 | logit_rec_prop_est | -0.500 | 0.127 |
| :--- | :--- | ---: | :--- | :--- | :--- | :--- | :--- |
| 238 | rec_dev_est | 0.324 | 0.178 | 288 | logit_rec_prop_est | -0.388 | 0.410 |
| 239 | rec_dev_est | -0.068 | 0.194 | 289 | logit_rec_prop_est | -0.004 | 0.431 |
| 240 | rec_dev_est | -1.509 | 0.352 | 290 | logit_recprop_est | -0.365 | 0.142 |
| 241 | rec_dev_est | -0.460 | 0.208 | 291 | logit_recprop_est | -0.126 | 0.225 |
| 242 | rec_dev_est | -1.601 | 0.427 | 292 | logit_rec_prop_est | 0.451 | 0.280 |
| 293 | logit_rec_prop_est | -0.106 | 0.365 |  |  |  |  |
| 294 | logit_rec_prop_est | -0.451 | 0.347 |  |  |  |  |
| 295 | logit_rec_prop_est | -0.652 | 0.197 |  |  |  |  |
| 296 | logit_rec_prop_est | -0.490 | 0.292 |  |  |  |  |
| 297 | logit_rec_prop_est | -0.440 | 0.346 |  |  |  |  |
| 298 | logit_rec_prop_est | -0.207 | 0.332 |  |  |  |  |
| 299 | logit_rec_prop_est | -0.307 | 0.424 |  |  |  |  |
| 300 | logit_rec_prop_est | -0.300 | 0.312 |  |  |  |  |
| 301 | logit_rec_prop_est | 0.319 | 0.211 |  |  |  |  |
| 302 | logit_rec_prop_est | 0.553 | 0.453 |  |  |  |  |
| 303 | logit_rec_prop_est | 0.792 | 0.301 |  |  |  |  |
| 304 | logit_rec_prop_est | 0.148 | 0.454 |  |  |  |  |
| 305 | logit_rec_prop_est | 0.536 | 0.481 |  |  |  |  |
| 306 | logit_rec_prop_est | 0.171 | 0.448 |  |  |  |  |
| 307 | logit_rec_prop_est | -0.967 | 0.552 |  |  |  |  |
| 308 | survey_q[1] | 0.922 | 0.028 |  |  |  |  |
| 309 | log_add_cv[2] | -0.971 | 0.289 |  |  |  |  |
|  |  |  |  |  |  |  |  |

Table 7. Natural mortality estimates for three model scenarios during different year blocks.

| Model | 1975-1979, |  | 1980-1984 | 1985-2022 |
| :---: | :---: | :---: | :---: | :---: |
|  | Sex | 1985-2022 |  |  |
| 21.1b | Males | 0.180 | 0.886 |  |
|  | Females | 0.238 | 1.174 |  |
| 22.0 | Males |  |  | 0.180 |
|  | Females |  |  | 0.232 |
| 22.0a | Males |  |  | 0.228 |
|  | Females |  |  | 0.261 |

Table 8. Area-swept estimates of mature female abundance (million crab) and model estimates of effective spawning biomass (ESB, Zheng et al. 1995b) (1000 t) during 2011-2022 for groundfish fisheries bycatch (PSC) calculation. (*mature female abundance in 2020 is the model projected value). Note that PSC limits apply to previous-year ESB.

| Year | Mature female <br> abundance | Effective spawning <br> biomass |
| :---: | :---: | :---: |
| 2011 | 28.520 | 19.541 |
| 2012 | 21.121 | 20.029 |
| 2013 | 15.694 | 22.382 |
| 2014 | 38.580 | 23.272 |
| 2015 | 18.666 | 21.098 |
| 2016 | 22.633 | 19.147 |
| 2017 | 18.497 | 18.042 |
| 2018 | 9.106 | 15.093 |
| 2019 | 8.587 | 12.705 |
| 2020 | $9.668^{*}$ | 11.394 |
| 2021 | 6.432 | 9.463 |
| 2022 | 8.004 | 8.894 |

Table 9a. Annual abundance estimates (million crab), mature male biomass (MMB, 1000 t ), and total survey biomass (1000 t) for red king crab in Bristol Bay estimated by length-based model 21.1b during 1975-2022. MMB for year $t$ is on Feb. 15, year $t+1$. Size measurements are mm carapace length.

| Year (t) | Males |  |  |  | FemalesMature$(>89 \mathrm{~mm})$ | Total Recruits | Total Survey Biomass |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\begin{gathered} \text { Mature } \\ (>119 \mathrm{~mm}) \end{gathered}$ | $\begin{gathered} \text { Legal } \\ (>134 \mathrm{~mm}) \end{gathered}$ | $\begin{gathered} \text { MMB } \\ (>119 \mathrm{~mm}) \end{gathered}$ | SD MMB |  |  | Model Est. (>64 mm) | Area-Swept (>64 mm) |
| 1975 | 55.540 | 28.226 | 83.213 | 0.050 | 54.794 | NA | 235.861 | 199.643 |
| 1976 | 65.221 | 35.521 | 99.070 | 3.381 | 83.134 | 63.885 | 275.709 | 327.615 |
| 1977 | 72.440 | 41.304 | 113.031 | 3.817 | 110.373 | 41.197 | 297.013 | 371.223 |
| 1978 | 77.709 | 46.495 | 119.799 | 3.719 | 114.468 | 64.349 | 300.165 | 343.189 |
| 1979 | 68.280 | 47.420 | 99.943 | 3.121 | 109.505 | 115.470 | 288.724 | 165.449 |
| 1980 | 50.101 | 37.749 | 30.332 | 1.572 | 111.469 | 150.262 | 273.367 | 247.226 |
| 1981 | 14.471 | 8.026 | 6.551 | 1.017 | 48.863 | 67.936 | 109.145 | 131.145 |
| 1982 | 6.784 | 2.164 | 6.563 | 0.850 | 21.395 | 241.558 | 65.174 | 141.898 |
| 1983 | 6.175 | 2.171 | 7.407 | 0.590 | 14.199 | 93.067 | 57.646 | 48.476 |
| 1984 | 6.151 | 2.291 | 5.230 | 0.351 | 13.980 | 64.266 | 50.624 | 152.607 |
| 1985 | 7.558 | 1.884 | 9.676 | 0.376 | 9.730 | 10.204 | 34.842 | 34.138 |
| 1986 | 12.174 | 4.646 | 15.064 | 0.558 | 13.636 | 30.361 | 45.545 | 47.434 |
| 1987 | 14.360 | 6.686 | 20.405 | 0.701 | 17.029 | 9.473 | 51.326 | 69.245 |
| 1988 | 14.454 | 8.458 | 25.123 | 0.724 | 21.453 | 6.184 | 54.700 | 54.597 |
| 1989 | 15.546 | 9.750 | 27.979 | 0.701 | 20.233 | 8.033 | 57.355 | 55.136 |
| 1990 | 15.014 | 10.437 | 24.139 | 0.724 | 18.058 | 20.848 | 57.416 | 59.451 |
| 1991 | 11.547 | 8.652 | 18.449 | 0.760 | 17.435 | 13.111 | 52.333 | 83.892 |
| 1992 | 9.280 | 6.463 | 17.241 | 0.750 | 18.579 | 3.023 | 47.662 | 37.334 |
| 1993 | 10.499 | 6.160 | 15.852 | 0.826 | 17.269 | 9.070 | 47.203 | 52.906 |
| 1994 | 10.350 | 6.016 | 21.696 | 0.806 | 14.681 | 2.910 | 42.628 | 32.104 |
| 1995 | 10.862 | 7.892 | 24.839 | 0.730 | 13.574 | 59.439 | 48.696 | 38.068 |
| 1996 | 11.145 | 8.548 | 23.323 | 0.698 | 19.693 | 8.630 | 58.051 | 43.959 |
| 1997 | 10.587 | 7.787 | 22.063 | 0.688 | 28.739 | 4.446 | 64.180 | 84.030 |
| 1998 | 15.903 | 7.751 | 24.944 | 0.811 | 25.234 | 12.405 | 68.062 | 84.101 |
| 1999 | 16.967 | 9.732 | 28.678 | 0.875 | 21.317 | 33.581 | 66.662 | 64.754 |
| 2000 | 14.654 | 10.648 | 28.897 | 0.869 | 22.735 | 12.468 | 68.283 | 67.381 |
| 2001 | 14.471 | 10.216 | 29.284 | 0.838 | 25.762 | 12.893 | 71.897 | 52.455 |
| 2002 | 17.318 | 10.409 | 33.344 | 0.802 | 25.010 | 51.478 | 76.960 | 69.086 |
| 2003 | 18.144 | 12.050 | 32.898 | 0.816 | 30.661 | 11.755 | 83.108 | 115.760 |
| 2004 | 16.357 | 11.629 | 30.409 | 0.817 | 37.823 | 10.951 | 84.595 | 130.556 |
| 2005 | 18.229 | 10.845 | 30.944 | 0.821 | 35.026 | 39.536 | 85.554 | 105.727 |
| 2006 | 17.344 | 11.425 | 31.316 | 0.808 | 35.327 | 18.930 | 85.403 | 94.477 |
| 2007 | 15.638 | 11.175 | 26.310 | 0.828 | 39.205 | 12.793 | 86.951 | 103.327 |
| 2008 | 16.023 | 9.509 | 24.990 | 0.876 | 36.751 | 7.149 | 83.339 | 113.082 |
| 2009 | 15.870 | 9.465 | 25.857 | 0.906 | 32.183 | 8.156 | 77.330 | 90.547 |
| 2010 | 14.749 | 9.692 | 25.141 | 0.887 | 28.167 | 21.789 | 72.264 | 80.501 |
| 2011 | 12.486 | 9.152 | 24.850 | 0.822 | 27.729 | 12.611 | 67.802 | 66.408 |
| 2012 | 11.131 | 8.608 | 23.277 | 0.758 | 29.488 | 7.303 | 66.173 | 60.697 |
| 2013 | 11.042 | 7.865 | 22.213 | 0.704 | 27.879 | 5.315 | 63.453 | 62.217 |
| 2014 | 10.759 | 7.568 | 20.208 | 0.674 | 24.647 | 3.324 | 58.710 | 113.135 |
| 2015 | 9.238 | 6.891 | 17.223 | 0.664 | 21.092 | 5.278 | 51.909 | 64.175 |
| 2016 | 7.483 | 5.797 | 14.188 | 0.668 | 17.965 | 10.332 | 45.238 | 60.958 |
| 2017 | 5.936 | 4.699 | 11.595 | 0.673 | 16.325 | 4.358 | 40.299 | 52.935 |
| 2018 | 5.173 | 3.797 | 10.342 | 0.679 | 14.911 | 9.732 | 37.304 | 28.805 |
| 2019 | 5.919 | 3.519 | 11.240 | 0.742 | 13.124 | 5.294 | 36.024 | 28.539 |
| 2020 | 6.489 | 4.032 | 12.859 | 0.804 | 12.066 | 5.793 |  |  |
| 2021 | 7.671 | 4.652 | 16.637 | 0.830 | 10.982 | 6.606 | 36.141 | 28.476 |
| 2022 | 8.658 | 5.963 | 16.953 | 1.180 | 10.201 | 9.006 | 38.016 | 36.198 |

Table 9b. Annual abundance estimates (million crab), mature male biomass (MMB, 1000 t ), and total survey biomass ( 1000 t ) for red king crab in Bristol Bay estimated by length-based model 22.0 during 19752022. MMB for year $t$ is on Feb. 15, year $t+1$. Size measurements are mm carapace length.

| Year (t) | Males |  |  |  | Females | Total Recruits | Total Survey Biomass |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\begin{gathered} \text { Mature } \\ (>119 \mathrm{~mm}) \end{gathered}$ | $\begin{gathered} \text { Legal } \\ (>134 \mathrm{~mm}) \end{gathered}$ | $\begin{gathered} \text { MMB } \\ (>119 \mathrm{~mm}) \end{gathered}$ | SD MMB | $\begin{gathered} \text { Mature } \\ (>89 \mathrm{~mm}) \end{gathered}$ |  | Model Est. <br> ( $>64 \mathrm{~mm}$ ) | Area-Swept (>64 mm) |
| 1985 | 8.557 | 2.350 | 11.651 | 0.007 | 8.440 | NA | 34.983 | 34.138 |
| 1986 | 12.986 | 5.351 | 17.143 | 0.505 | 11.908 | 31.070 | 44.854 | 47.434 |
| 1987 | 14.402 | 7.309 | 21.340 | 0.677 | 15.698 | 9.169 | 50.338 | 69.245 |
| 1988 | 14.418 | 8.689 | 25.552 | 0.704 | 20.545 | 6.011 | 53.589 | 54.597 |
| 1989 | 15.533 | 9.777 | 28.169 | 0.696 | 19.574 | 8.255 | 56.086 | 55.136 |
| 1990 | 15.089 | 10.459 | 24.356 | 0.739 | 17.604 | 20.523 | 56.080 | 59.451 |
| 1991 | 11.608 | 8.711 | 18.620 | 0.785 | 17.115 | 12.964 | 51.198 | 83.892 |
| 1992 | 9.366 | 6.515 | 17.426 | 0.776 | 18.263 | 3.043 | 46.836 | 37.334 |
| 1993 | 10.616 | 6.219 | 16.093 | 0.855 | 17.056 | 9.144 | 46.525 | 52.906 |
| 1994 | 10.513 | 6.101 | 22.010 | 0.830 | 14.609 | 3.161 | 42.186 | 32.104 |
| 1995 | 10.986 | 8.007 | 25.120 | 0.748 | 13.638 | 58.618 | 47.973 | 38.068 |
| 1996 | 11.305 | 8.643 | 23.651 | 0.719 | 19.628 | 8.485 | 57.278 | 43.959 |
| 1997 | 10.662 | 7.906 | 22.290 | 0.708 | 28.458 | 4.532 | 63.631 | 84.030 |
| 1998 | 16.156 | 7.799 | 25.401 | 0.838 | 25.175 | 12.459 | 67.618 | 84.101 |
| 1999 | 17.270 | 9.916 | 29.280 | 0.907 | 21.402 | 33.141 | 66.212 | 64.754 |
| 2000 | 14.880 | 10.863 | 29.431 | 0.900 | 22.783 | 12.372 | 67.855 | 67.381 |
| 2001 | 14.670 | 10.391 | 29.770 | 0.867 | 25.786 | 13.073 | 71.515 | 52.455 |
| 2002 | 17.558 | 10.556 | 33.871 | 0.830 | 25.156 | 49.985 | 76.328 | 69.086 |
| 2003 | 18.387 | 12.227 | 33.442 | 0.845 | 30.609 | 11.816 | 82.352 | 115.760 |
| 2004 | 16.544 | 11.810 | 30.880 | 0.846 | 37.556 | 10.964 | 83.936 | 130.556 |
| 2005 | 18.450 | 10.978 | 31.431 | 0.854 | 35.019 | 38.925 | 84.824 | 105.727 |
| 2006 | 17.553 | 11.590 | 31.797 | 0.842 | 35.322 | 18.643 | 84.702 | 94.477 |
| 2007 | 15.802 | 11.330 | 26.723 | 0.864 | 39.147 | 12.975 | 86.364 | 103.327 |
| 2008 | 16.242 | 9.630 | 25.461 | 0.918 | 36.866 | 7.063 | 83.001 | 113.082 |
| 2009 | 16.095 | 9.624 | 26.353 | 0.951 | 32.483 | 8.334 | 77.219 | 90.547 |
| 2010 | 14.996 | 9.858 | 25.679 | 0.933 | 28.616 | 21.279 | 72.204 | 80.501 |
| 2011 | 12.713 | 9.340 | 25.376 | 0.865 | 28.161 | 12.983 | 67.831 | 66.408 |
| 2012 | 11.310 | 8.780 | 23.735 | 0.798 | 29.908 | 7.126 | 66.312 | 60.697 |
| 2013 | 11.238 | 8.006 | 22.677 | 0.742 | 28.489 | 5.404 | 63.718 | 62.217 |
| 2014 | 10.954 | 7.718 | 20.672 | 0.713 | 25.260 | 3.443 | 59.092 | 113.135 |
| 2015 | 9.414 | 7.039 | 17.660 | 0.705 | 21.759 | 5.334 | 52.412 | 64.175 |
| 2016 | 7.648 | 5.935 | 14.603 | 0.710 | 18.643 | 10.655 | 45.840 | 60.958 |
| 2017 | 6.089 | 4.830 | 11.990 | 0.715 | 17.006 | 4.536 | 41.040 | 52.935 |
| 2018 | 5.319 | 3.918 | 10.721 | 0.719 | 15.634 | 9.940 | 38.162 | 28.805 |
| 2019 | 6.122 | 3.635 | 11.695 | 0.781 | 13.838 | 5.520 | 36.977 | 28.539 |
| 2020 | 6.713 | 4.187 | 13.370 | 0.840 | 12.782 | 5.916 |  |  |
| 2021 | 7.939 | 4.820 | 17.230 | 0.862 | 11.737 | 6.793 | 37.146 | 28.476 |
| 2022 | 8.922 | 6.167 | 17.158 | 1.204 | 10.992 | 9.513 | 38.940 | 36.198 |

Table 9c. Annual abundance estimates (million crab), mature male biomass (MMB, 1000 t ), and total survey biomass (1000 t) for red king crab in Bristol Bay estimated by length-based model 22.0a during 1975-2022. MMB for year $t$ is on Feb. 15, year $t+1$. Size measurements are mm carapace length.

| Year (t) | Males |  |  |  | Females | Total Recruits | Total Survey Biomass |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\begin{gathered} \text { Mature } \\ (>119 \mathrm{~mm}) \end{gathered}$ | $\begin{gathered} \text { Legal } \\ (>134 \mathrm{~mm}) \end{gathered}$ | $\begin{gathered} \text { MMB } \\ (>119 \mathrm{~mm}) \end{gathered}$ | SD MMB | $\begin{aligned} & \text { Mature } \\ & (>89 \mathrm{~mm}) \end{aligned}$ |  | Model Est. <br> (>64 mm) | Area-Swept (>64 mm) |
| 1985 | 8.837 | 2.375 | 11.771 | 0.007 | 10.572 | NA | 34.058 | 34.138 |
| 1986 | 13.669 | 5.465 | 17.699 | 0.527 | 14.890 | 42.936 | 44.651 | 47.434 |
| 1987 | 15.464 | 7.611 | 22.496 | 0.717 | 19.678 | 13.167 | 50.907 | 69.245 |
| 1988 | 15.613 | 9.197 | 27.056 | 0.749 | 25.738 | 8.523 | 54.866 | 54.597 |
| 1989 | 16.905 | 10.330 | 29.968 | 0.755 | 24.215 | 11.387 | 57.743 | 55.136 |
| 1990 | 16.338 | 11.099 | 26.071 | 0.821 | 21.462 | 27.952 | 57.729 | 59.451 |
| 1991 | 12.521 | 9.272 | 19.972 | 0.870 | 20.935 | 18.883 | 52.802 | 83.892 |
| 1992 | 10.235 | 6.932 | 18.686 | 0.858 | 22.567 | 4.469 | 48.624 | 37.334 |
| 1993 | 11.773 | 6.649 | 17.712 | 0.958 | 21.033 | 12.766 | 48.605 | 52.906 |
| 1994 | 11.889 | 6.737 | 24.062 | 0.954 | 17.867 | 4.077 | 44.562 | 32.104 |
| 1995 | 12.148 | 8.741 | 26.941 | 0.871 | 16.446 | 79.363 | 50.163 | 38.068 |
| 1996 | 12.243 | 9.191 | 25.037 | 0.820 | 24.647 | 13.010 | 58.696 | 43.959 |
| 1997 | 11.440 | 8.283 | 23.274 | 0.786 | 35.915 | 6.420 | 65.015 | 84.030 |
| 1998 | 17.674 | 8.133 | 27.168 | 0.948 | 31.183 | 17.044 | 69.465 | 84.101 |
| 1999 | 18.935 | 10.690 | 31.490 | 1.034 | 26.102 | 45.925 | 68.566 | 64.754 |
| 2000 | 16.129 | 11.684 | 31.266 | 1.012 | 28.030 | 17.680 | 70.155 | 67.381 |
| 2001 | 15.878 | 10.945 | 31.406 | 0.978 | 31.953 | 17.060 | 73.747 | 52.455 |
| 2002 | 19.212 | 11.096 | 35.944 | 0.977 | 30.762 | 69.824 | 78.686 | 69.086 |
| 2003 | 19.968 | 12.999 | 35.497 | 0.997 | 38.074 | 16.776 | 84.520 | 115.760 |
| 2004 | 17.833 | 12.511 | 32.579 | 0.986 | 47.217 | 15.214 | 86.324 | 130.556 |
| 2005 | 20.144 | 11.557 | 33.583 | 1.031 | 43.367 | 51.883 | 87.429 | 105.727 |
| 2006 | 19.081 | 12.405 | 33.875 | 1.029 | 43.515 | 27.054 | 87.195 | 94.477 |
| 2007 | 17.021 | 12.023 | 28.418 | 1.029 | 48.035 | 18.089 | 88.725 | 103.327 |
| 2008 | 17.708 | 10.193 | 27.435 | 1.107 | 44.885 | 10.079 | 85.576 | 113.082 |
| 2009 | 17.732 | 10.371 | 28.684 | 1.172 | 38.807 | 11.759 | 80.075 | 90.547 |
| 2010 | 16.634 | 10.721 | 28.194 | 1.174 | 33.698 | 28.137 | 74.998 | 80.501 |
| 2011 | 14.082 | 10.203 | 27.633 | 1.103 | 33.038 | 18.150 | 70.140 | 66.408 |
| 2012 | 12.392 | 9.463 | 25.515 | 1.013 | 34.946 | 9.825 | 67.981 | 60.697 |
| 2013 | 12.312 | 8.516 | 24.274 | 0.952 | 33.078 | 7.208 | 64.879 | 62.217 |
| 2014 | 11.975 | 8.223 | 22.134 | 0.903 | 28.918 | 4.484 | 59.788 | 113.135 |
| 2015 | 10.200 | 7.512 | 18.804 | 0.845 | 24.450 | 6.930 | 52.626 | 64.175 |
| 2016 | 8.174 | 6.276 | 15.345 | 0.792 | 20.625 | 13.442 | 45.504 | 60.958 |
| 2017 | 6.380 | 5.022 | 12.314 | 0.741 | 18.657 | 5.932 | 40.089 | 52.935 |
| 2018 | 5.500 | 3.975 | 10.766 | 0.711 | 16.967 | 12.628 | 36.765 | 28.805 |
| 2019 | 6.325 | 3.631 | 11.645 | 0.755 | 14.864 | 6.919 | 35.247 | 28.539 |
| 2020 | 6.883 | 4.181 | 13.195 | 0.794 | 13.645 | 7.788 |  |  |
| 2021 | 8.101 | 4.774 | 16.927 | 0.804 | 12.461 | 8.740 | 34.992 | 28.476 |
| 2022 | 8.912 | 6.060 | 15.714 | 1.051 | 11.688 | 12.138 | 36.458 | 36.198 |

## Figures



Figure 1. Current harvest rate strategy (line) for the Bristol Bay red king crab fishery and the associated annual prohibited species catch (PSC) limits (numbers of crab) of Bristol Bay red king crab in the groundfish fisheries in zone 1 in the eastern Bering Sea. Harvest rates are based on current-year estimates of effective spawning biomass (ESB, Zheng et al. 1995b), whereas PSC limits apply to previous-year ESB.

## Data by type and year



Figure 2. Data types and ranges used for the stock assessment.


Figure 3. Retained catch biomass and bycatch mortality biomass ( $t$ ) for Bristol Bay red king crab from 1953 to 2021. Directed pot bycatch data were not available from the observer program before 1990 and are not included in this figure.


Figure 4. Comparison of survey legal male abundances and catches per unit effort for Bristol Bay red king crab from 1968 to 2022.


Figure 5a. Survey abundances by 5-mm carapace length bin for male Bristol Bay red king crab from 1975 to 2022.


Figure 5 b. Survey abundances by 5 mm carapace length bin for female Bristol Bay red king crab from 1975 to 2022.
Figure 6. Holding spot for VAST results in future assessments.


Figure 7a. Comparison of NMFS survey abundance proportions of total NMFS and BSFRF side-byside trawl surveys during 2013-2016 for Bristol Bay red king crab. Sizes of circles are proportional to total abundances.


Figure 7b. Comparison of ratios of NMFS survey abundances to BSFRF side-by-side survey abundances during 2013-2016 for Bristol Bay red king crab. Sizes of circles are proportional to total abundances.


Figure 7c. Comparison of ratios of NMFS survey abundances to BSFRF side-by-side survey abundances during 2013-2016 for Bristol Bay red king crab. Sizes of circles are proportional to total abundances. The abundance-weighted average ratio is 0.891 for $\mathrm{crab} \geq 135 \mathrm{~mm}$ carapace length from all four years of data. The approach to compute this overall ratio is documented in section D. Data, 4. Bering Sea Fisheries Research Foundation Survey Data.


Figure 8a. Estimated NMFS trawl survey selectivities under models 21.1b, 22.0, and 22.0a


Figure 8b. Estimated total pot fishery selectivities and retained proportions and groundfish fisheries bycatch selectivities under model 21.1b.


Figure 8c. Estimated total pot fishery selectivities and retained proportions and groundfish fisheries bycatch selectivities under model 22.0.


Figure 8d. Estimated total pot fishery selectivities and retained proportions and groundfish fisheries bycatch selectivities under model 22.0a.


Figure 9a. Comparison of estimated probabilities of molting of male red king crab in Bristol Bay for different periods with model 21.1b. Molting probabilities for periods 1954-1961 and 19661969 were estimated by Balsiger (1974) from tagging data. Molting probabilities for 1975-1979 and 1980-2022 were estimated with a length-based model.


Figure 9 b . Comparison of estimated probabilities of molting of male red king crab in Bristol Bay for different periods with models 22.0 and 22.0a during 1985-2022. Molting probabilities for periods 1954-1961 and 1966-1969 were estimated by Balsiger (1974) from tagging data.


Figure 10a. Comparisons of area-swept estimates of total NMFS survey biomass and model prediction for model estimates in 2022 under models 21.1, 22.0, and 22.0a. The error bars are plus and minus 2 standard deviations of model 21.1b.


Figure 10b. Comparisons of survey biomass estimates by sex (upper plot for males and lower plot for females) by the BSFRF survey and the model for model estimates in 2022 (models 21.1b, 22.0, 22.0a). The error bars are plus and minus 2 standard deviations of model 21.1b. The BSFRF survey catchability is assumed to be 1.0 for all models.


Figure 10c. Comparisons of estimated BSFRF survey selectivities with models 21.1b, 22.0, and 22.0a. The BSFRF survey catchability is assumed to be 1.0 for all models.


Figure 10d. Comparisons of length compositions by the BSFRF survey and the model estimates during 2007-2008 and 2013-2016 with models 21.1b, 22.0, and 22.0a.


Figure 11a. Estimated absolute mature male biomasses during 1975-2022 for the base model (21.1b) in 2021 and 2022 model years. Mature male biomass is estimated on Feb. 15, year+1.


Figure 11b. Estimated absolute mature male biomasses during 1975-2022 for models 21.1b, 22.0, and 22.0a. Mature male biomass is estimated on Feb. 15, year+1.


Figure 11c. Estimated absolute mature male biomasses during 1985-2022 for models 21.1b, 22.0, and 22.0a. Mature male biomass is estimated on Feb. 15, year+1.


Figure 12a. Estimated recruitment time series during 1976-2022 with models 21.1b, 22.0 and 22.0a. Mean male recruits during 1984-2021 was used to estimate $B_{35 \%}$. Recruitment estimates in the terminal year (2022) are unreliable.


Figure 12b. Estimated recruitment length distributions with models 21.1b, 22.0, and 22.0a.


Figure 13a. Relationships between full fishing mortalities for the directed pot fishery and mature male biomass on Feb. 15 during 1975-2021 under model 21.1b. Average of recruitment from 1984 to 2020 was used to estimate $B_{35 \%}$.


Figure 13b. Relationships between full fishing mortalities for the directed pot fishery and mature male biomass on Feb. 15 during 1985-2021 under model 22.0. Average of recruitment from 1984 to 2020 was used to estimate $B_{35 \%}$.


Figure 13c. Relationships between full fishing mortalities for the directed pot fishery and mature male biomass on Feb. 15 during 1985-2021 under model 22.0a. Average of recruitment from 1986 to 2021 was used to estimate $B_{35 \%}$.


Figure 13d. Comparison of estimated natural mortality and directed pot fishing mortality for models 21.1b, 22.0, and 22.0a.


Figure 14a. Relationships between mature male biomass on Feb. 15 and total recruits at age 5 (i.e., 6 -year time lag) for Bristol Bay red king crab under model 21.1b. Numerical labels are years of mating, and the vertical dotted line is the estimated $B_{35 \%}$ based on the mean recruitment level during 1984 to 2021.


Figure 14b. Relationships between log recruitment per mature male biomass and mature male biomass on Feb. 15 for Bristol Bay red king crab under model 21.1b. Numerical labels are years of mating, and the line is the regression line for data of 1978-2015.


Figure 15a. Average clutch fullness and proportion of empty clutches of newshell (shell conditions 1 and 2) mature female crab $>89 \mathrm{~mm}$ CL from 1975 to 2022 from survey data. Oldshell females were excluded. The blue dashed line is the mean clutch fullness during two periods before 1992 and after 1991.


Figure 15b. Clutch fullness distribution of newshell (shell conditions 1 and 2) mature female crab $>89 \mathrm{~mm}$ CL from 1975 to 2022 from survey data. Oldshell females were excluded.


Figure 16a. Observed (dots) and predicted (lines) RKC catch and bycatch biomass under models 21.1b, 22.0, and 22.0a.


Figure 16b. Observed (dots) and predicted (lines) RKC bycatch biomass from groundfish fisheries and the Tanner crab fishery under models 21.1b, 22.0, and 22.0a. Trawl bycatch biomass was 0 before 1976.


Figure 17. Standardized residuals of NMFS survey biomass under model 21.1b, 22.0 and 22.0a.


Mid-point of size-class (mm)
Figure 18. Comparison of area-swept and model estimated NMFS survey length frequencies of Bristol Bay male red king crab by year under models 21.1b, 22.0, and 22.0a.


Mid-point of size-class (mm)
Figure 19. Comparison of area-swept and model estimated NMFS survey length frequencies of Bristol Bay female red king crab by year under models 21.1b, 22.0, and 22.0a.


Mid-point of size-class (mm)
Figure 20. Comparison of observed and model estimated retained length frequencies of Bristol Bay male red king crab by year in the directed pot fishery under models 21.1b, 22.0, and 22.0a.


Figure 21. Comparison of observer and model estimated total observer length frequencies of Bristol Bay male red king crab by year in the directed pot fishery under models 21.1b, 22.0, and 22.0a.


Figure 22. Comparison of observer and model estimated discarded length frequencies of Bristol Bay female red king crab by year in the directed pot fishery under models $21.1 \mathrm{~b}, 22.0$, and 22.0 aretro.


Mid-point of size-class (mm)
Figure 23a. Comparison of observer and model estimated discarded length frequencies of Bristol Bay male red king crab by year in the groundfish trawl fisheries under models 21.1b, 22.0, and 22.0a.


Mid-point of size-class (mm)
Figure 23b. Comparison of observer and model estimated discarded length frequencies of Bristol Bay female red king crab by year in the groundfish trawl fisheries under models 21.1b, 22.0, and 22.0a.


Figure 24a. Comparison of observer and model estimated discarded length frequencies of Bristol Bay male red king crab by year in the groundfish fixed gear fisheries under models 21.1b, 22.0, and 22.01.


Mid-point of size-class (mm)
Figure 24b. Comparison of observer and model estimated discarded length frequencies of Bristol Bay female red king crab by year in the groundfish fixed gear fisheries under models 21.1b, 22.0, and 22.0a.


Figure 24c. Comparison of observer and model estimated discarded length frequencies of Bristol Bay red king crab by year in the Tanner crab fishery under models 21.1b, 22.0, and 22.0a. Length composition data during 1994-2009 were not used before 2021.

Model 21.1b, Survey Males

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Figure 25a. Residuals of proportions of NMFS survey male red king crab by year and carapace length (mm) under model 21.1b. Green circles are positive residuals, and red circles are negative residuals.

## Model 22.0, Survey Males



Figure 25b. Residuals of proportions of NMFS survey male red king crab by year and carapace length (mm) under model 22.0. Green circles are positive residuals, and red circles are negative residuals.

## Model 22.0a, Survey Males



Figure 25c. Residuals of proportions of NMFS survey male red king crab by year and carapace length (mm) under model 22.0a. Green circles are positive residuals, and red circles are negative residuals.

## Model 21.1b, Survey Females



Figure 26a. Residuals of proportions of NMFS survey female red king crab by year and carapace length ( mm ) under model 21.1 b . Green circles are positive residuals, and red circles are negative residuals.

## Model 22.0, Survey Females



Figure 26b. Residuals of proportions of NMFS survey female red king crab by year and carapace length (mm) under model 22.0. Green circles are positive residuals, and red circles are negative residuals.

## Model 22.0a, Survey Females



Figure 26c. Residuals of proportions of NMFS survey female red king crab by year and carapace length ( mm ) under model 22.0a. Green circles are positive residuals, and red circles are negative residuals.


Figure 27a. Comparison of hindcast estimates of mature male biomass on Feb. 15 of Bristol Bay red king crab from 1975 to 2021 made with terminal years 2012-2022 with model 21.1b. These are results of the 2022 model. Legend shows the terminal year.


Figure 27b. Comparison of hindcast estimates of mature male biomass on Feb. 15 of Bristol Bay red king crab from 1975 to 2021 made with terminal years 2013-2022 with model 22.0. These are results of the 2022 model. Legend shows the terminal year.


Figure 27c. Comparison of hindcast estimates of mature male biomass on Feb. 15 of Bristol Bay red king crab from 1975 to 2021 made with terminal years 2013 to 2022 with model 22.0a. These are results of the 2022 model. Legend shows the terminal year.


Figure 28a. Comparison of hindcast estimates of total recruitment for model 21.1b of Bristol Bay red king crab from 1976 to 2022 made with terminal years 2012-2022. These are results of the model 21.1b. Legend shows the terminal year.


Figure 28b. Evaluation of Bristol Bay red king crab retrospective errors on recruitment estimates as a function of the number of years in the model for model 21.1b.


Figure 28c. Mean ratios of retrospective estimates of recruitments to those estimated in the most recent year (2022) and standard deviations (red line) of the ratios as a function of the number of years in the model for model 21.1b.


Figure 29a. Comparison of estimates of legal male abundance (top) and mature males (bottom) of Bristol Bay red king crab from 1968 to 2021 made with terminal years 2004-2021 with the base models. Model 21.1b is used for 2021. These are results of historical assessments. Legend shows the year in which the assessment was conducted.


Figure 30a. Histogram of estimated mature male biomass on Feb. 15, 2022, under model 21.1b with the MCMC approach.


Figure 30b. Histogram of the 2021 estimated OFL under model 21.1b with the MCMC approach.


Figure 31. Cumulative probabilities of estimated ratios of MMB on Feb. 15, 2023, to corresponding estimated $B_{35 \%}$ values under model 21.1 b with the MCMC approach. Zero probability is below the estimated minimum thresholds.


Figure 32. Projected mature male biomass on Feb. 15 with four fishing mortalities in the directed fishery $-F=0, \mathrm{~F}=0.083, \mathrm{~F}=0.167$, and $\mathrm{F}=0.25$, during 2022-2032. Input parameter estimates are based on model 21.1b. Crab year "2022" represents Feb. 15, 2023. Shaded areas represent a 5\% to 95\% limits.


Figure 33. Cumulative probabilities of estimated ratios of MMB during 2022-2025 to corresponding estimated $B_{35 \%}$ values under model 21.1b with the MCMC approach and four fishing mortality values. Crab year "2022" represents Feb. 15, 2023.


Figure 34. Length frequency distributions of male (top panel) and female (bottom panel) red king crab in Bristol Bay from NMFS trawl surveys during 2016-2022. For purposes of these graphs, abundance estimates are based on area-swept methods.



Figure 35b. Comparisons of NMFS survey area-swept estimates of mature and legal male abundances in Bristol Bay area (BB) and north of Bristol Bay area (North) during 1985-2022.

## Appendix A. Description of GMACS with Bristol Bay Red King Crab Options (mainly from the GMACS document)

## A. Model Description

## a. Population model

The basic dynamics account for growth, mortality, maturity state and shell condition (although most of the equations below do not explicitly refer to maturity state and shell condition). For the case in which shell condition is not distinguished:

$$
\begin{equation*}
\underline{N}_{y, t}^{g}=\left(\left(\mathbf{I}-\mathbf{P}_{y, t-1}^{g}\right)+\mathbf{X}_{y, t-1}^{g} \mathbf{P}_{y, t-1}^{g}\right) \mathbf{S}_{y, t-1}^{g} \underline{N}_{y, t-1}^{g}+\underline{\tilde{R}}_{y, t}^{g} \tag{A.1}
\end{equation*}
$$

where $\underline{N}_{y, t}^{g}$ is the number of animals by size-class of gender $g$ at the start of season $t$ of year $y$, $\mathbf{P}_{y, t}^{g}$ is a matrix with diagonals given by vector of molting probabilities for animals of gender $g$ at the start of season $t$ of year $y, \mathbf{S}_{y, t}^{g}$ is a matrix with diagonals given by the vector of probabilities of surviving for animals of gender $g$ during time-step $t$ of year $y$ (which may be of zero duration):

$$
\begin{equation*}
S_{y, t, l, l}^{g}=\exp \left(-Z_{y, t, l}^{g}\right) \tag{A.2}
\end{equation*}
$$

$\mathbf{X}_{y, t}^{g}$ is the size-transition matrix (probability of growing from one size-class to each of the other size-classes or remains in the same size class) for animals of gender $g$ during season $t$ of year $y$, $\underline{\underline{R}}_{y, t}^{g}$ is the recruitment (by size-class) to gear $g$ during season $t$ of year $y$ (which will be zero except for one season - the recruitment season), and $Z_{y, t, l}^{g}$ is the total mortality for animals of gender $g$ in size-class $l$ during season $t$ of year $y$. Note that mortality is continuous across a time-step.
The initial conditions for the model (i.e., the numbers-at-size at the start of the first year, $y_{1}$ ) is specified with an overall total recruitment multiplied by offsets for each size-class, i.e.:

$$
\begin{equation*}
N_{y_{1}, l}^{g}=R_{\mathrm{Init}} e^{\delta_{y_{1}, l}^{g}} / \sum_{g^{\prime}} \sum_{l^{\prime}} e^{\delta_{y_{1}, l}^{g^{\prime}}} \tag{A.3}
\end{equation*}
$$

The minimum carapace length for both males and females is set at 65 mm , and crab abundance is modeled with a length-class interval of 5 mm . The last length class includes all crab $\geq 160-\mathrm{mm}$ CL for males and $\geq 140-\mathrm{mm}$ CL for females. Thus, length classes/groups are 20 for males and 16 for females.

## b. Recruitment

Recruitment occurs once during each year. Recruitment by sex and size-class is the product of total recruitment, the split of the total recruitment to sex and the assignment of sex-specific recruitment to size-classes, i.e.:

$$
\tilde{R}_{y, t, l}^{g}=\bar{R} e^{\varepsilon_{y}} \begin{cases}\left(1+e^{\phi_{y}}\right)^{-1} p_{l}^{\mathrm{r}, \text { mal }} & \text { if } g=\text { males }  \tag{A.4}\\ \phi_{y}\left(1+e^{\phi_{y}}\right)^{-1} p_{l}^{\mathrm{r} \text { fem }} & \text { if } g=\text { females }\end{cases}
$$

where $\bar{R}$ is median recruitment, $\phi_{y}$ determines the sex ratio of recruitment during year $y$, and $p_{l}^{r, g}$ is the proportion of the recruitment (by gender and year) that recruits to size-class $l$ :

$$
\begin{equation*}
p_{l}^{r, g}=\int_{L_{l}^{\text {ou }}}^{L_{1}^{\text {Li }}} \frac{1}{\Gamma\left(\alpha^{r, g} / \beta^{r, g}\right)}\left(l / \beta^{r, g}\right)^{\left(\left(\alpha^{\left.\left.r, s / / \beta^{r, g}\right)-1\right)}\right.\right.} e^{-l / \beta^{r, g}} d l \tag{A.5}
\end{equation*}
$$

where $\alpha^{r, g}$ and $\beta^{r, g}$ are the parameters that define a gamma function for the distribution of recruits to size-class. Equation A. 5 can be restricted to a subset of size-classes, in which case the results from Equation A. 5 are normalized to sum to 1 over the selected size-classes.

## c. Total mortality / probability of encountering the gear

Total mortality is the sum of fishing mortality and natural mortality, i.e.:

$$
\begin{equation*}
Z_{y, t, l}^{g}=\rho_{y, t}^{\mathrm{M}} M_{y}^{g} \tilde{M}_{l}+\sum_{f} S_{y, t, l}^{f, g}\left(\lambda_{y, t, l}^{f, g}+\Omega_{y, t, l}^{f, g}\left(1-\lambda_{y, t, l}^{f, g}\right)\right) F_{y, t}^{f, g} \tag{A.6}
\end{equation*}
$$

where $\rho_{y, t}^{\mathrm{M}}$ is the proportion of natural mortality that occurs during season $t$ for year $y, M_{y}^{g}$ is the rate of natural mortality for year $y$ for animals of gender $g$ (applies to animals for which $\tilde{M}_{l}=1$ ), $\tilde{M}_{l}$ is the relative natural mortality for size-class $l, S_{y, t l l}^{f, g}$ is the (capture) selectivity for animals of gender $g$ in size-class $l$ by fleet $f$ during season $t$ of year $y, \lambda_{y, t, l}^{f, g}$ is the probability of retention for animals of gender $g$ in size-class $l$ by fleet $f$ during season $t$ of year $y, \Omega_{y, t, l}^{f, g}$ is the mortality rate for discards of gender $g$ in size-class $l$ by fleet $f$ during season $t$ of year $y$, and $F_{y, t}^{f, g}$ is the fullyselected fishing mortality for animals of gender $g$ by fleet $f$ during season $t$ of year $y$.
The probability of encountering the gear (occurs instantaneously) is given by:

$$
\begin{equation*}
\tilde{Z}_{y, t, l}^{g}=\sum S_{y, t, l}^{f, g} F_{y, t}^{f, g} \tag{A.7}
\end{equation*}
$$

Note that Equation A. 7 is computed under the premise that fishing is instantaneous and hence that there is no natural mortality during season $t$ of year $y$. The logarithms of the fully-selected fishing mortalities by season are modelled as:

$$
\begin{gather*}
\ell \mathrm{n} F_{y, t}^{f, \mathrm{mal}}=\ell \mathrm{n} F^{f, \mathrm{mal}}+\xi_{y, t}^{f, \mathrm{mal}}  \tag{A.8}\\
\ell \mathrm{n} F_{y, t}^{f, \mathrm{fem}}=\ell \mathrm{n} F_{y, t}^{f, \mathrm{mal}}+\phi^{f}+\xi_{y, t}^{f, \mathrm{fem}} \tag{A.9}
\end{gather*}
$$

where $F^{f, \text { mal }}$ is the reference fully-selected fishing mortality rate for fleet $f, \phi^{f}$ is the offset between female and male fully-selected fishing mortality for fleet $f$, and $\xi_{y, t}^{f, g}$ are the annual deviation of fully-selected fishing mortality for fleet $f$ (by gender).
Natural mortality can depend on time with blocked natural mortality (individual parameters). This option estimates natural mortality as parameters by block, i.e.:

$$
\begin{equation*}
M_{y}^{g}=e^{\psi_{y}^{g}} \tag{A.10}
\end{equation*}
$$

where $M_{y_{1}}^{g}$ is the rate of natural mortality for gender $g$ for the first year of the model, and $\psi_{y}^{g}$ is the annual change in natural mortality and changes in blocks of years.

It is possible to 'mirror' the values for the $\psi_{y}^{8}$ parameters (between genders and between blocks), which allows male and female natural mortality to be the same, and for natural mortality to be the same for discontinuous blocks (based on Equation A.10). It is also possible to estimate a ratio of natural mortality between genders. The deviations in natural mortality can also be penalized to avoid unrealistic changes in natural mortality to fit 'quirks' in the data.

## d. Landings, discards, total catch

The model keeps track of (and can be fitted to) landings, discards, total catch by fleet in season with continuous mortality:

Landed catch

$$
\begin{gather*}
C_{y, t, l}^{\mathrm{Land}, f, g}=\frac{\lambda_{y, t, l}^{f, g} S_{y, t, l}^{f, g} F_{y, t}^{f, g}}{Z_{y, t, l}^{g}} N_{y, t, l}^{f, g}\left(1-e^{-Z_{y, t, l}^{g}}\right)  \tag{A.11}\\
C_{y, t, l}^{\text {Disc, } f, g}=\frac{\left(1-\lambda_{y, t, l}^{f, g}\right) S_{y, t, l}^{f, g} F_{y, t}^{f, g}}{Z_{y, t, l}^{g}} N_{y, t, l}^{f, g}\left(1-e^{-Z_{y, t, l}^{g}}\right) \tag{A.12}
\end{gather*}
$$

Discards

Total catch

$$
\begin{equation*}
C_{y, t, l}^{\mathrm{Total}, f, g}=\frac{S_{y, t, l}^{f, g} F_{y, t}^{f, g}}{Z_{y, t, l}^{g}} N_{y, t, l}^{f, g}\left(1-e^{-Z_{y, t, l}^{g}}\right) \tag{A.13}
\end{equation*}
$$

Landings, discards, and total catches by fleet can be aggregated over gender (e.g., when fitting to removals reported as gender-combined). Equations A.11-13 are extended naturally for the case in which the population is represented by shell condition and/or maturity status (given the assumption that fishing mortality, retention and discard mortality depend on gender and time, but not on shell condition nor maturity status).

Landings, discards, and total catches by fleet can be reported in numbers (Equations A.11-13) or in terms of weight. For example, the landings, discards, and total catches by fleet, season, year, and gender for the total (over size-class) removals are computed as:

$$
\begin{equation*}
C_{y, t}^{\mathrm{Land}, g, f}=\sum_{l} C_{y, t, l}^{\mathrm{Land}, g, f} w_{y, l}^{g} ; C_{y, t}^{\mathrm{Disc}, g, f}=\sum_{l} C_{y, t, l}^{\mathrm{Disc}, g, f} w_{y, l}^{g} ; C_{y, t}^{\mathrm{Total}, g, f}=\sum_{l} C_{y, t, l}^{\mathrm{Total}, g, f} w_{y, l}^{g} \tag{A.14}
\end{equation*}
$$

where $C_{y, t}^{\mathrm{Land}, g, f}, C_{y, t}^{\mathrm{Disc}, g, f}$, and $C_{y, t}^{\text {Total, }, f, f}$ are respectively the landings, discards, and total catches in weight by fleet, season, year, and gender for the total (over size-class) removals, and $w_{y, l}^{g}$ is the weight of an animal of gender $g$ in size-class $l$ during year $y$.

## e. Selectivity / retention

Selectivity (the probability of encountering the gear) and retention (the probability of being landed given being captured) are logistic function:

$$
\begin{equation*}
\left.S_{l}=1-\left(1+\frac{\exp \left(\left(\bar{L}_{l}-S_{50}\right)\right.}{\sigma^{S}}\right)\right)^{-1} \tag{A.15}
\end{equation*}
$$

where $S_{50}$ is the size corresponding to $50 \%$ selectivity, $\sigma^{s}$ is the "standard deviation" of the selectivity curve, and $\bar{L}_{l}$ is the midpoint of size-class $l$.

It is possible to assume that selectivity for one fleet is the product of two of the selectivity patterns. This option is used to model cases in which one survey (NMFS trawl survey) is located within the footprint of another survey (BSFRF trawl survey).
The options to model retention are the same as those for selectivity, except that it is possible to estimate an asymptotic parameter, which allows discard of animals that would be "fully retained" according to the standard options for (capture) selectivity.

Selectivity and retention can be defined for blocks of contiguous years. Two blocks are used for NMFS survey selectivity (before 1982 and after 1981) due to gear modifications and two blocks are used for the directed pot fishery retention (before 2005 and after 2004) due to the fishery rationalization.

## f. Growth

Growth is a key component of any size-structured model. It is modelled in terms of molt probability and the size-transition matrix (the probability of growing from each size-class to each of the other size-classes, constrained to be zero for sizes less than the current size). Note that the size-transition matrix has entries on its diagonal, which represent animals that molt but do not change size-classes.

## (1) Molt probability

There are two options for modelling the probability of molting as a function of size, $P_{l, l}$ :

- Constant probability ( 1 for females)
- Logistic probability (for males), i.e.:

$$
\begin{equation*}
P_{l, l}=1-\left(1+\exp \left(\left(\bar{L}_{l}-P_{50}\right) / \sigma^{P}\right)\right)^{-1} \tag{A.16}
\end{equation*}
$$

where $P_{50}$ is the size at which the probability of molting is 0.5 , and $\sigma^{s}$ is the "standard deviation" of the molt probability function.

Molt probability is specified by gender and can change in blocks (one block before 1981 and one block after 1980 for males).

## (2) Size-transition

The proportion of animals in size-class $j$ that grow to be in size-class $i\left(X_{i, j}\right)$ can be pre-specified as gamma-distributed size-increments:

$$
\begin{equation*}
X_{i, j}=\int_{L_{j}^{\mathrm{Low}}}^{L_{j i}^{\mathrm{i}}} \frac{1}{\Gamma\left(I_{i} / \tilde{\beta}\right)}\left(\left(l-\bar{L}_{i}\right) / \tilde{\beta}\right)^{\left(I_{i} / \tilde{\beta}\right)-1} e^{-\left(l-\bar{L}_{i}\right) / \tilde{\beta}} d l \tag{A.17}
\end{equation*}
$$

where $I_{i}$ is the 'expected' growth increment for an animal in size-class $i$ (a linear function of the mid-point of size-class $i$ ), $\tilde{\beta}$ determines the variation in growth among individuals, and $L_{j}^{\text {low }}$ and $L_{j}^{\text {li }}$ are respectively the lower and upper bounds of size-class $j$.

The size-transition matrix is specified by gender and can change in blocks (one block for males and three blocks for females (1975-1982, 1983-1993, and 1994-present based on changes in sizes at maturity).

## B. Outputs, Projections and OFL Calculation

## a. Core model outputs

The core model outputs are the N-matrix, the matrix of fully-selected fishing mortalities, the timeseries of spawning stock biomass, mature male biomass (SSB), the values for the model parameters, and the predictions related to the observations. The spawning stock biomass (and hence mature male biomass) is defined according to:

$$
\begin{equation*}
S S B_{y}=\sum_{g} p^{\mathrm{SSB}, g} \sum_{l} N_{y, t^{*}, l}^{g} \tag{A.18}
\end{equation*}
$$

where $p^{\mathrm{SSB}, g}$ is the relative contribution of gender $g$ to spawning biomass ( $p^{\mathrm{SSB}, \mathrm{mal}}=1 ; p^{\mathrm{SSB}, \mathrm{fem}}=0$ corresponds to spawning stock biomass equating to mature male biomass), and $t^{*}$ is the season in which spawning takes place (spawning occurs at the start of the season).
Definition of model outputs:
(1) Biomass: two population biomass measurements are used in this report: total survey biomass (crab >64 mm CL) and mature male biomass (males >119 mm CL). Mating time is assumed to Feb. 15.
(2) Recruitment: new entry of number of males in the 1 st seven length classes ( $65-99 \mathrm{~mm}$ CL) and new entry of number of females in the 1st five length classes ( $65-89 \mathrm{~mm} \mathrm{CL}$ ).
(3) Fishing mortality: full-selected instantaneous annual fishing mortality rate at the time of fishery.

## b. Biological reference points

The key biological reference points are the proxy for $F_{\text {MSY }}$, the proxy for $B_{\text {MSY }}$ and the Overfishing Level (OFL).

## (1) The proxy for $F_{M S Y}$

The specification for the proxy for $F_{\text {MSY }}$ depends on the tier in which the stock is placed. BBRKC belongs to Tier 3, and the proxy for $F_{\mathrm{MSY}}$ is $F_{35 \%}$, the value of a multiplier on the fully-selected fishing mortality rates for directed fisheries in the final year of the assessment such that spawning biomass-per-recruit is $35 \%$ of the unfished level. The fully-selected fishing mortality rates for nondirected fisheries are set to recent averages (recent 5 years for BBRKC). The unfished spawning biomass-per-recruit, $\operatorname{SSBPR}(\underline{0})$, is calculated by projecting the population model forward where fishing mortality is zero for all fleets, and recruitment is constant (and ideally equal to 1 ). $F_{35 \%}$ is then computed (using Newtons' method) such that:

$$
\begin{equation*}
\operatorname{SSBPR}(\underline{\alpha} \overline{\tilde{F}})=0.35 \operatorname{SSBPR}(\underline{0}) \tag{A.19}
\end{equation*}
$$

where $\underline{\bar{F}}$ is the vector of recent average fully-selected fishing mortalities, and $\underline{\alpha}$ is a vector with 1 for the non-directed fisheries and a calculated constant for the directed fisheries.

## (2) The proxy for $B_{M S Y}$

The specification for the proxy for $B_{\text {MSY }}$ depends on the tier in which the stock is placed. For stocks in Tier 4, the proxy for $B_{\text {MSY }}$ is the average spawning stock biomass over a pre-specified number of years. For Tier 3 , the proxy for $B_{\mathrm{MSY}}$ is $0.35 \operatorname{SSBPR(\underline {0})\text {multipliedbythemeanrecruitmentover}}$ a pre-specified number of years. GMACS estimates annual recruitments by sex through estimating annual recruitment deviations and annual recruitment proportions by sex. Pre-specified numbers of years are needed in the control file for recruitment average and for mean recruitment sex ratio, respectively.

## (3) Calculating the OFL

The OFL is the total catch (in weight) encountered by the gear that dies either due to being landed or due to being discarded when fully-selected fishing mortality is computed using the OFL control rule. The total catch

$$
\begin{equation*}
O F L=\sum_{g} \sum_{t} w_{y_{2}, l}^{g} \frac{S_{y_{2}, t, l}^{f, g}\left(\lambda_{y_{2}, t, l}^{f, g}+\Omega_{y_{2}, t, l}^{f, g}\left(1-\lambda_{y_{2}, t, l}^{f, g}\right) S_{y_{2}, t, l}^{f, g}\right) \alpha^{*, f} \bar{F}_{t}^{f, g}}{Z_{y_{2}+1, t, l}^{g}} N_{y_{2}+1, t, l}^{f, g}\left(1-e^{-Z_{y_{2}+1, l, l}^{g}}\right) \tag{A.20}
\end{equation*}
$$

where $y_{2}$ is the final year of the assessment, $\alpha^{*, f}$ is the multiplier on average fully-selected fishing mortality for fleet $f$ ( 1 for non-directed fisheries and a value computed from the OFL control rule for the directed fisheries), $\bar{F}_{t}^{f, g}$ is recent average fully-selected fishing mortality for fleet $f$ and gender $g$ during season $t$, and $Z_{y_{2}+1, t, l}^{g}$ is the total mortality on animals of gender $g$ in size-class $l$ during season $t$ of year $y_{2}+1$ :

$$
\begin{equation*}
Z_{y_{2}+1, t, l}^{g}=\rho_{y_{2}, t}^{\mathrm{M}} M_{y_{2}}^{g} \tilde{M}_{l}+\sum_{f} S_{y_{2}, t, l}^{f, g}\left(\lambda_{y_{2}, t, l}^{f, g}+\Omega_{y_{2}, t, l}^{f, g}\left(1-\lambda_{y_{2}, t, l}^{f, g}\right)\right) \alpha^{*, f} \bar{F}_{t}^{f, g} \tag{A.21}
\end{equation*}
$$

The values for entries of the vector $\alpha^{*}$ for the directed fisheries are determined using the OFL control rule:

- If the projected spawning stock biomass in year $\mathrm{y}_{2}+1$ when $\underline{\alpha}^{*}=\underline{\alpha}$ exceeds the proxy for $B_{\mathrm{MSY}}$, then $\alpha^{*, f}=\alpha^{f}$.
- If the projected spawning stock biomass in year $\mathrm{y}_{2}+1$ when $\underline{\alpha^{*}}=\underline{\alpha}$ is less than $25 \%$ of the proxy for $B_{\mathrm{MSY}}$, then $\alpha^{*, f}=0$.
- If the projected spawning stock biomass in year $y_{2}+1, S S B_{y_{2}}^{*}$ when $\underline{\alpha^{*}}=\underline{\alpha}$ lies between less than $25 \%$ and $100 \%$ of the proxy for $B_{\mathrm{MSY}}$, then $\alpha^{*, f}$ is tuned according to $\alpha^{*, f}=\frac{\alpha^{f}\left(\frac{S S B_{y_{2}}^{*}-0.1}{B_{M S Y}}\right)}{0.9}$ until convergence.


## c. Projections

The specifications for the projections relate to:

- The duration of the projection.
- Whether the fully-selected fishing mortalities for the non-directed fisheries are set to zero or to recent averages by fleet.
- The way in which future recruitment is generated. The options available are:
- Select a recruitment from a set of historical recruitments at random.
- Generate a future recruitment from a Ricker stock-recruitment relationship, i.e.:

$$
\begin{equation*}
R_{y}^{g}=S S B_{y-a^{*}} / S S B_{0} e^{-1.25 \ln h\left(S S B_{y-a^{*}} / S S B_{0}-1\right)} e^{\varepsilon_{y}-\sigma_{k}^{2} / 2} ; \varepsilon_{y} \sim N\left(0 ; \sigma^{2}\right) \tag{A.22}
\end{equation*}
$$

where $a^{*}$ is the time-lag between spawning and entering the first size-class in the model, $S S B_{0}$ is unfished spawning stock biomass, $h$ is the steepness of the stockrecruitment relationship, $\sigma_{R}$ is the variation in recruitment about the stockrecruitment relationship.

- Generate a future recruitment from a Beverton-Holt stock-recruitment relationship, i.e.:

$$
\begin{equation*}
R_{y}^{g}=\frac{4 R_{0} S S B_{y-a^{*}} / S S B_{0}}{(1-h)+(5 h-1) S S B_{y-a^{*}} / S S B_{0}} e^{\varepsilon_{y}-\sigma_{R}^{2} / 2} \quad \varepsilon_{y} \sim N\left(0 ; \sigma^{2}\right) \tag{A.23}
\end{equation*}
$$

where $R_{0}$ is unfished recruitment (i.e.. $S S B_{0} / \operatorname{SSBPR}(\underline{0})$ ).

- The control rule used to set fully-selected fishing mortality for the directed fisheries. The options are available
- Pre-specified values for fully-selected fishing mortality for each fishery.
- Pre-specified values subject to the dead catch not exceeding that corresponding to the OFL.
- Pre-specified values subject to the dead catch not exceeding that corresponding to the OFL and the landed catch not exceeding that corresponding to the State of Alaska harvest control rule.

The value for the steepness of the stock-recruitment relationship is computed such that the maximum sustainable yield occurs at $F_{35 \%}$, i.e.:

$$
\begin{equation*}
\left.\frac{d C(\underline{F})}{d F}\right|_{\underline{\underline{E}=\alpha^{*} \bar{F}}} \tag{A.24}
\end{equation*}
$$

where $C(\underline{F})$ is the equilibrium landed catch when the population model is projected forward deterministically under one of the two stock-recruitment relationships.

## C. Parameter Estimation

## a. Estimating Bycatch Fishing Mortalities for Years without Observer Data

Observer data are not available for the directed pot fishery before 1990 and the Tanner crab fishery before 1991. There are also extremely low observed bycatches in the Tanner crab fishery in 1994 and during 2006-2009. Bycatch fishing mortalities for male and females during 1975-1989 in the directed pot fishery were estimated as
$F_{t}^{d i s c, s}=r^{s} F_{t}^{d i r}$
where $r^{s}$ is the mean ratio of estimated bycatch discard fishing mortalities to the estimated directed pot fishing mortalities during 1990-2004 for sex $s$. Directed pot fishing practice has changed after 2004 due to fishery rationalization.

We used pot fishing effort (potlifts) east of $163^{\circ} \mathrm{W}$ in the Tanner crab fishery to estimate red king crab bycatch discard fishing mortalities in that fishery when observer data are not available (19751990, 1994, 2006-2009):
$F_{t}^{\text {Tanner }, s}=a^{s} E_{t}$
where $a^{s}$ is the mean ratio of estimated Tanner crab fishery bycatch fishing mortalities to fishing efforts during 1991-1993 for sex s, and $E_{t}$ is Tanner crab fishery fishing efforts east of $163^{\circ} \mathrm{W}$ in year $t$. Due to fishery rationalization after 2004, we used the data only during 1991-1993 to estimate the ratio.

## b. Likelihood Components

A maximum likelihood approach was used to estimate parameters. For length compositions ( $p_{l, t, s, s h}$ ), the likelihood functions are :
$R f=\prod_{l=1}^{L} \prod_{t=1}^{T} \coprod_{s=1}^{2} \prod_{s h=1}^{2} \frac{\left\{\exp \left[-\frac{\left(p_{l, t, s, s h}-\hat{p}_{l, t, s, s h}\right)^{2}}{2 \sigma_{l, t, s, s h}^{2}}\right]+0.01\right\}}{\sqrt{2 \pi \sigma_{l, t, s, s h}^{2}}}$
$\sigma_{l, t, s, s h}^{2}=\frac{\left[p_{l, t, s, s h}\left(1-p_{l, t, s, s h}\right)+\frac{0.1}{L}\right]}{n_{t}}$
where $L$ is the number of length groups, $T$ the number of years, and $n_{t}$ the effective sample size in year $t$, which was estimated for trawl survey, pot retained catch, total directed pot male catch, directed pot female discard, groundfish trawl discard, groundfish fixed gear discard, and Tanner crab fishery discard length composition data. $p_{l, t, s, s h}$ is the observed proportion of crab in lengthclass $l$, year $t$, sex $s$ and shell condition $s h$, and $\hat{p}_{l, t, s, s h}$ is the model-estimate corresponding to $p_{l, t, s, s h}$.

The weighted negative log likelihood functions are:

$$
\begin{gathered}
\text { Length compositions: }-\sum \ln \left(R f_{i}\right) \\
\text { Catch and bycatch biomasses: } \sum\left[\ln \left(\frac{C_{t}}{\bar{C}_{t}}\right)^{2} /\left(2 \ln \left(c v_{t}^{2}+1\right)\right)\right] \\
\text { NMFS survey biomass: } \sum\left[\ln \left(\ln \left(C V_{t}^{2}+1\right)\right)^{0.5}+\frac{\ln \left(\frac{B_{t}}{\bar{B}_{t}}\right)^{2}}{\left(2 \ln \left(C V_{t}^{2}+1\right)\right)}\right] \\
\text { BSFRF survey biomass: } \sum\left[\ln \left(\ln \left(C V_{t}^{2}+A V^{2}+1\right)\right)^{0.5}+\frac{\ln \left(\frac{B_{t}}{\bar{B}_{t}}\right)^{2}}{\left(2 \ln \left(C V_{t}^{2}+A V^{2}+1\right)\right)}\right] \\
R \text { variation: } \lambda_{R} \sum\left[\ln \left(\frac{R_{t}}{\bar{R}}\right)^{2}\right] \\
R \text { sex ratio: } \lambda_{s} \sum\left[\ln \left(\frac{\bar{R}_{M}}{\bar{R}_{F}}\right)^{2}\right] \\
\text { Groundfish bycatch fishing mortalities: } \lambda_{t} \sum\left[\ln \left(\frac{F_{t, g f}}{\overline{F_{g f}}}\right)^{2}\right] \\
\text { Pot female bycatch fishing mortalities: } \lambda_{p} \sum\left[\ln \left(\frac{F_{t, f}}{\overline{\bar{F}_{f}}}\right)^{2}\right] \\
\text { Trawl survey catchability: } \frac{(Q-\hat{Q})^{2}}{2 \sigma^{2}}
\end{gathered}
$$

where $R_{t}$ is the recruitment in year $t, \bar{R}$ the mean recruitment, $\bar{R}_{M}$ the mean male recruitment, $\bar{R}_{F}$ the mean female recruitment, $A V$ is additional $C V$ and estimated in the model, $\bar{F}_{g f}$ the mean groundfish bycatch fishing mortality (this is separated into trawl and fixed gear fishery bycatch), $\bar{F}_{f}$ the mean pot female bycatch fishing mortality, $Q$ summer trawl survey catchability, and $\sigma$ the estimated standard deviation of $Q$ (all models).

Weights $\lambda_{j}$ are assumed to be 2 for recruitment variation, 10 for recruitment sex ratio, 0.2 for pot female bycatch fishing mortality, and 0.1 for trawl bycatch fishing mortality. These $\lambda_{j}$ values correspond to CV values of $0.53,0.23,3.34$, and 12.14 , respectively.

## c. Population State in Year 1.

The total abundance and proportions for the first year are estimated in the model.

## d. Parameter estimation framework:

(1) Parameters estimated independently

Basic natural mortality, length-weight relationships, and mean growth increments per molt were estimated independently outside of the model. Mean length of recruits to the model depends on growth and was assumed to be 72.5 for both males and females. Handling mortality rates were set to 0.2 for the directed pot fishery, 0.25 for the Tanner crab fishery, 0.5 for the groundfish fixed gear fishery, and 0.8 for the groundfish trawl fishery.

## i. Natural Mortality

Based on an assumed maximum age of 25 years and the $1 \%$ rule (Zheng 2005), a base $M$ was estimated to be 0.18 for males.

## ii. Length-weight Relationship

Length-weight relationships for males and females were as follows:

$$
\begin{array}{ll}
\text { Immature Females: } & W=0.000408 L^{3.127956} \\
\text { Ovigerous Females: } & W=0.003593 L^{2.666076}  \tag{A.29}\\
\text { Males: } & W=0.0004031 L^{3.141334}
\end{array}
$$

where $W$ is weight in grams, and $L$ CL in mm.

## iii. Growth Increment per Molt

A variety of data are available to estimate male mean growth increment per molt for Bristol Bay RKC. Tagging studies were conducted during the 1950s, 1960s and 1990s, and mean growth increment per molt data from these tagging studies in the 1950s and 1960s were analyzed by Weber and Miyahara (1962) and Balsiger (1974). Modal analyses were conducted for the data during 1957-1961 and the 1990s (Weber 1967; Loher et al. 2001). Mean growth increment per molt may be a function of body size and shell condition and vary over time (Balsiger 1974; McCaughran and Powell 1977); however, for simplicity, mean growth increment per molt was assumed to be only a function of body size in the models. Tagging data were used to estimate mean growth increment per molt as a function of pre-molt length for males (Figure A2). The results from modal analyses of 1957-1961 and the 1990s were used to estimate mean growth increment per molt for immature females during 1975-1993 and 1994-2020, respectively, and the data presented in Gray (1963) were used to estimate those for mature females for model scenarios (Figure A2). To make a smooth transition of growth increment per molt from immature to mature females, weighted growth increment averages of $70 \%$ and $30 \%$ at 92.5 mm CL pre-molt length and $90 \%$ and $10 \%$ at 97.5 mm CL were used, respectively, for mature and immature females during 1983-1993. These percentages are roughly close to the composition of maturity. During 1975-1982, females matured at a smaller size, so the growth increment per molt as a function of length was shifted to smaller increments. Likewise, during 1994-2021, females matured at a slightly higher size, so the growth increment per molt was shifted to high increments for immature crab (Figure A2). Once mature, the growth increment per molt for male crab decreases slightly and annual molting probability decreases, whereas the growth increment for female crab decreases dramatically but annual molting probability remains constant at 1.0 (Powell 1967).

## iv. Sizes at Maturity for Females

The NMFS collected female reproductive condition data during the summer trawl surveys. Mature females are separated from immature females by a presence of egg clutches or egg cases. Proportions of mature females at $5-\mathrm{mm}$ length intervals were summarized and a
logistic curve was fitted to the data each year to estimate sizes at $50 \%$ maturity. Sizes at $50 \%$ maturity are illustrated in Figure A3 with mean values for three different periods (1975-82, 1983-93, and 1994-2021).

## v. Sizes at Maturity for Males

Although size at sexual maturity for Bristol Bay red king crab males has been estimated (Paul et al. 1991), there are no data for estimating size of functional maturity collected in the natural environment. Sizes at functional maturity for Bristol Bay male RKC have been assumed to be 120 mm CL (Schmidt and Pengilly 1990). This is based on mating pair data collected off Kodiak Island (Figure A4). Sizes at maturity for Bristol Bay female RKC are about 90 mm CL, about 15 mm CL less than Kodiak female RKC (Pengilly et al. 2002). The size ratio of mature males to females is 1.3333 at sizes at maturity for Bristol Bay RKC, and since mature males grow at much larger increments than mature females, the mean size ratio of mature males to females is most likely larger than this ratio. Size ratios of the large majority of Kodiak mating pairs were less than 1.3333, and in some bays, only a small proportion of mating pairs had size ratios above 1.3333 (Figure A4).

In the laboratory, male RKC as small as 80 mm CL from Kodiak and Southeast Alaska can successfully mate with females (Paul and Paul 1990). But few males less than 100 mm CL were observed to mate with females in the wild. Based on the size ratios of males to females in the Kodiak mating pair data, setting 120 mm CL as a minimum size of functional maturity for Bristol Bay male RKC is proper in terms of managing the fishery.

## vi. Potential Reasons for High Mortality during the Early 1980s

Bristol Bay red king crab abundance had declined sharply during the early 1980s. Many factors have been speculated for this decline: (i) completely wiped out by fishing: the directed pot fishery, the other directed pot fishery (Tanner crab fishery), and bottom trawling; and (ii) high fishing and natural mortality. With the survey abundance, harvest rates in 1980 and 1981 were among the highest, thus the directed fishing definitely had a big impact on the stock decline, especially legal and mature males. However, for the sharp decline during 1980-1984 for males, 3 out of 5 years had low mature harvest rates. During the 1981-1984 decline for females, 3 out of 4 years had low mature harvest rates. Also pot catchability for females and immature males are generally much lower than for legal males, so the directed pot fishing alone cannot explain the sharp decline for all segments of the stock during the early 1980s.
Red king crab bycatch in the eastern Bering Sea Tanner crab fishery is another potential factor (Griffin et al. 1983). The main overlap between Tanner crab and Bristol Bay red king crab is east of $163^{\circ} \mathrm{W}$. No absolute red king crab bycatch estimates are available until 1991. So there are insufficient data to fully evaluate the impact. Tanner crab retained catch and potlifts from the eastern Bering Sea Tanner crab fishery are illustrated in Figure A5. The observed red king crab bycatch in the Tanner crab fishery during 1991-2015 and total potlifts east of $166^{\circ} \mathrm{W}$ during 1975 to 2015 were used to estimate the bycatch mortality in the current model. Because winter sea surface temperatures and air temperatures were warmer (which means a lower handling mortality rate) and there were fewer potlifts during
the early 1980s than during the early 1990s, bycatch in the Tanner crab fishery is unlikely to have been a main factor for the sharp decline of Bristol Bay red king crab.
Several factors may have caused increases in natural mortality. Crab diseases in the early 1980s were documented by Sparks and Morado (1985), but inadequate data were collected to examine their effects on the stock. Stevens (1990) speculated that senescence may be a factor because many crab in the early 1980s were very old due to low temperatures in the 1960s and early 1970s. The biomass of the main crab predator, Pacific cod, increased about 10 times during the late 1970s and early 1980s. Yellowfin sole biomass also increased substantially during this period. Predation is primarily on juvenile and molting/softshell crab. But we lack stomach samples in shallow waters (juvenile habitat) and during the period when red king crab molt. Also cannibalism occurs during molting periods for red king crab. High crab abundance in the late 1970s and early 1980s may have increased the occurrence of cannibalism.

Overall, the likely causes for the sharp decline in the early 1980s are combinations of the above factors, such as pot fisheries on legal males, bycatch, and predation on females and juvenile and sublegal males, senescence for older crab, and disease for all crab. In our model, we estimated one mortality parameter for males and another for females during 1980-1984. We also estimated a mortality parameter for females during 1976-1979 and 1985-1993. These three mortality parameters are additional to the basic natural mortality of $0.18 \mathrm{yr}^{-1}$, all directed fishing mortality, and non-directed fishing mortality. These three mortality parameters could be attributed to natural mortality as well as undocumented nondirected fishing mortality. The model fit the data much better with these three parameters than without them.
(2) Parameters estimated conditionally

The following model parameters were estimated for male and female crab: total recruits for each year (year class strength $R_{t}$ for $t=1975$ to 2020), total abundance in the first year (1975), growth parameter $\beta$, and recruitment parameter $\beta_{r}$ for males and females separately. Molting probability parameters $\beta$ and $L_{50}$ were also estimated for male crab. Estimated parameters also include different sets of $\beta$ and $L_{50}$ for total selectivity and retained proportions, $\beta$ and $L_{50}$ for pot-discarded female selectivity, $\beta$ and $L_{50}$ for potdiscarded male and female selectivities from the eastern Bering Sea Tanner crab fishery, $\beta$ and $L_{50}$ for groundfish trawl and fixed gear discarded selectivities, and different sets of $\beta$ and $L_{50}$ for NMFS trawl survey male and female selectivities separately. The NMFS survey catchabilities $Q$ for some models were also estimated. Different sets of $\beta$ and $L_{50}$ for selectivity parameters were estimated for the survey data from the Bering Fisheries Research Foundation. Annual fishing mortalities were also estimated for the directed pot fishery for males (1975-2020), pot-discarded females from the directed fishery (19902020), pot-discarded males and females from the eastern Bering Sea Tanner crab fishery (1991-93, 2013-15), groundfish trawl discarded males and females (1976-2020), and groundfish fixed gear discarded males and females (1996-2020). One additional mortality parameter for years 1980-1984 for males and a constant to multiply male natural mortality
for estimating female natural mortality were also estimated. Some estimated parameters were constrained in the model. For example, male and female recruitment estimates were forced to be close to each other for a given year.


Figure A1. Estimated capture probabilities for NMFS Bristol Bay red king crab trawl surveys by Weinberg et al. (2004) and the Bering Sea Fisheries Research Foundation surveys.


Figure A2. Mean growth increments per molt for Bristol Bay red king crab. Note: "tagging"---based on tagging data; "mode"---based on modal analysis. The female growth increments per molt are for different model scenarios.


Figure A3. Estimated sizes at $50 \%$ maturity for Bristol Bay female red king crab from 1975 to 2021. Averages for three periods (1975-82, 1983-93, and 1994-2021) are plotted with a line.


Figure A4. Histograms of carapace lengths (CL) and CL ratios of males to females for male shell ages $\leq 13$ months of red king crab males in grasping pairs; Powell's Kodiak data. Upper plot: all locations and years pooled; middle plot: location 11; lower plot: locations 4 and 13. Sizes at maturity for Kodiak red king crab are about 15 mm larger than those for Bristol Bay red king crab. (Doug Pengilly, ADF\&G, pers. comm.).


Figure A5. Tanner crab retained catch and potlifts for total eastern Bering Sea Tanner crab fishery (upper plot) and the Tanner crab fishery east of $163^{\circ} \mathrm{W}$ (bottom).

## Appendix B. Data files for model 21.1b

## Model 21.1b data file for 2022

\#updating trawl and fixed gear bycatch data for length comp. during 1986-2022 and biomass 2009-2021
\# base data file 21.1b model fall 2022




| 2019 | 3 | 1 | 2 | 218.0 | 0.07 | 0 | 1 | 0 | 0.2 |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2020 | 3 | 1 | 2 | 76.1 | 0.07 | 01 | 1 | 0 | 0.2 |  |  |  |
| 2021 | 3 | 1 | 2 | 29.4 | 0.07 | 0 | 1 | 0 | 0.2 \#update annually |  |  |  |
| \#\# Trawl fishery discards (t, without applying to handling morta |  |  |  |  |  |  |  |  |  |  |  |  |
| \#year | seas |  | fleet sex |  | obs cv | type | units |  | mult | effort discard_mortality |  |  |
| 1976 | 5 | 2 | 0 | 853.494 | 0.10 | 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1977 | 5 | 2 | 0 | 1562.313 | $3 \quad 0.10$ | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1978 | 5 | 2 | 0 | 1650.775 | -0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1979 | 5 | 2 | 0 | 1664.925 | $5 \quad 0.10$ | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1980 | 5 | 2 | 0 | 1295.625 | $5 \quad 0.10$ | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1981 | 5 | 2 | 0 | 274.229 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1982 | 5 | 2 | 0 | 718.610 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1983 | 5 | 2 | 0 | 525.554 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1984 | 5 | 2 | 0 | 1367.550 | $0 \quad 0.10$ | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1985 | 5 | 2 | 0 | 487.576 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1986 | 5 | 2 | 0 | 250.758 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1987 | 5 | 2 | 0 | 233.045 | 0.10 | 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1988 | 5 | 2 | 0 | 747.996 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1989 | 5 | 2 | 0 | 219.023 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1990 | 5 | 2 | 0 | 324.883 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1991 | 5 | 2 | 0 | 436.783 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1992 | 5 | 2 | 0 | 366.816 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1993 | 5 | 2 | 0 | 501.770 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1994 | 5 | 2 | 0 | 109.129 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1995 | 5 | 2 | 0 | 102.623 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1996 | 5 | 2 | 0 | 113.495 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1997 | 5 | 2 | 0 | 71.862 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1998 | 5 | 2 | 0 | 232.580 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 1999 | 5 | 2 | 0 | 188.101 | - 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2000 | 5 | 2 | 0 | 102.161 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2001 | 5 | 2 | 0 | 241.011 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2002 | 5 | 2 | 0 | 189.018 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2003 | 5 | 2 | 0 | 171.114 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2004 | 5 | 2 | 0 | 216.889 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2005 | 5 | 2 | 0 | 155.924 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2006 | 5 | 2 | 0 | 189.660 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2007 | 5 | 2 | 0 | 192.571 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2008 | 5 | 2 | 0 | 170.561 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2009 | 5 | 2 | 0 | 118.672 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2010 | 5 | 2 | 0 | 104.005 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2011 | 5 | 2 | 0 | 70.286 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2012 | 5 | 2 | 0 | 42.641 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2013 | 5 | 2 | 0 | 83.613 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2014 | 5 | 2 | 0 | 43.129 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2015 | 5 | 2 | 0 | 56.410 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2016 | 5 | 2 | 0 | 84.127 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2017 | 5 | 2 | 0 | 114.624 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2018 | 5 | 2 | 0 | 97.561 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2019 | 5 | 2 | 0 | 100.915 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2020 | 5 | 2 | 0 | 100.842 | 0.10 | - 2 | 1 | 1 | 0 | 0.8 |  |  |
| 2021 | 5 | 2 | 0 | 42.990 | 0.10 |  | 1 | 1 | 0 | 0.8 \# update | nually - gf_we | ght? |
| \# Tanner crab fishery discards males |  |  |  |  |  |  |  |  |  |  |  |  |
| \#year | seas |  | fleet sex |  | x obs cv | type | units |  | mult | potlifts | discard_mortality |  |
| 1975 | 5 | 3 | 1 | $0 \quad 0.07$ | (1) | 11 |  | . 445 |  | 0.25 |  |  |
| 1976 | 5 | 3 | 1 | $0 \quad 0.07$ |  | 11 | 233.667 |  |  | 0.25 |  |  |
| 1977 | 5 | 3 | 1 | $0 \quad 0.07$ | . 07 | 11 | 408.437 |  |  | 0.25 |  |  |
| 1978 | 5 | 3 | 1 | $0 \quad 0.07$ | . 07 | 11 | 356.594 |  |  | 0.25 |  |  |
| 1979 | 5 | 3 | 1 | $0 \quad 0.07$ | . 07 | 11 | 476.410 |  |  | 0.25 |  |  |
| 1980 | 5 | 3 | 1 | $0 \quad 0.07$ | 072 | 11 | 496.751 |  |  | 0.25 |  |  |
| 1981 | 5 | 3 | 1 | $0 \quad 0.07$ | 072 | 11 | 322.634 |  |  | 0.25 |  |  |
| 1982 | 5 | 3 | 1 | $0 \quad 0.07$ | 072 | 11 | 192.538 |  |  | 0.25 |  |  |
| 1983 | 5 | 3 | 1 | $0 \quad 0.07$ | . 07 | 11 |  | 546 |  | 0.25 |  |  |
| 1984 | 5 | 3 | 1 | $0 \quad 0.07$ | 72 | 11 |  | 037 |  | 0.25 |  |  |
| \#1985 | 5 | 3 | 1 | $0 \quad 0.07$ | 72 | 11 |  | 001 |  | 0.25 |  |  |
| \#1986 | 5 | 3 | 1 | $0 \quad 0.07$ | 72 | 11 |  | 001 |  | 0.25 |  |  |
| 1987 | 5 | 3 | 1 | $0 \quad 0.07$ | 72 | 11 |  | 827 |  | 0.25 |  |  |
| 1988 | 5 | 3 | 1 | $0 \quad 0.07$ | 72 | 11 |  | 551 |  | 0.25 |  |  |
| 1989 | 5 | 3 | 1 | $0 \quad 0.07$ | 72 | 11 |  | . 175 |  | 0.25 |  |  |
| 1990 | 5 | 3 | 1 | 0.000 | 0.07 | 21 | 1 |  | . 82 | 0.25 |  |  |
| 1991 | 5 | 3 | 1 | 1890.540 | 0.07 | 21 | 1 |  | . 864 | 0.25 |  |  |
| 1992 | 5 | 3 | 1 | 263.854 | 0.07 | 21 | 1 |  | . 922 | 0.25 |  |  |
| 1993 | 5 | 3 | 1 | 118.614 | 0.07 | 21 | 1 |  | . 62 | 0.25 |  |  |


| 1994 | 5 | 3 | 1 | 38.907 | 0.07 | 2 | 1 | 1 | 228.254 | 0.25 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| $\# 1995$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | 201.988 | 0.25 |
| $\# 1996$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | 64.989 | 0.25 |
| $\# 1997$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 1998$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 1999$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 2000$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 2001$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 2002$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 2003$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 2004$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 2005$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| 2006 | 5 | 3 | 1 | 14.334 | 0.07 | 2 | 1 | 1 | 15.273 | 0.25 |
| 2007 | 5 | 3 | 1 | 5.536 | 0.07 | 2 | 1 | 1 | 26.441 | 0.25 |
| 2008 | 5 | 3 | 1 | 9.245 | 0.07 | 2 | 1 | 1 | 19.401 | 0.25 |
| 2009 | 5 | 3 | 1 | 3.089 | 0.07 | 2 | 1 | 1 | 6.635 | 0.25 |
| $\# 2010$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 2011$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 2012$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| 2013 | 5 | 3 | 1 | 37.426 | 0.07 | 2 | 1 | 1 | 16.633 | 0.25 |
| 2014 | 5 | 3 | 1 | 68.588 | 0.07 | 2 | 1 | 1 | 72.768 | 0.25 |
| 2015 | 5 | 3 | 1 | 189.229 | 0.07 | 2 | 1 | 1 | 130.302 | 0.25 |
| $\# 2016$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 2017$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 2018$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 2019$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 2020$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 |
| $\# 2021$ | 5 | 3 | 1 | 0.000 | 0.07 | 2 | 1 | 1 | $1 \mathrm{e}-4$ | 0.25 \# update annually |
|  |  |  |  |  |  |  |  |  |  |  |




\#\# relative abundance data
\#\# Units of Abundance: 1 = biomass, 2 = numbers
\#\# TODO: add column for maturity for terminal molt life-histories
\#\# for BBRKC Units are in 1000 mt.

\#\# Number of relative abundance indicies
2
\# Data type (1=total selectivity; 2=retention*selectivity)
11
\#\# Number of rows in each index
106
\# Survey data (abundance indices, units are 1000 mt
\#Index Year Season Fleet Sex Abundance CV Units

| 1975 | 1 | 5 | 1 | 0 | 133084.0 | 0.193 | 1 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |


| 1976 | 1 | 5 | 1 | 0 | 256362.2 | 0.207 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |


| 1977 | 1 | 5 | 1 | 0 | 232538.7 | 0.144 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

$1978 \quad 1 \quad 5 \quad 1 \quad 0 \quad 199542.2 \quad 0.152 \quad 1 \quad 0$

| 1979 | 1 | 5 | 1 | 0 | 102448.2 | 0.164 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

$1980 \quad 1 \quad 5 \quad 1 \quad 0 \quad 166524.3 \quad 0.221 \quad 1 \quad 0$

| 1981 | 1 | 5 | 1 | 0 | 68294.4 | 0.190 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

19821 | 19 | 1 | 0 | 72296.3 | 0.251 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

| 1983 | 1 | 5 | 1 | 0 | 34761.9 | 0.214 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |


| 1984 | 1 | 5 | 1 | 0 | 96418.3 | 0.606 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

$1985 \quad 1 \quad 5 \quad 1 \quad 0 \quad 26819.4 \quad 0.159 \quad 1 \quad 0$

| 1986 | 1 | 5 | 1 | 0 | 40549.3 | 0.420 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

$1987 \quad 1 \quad 5 \quad 1 \quad 0 \quad 46769.1 \quad 0.209 \quad 1 \quad 0$

| 1988 | 1 | 5 | 1 | 0 | 35373.6 | 0.228 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

$1989 \quad 1 \quad 5 \quad 1 \quad 0 \quad 42357.7 \quad 0.232 \quad 1$

| 1990 | 1 | 5 | 1 | 0 | 38727.8 | 0.242 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

$1991 \quad 1 \quad 5 \quad 1 \quad 0 \quad 66528.0 \quad 0.443 \quad 1$

1992 | 1 | 5 | 1 | 0 | 25096.2 | 0.176 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

| 1993 | 1 | 5 | 1 | 0 | 35670.6 | 0.198 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

$1994 \quad 1 \quad 5 \quad 1 \quad 0 \quad 23002.5 \quad 0.174 \quad 1 \quad 0$
$1995 \quad 1 \quad 5 \quad 1 \quad 0 \quad 27251.9 \quad 0.266 \quad 1 \quad 0$

| 1996 | 1 | 5 | 1 | 0 | 26815.7 | 0.203 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 1997 | 1 | 5 | 1 | 0 | 59638.3 | 0.264 | 1 | 0 |

$1998 \quad 1 \quad 5 \quad 1 \quad 0 \quad 46208.6 \quad 0.182 \quad 1$


| 2 | 2008 | 1 | 6 | 1 | 0 | 67569 | 0.094 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 2 | 2013 | 1 | 6 | 1 | 0 | 68384 | 0.209 | 1 | 0 |
| 2 | 2014 | 1 | 6 | 1 | 0 | 62327 | 0.192 | 1 | 0 |
| 2 | 2015 | 1 | 6 | 1 | 0 | 63709 | 0.161 | 1 | 0 |
| 2 | 2016 | 1 | 6 | 1 | 0 | 34417 | 0.22 | 1 | 0 |
| 2 | 2007 | 1 | 6 | 2 | 0 | 50811 | 0.116 | 1 | 0 |
| 2 | 2008 | 1 | 6 | 2 | 0 | 38472 | 0.094 | 1 | 0 |
| 2 | 2013 | 1 | 6 | 2 | 0 | 26633 | 0.209 | 1 | 0 |
| 2 | 2014 | 1 | 6 | 2 | 0 | 49414 | 0.192 | 1 | 0 |
| 2 | 2015 | 1 | 6 | 2 | 0 | 35244 | 0.161 | 1 | 0 |
| 2 | 2016 | 1 | 6 | 2 | 0 | 43399 | 0.22 | 1 | 0 |


\#Total males

| \#Year | Season | Fleet | Sex | Type | Shell | Maturity | Nsamp | DataVec |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1990 | 3 | 1 | 1 | 0 | 0 | 0 | 127.2 | 0 | 0 | 0.0004 | 0.0028 | 0.0016 | 0.0043 | 0.0024 | 0.013 | 0.0173 | 0.0263 | 0 |
| 1991 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0009 | 0.0038 | 0.0075 | 0.0081 | 0.0092 | 0.0149 | 0.0124 | 0.0241 | 0.0236 | 0.0262 | 0 |
| 1992 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0 | 0.0006 | 0.0008 | 0.0075 | 0.0151 | 0.0375 | 0.0591 | 0.0777 | 0.0806 | 0.0838 | 0 |
| 1993 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0009 | 0.0025 | 0.0032 | 0.0028 | 0.0035 | 0.007 | 0.0177 | 0.0325 | 0.0445 | 0.0615 | 0 |
| 1996 | 3 | 1 | 1 | 0 | 0 | 0 | 32.1 | 0 | 0 | 0 | 0.0047 | 0.0187 | 0.0296 | 0.0265 | 0.0109 | 0.0171 | 0.0249 | 0 |
| 1997 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0 | 0.0001 | 0.0002 | 0.0003 | 0.0006 | 0.0081 | 0.0227 | 0.0446 | 0.0519 | 0.0534 | 0 . |
| 1998 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0001 | 0.0002 | 0.0004 | 0.0021 | 0.0037 | 0.0054 | 0.0056 | 0.0104 | 0.0246 | 0.0588 | 0 |
| 1999 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0 | 0 | 0 | 0.0013 | 0.0013 | 0.0006 | 0.0017 | 0.0013 | 0.0025 | 0.0059 | 0 |
| 2000 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0002 | 0.002 | 0.0071 | 0.0185 | 0.0234 | 0.0242 | 0.0256 | 0.0262 | 0.0254 | 0.0291 | 0 |
| 2001 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0004 | 0.0023 | 0.0037 | 0.005 | 0.0066 | 0.0139 | 0.0249 | 0.0381 | 0.0447 | 0.0539 | 0 |
| 2002 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0017 | 0.0046 | 0.0044 | 0.0051 | 0.0043 | 0.0054 | 0.0066 | 0.0151 | 0.0272 | 0.0504 | 0 |
| 2003 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0034 | 0.0053 | 0.0065 | 0.0144 | 0.0257 | 0.0323 | 0.0355 | 0.0335 | 0.0315 | 0.0322 | 0 |
| 2004 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0001 | 0.0019 | 0.0061 | 0.016 | 0.021 | 0.0231 | 0.0316 | 0.0519 | 0.0613 | 0.0616 | 0 |
| 2005 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0001 | 0.0005 | 0.0008 | 0.0017 | 0.0044 | 0.0128 | 0.0199 | 0.0243 | 0.0264 | 0.0383 | 0 |
| 2006 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0001 | 0.0006 | 0.0019 | 0.0065 | 0.014 | 0.0171 | 0.0166 | 0.0154 | 0.02 | 0.0333 | 0 |
| 2007 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0006 | 0.0021 | 0.0034 | 0.0051 | 0.0089 | 0.0191 | 0.034 | 0.044 | 0.0477 | 0.044 | 0 . |
| 2008 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0001 | 0.0002 | 0.0007 | 0.0026 | 0.0059 | 0.0078 | 0.0088 | 0.0118 | 0.0243 | 0.0445 | 0 |
| 2009 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0002 | 0.0005 | 0.0009 | 0.0016 | 0.0021 | 0.0038 | 0.0093 | 0.0213 | 0.033 | 0.0371 | 0 |
| 2010 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0004 | 0.0006 | 0.0013 | 0.0028 | 0.0044 | 0.0061 | 0.0077 | 0.0113 | 0.0179 | 0.0286 | 0 |
| 2011 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0008 | 0.0031 | 0.0055 | 0.0097 | 0.01 | 0.0089 | 0.0129 | 0.0147 | 0.0193 | 0.0265 | 0 |
| 2012 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0002 | 0.0003 | 0.0008 | 0.0014 | 0.0037 | 0.0088 | 0.0141 | 0.0189 | 0.018 | 0.0192 | 0 |
| 2013 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0001 | 0.0007 | 0.0017 | 0.0022 | 0.0047 | 0.0058 | 0.0096 | 0.015 | 0.0257 | 0.0378 | 0 |
| 2014 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0003 | 0.0006 | 0.0008 | 0.0012 | 0.0017 | 0.0038 | 0.0063 | 0.0111 | 0.0155 | 0.0206 | 0 |
| 2015 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0001 | 0.0002 | 0.0008 | 0.0017 | 0.0038 | 0.0059 | 0.0063 | 0.007 | 0.012 | 0.0271 | 0 |
| 2016 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0001 | 0.0002 | 0.0015 | 0.0034 | 0.0046 | 0.0064 | 0.0111 | 0.0188 | 0.0225 | 0.0279 | 0 |
| 2017 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0002 | 0.0006 | 0.0031 | 0.0115 | 0.0241 | 0.0341 | 0.0294 | 0.0235 | 0.0197 | 0.0248 | 0 |
| 2018 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0.0004 | 0.0027 | 0.0072 | 0.0082 | 0.0067 | 0.011 | 0.0232 | 0.0432 | 0.0643 | 0.0723 | 0 |
| 2019 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0 | 0.0001 | 0.0002 | 0.0019 | 0.0084 | 0.017 | 0.0218 | 0.0194 | 0.0196 | 0.0356 | 0 |
| 2020 | 3 | 1 | 1 | 0 | 0 | 0 | 150 | 0 | 0.0007 | 0.0034 | 0.0075 | 0.0101 | 0.0142 | 0.0177 | 0.03 | 0.0426 | 0.0589 | 0 |
| 2021 | 3 | 1 | 1 | 0 | 0 | 0 | 55 | 0.0009 | 0.0000 | 0.0082 | 0.0127 | 0.0073 | 0.0127 | 0.0145 | 0.0200 | 0.0218 | 0.0418 | 0 |


\#Trawl bycatch male
\#Year Season Fleet Sex Type Shell Maturity Nsamp DataVec




| 2004 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 2004 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0000 | 0.0016 | 0.0016 | 0.0142 | 0.0299 | 0.0377 | 0.0393 | 0.0299 | 0.0535 | 0.0330 | 0.0409 | 0.0220 |
| 2006 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0010 | 0.0058 | 0.0077 | 0.0048 | 0.0087 | 0.0212 | 0.0346 | 0.0500 | 0.0673 | 0.0596 | 0.0260 | 0.0308 | 0.0221 | 0.0192 |
| 2007 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0009 | 0.0009 | 0.0043 | 0.0094 | 0.0068 | 0.0103 | 0.0154 | 0.0231 | 0.0300 | 0.0308 | 0.0342 | 0.0257 |
| 2008 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0016 | 0.0016 | 0.0033 | 0.0139 | 0.0269 | 0.0359 | 0.0359 | 0.0376 | 0.0465 | 0.0563 | 0.0522 | 0.0506 |
| 2009 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0000 | 0.0006 | 0.0063 | 0.0044 | 0.0075 | 0.0169 | 0.0307 | 0.0269 | 0.0263 | 0.0269 | 0.0338 | 0.0194 |
| 2010 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0017 | 0.0120 | 0.0231 | 0.0453 | 0.0427 | 0.0256 | 0.0274 | 0.0342 | 0.0436 | 0.0393 |
| 2011 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0011 | 0.0011 | 0.0011 | 0.0011 | 0.0044 | 0.0122 | 0.0244 | 0.0322 | 0.0322 | 0.0433 | 0.0599 | 0.0511 | 0.0488 | 0.0433 |
| 2012 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0046 | 0.0137 | 0.0091 | 0.0068 | 0.0342 | 0.0091 | 0.0205 | 0.0228 | 0.0433 | 0.0456 | 0.0524 | 0.0342 |
| 2013 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0036 | 0.0320 | 0.0214 | 0.0463 | 0.0142 | 0.0356 | 0.0320 | 0.0285 | 0.0569 | 0.0463 |
| 2014 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0038 | 0.0038 | 0.0038 | 0.0077 | 0.0268 | 0.0153 | 0.0460 | 0.0307 | 0.0268 | 0.0153 | 0.0115 | 0.0115 |
| 2015 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0024 | 0.0024 | 0.0073 | 0.0342 | 0.0293 | 0.0465 | 0.0538 | 0.0318 | 0.0465 | 0.0367 | 0.0293 | 0.0293 | 0.0220 |
| 2016 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0065 | 0.0049 | 0.0016 | 0.0081 | 0.0097 | 0.0097 | 0.0097 | 0.0227 | 0.0373 | 0.0324 | 0.0340 | 0.0243 |
| 2017 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0028 | 0.0028 | 0.0181 | 0.0056 | 0.0070 | 0.0028 | 0.0056 | 0.0070 | 0.0097 | 0.0153 | 0.0153 | 0.0125 |
| 2018 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0045 | 0.0067 | 0.0112 | 0.0078 | 0.0112 | 0.0157 | 0.0347 | 0.0168 | 0.0202 | 0.0246 | 0.0291 | 0.0314 | 0.0325 |
| 2019 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0024 | 0.0024 | 0.0097 | 0.0036 | 0.0085 | 0.0194 | 0.0073 | 0.0109 | 0.0122 | 0.0170 | 0.0182 | 0.0170 | 0.0207 | 0.0182 |
| 2020 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0026 | 0.0026 | 0.0026 | 0.0092 | 0.0052 | 0.0105 | 0.0079 | 0.0131 | 0.0105 | 0.0065 | 0.0131 | 0.0209 | 0.0157 |
| 2021 | 5 | 2 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0020 | 0.0000 | 0.000 | 0.0040 | 0.0060 | 0.0099 | 0.0080 | 0.0219 | 0.0040 | 0.0080 | 0.0119 | 0.0159 | 0.0219 |


| \#Tanner | crab | bycatch | Male |  |  |  | combi |  | it | e norm | ized to | ( be 1) |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| \#Year | Season | Fleet | Sex | Type |  | Shell Mat | turity | Nsamp | DataVec |  |  |  |  |  |  |  |  |  |  |
| 1991 | 5 | 1 | 1 |  | 0 | 0 | 0 | 50 | 0.0026 | 0.0048 | 0.0029 | 0.0042 | 0.0051 | 0.0042 | 0.0102 | 0.0141 | 0.0144 | 0.0112 | 0 |
| 1992 | 5 | 1 | 1 |  | 0 | 0 | 0 | 48.25 | 0.0000 | 0.0000 | 0.0010 | 0.0031 | 0.0114 | 0.0166 | 0.0259 | 0.0238 | 0.0259 | 0.0301 | 0 |
| 1993 | 5 | 1 | 1 |  | 0 | 0 | 0 | 24.85 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0040 | 0.0020 | 0.0262 | 0.0483 | 0.0584 | 0.0664 | 0 |
| 1994 | 5 | 1 | 1 |  | 0 | 0 | 0 | 0.85 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0588 | 0 |
| 2006 | 5 | 1 | 1 |  | 0 | 0 | 0 | 7 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0214 | 0.0500 | 0.0429 | 0.0500 | 0.0786 | 0.0857 | 0 |
| 2007 | 5 | 1 | 1 |  | 0 | 0 | 0 | 2.65 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0189 | 0.0189 | 0 |
| 2008 | 5 | 1 | 1 |  | 0 | 0 | 0 | 7.25 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0069 | 0.0138 | 0.0276 | 0.0897 | 0 |
| 2009 | 5 | 1 | 1 |  | 0 | 0 | 0 | 9.65 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0052 | 0.0155 | 0.0207 |  |
| 2013 | 5 | 1 | 1 |  | 0 | 0 | 0 | 40.7 | 0.0000 | 0.0012 | 0.0000 | 0.0000 | 0.0000 | 0.0086 | 0.0074 | 0.0135 | 0.0184 | 0.0393 | 0 |
| 2014 | 5 | 1 | 1 |  | 0 | 0 | 0 | 31.55 | 0.0000 | 0.0000 | 0.0016 | 0.0000 | 0.0079 | 0.0079 | 0.0127 | 0.0190 | 0.0158 | 0.0317 | 0 |
| 2015 | 5 | 1 | 1 |  | 0 | 0 | 0 | 50 | 0.0017 | 0.0038 | 0.0017 | 0.0024 | 0.0181 | 0.0247 | 0.0178 | 0.0115 | 0.0153 | 0.0205 | 0 |
| \#Tanne | crab | bycatch | $f$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| \#Year | Season | Fleet | Sex | Type |  | Shell Mat | turity | Nsamp | DataVec |  |  |  |  |  |  |  |  |  |  |
| 1991 | 5 | 1 | 2 |  | 0 | 0 | 0 | 0 | 0.0051 | 0.0105 | 0.0096 | 0.0102 | 0.0240 | 0.0326 | 0.0565 | 0.0466 | 0.0827 | 0.1150 | 0 |
| 1992 | 5 | 1 | 2 |  | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0010 | 0.0062 | 0.0228 | 0.0456 | 0.0819 | 0.0933 | 0.0870 | 0.0539 | 0 |
| 1993 | 5 | 1 | 2 |  | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0040 | 0.0342 | 0.0825 | 0.1127 | 0.0805 | 0.0362 | 0 |
| 1994 | 5 | 1 | 2 |  | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0588 | 0.0000 | 0 |
| 2006 | 5 | 1 | 2 |  | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0000 | 0.0071 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0143 | 0 |
| 2007 | 5 | 1 | 2 |  | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0189 | 0.1698 | 0.2264 | 0.2453 | 0.1321 | 0 |
| 2008 | 5 | 1 | 2 |  | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0069 | 0.0069 | 0.0138 | 0.0621 | 0.0552 |  |
| 2009 | 5 | 1 | 2 |  | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0155 | 0.0622 | 0.0674 | 0.0518 | 0 |
| 2013 | 5 | 1 | 2 |  | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0221 | 0.0504 | 0.1806 | 0.1437 | 0.0774 | 0 |
| 2014 | 5 | 1 | 2 |  | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0016 | 0.0032 | 0.0111 | 0.0222 | 0.0475 | 0.0539 | 0.1442 | 0.1537 |  |
| 2015 | 5 | 1 | 2 |  | 0 | 0 | 0 | 0 | 0.0003 | 0.0014 | 0.0028 | 0.0052 | 0.0240 | 0.0348 | 0.0637 | 0.1031 | 0.1445 | 0.1114 | 0 |
| \# Fixed | gear | crab | byc |  | Male |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| \#Year | season | Fleet | Sex | Type |  | Shell Mat | turity | Nsamp | DataVec |  |  |  |  |  |  |  |  |  |  |
| 1996 | 54 | 10 | 0 | 03 | 37.8 | 0.0026 | 0.0013 | 0.0066 | 0.0053 | 0.0026 | 0.0053 | 0.0132 | 0.0132 | 0.0079 | 0.0146 | 0.0146 | 0.0079 | 0.0146 | 0 |
| 1997 | 54 | 10 | 0 | 05 | 50 | 0.0000 | 0.0000 | 0.0024 | 0.0024 | 0.0134 | 0.0284 | 0.0504 | 0.0686 | 0.0654 | 0.0607 | 0.0496 | 0.0315 | 0.0347 | 0 |
| 1998 | 54 | 10 | 0 | 05 | 50 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0019 | 0.0019 | 0.0039 | 0.0077 | 0.0125 | 0.0251 | 0.0367 | 0.0521 | 0 |
| 1999 | 54 | 10 | 0 | 05 | 50 | 0.0031 | 0.0006 | 0.0019 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0025 | 0.0094 | 0.0218 | 0.0524 |  |
| 2000 | 54 | 10 | 0 | 02 | 29.55 | 50.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0085 | 0.0169 | 0.0321 | 0.0271 | 0.0761 | 0.0508 | 0 |
| 2001 | 54 | 10 | 0 | 05 | 50 | 0.0000 | 0.0002 | 0.0006 | 0.0004 | 0.0016 | 0.0044 | 0.0074 | 0.0111 | 0.0201 | 0.0221 | 0.0239 | 0.0233 | 0.0257 | 0 |
| 2002 | 54 | 10 | 0 | 05 | 50 | 0.0000 | 0.0000 | 0.0000 | 0.0003 | 0.0009 | 0.0017 | 0.0003 | 0.0020 | 0.0049 | 0.0111 | 0.0151 | 0.0220 | 0.0305 | 0 |
| 2003 | 54 | 10 | 0 | 05 | 50 | 0.0011 | 0.0000 | 0.0032 | 0.0118 | 0.0150 | 0.0171 | 0.0235 | 0.0107 | 0.0075 | 0.0118 | 0.0128 | 0.0299 | 0.0310 | 0 |
| 2004 | 54 | 10 | 0 | 05 | 50 | 0.0000 | 0.0005 | 0.0018 | 0.0055 | 0.0037 | 0.0092 | 0.0114 | 0.0284 | 0.0307 | 0.0288 | 0.0343 | 0.0247 | 0.0243 |  |
| 2005 | 54 | 10 | 0 | 05 | 50 | 0.0005 | 0.0000 | 0.0014 | 0.0000 | 0.0005 | 0.0042 | 0.0009 | 0.0116 | 0.0075 | 0.0075 | 0.0205 | 0.0266 | 0.0266 | 0 |
| 2006 | 54 | 10 | 0 | 05 | 50 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0005 | 0.0027 | 0.0016 | 0.0070 | 0.0070 | 0.0107 | 0.0161 | 0.0155 | 0 |
| 2007 | 54 | 10 | 0 | 03 | 39.25 | 50.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0013 | 0.0000 | 0.0000 | 0.0013 | 0.0025 | 0.0051 | 0.0051 | 0.0140 | 0.0051 | 0 |
| 2008 | 54 | 10 | 0 | 05 | 50 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0026 | 0.0069 | 0.0172 | 0.0232 | 0.0369 | 0.0378 | 0.0464 | 0 |
| 2009 | 54 | 10 | 0 | 05 | 50 | 0.0000 | 0.0000 | 0.0000 | 0.0000 | 0.0009 | 0.0009 | 0.0009 | 0.0101 | 0.0129 | 0.0129 | 0.0129 | 0.0202 | 0.0395 |  |
| 2010 | 54 | 10 | 0 | 02 | 25.65 | $5 \quad 0.0000$ | 0.0000 | 0.0019 | 0.0000 | 0.0000 | 0.0019 | 0.0019 | 0.0000 | 0.0019 | 0.0039 | 0.0117 | 0.0136 | 0.0273 |  |
| 2011 | 54 | 10 | 0 | 05 | 50 | 0.0000 | 0.0000 | 0.0008 | 0.0017 | 0.0000 | 0.0025 | 0.0017 | 0.0025 | 0.0042 | 0.0025 | 0.0050 | 0.0067 | 0.0076 |  |
| 2012 | 54 | 10 | 0 | 05 | 50 | 0.0000 | 0.0000 | 0.0003 | 0.0007 | 0.0013 | 0.0010 | 0.0047 | 0.0074 | 0.0114 | 0.0138 | 0.0225 | 0.0269 | 0.0316 | 0 |
| 2013 | 54 | 10 | 0 | 05 | 50 | 0.0073 | 0.0097 | 0.0153 | 0.0253 | 0.0210 | 0.0185 | 0.0211 | 0.0215 | 0.0232 | 0.0264 | 0.0275 | 0.0327 | 0.0340 | 0 |
| 2014 | 54 | 10 | 0 | 05 | 50 | 0.0019 | 0.0026 | 0.0040 | 0.0026 | 0.0033 | 0.0054 | 0.0089 | 0.0128 | 0.0121 | 0.0145 | 0.0191 | 0.0238 | 0.0285 | 0 |
| 2015 | 54 | 10 | 0 | 05 | 50 | 0.0007 | 0.0011 | 0.0007 | 0.0022 | 0.0063 | 0.0098 | 0.0107 | 0.0130 | 0.0125 | 0.0192 | 0.0177 | 0.0170 | 0.0150 | 0 |
| 2016 | 54 | 10 | 0 | 05 | 50 | 0.0018 | 0.0032 | 0.0062 | 0.0090 | 0.0192 | 0.0210 | 0.0240 | 0.0291 | 0.0261 | 0.0229 | 0.0247 | 0.0189 | 0.0155 | 0 |
| 2017 | 54 | 10 | 0 | 05 | 50 | 0.0000 | 0.0014 | 0.0000 | 0.0071 | 0.0141 | 0.0148 | 0.0163 | 0.0120 | 0.0071 | 0.0163 | 0.0085 | 0.0120 | 0.0078 | 0 |
| 2018 | 54 | 10 | 0 | 05 | 50 | 0.0009 | 0.0020 | 0.0041 | 0.0080 | 0.0045 | 0.0126 | 0.0242 | 0.0392 | 0.0399 | 0.0470 | 0.0385 | 0.0255 | 0.0201 |  |
| 2019 | 54 | 10 | 0 | $0 \quad 4$ | 43.15 | 50.0000 | 0.0023 | $0.0046 \quad 0$ | 0.01040 | 0.01850 | 0.01970 | 0.02550 | 0.0209 | 0.0209 | 0.01970 | . 0070 | . 01390 | . 0139 |  |
| 2020 | 54 | 10 | 0 | 012. | . 30 | $0.0122 \quad 0.00$ | $0000 \quad 0.0$ | 0.00410 .0 | $0.0000 \quad 0.0$ | 00000.0 | . 0000 0.0 | $0.0000 \quad 0.0$ | 0.0410 .00 | 00810.00 | 0.00 | 0810.0 | 030.0 | 1220.0 |  |

\# Fixed gear crab bycatch female
\#Year Season Fleet Sex Type Shell Maturity Nsamp DataVec

| 1996 | 5 | 4 | 2 | 0 | 0 | 0 | 0 | 0.0066 | 0.0013 | 0.0053 | 0.0040 | 0.0159 | 0.0079 | 0.0238 | 0.0423 | 0.0556 | 0.0860 | 0.1270 | 0.1230 | 0.0847 | 0.0741 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |


| 1997 | 5 | 4 | 2 | 0 | 0 | 0 | 0 | 0.0000 | 0.0000 | 0.0008 | 0.0008 | 0.0047 | 0.0126 | 0.0299 | 0.0260 | 0.0339 | 0.0252 | 0.0165 | 0.0126 | 0.0071 | 0.0071 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

1999
2000
2002
2004






2011
2012
2013


2015

2017

2019
$\begin{array}{lllll}2018 & 5 & 4 & 2 & 0 \\ 2020 & 5 & 4 & 2 & 0 \\ 2021 & 5 & 4 & 2 & 0\end{array}$
\#NMFS males combined

| \#NMFS | males |  | combined |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| \#Year | Season |  | Fleet |  | Type |  |
| 1975 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1976 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1977 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1978 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1979 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1980 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1981 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1982 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1983 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1984 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1985 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1986 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1987 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1988 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1989 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1990 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1991 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1992 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1993 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1994 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1995 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1996 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1997 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1998 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 1999 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2000 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2001 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2002 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2003 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2004 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2005 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2006 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2007 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2008 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2009 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2010 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2011 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2012 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2013 | 1 | 5 | 1 | 0.000 | 0 | 0 |


| 2014 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 2015 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2016 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2017 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2018 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2019 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2021 | 1 | 5 | 1 | 0.000 | 0 | 0 |
| 2022 | 1 | 5 | 1 | 0.0000 | 0 | 0 |

\#NMFS female
\#Year Season Fleet Sex Type Shell Maturity Nsamp DataVec

1978
1979
1980
1981
1982
1983
1984
1985
1986
1987
1988
1989
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1992
1993
1994
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1997
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1999
2000
2001
2002
2003
2005
2006
2007
2008
2010
2011
2012
2013
2015
2016
2017
2018
2019
$\begin{array}{lllllll}2022 & 1 & 5 & 2 & 0.000 & 0 & 0\end{array}$

| 200 | 0.0000 | 0.0005 | 0.0026 | 0.0030 | 0.0160 | 0.0313 | 0.0437 | 0.0348 | 0.0313 | 0.0192 | 0.0231 | 0.0326 | 0.0336 | 0. |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 200 | 0.0105 | 0.0207 | 0.0103 | 0.0093 | 0.0047 | 0.0110 | 0.0158 | 0.0149 | 0.0244 | 0.0187 | 0.0285 | 0.0203 | 0.0235 | 0. |
| 200 | 0.0066 | 0.0009 | 0.0026 | 0.0032 | 0.0041 | 0.0043 | 0.0034 | 0.0083 | 0.0069 | 0.0129 | 0.0085 | 0.0145 | 0.0127 | 0. |
| 200 | 0.0032 | 0.0011 | 0.0029 | 0.0095 | 0.0243 | 0.0199 | 0.0135 | 0.0068 | 0.0083 | 0.0077 | 0.0086 | 0.0134 | 0.0064 | 0. |
| 161 | 0.0051 | 0.0173 | 0.0173 | 0.0153 | 0.0093 | 0.0161 | 0.0144 | 0.0174 | 0.0367 | 0.0160 | 0.0334 | 0.0210 | 0.0033 | 0. |
| 143 | 0.0017 | 0.0036 | 0.0106 | 0.0071 | 0.0071 | 0.0314 | 0.0157 | 0.0244 | 0.0231 | 0.0336 | 0.0299 | 0.0436 | 0.0424 | 0. |
| 142.75 | 0.0038 | 0.0187 | 0.0136 | 0.0140 | 0.0248 | 0.0129 | 0.0183 | 0.0211 | 0.0198 | 0.0245 | 0.0236 | 0.0229 | 0.0384 |  |
| 180 | 0.0000 | 0.0044 | 0.0176 | 0.0143 | 0.0197 | 0.0098 | 0.0256 | 0.0319 | 0.0209 | 0.0241 | 0.0168 | 0.0295 | 0.0402 |  |

$\begin{array}{ccccccccccccccc}200 & 0.0000 & 0.0005 & 0.0026 & 0.0030 & 0.0160 & 0.0313 & 0.0437 & 0.0348 & 0.0313 & 0.0192 & 0.0231 & 0.0326 & 0.0336 & 0 . \\ 200 & 0.0105 & 0.0207 & 0.0103 & 0.0093 & 0.0047 & 0.0110 & 0.0158 & 0.0149 & 0.0244 & 0.0187 & 0.0285 & 0.0203 & 0.0235 & 0 . \\ 200 & 0.0066 & 0.0009 & 0.0026 & 0.0032 & 0.0041 & 0.0043 & 0.0034 & 0.0083 & 0.0069 & 0.0129 & 0.0085 & 0.0145 & 0.0127 & 0 . \\ 200 & 0.0032 & 0.0011 & 0.0029 & 0.0095 & 0.0243 & 0.0199 & 0.0135 & 0.0068 & 0.0083 & 0.0077 & 0.0086 & 0.0134 & 0.0064 & 0 . \\ 161 & 0.0051 & 0.0173 & 0.0173 & 0.0153 & 0.0093 & 0.0161 & 0.0144 & 0.0174 & 0.0367 & 0.0160 & 0.0334 & 0.0210 & 0.0033 & 0 . \\ 143 & 0.0017 & 0.0036 & 0.0106 & 0.0071 & 0.0071 & 0.0314 & 0.0157 & 0.0244 & 0.0231 & 0.0336 & 0.0299 & 0.0436 & 0.0424 & 0 . \\ 142.75 & 0.0038 & 0.0187 & 0.0136 & 0.0140 & 0.0248 & 0.0129 & 0.0183 & 0.0211 & 0.0198 & 0.0245 & 0.0236 & 0.0229 & 0.0384 \\ 180 & 0.0000 & 0.0044 & 0.0176 & 0.0143 & 0.0197 & 0.0098 & 0.0256 & 0.0319 & 0.0209 & 0.0241 & 0.0168 & 0.0295 & 0.0402\end{array}$

$\begin{array}{lllllllllllllllllllllll}1975 & 1 & 5 & 2 & 0.000 & 0 & 0 & 0 & 0.0331 & 0.0401 & 0.0481 & 0.0494 & 0.0564 & 0.0439 & 0.0444 & 0.0454 & 0.0326 & 0.0289 & 0.0162 & 0.0158 & 0.0116 & 0 .\end{array}$ $\begin{array}{lllllllllllllllllllllllll}1976 & 1 & 5 & 2 & 0.000 & 0 & 0 & 0 & 0.0029 & 0.0092 & 0.0313 & 0.0563 & 0.0688 & 0.0628 & 0.0494 & 0.0269 & 0.0121 & 0.0137 & 0.0066 & 0.0049 & 0.0023 & 0\end{array}$ | 1977 | 1 | 5 | 2 | 0.000 | 0 | 0 | 0 | 0.0026 | 0.0068 | 0.0079 | 0.0193 | 0.0337 | 0.0701 | 0.0808 | 0.0715 | 0.0453 | 0.0435 | 0.0415 | 0.0316 | 0.0151 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | $\begin{array}{llllllllllllll}0.0026 & 0.0068 & 0.0079 & 0.0193 & 0.0337 & 0.0701 & 0.0808 & 0.0715 & 0.0453 & 0.0435 & 0.0415 & 0.0316 & 0.0151 & 0.0 \\ 0.0060 & 0.0111 & 0.0187 & 0.0201 & 0.0233 & 0.0418 & 0.0920 & 0.1212 & 0.0791 & 0.0440 & 0.0301 & 0.0267 & 0.0176 & 0 .\end{array}$ $\begin{array}{lllllllllllllll}0.0286 & 0.0154 & 0.0121 & 0.0147 & 0.0148 & 0.0230 & 0.0381 & 0.0734 & 0.0922 & 0.0876 & 0.0565 & 0.0336 & 0.0215 & 0 .\end{array}$ $\begin{array}{lllllllllllll}0.0048 & 0.0219 & 0.0322 & 0.0292 & 0.0597 & 0.0820 & 0.0487 & 0.0581 & 0.0540 & 0.0424 & 0.0315 & 0.0130 & 0.0110 \\ 0.0152 & 0.0113 & 0.0151 & 0.0190 & 0.0366 & 0.0456 & 0.0443 & 0.0472 & 0.0600 & 0.0774 & 0.0804 & 0.0510 & 0.0252\end{array} \quad 0$. $\begin{array}{lllllllllllll}0.0152 & 0.0113 & 0.0151 & 0.0190 & 0.0366 & 0.0456 & 0.0443 & 0.0472 & 0.0600 & 0.0774 & 0.0804 & 0.0510 & 0.0252 \\ 0.0536 & 0.0954 & 0.0603 & 0.0378 & 0.0423 & 0.0482 & 0.0398 & 0.0232 & 0.0190 & 0.0257 & 0.0281 & 0.0203 & 0.0114\end{array} 0$. $\begin{array}{lllllllllllll}0.0536 & 0.0954 & 0.0603 & 0.0378 & 0.0423 & 0.0482 & 0.0398 & 0.0232 & 0.0190 & 0.0257 & 0.0281 & 0.0203 & 0.0114 \\ 0.0174 & 0.0383 & 0.0475 & 0.0629 & 0.0647 & 0.0398 & 0.0341 & 0.0152 & 0.0107 & 0.0042 & 0.0090 & 0.0056 & 0.0061\end{array} \quad 0$. $\begin{array}{llllllllllllll}0.0174 & 0.0585 & 0.1229 & 0.1105 & 0.0647 & 0.0325 & 0.0159 & 0.0119 & 0.0038 & 0.0017 & 0.0000 & 0.0004 & 0.0001 & 0 .\end{array}$ $\begin{array}{lllllllllllll}0.0174 & 0.0585 & 0.1229 & 0.115 & 0.0647 & 0.0325 & 0.015 & 0.0119 & 0.0038 & 0.0017 & 0.0000 & 0.002 & 0.0155 \\ 0.0009 & 0.0377 & 0.0521 & 0.0643 & 0.0555 & 0.0516 & 0.0397 & 0.0161 & 0.0068 & 0.0000 & 0.0000 & 0.0015 & 0\end{array}$ $\begin{array}{llllllllllllll}0.0124 & 0.0224 & 0.0355 & 0.0274 & 0.0263 & 0.0313 & 0.0362 & 0.0388 & 0.0274 & 0.0113 & 0.0072 & 0.0008 & 0.0000 & 0.0 \\ 0.0013 & 0.0124 & 0.0525 & 0.0918 & 0.0761 & 0.0462 & 0.0445 & 0.0569 & 0.0414 & 0.0292 & 0.0179 & 0.0079 & 0.0018 & 0 .\end{array}$ $\begin{array}{llllllllllllll}0.0006 & 0.0076 & 0.0064 & 0.0062 & 0.0139 & 0.0695 & 0.0910 & 0.0979 & 0.0697 & 0.0600 & 0.0407 & 0.0184 & 0.0077 & 0 \\ 0.0017 & 0.0000 & 0.0017 & 0.0082 & 0.0310 & 0.0740 & 0.0646 & 0.0692 & 0.0531 & 0.0376 & 0.0315 & 0.0194 & 0.0064 & 0\end{array}$ $\begin{array}{llllllllllllll}0.0017 & 0.0000 & 0.0017 & 0.0082 & 0.0310 & 0.0740 & 0.0646 & 0.0692 & 0.0531 & 0.0376 & 0.0315 & 0.0194 & 0.0064 & 0.0 \\ 0.0041 & 0.0052 & 0.0235 & 0.0513 & 0.0525 & 0.0071 & 0.0256 & 0.0601 & 0.0732 & 0.0708 & 0.0633 & 0.0410 & 0.0215 & 0 .\end{array}$ $\begin{array}{lllllllllllll}0.0042 & 0.0115 & 0.0196 & 0.0320 & 0.0218 & 0.0344 & 0.0343 & 0.0310 & 0.0366 & 0.0329 & 0.0281 & 0.0431 & 0.0232 \\ 0.0000 & 0.0053 & 0.0074 & 0.0197 & 0.0364 & 0.0414 & 0.0625 & 0.0448 & 0.0353 & 0.0273 & 0.0450 & 0.0407 & 0.0265\end{array}$ $\begin{array}{lllllllllllll}0.0000 & 0.0053 & 0.0074 & 0.0197 & 0.0364 & 0.0414 & 0.0625 & 0.0448 & 0.0353 & 0.0273 & 0.0450 & 0.0407 & 0.0265 \\ 0.0066 & 0.0080 & 0.0175 & 0.0085 & 0.0131 & 0.0248 & 0.0437 & 0.0647 & 0.0639 & 0.0269 & 0.0300 & 0.0268 & 0.0271\end{array} \quad 0$.

 $\begin{array}{llllllllllllll}0.0294 & 0.0482 & 0.0316 & 0.0145 & 0.0139 & 0.0182 & 0.0163 & 0.0254 & 0.0234 & 0.0334 & 0.0272 & 0.0234 & 0.0240 & 0 . \\ 0.0260 & 0.0219 & 0.0436 & 0.0794 & 0.0796 & 0.0436 & 0.0226 & 0.0218 & 0.0245 & 0.0202 & 0.0161 & 0.0285 & 0.0244 & 0 .\end{array}$ $\begin{array}{lllllllllllllll}0.0260 & 0.0219 & 0.0436 & 0.0794 & 0.0796 & 0.0436 & 0.0226 & 0.0218 & 0.0245 & 0.0202 & 0.0161 & 0.0285 & 0.0244 & 0.0 \\ 0.0004 & 0.0037 & 0.0016 & 0.0020 & 0.0146 & 0.0791 & 0.0969 & 0.0616 & 0.0212 & 0.0137 & 0.0095 & 0.0146 & 0.0143 & 0 .\end{array}$ $\begin{array}{llllllllllllll}0.0145 & 0.0196 & 0.0101 & 0.0088 & 0.0111 & 0.0116 & 0.0303 & 0.1040 & 0.1153 & 0.0594 & 0.0303 & 0.0252 & 0.0225 & 0 . \\ 0.0243 & 0.0169 & 0.0125 & 0.0115 & 0.0044 & 0.0055 & 0.0093 & 0.0164 & 0.0512 & 0.0800 & 0.0583 & 0.0358 & 0.0340 & 0 .\end{array}$ $\begin{array}{llllllllllllll}0.0243 & 0.0169 & 0.0125 & 0.0115 & 0.0044 & 0.0055 & 0.0093 & 0.0164 & 0.0512 & 0.0800 & 0.0583 & 0.0358 & 0.0340 & 0 . \\ 0.0018 & 0.0067 & 0.0269 & 0.0403 & 0.0357 & 0.0272 & 0.0255 & 0.0226 & 0.0358 & 0.0524 & 0.0676 & 0.0603 & 0.0419 & 0 .\end{array}$
 $\begin{array}{lllllllllllll}0.0506 & 0.0769 & 0.0485 & 0.0247 & 0.0222 & 0.0176 & 0.0225 & 0.0520 & 0.0399 & 0.0296 & 0.0163 & 0.0206 & 0.0205 \\ 0.0163 & 0.0059 & 0.0143 & 0.0314 & 0.0414 & 0.0464 & 0.0239 & 0.0292 & 0.0351 & 0.0533 & 0.0526 & 0.0356 & 0.0219\end{array} \quad 0$. $\begin{array}{llllllllllllll}0.0163 & 0.0059 & 0.0143 & 0.0314 & 0.0414 & 0.0464 & 0.0239 & 0.0292 & 0.0351 & 0.0533 & 0.0526 & 0.0356 & 0.0219 & 0 . \\ 0.0279 & 0.0327 & 0.0194 & 0.0132 & 0.0199 & 0.0369 & 0.0577 & 0.0514 & 0.0334 & 0.0204 & 0.0196 & 0.0232 & 0.0184 & 0 \text {. }\end{array}$ $\begin{array}{llllllllllllll}0.0405 & 0.0561 & 0.0457 & 0.0116 & 0.0099 & 0.0336 & 0.0386 & 0.0521 & 0.0567 & 0.0468 & 0.0336 & 0.0383 & 0.0347 & 0 . \\ 0.0143 & 0.0139 & 0.0198 & 0.0425 & 0.0615 & 0.0462 & 0.0254 & 0.0259 & 0.0481 & 0.0656 & 0.0619 & 0.0415 & 0.0301 & 0 .\end{array}$ $\begin{array}{llllllllllllll}0.0015 & 0.0023 & 0.0064 & 0.0078 & 0.0155 & 0.0356 & 0.0574 & 0.0560 & 0.0325 & 0.0570 & 0.0614 & 0.0641 & 0.0459 & 0 .\end{array}$ $\begin{array}{lllllllllllll}0.0015 & 0.0023 & 0.0064 & 0.0078 & 0.0155 & 0.0356 & 0.0574 & 0.0560 & 0.0325 & 0.0570 & 0.0614 & 0.0641 & 0.0459 \\ 0.0000 & 0.0027 & 0.0054 & 0.0136 & 0.0116 & 0.0167 & 0.0303 & 0.0570 & 0.0724 & 0.0560 & 0.0555 & 0.0562 & 0.0575 \\ 0.0\end{array}$. $\begin{array}{llllllllllllll}0.0005 & 0.0019 & 0.0050 & 0.0055 & 0.0081 & 0.0122 & 0.0206 & 0.0466 & 0.0656 & 0.0866 & 0.0645 & 0.0603 & 0.0523 & 0 \\ 0.0018 & 0.0006 & 0.0037 & 0.0048 & 0.0069 & 0.0116 & 0.0213 & 0.0365 & 0.0565 & 0.0927 & 0.0955 & 0.0700 & 0.0509 & 0\end{array}$ $\begin{array}{lllllllllllll}0.0058 & 0.0085 & 0.0092 & 0.0141 & 0.0284 & 0.0310 & 0.0384 & 0.0484 & 0.0299 & 0.0530 & 0.0637 & 0.0905 & 0.0635\end{array} 0$ $\begin{array}{llllllllllllll}0.0293 & 0.0180 & 0.0191 & 0.0250 & 0.0281 & 0.0461 & 0.0351 & 0.0220 & 0.0331 & 0.0355 & 0.0365 & 0.0461 & 0.0663 & 0 . \\ 0.0008 & 0.0027 & 0.0093 & 0.0112 & 0.0067 & 0.0125 & 0.0202 & 0.0384 & 0.0429 & 0.0450 & 0.0304 & 0.0302 & 0.0455 & 0 .\end{array}$ $\begin{array}{lllllllllllllll}0.0000 & 0.0000 & 0.0012 & 0.0040 & 0.0091 & 0.0258 & 0.0219 & 0.0320 & 0.0499 & 0.0770 & 0.0569 & 0.0456 & 0.0307 & 0\end{array}$ $\begin{array}{llllllllllllll}0.0074 & 0.0129 & 0.0110 & 0.0055 & 0.0120 & 0.0114 & 0.0107 & 0.0234 & 0.0408 & 0.0461 & 0.0616 & 0.0668 & 0.0531 & 0 .\end{array}$ $\begin{array}{llllllllllllll}0.0120 & 0.0019 & 0.0036 & 0.0043 & 0.0026 & 0.0051 & 0.0143 & 0.0141 & 0.0390 & 0.0714 & 0.0782 & 0.1023 & 0.0737 & 0.0 \\ 0.0010 & 0.0028 & 0.0030 & 0.0126 & 0.0258 & 0.0248 & 0.0167 & 0.0188 & 0.0214 & 0.0511 & 0.0665 & 0.0804 & 0.0885 & 0 .\end{array}$ $\begin{array}{llllllllllllll}0.0031 & 0.0109 & 0.0172 & 0.0186 & 0.0094 & 0.0198 & 0.0516 & 0.0362 & 0.0421 & 0.0296 & 0.0254 & 0.0652 & 0.0462 & 0 .\end{array}$ $\begin{array}{lllllllllllll}0.0017 & 0.0105 & 0.0018 & 0.0070 & 0.0070 & 0.0140 & 0.0143 & 0.0174 & 0.0312 & 0.0355 & 0.0335 & 0.0279 & 0.0515 \\ 0.0107 & 0.0051 & 0.0100 & 0.0121 & 0.0033 & 0.0000 & 0.0120 & 0.0356 & 0.0296 & 0.0189 & 0.0224 & 0.0309 & 0.0446\end{array} 0$. $\begin{array}{llllllllllllll}0.0107 & 0.0051 & 0.0100 & 0.0121 & 0.0033 & 0.0000 & 0.0120 & 0.0356 & 0.0296 & 0.0189 & 0.0224 & 0.0309 & 0.0446 & 0 . \\ 0.0100 & 0.0146 & 0.0170 & 0.0288 & 0.0125 & 0.0222 & 0.0360 & 0.0418 & 0.0440 & 0.0257 & 0.0182 & 0.0310 & 0.0236 & 0 .\end{array}$
\#BSFRF males
\#Year Season Fleet Sex Type Shell Maturity Nsamp DataVec


 2014 $105.750 \quad 0 \quad 0.003 \quad 0.0101 \quad 0.0118 \quad 0.0448 \quad 0.0546 \quad 0.0423 \quad 0.047 \quad 0.0164 \quad 0.0221 \quad 0.0321 \quad 0.0226 \quad 0.0369 \quad 0$.

\#BSFRF females
\#Year Season Fleet Sex Type Shell Maturity Nsamp DataVec







```
## Growth data
# Type of growth increment (1=growth increment with a CV;2=size-at-release; size-at)
O
# nobs_growth
## Note SM used loewss regression for males BBRKC data
## and cubic spine to interpolate 3 sets of female BBRKC data
# MidPoint Sex Increment CV
#67.5 2 14.766667 1000000000000000000000
# MidPoint Sex MidPoint Time-at-liberty Size-trans matrix Number of points
# Release Recapture
## eof
9999
```


## Model 21.1b control file for 2022



| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 5 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 6 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 7 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 8 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 9 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 10 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 | \# | \# Deviation for size-class 11 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 12 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 13 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 | \# | Deviation for size-class 14 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 15 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 16 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 | \# | \# Deviation for size-class 17 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 | \# | Deviation for size-class 18 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 19 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 20 |
| 0.425 | 2053 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 1 |
| 2.268 | 2660 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 2 |
| 1.810 | 3080 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 3 |
| 1.370 | 5111 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 4 |
| 1.158 | 87990 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 5 |
| 0.596 | 84439 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 6 |
| 0.225 | 1257 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 7 |
| -0.02 | 7565368 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 8 |
| -0.21 | 895269 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 9 |
| -0.56 | 77780 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 10 |
| -0.97 | 300021 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 11 |
| -1.24 | 2031 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 12 |
| -1.49 | 7450 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 13 |
| -1.94 | 1253 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 14 |
| -2.05 | 60679 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 15 |
| -1.94 | 06430 | -10 | 4 | 9 |  | 10.0 | 20.00 | \# Deviation for size-class 16 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 17 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 18 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 19 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 20 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 1 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 2 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 3 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 4 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 5 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 6 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 7 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 8 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 9 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 10 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 11 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 | \# | \# Deviation for size-class 12 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 13 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | Deviation for size-class 14 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 15 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 16 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 17 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 18 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 19 |
| -100.00 | -101 | 5 | -2 | 0 | 10.0 | 20.00 |  | \# Deviation for size-class 20 |

\# weight-at-length input method (1 = allometry [w_1 = a*l^b], $2=$ vector by sex)
2
\#\# Males
$\begin{array}{lllllllllllllllllllll}0.000224781 & 0.000281351 & 0.000346923 & 0.000422209 & 0.000507927 & 0.000604802 & 0.000713564 & 0.00083495 & 0.0009697 & 0.00111856 & 0.00128229 & 0.00146163 & 0 .\end{array}$
\#\# Females
$\begin{array}{llllllllllllllll}0.0002151 & 0.00026898 & 0.00033137 & 0.00040294 & 0.00048437 & 0.00062711 & 0.0007216 & 0.00082452 & 0.00093615 & 0.00105678 & 0.00118669 & 0.00132613 & 0 .\end{array}$
\# Proportion mature by sex

| 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 0 | 0 | 0 | 0 | 0 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
| \# | Proportion | legal | by | sex |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
| 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |



```
## GROWTH PARAMETER CONTROLS
##
## Two lines for each parameter if split sex, one line if not ##
```




```
3
# growth increment model (1=alpha/beta; 2=estimated by size-class;3=pre-specified/emprical)
3
# molt probability function (0=pre-specified; 1=flat;2=declining logistic)
2
# Maximum size-class for recruitment(males then females)
75
## number of size-increment periods
1 3
## Year(s) size-incremnt period changes (blank if no changes)
1983 1994
## number of molt periods
22
## Year(s) molt period changes (blank if no changes)
1980 1980
## Beta parameters are relative (1=Yes;0=no)
1
```





| 12.2 | 0 | 20 | -33 | 0 | 0 | 999 |  | \# Females |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 10.5 | 0 | 20 | -33 | 0 | 0 | 999 |  | \# Females |  |
| 8.90 | 20 | -33 | 0 | 0 | 999 |  | \# F | Females |  |
| 7.90 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 7.20 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 6.60 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 6.10 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 5.60 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 5.10 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 4.60 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 4.10 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 3.60 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 3.20 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 2.70 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 2.20 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 1.70 | 20 | -33 | 0 | 0 | 999 |  | \# F | Females |  |
| 1.20 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 0.70 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 0.0 | -1. |  | 1.0 | -7 | 0 | 0 | 999 | 9 \# Females | (beta) |
| \#1.38 |  | 0.5 | 3.7 | -7 | 0 | 0 | 999 | 9 \# Females | (beta) |
| 15.1 | 0 | 20 | -33 | 0 | 0 | 999 |  | \# Females |  |
| 140 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 12.9 | 0 | 20 | -33 | 0 | 0 | 999 |  | \# Females |  |
| 11.8 | 0 | 20 | -33 | 0 | 0 | 999 |  | \# Females |  |
| 10.6 | 0 | 20 | -33 | 0 | 0 | 999 |  | \# Females |  |
| 8.70 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 7.40 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 6.60 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 6.10 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 5.60 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 5.10 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 4.60 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 4.10 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 3.60 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 3.20 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 2.70 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 2.20 | 20 | -33 | 0 | 0 | 999 |  | \# F | Females |  |
| 1.70 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 1.20 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 0.70 | 20 | -33 | 0 | 0 | 999 |  |  | Females |  |
| 0.0 | -1. |  | 1.0 | -7 | 0 | 0 | 999 | 9 \# Females | (beta) |
| \#1.38 |  | 0.5 | 3.7 | -7 | 0 | 0 | 999 | 9 \# Females | (beta) |




| \#\# LEGEND |  |  |
| :--- | :--- | :--- |
| $\# \#$ | sel type: $0=$ parametric, $1=$ coefficients (NIY), $2=$ logistic, $3=$ logistic95, |  |
| \#\# | $4=$ double normal (NIY) | \#\# |
| \#\# | gear index: use +ve for selectivity, -ve for retention | \#\# |
| $\# \#$ | sex dep: 0 for sex-independent, 1 for sex-dependent |  |


\#\# Gear-1 Gear-2 Gear-3 Gear-4 Gear-5 Gear-6 \#\# PotFshry TrawlByc TCFshry FixedGr nMFS BSFRF



 \# Gear-1

|  | 1 | 1 | 1 | 1 | 125.0000 | 5 | 190 | 0 | 1 | 999 | 419 | 975 | 2021 \# | 4 |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 2 | 1 | 8.0 | 0.1 | 20 | 0 | 1 | 999 | 419 | 975 | 2021 \# | 4 |  |  |
| \# Gear-1 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| \# | 1 | 1 | 1 | 1 | 67.5 | 0 | 200 | 0 | 1 | 999 | -999 | 1975 | 2018 | \#4 |  | \#parameters for cubic spine |
| \# | 1 | 2 | 2 | 1 | 87.5 | 0 | 200 | 0 | 1 | 999 | -999 | 1975 | 2018 | \#4 |  |  |
| \# | 1 | 3 | 3 | 1 | 97.5 | 0 | 200 | 0 | 1 | 999 | -999 | 1975 | 2018 | \#4 |  |  |
| \# | 1 | 4 | 4 | 1 | 112.5 | 0 | 200 | 0 | 1 | 999 | -999 | 1975 | 2018 | \#4 |  |  |
| \# | 1 | 5 | 5 | 1 | 162.5 | 0 | 200 | 0 | 1 | 999 | -999 | 1975 | 2018 | \#4 |  |  |
| \# | 1 | 6 | 6 | 1 | 0.001 |  | 0001 | 0.99999 | 0 | 1 | 999 | 4 | 1975 | 2018 | \#4 |  |
| \# | 1 | 6 | 7 | 1 | 0.1 |  | 0001 | 0.99999 | 0 | 1 | 999 | 4 | 1975 | 2018 | \#4 |  |
| \# | 1 | 6 | 8 | 1 | 0.3 |  | 0001 | 0.99999 | 0 | 1 | 999 | 4 | 1975 | 2018 | \#4 |  |
| \# | 1 | 6 | 9 | 1 | 0.7 |  | 0001 | 0.99999 | 0 | 1 | 999 | 4 | 1975 | 2018 | \#4 |  |
| \# | 1 | 6 | 10 | 1 | 0.99999 | 0.0 | 0001 | 1.01 | 0 | 1 | 999 | -4 | 1975 | 2018 | \#4 |  |
|  | 1 | 3 | 1 | 2 | 84.00 | 5 | 150 | 0 | 1 | 999 | 419 | 975 | 2021 |  |  |  |
|  | 1 | 4 | 2 | 2 | 4.0000 | 0.1 | 20 | 0 | 1 | 999 | 419 | 975 | 2021 |  |  |  |
| \# Gear-2 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  | 2 | 5 | 1 | 0 | 165.0 | 5 | 190 | 0 | 1 | 999 | 4 | 1975 | 2021 |  |  |  |
|  | 2 | 6 | 2 | 0 | 15.0000 | 0.1 | 25 | 0 | 1 | 999 | 4 | 1975 | 2021 |  |  |  |
| \# Gear-3-9 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  | \# 3 | 7 | 1 | 1 | 103.275 | 5 | 190 | 1 |  | 103.275 | 20.66 | 4 | 1975 | 2019 |  |  |
|  | \# 3 | 8 | 2 | 1 | 8.834 | 0. | 125 | 1 |  | 8.834 | 1.76 | 4 | 1975 | 2019 |  |  |
|  | \# 3 | 9 | 1 | 2 | 91.178 | 5 | 190 | 1 |  | 91.178 | 18.24 | 4 | 1975 | 2019 |  |  |
|  | \# 3 | 10 | 2 | 2 | 2.5 | 0. | 125 | 1 |  | 2.5 | 0.50 | 4 | 1975 | 2019 |  |  |
|  | \#3 | 7 | 1 | 1 | 103.275 | 5 | 190 | 1 |  | 03.275 | 51.64 | 4 | 1975 | 2019 |  |  |
|  | \#3 | 8 | 2 | 1 | 8.834 | 0.1 | 25 | 1 |  | . 834 | 4.42 | 4 | 1975 | 2019 |  |  |
|  | \#3 | 9 | 1 | 2 | 91.178 | 5 | 190 | 1 |  | 1.178 | 45.59 | 4 | 1975 | 2019 |  |  |
|  | \#3 | 10 | 2 | 2 | 2.5 | 0.1 | 25 | 1 |  | . 5 | 1.25 | 4 | 1975 | 2019 |  |  |
|  | 3 | 7 | 1 | 1 | 103.275 | 5 | 190 | 1 |  | 03.275 | 30.98 | 4 | 1975 | 2021 |  |  |
|  | 3 | 8 | 2 | 1 | 8.834 | 0.1 | 25 | 1 |  | . 834 | 2.65 | 4 | 1975 | 2021 |  |  |
|  | 3 | 9 | 1 | 2 | 91.178 | 5 | 190 | 1 |  | 1.178 | 27.35 | 4 | 1975 | 2021 |  |  |
|  | 3 | 10 | 2 | 2 | 2.5 | 0.1 | 25 | 1 |  | . 5 | 0.75 | 4 | 1975 | 2021 |  |  |
| \# Gear-4 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  | 4 | 11 | 1 | 0 | 115.0 | 5 | 190 | 0 | 1 | 999 | 4 | 1975 | 2021 | \# dummy |  |  |
|  | 4 | 12 | 2 | 0 | 9.0 | 0.1 | 25 | 0 | 1 | 999 | 4 | 1975 | 2021 |  |  |  |
| \# Gear-5 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
|  | 5 | 13 | 1 | 0 | 75.0 | 30 | 190 | 0 | 1 | 999 | 5 | 1975 | 1981 | \#5 |  |  |
|  | 5 | 14 | 2 | 0 | 5.0 | 1 | 50 | 0 | 1 | 999 | 5 | 1975 | 1981 | \#5 |  |  |
|  | 5 | 15 | 1 | 0 | 80.0 | 30 | 190 | 0 | 1 | 999 | 5 | 1982 | 2022 | \#5 |  |  |
|  | 5 | 16 | 2 | 0 | 10.0 | 1 | 50 | 0 | 1 | 999 | 5 | 1982 | 2022 | \#5 |  |  |
|  | \# 5 | 17 | 1 | 2 | 70.0 | 30 | 180 | 0 | 1 | 999 | 5 | 1975 | 1981 | \#5 |  |  |


| \# 5 | 18 | 2 | 2 | 9.0 | 1 | 50 | 0 | 1 | 999 | 5 | 1975 | 1981 | \#5 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| \# 5 | 19 | 1 | 2 | 70.0 | 30 | 180 | 0 | 1 | 999 | 5 | 1982 | 2021 | \#5 |
| \# 5 | 20 | 2 | 2 | 4.00 | 1.0 | 50 | 0 | 1 | 999 | 5 | 1982 | 2021 | \#5 |
| \# Gear-6 |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 6 | 17 | 1 | 0 | 75.0 | 1 | 180 | 0 | 1 | 999 | 5 | 1975 | 2022 | \# 5 |
| 6 | 18 | 2 | 0 | 8.5 | 1 | 50 | 0 | 1 | 999 | 5 | 1975 | 2022 | \# 5 |
| \# 6 | 23 | 1 | 2 | 85.0 | 1 | 180 | 0 | 1 | 999 | 5 | 1975 | 2021 | \# 5 |
| \# 6 | 24 | 2 | 2 | 10.0 | 1 | 50 | 0 | 1 | 999 | 5 | 1975 | 2021 | \# 5 |




```
## PRIORS FOR CATCHABILITY
## If a uniform prior is selected for a parameter then the lb and ub are used (p1 ##
## and p2 are ignored). ival must be > 0 ##
```



```
## prior: 0 = uniform, 1 = normal, 2 = lognormal, 3 = beta, 4 = gamma ## 
```



```
\#\# ival lb ub phz prior p1 p2 Analytic? LAMBDA Emphasis
\begin{tabular}{lllllllllll} 
\# 0.896 & 0 & 2 & 6 & 1 & 0.843136 & 0.03 & 0 & 1 & 1 \\
0.896 & 0 & 2 & 6 & 1 & 0.896 & 0.03 & 0 & 1 & 1 & 1 \\
1.0 & 0 & 5 & -6 & 0 & 0.001 & 5.00 & 0 & 1 & 1
\end{tabular} \# BSFRF
```




```
## ADDITIONAL CV FOR SURVEYS/INDICES ## 
## If a uniform prior is selected for a parameter then the lb and ub are used (p1 ##
## and p2 are ignored). ival must be > 0 ##
## LEGEND
## prior type: 0 = uniform, 1 = normal, 2 = lognormal, 3 = beta, 4 = gamma ##
```



```
## ival lb ub phz prior p1 p2
```

| 0.0001 | 0.00001 | 10.0 | -4 | 4 | 1.0 | 100 | \# NMFS |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 0.25 | 0.00001 | 10.0 | 10 | 0 | 0.001 | 1.00 | \# BSFRF |

 \#\#\# Pointers to how the the additional CVs are used ( 0 ignore; >0 link to one of the parameters !!!NEW 1/2022
12

\#\# PENALTIES FOR AVERAGE FISHING MORTALITY RATE FOR EACH GEAR

\#\# Mean_F Female Offset STD_PHZ1 STD_PHZ2 PHZ_M PHZ_F





\#\# TIME VARYING NATURAL MORTALIIY RATES
\#\#
\#\# LEGEND
\#\# Type: $0=$ constant natural mortality
\#\#
\#\#
\#\# 1 = Random walk (deviates constrained by variance in M) \#\#
\#\# 2 = Cubic Spline (deviates constrained by nodes \& node-placement) \#\#
\#\# $\quad 3=$ Blocked changes (deviates constrained by variance at specific knots) \#\#
\#\# â€"â€"â€"â€"̂̂€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"â€"̂̂€"â€"â€"â€"â€"â€"â€"â€
\#\# Type
6
\#\# M is relative (YES=1; NO=0)
1
\#\# Phase of estimation
3
\#\# STDEV in m_dev for Random walk
0.25
\#\# Number of nodes for cubic spline or number of step-changes for option 3
2
2
\#\# Year position of the knots (vector must be equal to the number of nodes)
19801985
19801985
\# number of breakpoints in $M$ by size

```
0
## Specific initial values for the natural mortality devs (0-no, 1=yes)
1
```




```
\begin{tabular}{lcccc}
1.7342575 & 0 & 2 & 8 & 0 \\
0.000000 & -2 & 2 & -99 & 0 \\
\(\# 0.30\) & 0 & 2 & 8 & 0 \\
\(\# 0.000000\) & -2 & 2 & -99 & 0 \\
1.780586 & 0 & 2 & 8 & -1 \\
\(\# 9.262792\) & 0 & 2 & -8 & -3 \\
0.000000 & -2 & 2 & -99 & 0 \\
\(\# 0.400\) & 0 & 2 & 8 & -3 \\
\(\# 0.000000\) & -2 & 2 & -99 & 0 \\
\(\# 0.6\) & 0 & 2 & 8 & 0 \\
\(\# 0.804\) & 0 & 2 & 8 & 0 \\
\(\# 9.262792\) & 0 & 2 & 8 & -3 \\
\(\# 0.000000\) & -2 & 2 & -99 & 0
\end{tabular}
```



```
## ??????????????????????????????????????????????? ##
## TAGGING controls CONTROLS !!!NEW 1/2022
## ?????????????????????????????????????????????? ##
# emphasis on tagging data
```



```
## OTHER CONTROLS
```



```
1975 # First rec_dev
2021 # last rec_dev
    # # Estimated rec_dev phase
# # Estimated sex_ratio
0.5 # initial sex-ratio
    3 # Estimated rec_ini phase
    # VERBOSE FLAG (0 = off, 1 = on, 2 = objective func; 3 diagnostics)
    # Initial conditions ( }0=\mathrm{ Unfished, 1 = Steady-state fished, 2 = Free parameters, 3 = Free parameters (revised))
    # Lambda (proportion of mature male biomass for SPR reference points).
    # Stock-Recruit-Relationship (0 = none, 1 = Beverton-Holt)
    # Maximum phase (stop the estimation after this phase).
    # Maximum number of function calls
    # Calculate reference points (0=no) !!!NEW 1/2022
    200 # Year to compute equilibria !!!NEW 1/2022
```



```
## EMPHASIS FACTORS (CATCH)
```



```
#Ret_male Disc_male Disc_female Disc_trawl Disc_Tanner_male Disc_Tanner_female Disc_fixed
\begin{tabular}{rrrrrrr} 
& 1 & 1 & 1 & 1 & 1 & 1 \\
\(\#\) & 500 & 100 & 100 & 50 & 100 & 100
\end{tabular}
## EMPHASIS FACTORS (Priors) by fleet: fdev_total, Fdov_total, Fdev_year, Fdov_year !!!NEW 1/2022
1 1 0 0 # Pot fishery
1 1 0 0 # Trawl by-catch
1100 # Tanner by-catch
11 0 0 # Fixed by-catch
1 1 0 0 # Trawl survey
1 1 0 0 # BSFRF survey
```



```
## EMPHASIS FACTORS (Priors)
```



```
# Log_fdevs meanF Mdevs Rec_devs Initial_devs Fst_dif_dev Mean_sex-Ratio Fvecs Fdovs (!!!NEW for the last two 1/2022)
```



```
## EOF
9999
```


## Appendix C. Summary of the CIE Review of BBRKC in

## 2021

The virtual CIE review of the stock assessments for Bristol Bay red king crab and eastern Bering Sea snow crab was held online during March 22-26, 2021. The review was conducted by three independent experts: Drs. Yong Chen, Nick Caputi, and Billy Ernst. The review reports are at the end of this SAFE report. The followings are a brief summary of recommendations and plan to address these recommendations.

1. Identifying the possible sources of the large retrospective patterns and/or develop alternative approaches to provide catch advice if retrospective patterns persistent and biased errors are too large for the assessments to be considered reliable. Conducted more studies to identify temporal trends and/or time blocks of parameters, such as natural mortality and survey catchability, to be incorporated in future stock assessments.

Reply: Temporal changes in parameters may play an important role for the large retrospective patterns, and some data conflict between NMFS surveys and 20072008 and 2013-2016 BSFRF surveys also contributes to them. We used model 21.2 to add another time block (2018-2019) of natural mortality. The Mohn's rho value for mature male biomass decreases from 0.347 for model 19.3 d to 0.223 for model 21.2 . We will further examine the retrospective patterns and develop alternative model scenarios to reduce the retrospective patterns for the CPT meeting in January/May 2022. Potential changes in natural mortality over time play a big role for the large retrospective patterns during recent years, and additional time blocks of parameters for recent years will be further evaluated.
2. Survey performance/efficacy and selectivity curve evaluations in term of changes in distributions over time, and the stock area evaluation.

Reply: We would like to examine red king crab north of the management area of Bristol Bay sometime in the future to see whether they are part of the BBRKC stock. Hopefully, a tagging study can be conducted to examine the link between red king crab in these two areas. We have not seen the need for evaluating different kinds of survey selectivity curves now since large-size crab are generally inside the survey area. Some limited genetic and larval transport studies were conducted on the stock area in the past.
3. Surveying the red king crab juvenile crab abundance in nearshore locations may provide an estimate of younger juvenile abundance where the year-class is better defined.

Reply: We second this and have advocated this for a long time.
4. Examining VAST results on effects on the stock assessment model.

Reply: We will continue to use VAST results as a model scenario to compare it to the other model scenarios.
5. Evaluating commercial catch, effort, and CPUE for crab distributions, fishery performance, and population abundance relative to the trawl survey results and on impacts on survey timing and survey availability, and standardizing the CPUE for improvement, and conducting a depletion analysis.

Reply: Catch and bycatch are used in the model, the commercial CPUE is used to compare to the survey legal male abundance but not in the model, and fishing distributions and CPUE are often examined by ADF\&G. The fishing season has been very short in the most years, so the depletion analysis may not be much useful. Trawl surveys generally cover all red king crab distribution areas except for nearshore areas. We just started CPUE standardization work and will try to incorporate the standardized CPUE in the assessment model in 2022.
6. Extending estimates of sizes-at-50\% maturity for females and examining the impacts of changes on mature female biomass estimates. Conducting a sensitivity study to examine impacts of changes at sizes-at-maturity for males on mature male biomass estimates.

Reply: We will update the estimates of sizes-at-50\% maturity for females. Since the harvest strategy defines the sizes of mature females and males and the growth increments of males is not affected by changes in sizes-at-maturity, impacts of changes at sizes-at-maturity for males on mature male biomass estimates do not occur for the harvest strategy. The current defined size-atmaturity for males is for functional maturity and is much larger than the physiological mature sizes.
7. A model run just using data from 1985 to avoid high natural mortality during the early 1980s.

Reply: We have planned to do this in 2022.
8. Examining biological, environmental, and vessel performance data on the 2014 NMFS trawl survey to assess the survey abundance outlier and conducting a sensitivity study without the 2014 NMFS trawl survey data.

Reply: During the CIE review, we conducted this sensitivity study. The NMFS and BSFRF have examined biological, environmental, and vessel performance data on the 2014 NMFS trawl survey extensively. It is unlikely that we would drop this data point in the stock assessments since there are several data points in the survey time series that are as unexpected as the 2014 data.
9. Important to continue environmental SAFE reports.

Reply: We agree and hopefully it will be updated annually.
10. Besides overfished and overfishing, using MMB, recruitment, trends in commercial catch and CPUE, legal-size abundance and total survey biomass, and the projections and near future outlook to summarize the stock status.

Reply: We will add these in our summary of the stock status.
11. Modeling double bag experiment and BSFRF side-by-side survey data to improve the catchability prior.

Reply: This is a good suggestion. However, we do not use BSFRF side-by-side survey data to estimate the NMFS trawl catchability prior because we do not want to use these data twice since they are used in the model already.
12. Conducting new tagging study to update the outdated tagging/return data used in the assessments.

Reply: We agree with this recommendation. Hopefully, tagging study will be conducted for BBRKC in the future.

# Appendix D. Ecosystem and Socioeconomic Profile of the Bristol Bay Red King Crab stock - Report Card 

Erin Fedewa, Brian Garber-Yonts and Kalei Shotwell

September 2022


With Contributions from:
Matt Callahan, Curry Cunningham, Ben Daly, Jean Lee, Mike Litzow, Jens Nielsen, Katie Palof, Darren Pilcher, Jon Richar, Dale Robinson, Abigail Tyrell, Ellen Yasumiishi, Leah Zacher and Jie Zheng

## Current Year Update

The ecosystem and socioeconomic profile or ESP is a standardized framework for compiling and evaluating relevant stock-specific ecosystem and socioeconomic indicators and communicating linkages and potential drivers of the stock within the stock assessment process (Shotwell et al., In Review). The ESP process creates a traceable pathway from the initial development of indicators to management advice and serves as an on-ramp for developing ecosystem-linked stock assessments.
Please refer to the last full ESP document (Fedewa et al., 2020, Appendix E, pp. 172-204) which is available within the Bristol Bay red king crab (BBRKC) stock assessment and fishery evaluation or SAFE report for further information regarding the ecosystem and socioeconomic linkages for this stock.

## Management Considerations

The following are the summary considerations from current updates to the ecosystem and socioeconomic indicators evaluated for BBRKC:

- In 2022, bottom temperatures returned to near-average and the cold pool extended into the Bristol Bay management area. Results from the NOAA bottom trawl survey indicate that BBRKC female reproductive cycles were delayed due to relatively cold bottom temperatures. However, summer bottom temperatures were well-within the thermal range of juvenile and adult red king crab.
- Red king crab have experienced a steady decline in bottom water pH in the past two decades, reaching 7.89 in 2022. Continued declines to threshold pH levels of 7.8 could negatively affect juvenile red king crab growth, shell hardening and survival.
- BBRKC recruitment remains well below the long-term average. Concurrent declines in Pacific cod and benthic invertebrate densities in the past 7 years may suggest shared processes that drive productivity of Bristol Bay benthic communities.
- Spatial extent of mature male red king crab in Bristol Bay was above average in 2022, coinciding with increases in abundance. The relatively large spatial footprint of mature males in 2022 can be attributed to an increased use of nearshore habitats in Bristol Bay, and was likely driven by the return of cold waters $<2^{\circ} \mathrm{C}$ following a 2018-2019 heat wave.
- Closure of the BBRKC fishery for 2021/2022, which averaged nearly $\$ 55$ million in annual gross ex-vessel value over the previous five years, combined with economic effects of the historically low TAC for the EBS snow crab fishery, entail severe economic stresses affecting the EBS crab fleet and associated stakeholders during the current period. If persistent, this has potential to induce lasting structural changes in crab harvesting and processing sectors with associated changes from historical patterns of fleet fishing behavior.
- Incidental catch of BBRKC biomass in EBS groundfish fisheries during 2021 increased moderately from the previous year to slightly above average for the 2010-current period.


## Modeling Considerations

The following are the summary results from the intermediate and advanced stage monitoring analyses for BBRKC:

- The highest ranked predictor variables ( $>0.50$ inclusion probability) in the advanced stage monitoring analysis were: BBRKC recruit biomass, Pacific cod biomass, and the Arctic Oscillation. Due to concerns with autocorrelation in model-based estimates of mature male biomass, indicator importance tests in future BBRKC ESP updates will use recruitment estimates as a response variable.
- The advanced stage indicator analysis provides updates on developing research ecosystem linked models that are not yet included as a model alternative in the main stock assessment. We have not received updates on new research ecosystem linked models for BBRKC at this time.


## Assessment

## Ecosystem and Socioeconomic Processes

We summarize important processes that may be helpful for identifying productivity bottlenecks and dominant pressures on the stock in conceptual models detailing 1) ecosystem processes by RKC life history stage (Figure 1a) and 2) socioeconomic performance metrics (Figure 1b). The ecosystem conceptual model highlights abiotic and biotic processes identified by each life stage from the literature, process studies and laboratory rearing experiments. During early larval stages, RKC survival is dependent on spatiotemporal overlap with high densities of diatoms (Paul et al., 1989; Paul and Paul, 1980), optimal environmental conditions for development (Nakanishi, 1987) and dispersal to suitable settlement habitat (Daly et al., 2018). Specific habitat requirements for juvenile RKC include physical structure and high relief to both evade predators (Stoner, 2009; Pirtle et al., 2012) and provide increased foraging opportunities (Pirtle and Stoner, 2010). Late juvenile and adult RKC are less reliant on complex structure, and instead, spatial distributions and migration timing are driven by bottom temperatures (Loher and Armstrong, 2005; Zheng and Kruse, 2006; Zacher et al., 2018).

The socioeconomic conceptual model highlights fishery performance indicators that represent processes most directly involved in prosecution of the BBRKC fishery, and thus have the potential to differentially affect the condition of the stock, depending on how they influence the timing, spatial distribution, selectivity, and other aspects of fishing pressure. Implementation of the Crab Rationalization Program and the allocation of tradable crab harvest quota shares resulted in rapid consolidation of the BBRKC fleet and changed the timing of the fishery from short derby seasons to more extended seasons. These and other institutional changes continue to influence the geographic and inter-sectoral distribution of benefits produced by the BBRKC fleet.

## Indicator Suite

The following list of indicators for BBRKC is organized by categories: three for ecosystem indicators (physical, lower trophic, and upper trophic) and two for socioeconomic indicators (fishery performance and economic). A title, short description and contact name for the indicator contributor are provided. We also include the anticipated sign of the proposed relationship between the indicator and the stock population dynamics where relevant. Please refer to the last full ESP document for detailed information regarding the ecosystem and socioeconomic indicator descriptions and proposed mechanistic linkages for this stock (Fedewa et al., 2020). Time series of the ecosystem and socioeconomic indicators are provided in Figure 2a and Figure 2b, respectively. Summer pH values reported in past ESP products differ slightly from those reported in this document due to an updated ROMS model hindcast through August 2022, which simulates less large phytoplankton than the previous hindcast and results in comparatively lower surface pH values due to less photosynthesis, but greater bottom water pH due to less respiration. Also, please note that we have added back in two socioeconomic indicators that were presented in the full BBRKC ESP. Upon further evaluation, we have determined that these two indicators are useful for understanding health and condition of the stock. The two indicators are annual incidental catch of Bristol Bay red king crab in eastern Bering Sea trawl and fixed gear fisheries and TAC utilization in the Bristol Bay red king crab fishery, which we are monitoring for potential changes due to shifts in stock distribution.

## Ecosystem Indicators:

Physical Indicators (Figure 2a.a-e)
a.) The areal extent of the summer cold pool (EBS bottom trawl survey stations with bottom temperatures $<2^{\circ} \mathrm{C}$; contact: Erin Fedewa). Proposed sign of relationship is positive.
b.) Summer bottom temperatures in Bristol Bay from the AFSC eastern Bering Sea bottom trawl survey (contact: E. Fedewa). Proposed sign of relationship is positive.
c.) Winter-spring Arctic Oscillation index from the NOAA National Climate Data Center (contact: E. Fedewa). Proposed sign of relationship is positive.
d.) Spring pH index in Bristol Bay from the Bering10K ROMS model (Pilcher et al., 2019) (contact: D. Pilcher). Proposed sign of relationship is positive.
e.) Summer wind stress ( $\mathrm{m} / \mathrm{s}$ ) in Bristol Bay from NOAA/NCDC blended winds and MetopA ASCAT satellite (Zhang et al., 2006, NOAA/NESDIS, CoastWatch) (contact: D. Robinson). Proposed sign of relationship is negative.
Lower Trophic Indicators (Figure 2a.f)
f.) Spring chlorophyll-a biomass in Bristol Bay from MODIS satellites (contact: M. Callahan and J. Nielsen). Proposed sign of relationship is positive
Upper Trophic Indicators (Figure 2a.g-m)
g.) September juvenile sockeye salmon abundance in the EBS from the AFSC Bering Arctic Subarctic Integrated Survey (contact: E. Yasumiishi). Proposed sign of relationship is negative.
h.) Summer Pacific cod density in Bristol Bay from the AFSC eastern Bering Sea bottom trawl survey (contact: E. Fedewa). Proposed sign of relationship is negative.
i.) Summer benthic invertebrate density in Bristol Bay. Invertebrates include brittle stars, sea stars, sea cucumber, bivalves, non-commercial crab species, shrimp and polychaetes. (contact: Erin Fedewa). Proposed sign of relationship is positive.
j.) Annual red king crab recruit abundance ( $110-134 \mathrm{~mm}$ CL) in Bristol Bay from the AFSC eastern Bering Sea bottom trawl survey (contact: E. Fedewa). Proposed sign of relationship is positive.
k.) Summer mature male red king crab area occupied in Bristol Bay from the AFSC eastern Bering Sea bottom trawl survey (contact: E. Fedewa). Proposed sign of relationship is negative.
1.) Summer mature female red king crab area occupied in Bristol Bay from the AFSC eastern Bering Sea bottom trawl survey (contact: E. Fedewa). Proposed sign of relationship is negative.
m.) Annual male red king crab catch distance from shore in Bristol Bay during the fishery (contact: L. Zacher). Proposed sign of relationship is positive.

## Socioeconomic Indicators:

Fishery Performance Indicators (Figure 2b. a-d)
a.) Annual catch-per-unit-effort (CPUE), expressed as mean number of legal crabs per potlift, in the BBRKC fishery, representing relative efficiency of fishing effort (contact: B. Daly)
b.) Annual total potlifts in the BBRKC fishery, representing the level of fishing effort expended by the active fleet (contact: B. Daly)
c.) Annual number of active vessels in the Bristol Bay red king crab fishery, representing the level of fishing effort assigned to the fishery (contact: J. Lee)
d.) Estimated total incidental catch of BBRKC biomass ( kg ) in EBS groundfish fisheries (contact: J. Lee)
Economic Indicators (Figure 2b. e-h; all monetary values are inflation-adjusted to $\$ 2021$ value)
e.) Percentage of the annual BBRKC total allowable catch (TAC) (GHL prior to 2005) that was harvested by active vessels, including deadloss discarded at landing (contact: B. Garber-Yonts)
f.) Annual ex-vessel value (\$2021) of the BBRKC fishery landings, representing gross economic returns to the harvest sector, as a principal driver of fishery behavior (contact: J. Lee)
g.) Annual ex-vessel price per pound (\$2021) of BBRKC landings, representing per-unit gross economic returns to the harvest sector, as a principal driver of fishery behavior (contact: J. Lee)
h.) Annual ex-vessel revenue share, expressed as average proportion of total annual gross landings revenue from all fisheries earned from BBRKC landings by vessels active in the fishery (contact: J. Lee)

## Indicator Monitoring Analysis

There are up to three stages (beginning, intermediate, and advanced) of statistical analyses for monitoring the indicator suite listed in the previous section. The beginning stage is a relatively simple evaluation by traffic light scoring. This evaluates the current year trends relative to the mean of the whole time series, and provides a historical perspective on the utility of the whole indicator suite. The intermediate stage uses importance methods related to a stock assessment variable of interest (e.g., recruitment, biomass, catchability). These regression techniques provide a simple predictive performance for the variable of interest and are run separate from the stock assessment model. They provide the direction, magnitude, uncertainty of the effect, and an estimate of inclusion probability. The advanced stage is used for testing a research ecosystem linked model and output can be compared with the current operational model to understand information on retrospective patterns, prediction performance, and comparisons of other model output such as terminal spawning stock biomass or mean recruitment. This stage provides an onramp for introducing an alternative ecosystem linked stock assessment model to the current operational stock assessment model and can be used to understand the potential reduction in uncertainty by including the ecosystem information.

## Beginning Stage: Traffic Light Test

We use a simple scoring calculation for this beginning stage traffic light evaluation. Indicator status is evaluated based on being greater than ("high"), less than ("low"), or within ("neutral") one standard deviation of the long-term mean. A sign based on the anticipated relationship between the ecosystem indicators and the stock (generally shown in Figure 1a and specifically by indicator in the Indicator Suite, Ecosystem Indicators section) is also assigned to the indicator where possible. If a high value of an indicator generates good conditions for the stock and is also greater than one standard deviation above the mean, then that value receives a ' +1 ' score. If a high value generates poor conditions for the stock and is greater than one standard deviation above the mean, then that value receives a ' -1 ' score. All values less than or equal to one standard deviation from the long-term mean are average and receive a ' 0 ' score. The scores are summed by the three organizational categories within the ecosystem (physical, lower trophic, and upper trophic) or socioeconomic (fishery performance and economic performance) indicators and divided by the total number of indicators available in that category for a given year. The scores over time allow for comparison of the indicator performance and the history of stock productivity (Figure 3). We also provide five year indicator status tables with a color or text code for the relationship with the stock (Tables 1a,b) and evaluate the current year status in the historical indicator time series graphic (Figures $2 \mathrm{a}, \mathrm{b}$ ) for each ecosystem and socioeconomic indicator. Socioeconomic indicators representing the target fishery are reported, by calendar year, through 2020 (noting that virtually all active harvest activity occurs prior to January), the last year that the fishery was open (corresponding to the 2020-2021 crab season, and incidental catch is reported for the most recent full calendar year (2021).

We evaluate the status and trends of the ecosystem and socioeconomic indicators to understand the pressures on the BBRKC stock regarding recruitment, stock productivity, and stock health. We start with the physical indicators and proceed through the increasing trophic levels for the ecosystem indicators then evaluate the fishery performance and economic indicators as listed above. Here, we concentrate on updates since the last ESP. Overall, the physical and upper trophic indicators scored below average for 2022, while the lower trophic indicators were average (Figure 3). The fishery performance indicators
scored average for 2021, but this is based on only one indicator. The economic indicators were average for 2020. Compared to the previous data point, this is an increase from well below average for the physical indicators, the same for the lower trophic indicators, a decrease for the upper trophic indicators, an increase for the fishery performance indicators, and the same for the economic indicators.

Overall, trends in physical ecosystem indicators suggest a return to near-normal conditions in Bristol Bay with average bottom temperatures nearly $2^{\circ} \mathrm{C}$ colder than 2018-2019 heat conditions. A positive phase Arctic Oscillation index in winter 2022 may suggest favorable conditions for BBRKC productivity (Szuwalski et al., 2020), although continued declines in pH that are approaching a critical threshold for negative effects on growth and shell hardening remain concerning (Long et al., 2013). Results from the 2022 AFSC EBS bottom trawl survey indicate that reproductive cycles of mature female BBRKC were delayed due to relatively cold spring bottom temperatures in Bristol Bay (Zacher et al., in review). Delayed spring hatching of red king crab embryos relative to mid-May peak bloom timing may impact the spatiotemporal overlap between first-feeding larvae and preferred diatom prey, and larval retention may be reduced in relatively cold years (Daly et al., 2020). While recent year updates for juvenile sockeye salmon abundance were not yet available for this document, Bristol Bay's 2022 sockeye run was the largest on record and may be indicative of increased predation on larval RKC in recent years. However, near-average wind stress and chloropyll- $a$ biomass in Bristol Bay indicate suitable conditions for larval first-feeding success and survival.

Current-year values for upper trophic level Pacific cod and benthic invertebrate indicators are not yet available following the conclusion of the 2022 EBS bottom trawl survey. However, both indicators are on a downward trend and Pacific cod biomass has been below average since 2016 in Bristol Bay. BBRKC recruitment still remains well below average as well, and concurrent declines with Pacific cod and invertebrates may be suggestive of bottom-up forcing on benthic communities in Bristol Bay. Although inference on fall BBRKC distributions is limited due to a 2021/2022 fishery closure, mature male area occupied during the summer NOAA bottom trawl survey was above-average. This likely coincides with relatively high catches along the Alaska Peninsula (Zacher et al., in review), and may point to the importance of near-shore habitat in years when the cold pool extends south into the management area (Zacher et al., 2018).

Trends in fishery performance and economic indicators correspond to ongoing decline in TACs issued in the BBRKC fishery since the 2014/2015 season, with complete closure for the 2021/2022 season. Prior to the closure, effort in the fishery exhibited downward trends over recent years, as indicated by the number of active vessels and total number of potlifts, both of which reached the lowest points on record for active seasons during 2020/2021, while CPUE increased somewhat relative to the previous three seasons, but remained at a relative low compared to the post-rationalization period overall. Ex-vessel price declined slightly for the 2020/2021 season, but remained relatively high compared to the prior post-rationalization period; closure during 2021/2022 coincided with historically high ex-vessel and first wholesale prices reported in other BSAI crab fisheries during 2021 (Garber-Yonts et al, in prep). Consistent with the downward trend in TACs since 2014/2015, the $\$ 33.6$ million total annual gross ex-vessel value of BBRKC landings during 2021 continued the sharp declining trend observed over the period, with exvessel value and ex-vessel revenue share (the percentage share of BBRKC ex-vessel value relative to total annual ex-vessel value of landings, across all fisheries, by vessels active in BBRKC during the year) both reaching historical lows as of 2020. The loss of income associated with closure of the BBRKC fishery for $2021 / 2022$, which generated an average of nearly $\$ 55$ million in annual gross ex-vessel value over the previous five years, combined with the historically low TAC for the Eastern Bering Sea snow crab fishery, represent acute financial and economic stresses on the EBS crab industry and associated stakeholders. Apart from fishery management concerns regarding potentially substantial, long-term income and employment losses, if persistent, near-term financial stresses have potential to induce lasting structural changes in crab harvesting and processing sectors. While speculative, there is the potential for
structural changes to result in systematic changes in fleet fishing behavior, with implications for assessment modeling and in-season management across EBS crab fisheries.

## Intermediate Stage: Importance Test

We plan to update the second stage indicator analysis in 2023 and are exploring additional importance methods for BBRKC.

## Advanced Stage: Research Model Test

At this time we do not have any ecosystem research models to report for BBRKC.

## Data Gaps and Future Research Priorities

Environmental conditions are rapidly changing in the eastern Bering Sea and continued research is needed to identify temperature thresholds and characterize responses across BBRKC life stages to changes in bottom temperatures. Specifically, future laboratory and field research should focus on clarifying the range of optimal temperatures for embryo survival and successful settlement in juvenile nursery areas. In addition, potential climate-driven shifts in phenology and spatial distribution underscore the importance of assessing fishery interactions with trawl and pot gear relative to BBRKC migration patterns, molt-mate timing and spawning habitat.

Given the dramatic increase in Bristol Bay sockeye salmon coinciding with declines in BBRKC recruitment in recent years, we emphasize the importance of understanding predator-prey interactions and spatiotemporal overlap of major pelagic predators with BBRKC larval stages. Juvenile salmon diet studies conducted from 1984-1992 (Farley 2001, unpublished data) reported that juvenile sockeye salmon consumption of red king crab zoea exceeded $45 \%$ in several years, suggesting potential links between salmon predation and BBRKC recruitment. In more recent years, the Bering-Aleutian Salmon International Survey has taken place in late-September following peak settlement of BBRKC, and there appears to be no ongoing efforts to characterize diets of juvenile sockeye salmon in earlier summer months when BBRKC are likely important prey items. Furthermore, because the survey is biennial and occurs in September, data gaps across the time series preclude use of the indicator in monitoring analyses, and indicator updates are unavailable for the current-year ESP. Future efforts should focus on exploring additional larval predator datasets that are more timely and consistent. In addition, additional groundfish stomach data outside of the summer survey time series would inform predation mortality during the molt when RKC are highly vulnerable.

Low stock recruitment in the past decade also warrants a better understanding of early life history processes and bottlenecks to aid in developing meaningful larval indicators as early warning signs. Evaluating RKC phenology relative to spring bloom timing may be useful for predicting larval condition and subsequent survival to settlement. Additionally, evaluating larval drift patterns and identifying essential fish habitat for benthic juvenile RKC may support the development of a larval retention or settlement success indicator. Overall, we highlight the continued importance of developing a mechanistic understanding of driver-response relationships to facilitate the inclusion of ecosystem indicators in future management strategies for BBRKC.

BBRKC ESP developments for 2023 include: 1) updating the intermediate stage indicator analysis, 2) producing a Request for Indicators in January 2023 to highlight data gaps and propose new indicator contributions, 3) developing an indicator to quantify potential BBRKC-gear interactions using the Fishing Effects model, and 4) updating ecosystem and socioeconomic indicators and considerations prior to the 2023 Crab Plan Team meeting to inform BBRKC management.

We plan to further evaluate the information provided in the Economic SAFE and ACEPO report to determine what socioeconomic indicators could be provided in the ESP that are not redundant with those reports and related directly to stock health. This may result in a transition of socioeconomic indicators currently reported in this ESP to a different series of indicators. Additional consideration of the timing of the economic and community reports, which are lagged by 1-2 years (depending on the data source) from the annual stock assessment cycle, should also be undertaken. The Scientific and Statistical Committee (SSC) recently recommended that local knowledge, traditional knowledge, and subsistence information may be helpful for understanding recent fluctuations in stock health, shifts in stock distributions, or changes in size or condition of species in the fishery. We could include this information as supportive evidence and perspective on many indicators monitored within the ESP.

As indicators are improved or updated, they may replace those in the current set of indicators to allow for refinement of the BAS model and potential evaluation of performance and risk within the operational stock assessment model. The annual request for indicators (RFI) for the BBRKC ESP will include these data gaps and research priorities along with a list of potential new indicators that could be developed for the next full ESP assessment.

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## Tables

Table 1a. First stage ecosystem indicator analysis for BBRKC, including indicator title and the indicator status of the last five available years. The indicator status is designated with text, (greater than = "high", less than = "low", or within 1 standard deviation = "neutral" of time series mean). Fill color of the cell is based on the sign of the anticipated relationship between the indicator and the stock (blue or italicized text $=$ good conditions for the stock, red or bold text $=$ poor conditions, white $=$ average conditions). A gray fill and text = "NA" will appear if there were no data for that year.

| Indicator category | Indicator | 2018 <br> Status | 2019 <br> Status | $\begin{aligned} & 2020 \\ & \text { Status } \end{aligned}$ | 2021 <br> Status | 2022 <br> Status |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Physical | Summer Cold Pool SEBS BBRKC Survey | low | low | NA | low | neutral |
|  | Summer Temperature Bottom BBRKC Survey | high | high | NA | neutral | neutral |
|  | Winter Spring Arctic Oscillation Index Model | neutral | neutral | high | neutral | neutral |
|  | Spring pH BBRKC Model | low | low | low | low | low |
|  | Summer Wind Stress BBRKC Satellite | neutral | high | neutral | high | neutral |
| Lower <br> Trophic | Spring Chlorophylla Biomass SEBS Inner Shelf Satellite | neutral | neutral | neutral | neutral | neutral |
| Upper <br> Trophic | Summer Sockeye <br> Salmon Abundance EBS <br> Survey | high | NA | NA | NA | NA |
|  | Summer Pacific Cod Density BBRKC Survey | neutral | low | NA | neutral | NA |
|  | Summer Benthic Invertebrate Density BBRKC Survey | neutral | neutral | NA | neutral | NA |
|  | Annual Red King Crab Recruit Abundance BBRKC Survey | low | low | NA | low | low |
|  | Summer Red King Crab Male Area Occupied BBRKC Model | high | high | NA | neutral | high |


| Indicator <br> category | Indicator | $\mathbf{2 0 1 8}$ <br> Status | 2019 <br> Status | 2020 <br> Status | 2021 <br> Status | 2022 <br> Status |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
|  | Summer Red King Crab <br> Female Area Occupied <br> BBRKC Model | neutral | high | NA | high | neutral |
|  | Annual Red King Crab <br> Catch Distance Shore <br> BBRKC Fishery | neutral | high | neutral | NA | NA |

Table 1b. First stage socioeconomic indicator analysis for BBRKC, including indicator title and the indicator status of the last five available years. The indicator status is designated with text, (greater than $=$ "high", less than = "low", or within 1 standard deviation = "neutral" of time series mean). A gray fill and text = "NA" will appear if there were no data for that year.

| Indicator category | Indicator | $\begin{aligned} & 2017 \\ & \text { Status } \end{aligned}$ | 2018 <br> Status | $\begin{aligned} & 2019 \\ & \text { Status } \end{aligned}$ | 2020 <br> Status | 2021 <br> Status |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Fishery <br> Performance | Annual Red King Crab CPUE BBRKC Fishery | neutral | neutral | neutral | neutral | NA |
|  | Annual Red King Crab Total Potlift BBRKC Fishery | neutral | neutral | neutral | low | NA |
|  | Annual Red King Crab Active Vessels BBRKC Fishery | neutral | neutral | neutral | neutral | NA |
|  | Annual Red King Crab Incidental Catch EBS Fishery | high | neutral | neutral | neutral | neutral |
| Economic | Annual Red King Crab TAC Utilization BBRKC Fishery | neutral | neutral | neutral | neutral | NA |
|  | Annual Red King Crab Exvessel Value BBRKC Fishery | neutral | low | low | low | NA |
|  | Annual Red King Crab Exvessel Price BBRKC Fishery | neutral | high | high | high | NA |
|  | Annual Red King Crab Exvessel Revenue Share BBRKC Fishery | neutral | neutral | neutral | neutral | NA |

## Figures



Figure 1a: Life history conceptual model for BBRKC summarizing ecological information and key ecosystem processes affecting survival by life history stage. Thermal requirements by life history stage were determined from RKC laboratory studies. Red text means increases in process negatively affect survival, while blue text means increases in process positively affect survival.


Figure 1b: Conceptual model of socioeconomic performance metrics for BBRKC that may identify dominant pressures on the Bristol Bay red king crab stock.


Figure 2a. Selected ecosystem indicators for BBRKC with time series ranging from 1970 - present. Upper and lower solid green horizontal lines represent 1 standard deviation of the time series mean. Dotted green horizontal line is the mean of the time series. A symbol appears when current year data are available and follows the traffic light status table designations (triangle direction represents if above or below 1 standard deviation of the time series mean, color represents proposed relationship for stock, white circle for neutral).


Figure 2a (cont.). Selected ecosystem indicators for BBRKC with time series ranging from 1970 present. Upper and lower solid green horizontal lines represent 1 standard deviation of the time series mean. Dotted green horizontal line is the mean of the time series. A symbol appears when current year data are available and follows the traffic light status table designations (triangle direction represents if above or below 1 standard deviation of the time series mean, color represents proposed relationship for stock, white circle for neutral).


YEAR

Figure 2 a (cont.). Selected ecosystem indicators for BBRKC with time series ranging from 1970 present. Upper and lower solid green horizontal lines represent 1 standard deviation of the time series mean. Dotted green horizontal line is the mean of the time series. A symbol appears when current year data are available and follows the traffic light status table designations (triangle direction represents if above or below 1 standard deviation of the time series mean, color represents proposed relationship for stock, white circle for neutral).


Figure 2b. Selected socioeconomic indicators for BBRKC with time series ranging from 1966 - present. Upper and lower solid green horizontal lines represent 1 standard deviation of the time series mean. Dotted green horizontal line is the mean of the time series. A symbol appears when current year data are available and follows the traffic light status table designations (triangle direction represents if above or below 1 standard deviation from the time series mean, color represents proposed relationship for stock, white circle for neutral).


## YEAR

Figure 2b (cont.). Selected socioeconomic indicators for BBRKC with time series ranging from 1966 present. Upper and lower solid green horizontal lines represent 1 standard deviation of the time series mean. Dotted green horizontal line is the mean of the time series. A symbol appears when current year data are available and follows the traffic light status table designations (triangle direction represents if above or below 1 standard deviation from the time series mean, color represents proposed relationship for stock, white circle for neutral).

## Overall Stage 1 Score for Bristol Bay Red King Crab


$\rightarrow$ Physical $\quad+$ Fishery Performance

- Lower Trophic - E- Economic
- Upper Trophic

Figure 3: Simple summary traffic light score by category for ecosystem and socioeconomic indicators from 2000 to present.


[^0]:    "The SSC supports the BSFRF collaborative work with ADF\&G and NMFS to tag BBRKC."

