# 1A. Stock Assessment of Aleutian Islands Region Pollock 

Steve Barbeaux, James Ianelli, Sarah Gaichas, and Mark Wilkins<br>Alaska Fisheries Science Center<br>November 2006

## Executive Summary

Development of a detailed age-structured stock assessment for the Aleutian Islands Region pollock began in 2003 (Barbeaux et al. 2003) and has been further developed in 2004 and 2005 (Barbeaux et al. 2004, Barbeaux et al. 2005). In the initial study the near shore areas of the Aleutian chain island were isolated and identified as the Near, Rat, and Andreanof Island (NRA) sub-area. This sub-area was further refined to exclude the area east of $174^{\circ} \mathrm{W}$ to address data consistency issues. The Council supported this proposal and urged continuing development on an age-structured assessment model using data from the area west of $174^{\circ} \mathrm{W}$ (and omitting deep-water areas where survey data are unavailable).
Pollock fishery data collected near the eastern boundary of the Aleutian Islands region (between of $174^{\circ} \mathrm{W}$ and $170^{\circ} \mathrm{W}$ ) highlight stock structure uncertainty between the Aleutian Islands region, the Aleutian Islands Basin, and the EBS. Consequently, they are excluded from all of the age-structured assessment models presented below. We do, however, recognize that fluctuations in biomass observed from the summer Aleutian Islands bottom trawl (AIBT) survey data from this area do not indicate clear cut patterns and that substantial uncertainty in the stock structure exists. We have included model configurations which include all of the AIBT survey data and one model that excludes data from the eastern NRA area (NRA area east of $174^{\circ} \mathrm{W}$ ).
Spatial analyses of fishery, survey, and bycatch data using GIS methods reveal an important characteristic of pollock in the Aleutian Islands region: concentrations are highly variable and likely evolve quickly within seasons. These analyses underscore the challenge of evaluating stocks that: are highly mobile, spend variable time associated with the bottom, have patchy distributions, and are likely influenced by neighboring stocks.

## Summary of major changes

The model configuration differs from the 2005 assessment in that natural mortality is estimated within the reference model. The data for this year's assessment differ from last year in that age data from the 2006 Aleutian Islands Cooperative Acoustic Survey Study was available as a proxy for fisheries data and biomass estimates from the 2006 Aleutian Islands bottom trawl survey were incorporated.

## Changes in the assessment results

The estimate for natural mortality in the reference model was lower than that assumed in previous years, therefore biomass and F estimates are significantly lower than those from the 2005 assessment. The maximum permissible ABC for 2007 and 2008 under Tier 3a are 44,470 $t$ and $33,090 t$ respectively and the maximum permissible ABC under Tier 5 for both years using the lower natural mortality is $16,800 \mathrm{t}$.

## Response to SSC 2005 Comments

The SSC commented that they supported continued research of the Aleutians pollock stock. To that end we completed the 2006 Aleutian Islands cooperative acoustic survey study AICASS to ascertain whether it was possible to conduct acoustic surveys on pollock in the Central Aleutian Islands in the winter. This
project was a success and has provided not only verification of the technique, but valuable data on the biology and health of the Aleutian Islands pollock stock. Some the biological data such as age and length composition have been used directly in this assessment, other data such as pollock tissue samples await funding for further analysis. We propose further research of this type for 2007 with an expanded AICASS.

## Introduction

Walleye pollock (Theragra chalcogramma) are distributed throughout the Aleutian Islands with concentrations in areas and depths dependent on season. Generally, larger pollock occur in spawning aggregations during February - April. Three stocks of pollock inhabiting three regions in the Bering Sea Aleutian Islands (BSAI) are identified in the U.S. portion of the BSAI for management purposes. These stocks are: the eastern Bering Sea pollock occupying the eastern Bering Sea shelf from Unimak Pass to the U.S.-Russia Convention line; the Aleutian Islands Region pollock encompassing the Aleutian Islands shelf region from $170^{\circ} \mathrm{W}$ to the U.S.-Russia Convention line; and the Central Bering Sea-Bogoslof Island pollock. These three management stocks probably have some degree of exchange. The Central Bering Sea-Bogoslof stock is a group that forms a distinct spawning aggregation that has some connection with the deep water region of the Aleutian Basin. In the Russian Exclusive Economic Zone (EEZ), pollock are thought to form two stocks, a western Bering Sea stock centered in the Gulf of Olyutorski, and a northern stock located along the Navarin shelf from $171^{\circ} \mathrm{E}$ to the U.S.- Russia Convention line. The northern stock is believed to be a mixture of eastern and western Bering Sea pollock with the former predominant. Bailey et al. (1999) present a thorough review of the population structure of pollock throughout the north Pacific region. Recent genetic studies using mitochondrial DNA methods have found the largest differences to be between pollock from the eastern and western sides of the north Pacific.

Previously, Ianelli et al. (1997) developed a model for Aleutian Islands pollock and concluded that the spatial overlap and the nature of the fisheries precluded a clearly defined "stock" since much of the catch was removed very close to the eastern edge of the region and appeared continuous with catch further to the east. In some years a large portion of the pollock removed in the Aleutian Islands Region was from deep-water regions and appeared to be most aptly assigned as "Basin" pollock. This problem was confirmed in the 2003 Aleutian Islands pollock stock assessment (Barbeaux et al. 2003).

## Fishery

The nature of the pollock fishery in the Aleutian Islands Region has varied considerably since 1977 due to changes in the fleet makeup and in regulations. During the late 1970s through the 1980s the fishing fleet was primarily foreign. In 1989, the domestic fleet began operating in earnest and continued in the Aleutian Islands Region until 1999 when the North Pacific Fishery Management Council (NPFMC) recommended closing this region for directed pollock fishing due to concerns for Steller sea lion recovery. Table 1A. 1 provides a history of ABC, OFL, and catch for Aleutian Islands pollock since 1991. In 2005 the fishery was reopened with a $19,000 \mathrm{t}$ TAC. A directed pollock fishery was conducted in February 2005, but the vessels participating in the fishery failed to find commercially harvestable quantities outside of Steller sea lion critical habitat closure areas and removed less than 200 t of pollock. In addition, bycatch rates of Pacific Ocean perch were prohibitively high in areas where pollock aggregations were observed. The 2005 fishery is thought to have resulted in a net loss of revenue for participating vessels. Data on specific bycatch and discard rates for the 2005 fishery are not available due to issues of data confidentiality. In 2006 the Aleut Corporation, in partnership with the Alaska Fisheries Science Center, Adak Fisheries LLC and the owners and operators of the F/V Muir Milach, conducted the Aleutian Islands Cooperative Acoustic Survey Study (AICASS) to test the technical feasibility of conducting acoustic surveys of pollock in the Aleutian Islands using small ( $<32 \mathrm{~m}$ ) commercial fishing vessels (Barbeaux, in press). This work was supported under an exempted fishing permit that allowed
directed pollock fishing within Steller sea lion critical habitat. A total of 932 t of pollock were harvested during this study and biological data collected during this study are treated in the stock assessment as fishery data.

## Data

## Catch estimates

Estimates of pollock catch in the Aleutian Islands Region are derived from a variety of data sources (Table 1A.2). During the early period, the foreign-reported database (held at AFSC) is the main source of information and was used to derive the official catch statistics until about 1980 when the observer data were introduced to provide more reliable estimates. The foreign and joint-venture (JV) blend data takes into account observer data and reported catches and forms the basis of the official catch statistics until 1990. The NMFS Observer data are the raw observed catch estimates and provide an indication of the amount of catch observed relative to the current estimates from the blend data. Estimates of pollock discard levels have been available since 1990. During the years when directed fishing was allowed pollock discards represented a small fraction of the total catch (Table 1A.3).
For the period 1977-1984, the foreign reported catch database was used to partition catches while for 1985-2003, observer data were used. These proportions were then expanded to match the total catch (Table 1A.4; Fig. 1A.1).

The distribution of observed catch differed between the JV years (1977-1989) and the domestic fishery (1989-2002; Fig. 1A.2). In the early period, the JV fishery operated in the deep basin area extending westward to Bowers Ridge and in the eastern most portions of the Aleutian Islands. Some operations took place out to the west but observer coverage was limited. In the recent period (1989-1998, since the Aleutian Islands Region has been closed to directed pollock fishing since 1999) the fishery was more dispersed along the Aleutian Islands chain with no observed catches along Bowers Ridge and fewer operations in the deep basin area. Considering the spatial distribution of these fisheries, we recommended that the Aleutian Islands Region be broken into areas where apparent breaks existed (Fig. 1A.3). These breaks separate the northern "basin" area from the Aleutian Islands chain and split the eastern-most portion of the Aleutian Islands Region from the Aleutian Islands. Two regional partitions were developed, one called NRA (for Near, Rat, and Andreanof Island groups) extending to $170^{\circ} \mathrm{E}$, and another that excludes the eastern portion between $174^{\circ} \mathrm{W}$ and $170^{\circ} \mathrm{W}$. The time series of catch estimates for these two groups is shown in Table 1A.5. In the NRA area west of $174^{\circ} \mathrm{W}$ the fishery tended to concentrate in two distinct locations one on the north side of Atka Island around $174^{\circ} \mathrm{W}$ and the other near $177^{\circ} \mathrm{W}$ northwest of Adak Island. While the overall catch level was relatively low, the fishery moved far to the west in 1998 (Fig. 1A.4).

## Fishery length frequency

The number of hauls and length samples in the NRA region west of $174^{\circ} \mathrm{W}$ are quite small compared with the eastern and northern (basin) areas (Table 1A.6). The differences in the length frequencies appear to be substantial between regions (Barbeaux et al. 2004). During the early period, the region west of $174^{\circ} \mathrm{W}$ longitude was composed of smaller fish. This region also tended to have a broader range of lengths. The Basin region was similar to the eastern most region and the Bogoslof region (during the years when a fishery was allowed there). In the 2005 stock assessment we investigated whether the changes in length frequency distributions for the NRA region west of $174^{\circ} \mathrm{W}$ could be attributed to different seasonal concentrations of fishing. These investigations showed that before 1990, the fishery tended to be more concentrated later in the year, but inter-annually the fishery was consistent in the time between the eastern and western NRA (Barbeaux et al. 2005). We therefore concluded that differences in length distributions observed between these two regions could not be attributed to differences in the time of year in which the fishery was conducted. Intra-annual differences may show a trend that could be consistent with seasonality differences. The occurrence of larger fish later in the time series is likely due to the fishery
targeting on spawning pollock. Pollock average weights-at-age from the early period are lower than the recent period (Table 1A.7). As shown in the 2005 assessment the observed proportion of females in the catch appeared to show a slight decline over this period (Barbeaux et al. 2005).

## Fishery age composition

Catch-at-age composition estimates are made following Kimura (1989) and modified by Dorn (1992). Briefly, length-stratified age data are used to construct age-length keys for each stratum and sex. These keys are then applied to randomly sampled catch length frequency data. The stratum-specific age composition estimates are then weighted by the catch within each stratum to arrive at an overall age composition for each year. Data were collected through shore-side sampling and at-sea observers. The number of age samples and length samples was highly variable over this time period (Table 1A.8). This problem is exacerbated for samples collected from different areas and gears (Table 1A.9). The estimates for catch-age composition are shown in Table 1A.10. The age composition data collected in the 2006 AICASS was used as fishery data.

## Survey data

Bottom trawl survey effort in the Aleutian Islands region has not been as extensive as in the eastern Bering Sea. The National Marine Fisheries Service in conjunction with the Fisheries Agency of Japan completed bottom trawl surveys for the Aleutian Islands region (from $\sim 165^{\circ} \mathrm{W}$ to $\sim 170^{\circ} \mathrm{E}$ ) in 1980, 1983, and 1986. The Alaska Fisheries Science Center's Resource Assessment and Conservation Engineering Division (RACE) conducted bottom trawl surveys in this region in 1991, 1994, 1997, 2000, 2002, 2004 and 2006. Biomass estimates from the surveys conducted in the 1980s ranged between 309 and 779 thousand tons (mean 546). Biomass estimates from the five most recent RACE surveys ranged between 112 and 366 thousand tons (mean 225; Table 1A.11). The biomass estimates from the early surveys are not comparable with the biomass estimates obtained from the RACE trawl surveys because of differences in the net, fishing power of the vessels, and sampling design. In the early surveys, biomass estimates were computed using relative fishing power coefficients (RFPC) and were based on the most efficient trawl during each survey. Such methods will result in pollock biomass estimates that are higher than those obtained using standard methods employed in the RACE surveys. Plotted on a simple catch-per-tow basis, the relative distribution of pollock appears to be variable between years and areas (Fig. 1A.5).
RACE Aleutian Islands bottom trawl (AIBT) surveys prior to 2004 indicate that most of the pollock biomass has been located in the Eastern Aleutian Islands Area (Area 541) and along the north side of Unalaska-Umnak Islands in the eastern Bering Sea region ( $\sim 165^{\circ} \mathrm{W}$ and $170^{\circ} \mathrm{W}$ ). The 2004 Aleutian Islands trawl survey showed the greatest density and estimated biomass in the Unalaska-Umnak area in the eastern Bering Sea region, but only low densities of pollock were observed in the Unalaska-Umnak Islands area in the 2006 survey. If we ignore the biomass estimates from the Unalaska-Umnak area the 2004 and 2006 AIBT surveys are very similar and show a very different pattern of biomass abundance than the 2002 survey (Fig. 1A.5). Within the Aleutian Islands Region (Areas 541, 542, and 543) the 2002 AIBT survey indicated the highest densities and biomass were in the Central Aleutian Islands Area (Area 542) followed by the Eastern (Area 541) and Western areas (Area 543). In the 1991-2000 AIBT surveys the highest biomasses for the NRA Areas were estimated in Area 541 followed by Area 542 and Area 543. The earlier RACE AIBT surveys indicated a decline in pollock biomass in the portion of Area 541 east of $174^{\circ} \mathrm{W}$ longitude from a high of $53,865 \mathrm{t}$ in 1991 to a low of $28,985 \mathrm{t}$ in the 2000 survey. This trend was reversed in the 2002 survey with an estimate of $53,368 \mathrm{t}$, in 2004 with an estimate of $111,250 \mathrm{t}$, and 117,000 $t$ in 2006 (Table 1A.11). In the 1991-2002 surveys a number of large to medium sized tows were encountered throughout the Aleutians indicative of a fairly well distributed population. This is very different from the 2004 and 2006 survey estimates which indicated a low level of pollock abundance in both Area 542 and Area 543, and a much higher pollock density in Area 541. The 2004 survey revealed very few pollock throughout the NRA, except for a single large tow in Seguam pass. The distribution of pollock in the 2006 survey revealed a similar pattern to that of the 2004 survey with high CPUE in the

Seguam pass area. The 2006 survey found a higher concentration of pollock in the Delerof Islands that was not observed in 2004, but is consistent with aggregations observed in 2002. Like the 2004 survey there were very few pollock observed west of $180^{\circ}$ longitude. Since there has not been a substantial fishery in the Aleutians not has there been a substantial change in survey methodology or design, the continued decrease in pollock must be attributed to either a change in catchability due to vertical migration of pollock out of the reach of the bottom trawl, increased emigration of pollock out of the surveyed area, decreased recruitment, increased natural mortality exceeding recruitment, or some combination of these factors. Since the AIBT is limited to within the 500 m isobath the survey biomass estimates do not include mid-water pollock, nor do they include pollock located offshore of the 500 m isobath. These biomass estimates therefore represent an unknown portion of the total biomass. The biomass in this area may be greater if the on-bottom/off-bottom distribution is similar to that of the eastern Bering Sea (Ianelli et al. 2005). In addition, climatic and year class variation may cause a difference in the proportion of pollock available to the bottom trawl survey.

This year we looked at distribution patterns of pollock in relation to temperature and depth. We found that in comparison with pollock distribution observed in the 2004 Bering Sea and 2005 Gulf of Alaska bottom trawl surveys (BSBTS and GOABTS respectively), the distribution observed in the 2004 and 2006 Aleutian Islands bottom trawl surveys (AIBTS) was in a more limited temperature range and generally deeper (Fig. 1A.6). Overall the bottom temperature in the AIBTS was much less variable than in either the BSBTS or GOABTS at depth and ranged between the other surveys with the BSBTS generally cooler at depth and GOABTS warmer at depth. In the AIBTS the highest concentrations of pollock are encountered between 140 m and 300 m , while in the BSBTS the highest concentrations of pollock were above 100 m and above 150 m in the GOABTS. The 2006 AIBTS was colder at shallower depths than in 2004 and pollock concentrations appeared to shift towards deeper water (Fig. 1A.7). The shift of pollock distribution to deeper waters with colder bottom temperatures is consistent with a shift observed in the Bering Sea between 1999, a cold year, and 2004, a warm year (Fig. 1A.8).

## Survey Length Frequencies

There are apparent differences in pollock length-at-age between the Aleutian Island, Bering Sea, and Gulf of Alaska between ages 2 and 9 , with the Aleutian Islands pollock being largest, GOA next, then Bering Sea pollock the smallest at age (Fig. 1A.9). The pollock length frequency collection from the 2006 AIBTS showed the primary mode between 56 and 66 cm similar to previous years and is thought to be primarily composed of $2000 \mathrm{and} /$ or 1999 year-class fish (Fig. 1A.10). There was a small mode between 15 and 25 cm that would be consistent with 1 or 2 year old fish, but much fewer than observed in 2004. The 2004 AIBT survey found a large proportion of small fish (between 10 and 25 cm , indicative of 1 or 2 year old fish) in the NRA area west of $174^{\circ} \mathrm{W}$, but very few small fish east of $174^{\circ} \mathrm{W}$. The 2002 AIBT survey did not find very many small fish anywhere in the Aleutians. There were a large number of small fish observed in the 1994 and 2000 surveys throughout the NRA. The large numbers of 1 or 2 year old size pollock observed in the these surveys were assumed to have entered the fishable population in 1996 and 2002, respectively, and should have stabilized or increased pollock biomass in the Aleutian Islands in recent years.

## Other Surveys

In addition to the bottom trawl survey there has been one echo integration-trawl survey in a portion of the NRA. The R/V Kaiyo Maru conducted a survey between $170^{\circ} \mathrm{W}$ and $178^{\circ} \mathrm{W}$ longitude in the winter of 2002 after completing a survey of the Bogoslof region (Nishimura et al 2002; Fig. 1A.11). Due to difficulties in operating their large mid-water trawl on the steep slope area they determined that their catches in this area were insufficient for accurate species identification and biomass estimation. They did however come up with some preliminary biomass estimations. For the entire area from $170^{\circ} \mathrm{W}$ and $178^{\circ} \mathrm{W}$ longitudes they estimated a biomass of $93,000 \mathrm{t}$ of spawning pollock biomass with between $61,000 \mathrm{t}$ estimated in the NRA east of $173^{\circ} \mathrm{W}$ and $32,000 \mathrm{t}$ in the remainder of the survey area to $178^{\circ} \mathrm{W}$
longitude (Table 1A.12). The largest aggregations in the NRA area were observed at $174^{\circ} \mathrm{W}$ longitude north of Atka Island. Most of the pollock echo sign was observed along the slope of the Aleutian Islands relatively near shore.

In 2006 the Aleut corporation in conjunction with the Alaska Fisheries Science Center, Adak Fisheries LLC, and the F/V Muir Milach conducted the Aleutian Islands Cooperative Acoustic Survey Study (AICASS) to test the technical feasibility of conducting acoustic surveys of pollock in the Aleutian Islands using small commercial fishing vessels ( $<32 \mathrm{~m}$ ) (Barbeaux, in press). The study resulted in three surveys of a $180 \mathrm{~nm}^{2}$ area north of Atka Island between $174.5^{\circ} \mathrm{W}$ and $175.5^{\circ} \mathrm{W}$ longitude (Fig. 1A.12), two surveys of a $72 \mathrm{~nm}^{2}$ area within the larger area, and a $9 \mathrm{~nm}^{2}$ area within this area over a three week period. Since this survey was conducted in conjunction with a commercial fishery the catch-at-age data (Fig. 1A.13) from this survey was used as the Fishery catch at age data within the stock assessment model. Age structure and length frequency data collected during this study reveal that the 2000 and 1999 year classes were dominant in the studied population with strong 1992 and 1996 showing up as well. These year classes are also prevalent in the summer AIBTS and in the Eastern Bering Sea. The biomass within the large area started at $8,810 \mathrm{t}$ for the first survey, was at $7,980 \mathrm{t}$ for the second large survey (a statistically insignificant change), then dropped significantly to $3,639 \mathrm{t}$ for the final survey (Table 1A.13, Fig. 1A.14, and Fig. 1A.15 ). From the first to last survey there was a $59 \%$ decline in pollock abundance even though the fishery only had an $11 \%$ harvest rate (Fig. 1A.16). The reason for this decline, although undoubtedly in part due to fishing, is confounded with possible effects of spawning emigration from the survey area. Although, a trend in the maturity data showed that pollock began showing signs of active spawning only at the end of the study (Fig. 1A.17).

## Analytic Approach

The 2006 Aleutian Islands walleye pollock stock assessment uses the same modeling approach as last year's model; through the Assessment Model for Alaska (here referred to as AMAK). AMAK is a variation of the "Stock Assessment Toolbox" model presented to the plan team in the 2002 Atka mackerel stock assessment, with some small adjustments to the model and a user-friendly graphic interface.

The abundance, mortality, recruitment, and selectivity of the Aleutian Islands pollock were assessed with a stock assessment model constructed with AMAK as implemented using the ADMB software. The ADMB is a $\mathrm{C}++$ software language extension and automatic differentiation library. It allows for estimation of large numbers of parameters in non-linear models using automatic differentiation software developed into C++ libraries (Fournier 1998). The optimizer in ADMB is a quasi-Newton routine (Press et al. 1992). The model is determined to have converged when the maximum parameter gradient is less than a small constant (set to $1 \times 10^{-7}$ ). A feature of ADMB and AMAK is that it includes postconvergence routines to calculate standard errors (or likelihood profiles) for quantities of interest.

## Model structure

The AMAK model models catch-at-age with the standard Baranov catch equation. The population dynamics follows numbers-at-age over the period of catch history with natural and age-specific fishing mortality occurring throughout the 14-age-groups that are modeled (ages 2-15+). Age-2 recruitment in each year is estimated as deviations from a mean value expected from an underlying stock-recruitment curve. Deviations between the observations and the expected values are quantified with a specified error model and cast in terms of a penalized log-likelihood. This overall $\log$-likelihood $(L)$ is the weighted sum of the calculated log-likelihoods for each data component and model penalties. The component weights are inversely proportional to the specified (or in some cases, estimated) variances. Barbeaux et al. 2005, Appendix Tables 1-3 provide a description of the variables used, and the basic equations describing the population dynamics of Aleutian Islands pollock and likelihood equations. The model was modified from that of Barbeaux et al. (2003). These modifications include a feature that allows a user-specified agerange for which to apply the survey (or other abundance index) catchability. For example, specifying the
age-range of 6-10 (as was done for Aleutian Islands pollock) means that the average age-specific catchability of the survey is set to the parametric value (either specified as fixed, as in this assessment, or estimated). Also, in the 2003 assessment age- 1 pollock were explicitly modeled whereas in the work presented here, they were dropped from consideration because observations of age-1 pollock are irregular, and in trials where they were included, they were found to limit the flexibility to incorporate alternative model specifications such as parametric forms of selectivity functions. The quasi ${ }^{1}$ likelihood components and the distribution assumption of the error structure are given below:

| Likelihood Component | Distribution Assumption |
| :--- | :--- |
| Catch biomass | Lognormal |
| Catch age composition | Multinomial |
| Survey catch biomass | Lognormal |
| Survey catch age composition | Multinomial |
| Recruitment deviations | Lognormal |
| Stock recruitment curve | Lognormal |
| Selectivity smoothness (in age-coefficients, survey and fishery) | Lognormal |
| Selectivity change over time (fishery only) | Lognormal |
| Priors (where applicable) | Lognormal |

The age-composition components are heavily influenced by the sample size assumptions specified for the multinomial likelihood. Since sample variances of our catch-at-age estimates are available (Dorn 1992), "effective sample sizes" $\left(\dot{N}_{i, j}\right)$ can be derived as follows (where $i$ indexes year, and $j$ indexes age):

$$
\dot{N}_{i, j}=\frac{p_{i, j}\left(1-p_{i, j}\right)}{\operatorname{var}\left(p_{i, j}\right)}
$$

where $p_{i, j}$ is the proportion of pollock in age group $j$ in year $i$ plus an added constant of 0.01 to provide some robustness. The variance of $p_{i, j}$ was obtained from the estimates of variance in catch-at-age.
Thompson et al., (2003, p. 137) and Thompson (pers. comm.) show that the above is a random variable that has its own distribution. They show that the harmonic mean of this distribution is equal to the true sample size in the multinomial distribution. This property was used to obtain sample size estimates for the surveys and fishery numbers-at-age estimates:

| Fishery data | Year | 1978 | 1979 | 1980 | 1981 | 1982 | 1983 | 1984 | 1985 | 1987 |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
|  | $\dot{N}_{i, \bullet}$ | 246 | 170 | 119 | 215 | 553 | 81 | 296 | 225 | 150 |
|  | Year | 1990 | 1992 | 1993 | 1994 | 1995 | 1996 | 1997 | 1998 | 2006 |
|  | $\dot{N}_{i, \bullet}$ | 199 | 238 | 172 | 327 | 211 | 228 | 30 | 302 | 300 |
| Survey data |  |  |  |  |  |  |  |  |  |  |
|  | Year | 1991 | 1994 | 1997 | 2000 | 2002 | 2004 |  |  |  |
|  | $\dot{N}_{i, \bullet}$ | $1^{*}$ | 740 | 690 | 831 | 1124 | 774 |  |  |  |

*The 1991 value was down-weighted by a factor of 1,000 because the samples collected in that year were not representative of the region considered.

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## Parameters

Parameters estimated independently

## Natural Mortality

For two base models (Model 1 and Model 2A) a natural mortality value of 0.3 was used for comparison with last year's assessment. We started with a value of $M$ based on the studies of Wespestad and Terry (1984) for the Bering Sea. Wespestad and Terry (1984) provide estimates of $M=0.3$ for ages $3+$ (Table 1A.14). Currently, the assessment model does not allow for age-specific natural mortality rates. It should be noted that in general, a higher natural mortality rate for age 2 pollock may be more appropriate (e.g., Ianelli et al. 2003) and that this model differs from the Eastern Bering Sea model in this manner. In the future, we will be investigating methods to improve AMAK to include age varying natural mortality.

## Length and Weight at Age

We estimated length and weight-at-age separately for the survey and for the fishery. We obtained survey estimates from AIBT surveys and computed fishery estimates from observer data and the 2006 AICASS. For the time period between 1978 and 1990 the von Bertalanffy growth curve parameters and length weight regression parameters for length and weight at age estimates for surveys were estimated for the 1980, 1983, and 1986 AIBT surveys (Table 1A.15). For the time period between 1990 and 2006 we calculated the average length at age by weighted averages by age and calculated the length-weight relationships using linear regression analysis. Data for these analyses were retrieved from the Resource Assessment and Conservation Engineering Division's (RACE) survey database. Length and weight-at-age data were available for the 1991, 1994, 1997, 2000, 2002, and 2004 AIBT surveys. For years without survey length and weight-at-age data we used the mean values at age for the two nearest surveys (Table 1A.16). Fishery data east of $174^{\circ} \mathrm{W}$ longitude were excluded from the data set for calculating length and weight-at-age. For the fishery, we used year (when available) and age-specific estimates of average weights-at-age as computed from the fishery age and length sampling programs from data collected west of $174^{\circ} \mathrm{W}$. These values (Table 1A.17) are important for converting model estimated catch-at-age (in numbers) to estimated total annual harvests (by weight).

## Maturity at Age

The maturity at age schedule is based on the studies of Wespestad and Terry (1984; Table 1A.18). An updated analysis on maturity-at-age using more recent data was presented in the 2005 Bering Sea pollock stock assessment. However, the EBS data collected in 2002 and 2003 are in agreement with that observed by Westpestad and Terry (1984), and a change in model configuration is not warranted at this time.

## Parameters estimated conditionally

Deviations between the observations and the expected values are quantified with a specified error structure. Lognormal error is assumed for estimates of survey and fishery catch, and a multinomial error structure is assumed for analysis of the survey and fishery age compositions. These error structures are used to estimate the following parameters conditionally within the model.

## Fishing Mortality

Fishing mortality in all models was parameterized to be separable with both an age component (selectivity) and a year component. In all models selectivity is conditioned so that the mean value over all ages will be equal to one. To provide regularity in the age component, a penalty was imposed on sharp shifts in selectivity between ages using the sum of squared second differences. In addition, the age component parameters are assumed constant for the last 4 age groups (ages 12-15). Finally, selectivity was allowed to vary over time. The model was set with controls selecting the degree to which selectivity is allowed to change between ages and over time.

## Survey Catchability

For the bottom trawl survey, survey catchability-at-age follows the parameterization similar to the fishery selectivity-at-age presented above. The catchability-at-age relationship is modeled with a smoothed nonparametric relationship that can take on any shape (with penalties controlling the degree of change and curvature specified by the user). To provide regularity in the age component, a penalty was imposed on sharp shifts in catchability-at-age between ages using the sum of squared second differences. In addition, the age component parameters are assumed constant for the last 4 age groups (ages 12-15). As noted above, the model allows specification of the age-range over which the catchability parameter is applied. For Aleutian Islands pollock, ages 6-10 were selected to have the average catchability (factoring selectivity components) equal to the catchability parameter value.
In the 2004 Aleutian Islands pollock stock assessment, the focus of our analysis was to evaluate a key model assumption: the extent to which the NMFS summer bottom trawl survey catchability should be estimated by the available data (resulting in very high stock sizes) or constrained to be close to a value of 1.0 (implying that the area-swept survey method during the summer months reasonably applies to a fishery that will likely occur during the winter). We provided evidence that suggests that fixing the value of survey catchability to 1.0 is unreasonable. However, recognizing that no other information is available to "anchor" the assessment model to an absolute biomass level, the authors were reluctant to proceed with specifying influential prior distributions on catchability values. The effects of the fishery on the pollock population dynamics appear to be poorly determined given the available data. This could be due to a number of factors including: characteristics of Aleutian Islands pollock relative to adjacent regions, poor quality data, and the possibility that the fishing effects are minor relative to other factors. The latter point is likely to be true at least for the recent period where the fishery removals have been minor since 1999. We have therefore selected a fixed catchability value of 1.00 for our 2006 preferred alternative models.

## Recruitment

A reparameterized form of the Beverton-Holt stock recruitment relationship based on Francis (1992) was used. Values for the stock recruitment function parameters $\alpha$ and $\beta$ are calculated from the values of $R_{0}$ (the number of 0 -year-olds in the absence of exploitation and recruitment variability) and the "steepness" (h) of the stock-recruit relationship. The "steepness" parameter is the fraction of $R_{0}$ to be expected (in the absence of recruitment variability) when the mature biomass is reduced to $20 \%$ of its pristine level (Francis 1992). As an example, a value of $h=0.8$ implies that at $20 \%$ of the unfished spawning stock size will result in an expected value of $80 \%$ of the unfished recruitment level. The steepness parameter $(h)$ was estimated with a prior of 0.7 and CV of 0.2 , and sigma $r$ was set at 0.6 for all model runs.

## Natural Mortality

For Model 2B natural mortality (M) was estimated within the model using an uninformative prior starting with a value of 0.3 with a CV of 0.2. The addition of the 2006 catch-at-age data from the 2006 AICASS allowed for improved model stability while estimating natural mortality.

## Model evaluation

Three models were evaluated for this year's stock assessment (Model 1, Model 2A, and Model 2B). All three models are configured with a survey catchability of 1.0 , a stock recruitment steepness parameter centered on 0.7 with a CV of 0.2 as a (normal) prior distribution and sigma $r$ of 0.6 . The data configuration for the three models differ in that Model 1 contains only survey data from the NRA area west of $174^{\circ} \mathrm{W}$ longitude and Model 2A and Model 2B contain survey data from the entire NRA region. Model 2A and Model 2B differ in that natural mortality is set at 3.0 in Model 2A, but is estimated within Model 2B with a prior of 0.3 and CV of 0.2 . Models 1 and 2A have the same parameterization as Models 1 and 2 in the 2005 Aleutian Islands pollock stock assessment with the addition of the 2006 survey biomass estimate and the 2006 AICASS catch-at-age estimates.

Relative differences in model fits are shown in Table 1A. 19 and key results are presented in Table 1A.20. By including the survey biomass from the area east of $174^{\circ} \mathrm{W}$ Model 2A and 2B show a marked improvement in fit from Model 1 based on lowest quasi-likelihood. This is primarily due to a better fit to the survey index, but Models 2A and 2B provide a better fit to all of the data components. A better fit to the survey index by Models 2A and 2B can be attributed to the lower intra-annual variability in the NRAwide biomass estimate with a much smoother trend which allows for a better fit to the model. The results from the 2004 and 2006 summer bottom trawl surveys are not consistent with the assumed stock delineation proposed in Model 1, and more analyses need to be conducted to determine a tenable stock delineation. Including an internal estimate of natural mortality in Model 2B improves the model fit over Model 2A. In particular Model 2B provides an improved fit to the age composition data for both the survey and fishery.

For all three models the fit to the survey data is relatively poor, but not surprisingly so, given the estimates of variance for the individual survey point estimates and the high intra-annual variability of the estimates. For all models the fit to the survey age composition data was excellent, except for the 1991 data which, for sampling reasons, was given less weight than for the other years. Results of fits to the fishery age-composition data were much poorer, the high variability in the age data probably reflects the diversity in sampling locations for the fishery in different years. The time-varying selectivity patterns estimated by the models show only slight changes for the survey, but a relatively large shift (to older fish) after 1990 for the fishery data coinciding with the change from a foreign fishery to a domestic fishery targeting spawning aggregations. The estimated total biomass trends for the three models diverge considerably (Fig. 1A.18). Differences between Model 1 and Model 2A are due to differences in the survey estimates for the two areas, differences between Model 2A and Model 2B are due to the lower estimated natural mortality in Model 2B. Model 2B MCMC results show that the natural mortality (M) estimate is stable with a mean of 0.235 and $\mathrm{CV}=0.10$ (Table 1A.21, Fig. 1A.19, and Fig. 1A. 20 ). A lower estimate for natural mortality than that estimated for the Bering Sea pollock is supported by estimates M for Bogoslof area pollock conducted by Ianelli et al. (2005) resulting in an estimate of M of 0.26 and by Westpestad and Terry (1984) estimate of $\mathrm{M}=0.2$ for Aleutian Basin pollock.

Because of the improved fit (lowest negative quasi-likelihood) to the available data and stable estimate for M, Model 2B was chosen as the reference model. Fits to the survey index, survey age composition, fishery age composition, and selectivity trend are show in Fig. 1A.21, Fig. 1A.22, Fig. 1A.23, and Fig. 1A. 24 respectively. As stated above the Model 2B fit to the bottom trawl survey index is better than Model 1, but remains relatively poor, the model has difficulty fitting the low 1994 survey index in light of the high level of catch observed in 1995 and the increase in the survey index in 1996. The model predicts a decline in abundance that corresponds with the high fishing mortality observed in the mid- to late1990s. Model 2B fits to the survey and fishery age composition data are for the most part very good.

## Results

## Abundance and exploitation trends

As indicated in the 2004 stock assessment analysis (Barbeaux et al, 2004), the abundance trend is highly conditioned on the assumptions made about the area-swept survey trawl assumptions on catchability. Even with catchability fixed at 1.0 , the uncertainty in the trend and level is very high. Bearing in mind the high degree of uncertainty, the total biomass trend (Table 1A. 22 and Fig. 1A.25) appears to have increased from 1999 to 2003 after cessation of directed fishing in the area, but from 2003 to 2006 was stable to decreasing. In this assessment total biomass is pollock at age 2 and above.

Spawning biomass appears to have been greatly influence by the high exploitation in the late 1990s (Fig. 1A.26). The highest fishing mortality occurred in 1995 ( $\mathrm{F}=0.71$ and Catch/biomass $=0.20$ ) when the fishery harvested more than $75 \%$ of the 1994 survey biomass estimate (Table 1A.23). The reference
model shows continued higher than average exploitation in 1997 and 1998 with $\mathrm{F}=0.67$ for both years. The spawning biomass has been increasing since 2000.

## Recruitment

Estimates of recruitment (at age 2) are estimated with high variance (Table 1A. 24 and Table 1A. 25 , Fig. 1A.27). The 1978 year-class is the largest ( 308 million age 2 recruits). The 1989 year class is the second largest ( 137 million age 2 recruits), and the 2000 the third ( 107 million age 2 recruits). Whether AI pollock recruitment is synchronous with other areas is an open question (e.g., the 1978, 1989, and 2000 year classes are also strong in the EBS region, Ianelli et al. 2005). An alternative explanation is that movement between other areas may affect year-class abundance. The extent to which adjacent stocks interact is an active area of research.

## Projections and harvest alternatives

For projection purposes we use the yield projections estimated for reference Model 2B. Because a directed fishery on pollock was banned between 1999 and 2004, and because the 2005 and 2006 fisheries were greatly limited, we do not believe the 2006 AI pollock selectivity-at-age assumed in these models would be relevant to a fully utilized directed fishery. For projections we used the selectivity-at-age derived from the 2005 EBS pollock assessment (Ianelli, et al 2005), because a current estimate for selectivity-at-age for a directed pollock fishery in the Aleutians is not available (Table 1A.26). The selectivity-at-age for the EBS pollock would be applicable if an Aleutian Islands Pollock fishery was prosecuted by EBS pollock fishing vessels. Catchability within the reference model (Model 2B) is fixed at 1.0. The reference model excludes fishery data from east of $174^{\circ} \mathrm{W}$ longitude, but includes all AIBTS data from west of $170^{\circ} \mathrm{W}$ longitude.

## Reference fishing mortality rates and yields

Amendment 56 to the BSAI Groundfish Fishery Management Plan (FMP) defines "overfishing level" (OFL), the fishing mortality rate used to set OFL ( $F_{O F L}$ ), the maximum permissible ABC , and the fishing mortality rate used to set the maximum permissible $\mathrm{ABC}\left(\max F_{A B C}\right)$. The fishing mortality rate used to set $\mathrm{ABC}\left(F_{\text {ABC }}\right)$ may be less than or equal to this maximum permissible level. The overfishing and maximum allowable ABC fishing mortality rates are given in terms of percentages of unfished female spawning biomass ( $F_{S P R \%}$ ), on fully selected age groups. The associated long-term average female spawner biomasses that would be expected under average estimated recruitment from 1983-2006 for Model 2B would be 67.49 million age 2 fish and $F$ equal to $F_{40 \%}$ and $F_{35 \%}$ are denoted $B_{40 \%}$ and $B_{35 \%}$, respectively. We chose to exclude the 1978 extreme recruitment event, treating it as an anomalous event therefore allowing a more conservative estimate of future recruitment. The Tiers require reference point estimates for biomass level determinations. We present the following reference points for NRA pollock for Tier 3 of Amendment 56. For our analyses, we estimated the following values from Model 2B:

| Female spawning biomass | Model 2B |
| ---: | ---: |
| $B_{100 \%}$ | $100,945 \mathrm{t}$ |
| $B_{40 \%}$ | $40,378 \mathrm{t}$ |
| $B_{35 \%}$ | $35,331 \mathrm{t}$ |
| $B_{07}$ | $82,210 \mathrm{t}$ |

## Specification of OFL and Maximum Permissible ABC

For Model 2B, the projected year 2007 female spawning biomass $\left(S B_{07}\right)$ is estimated to be $82,210 \mathrm{t}$, above the $B_{40 \%}$ value of $40,378 \mathrm{t}$ placing NRA pollock in Tier 3a. The maximum permissible ABC and OFL values under Tier 3a are:

Model 2B Tier 3a:

| Harvest Strategy | FSPR\% | Fishing Mortality Rate | 2007 Projected yield $(\mathrm{t})$ |
| :---: | :---: | :---: | :---: |
| max $F_{A B C}$ | $F_{40 \%}$ | 0.29 | $44,470 \mathrm{t}$ |
| $F_{\text {OFL }}$ | $F_{35 \%}$ | 0.38 | $54,540 \mathrm{t}$ |

Under Tier 5 with new model estimated natural mortality of 0.235 the 2007 ABC would be $16,800 \mathrm{t}$ $(94,992 \mathrm{tx} 0.75 \times 0.235=2 \mathrm{t})$ and under Tier 5 with an assumed natural mortality of 0.3 the 2007 ABC would be 21,370 t .

## ABC Considerations and Recommendation

## ABC Considerations

There remains considerable uncertainty in the Aleutian Islands pollock assessment. We've noted some concerns below:

1) The amount of interaction between the Aleutian stock and the Eastern Bering Sea stock is unknown. It is evident that some interaction does occur and that the abundance and composition of the eastern portion of the Aleutian Islands stock is highly confounded with that of the Eastern Bering Sea stock. Overestimation of the Aleutian Islands pollock stock productivity due to an influx of Eastern Bering Sea stock is a significant risk.
2) As assessed in the 2004 AI pollock stock assessment (Barbeaux et al. 2004), AIBT survey catchability is probably less than 1.0 , but we have no data to concretely anchor the value at anywhere less than 1.0. We therefore employed a default value for catchability of 1.00 (a conservative estimate). This provides a conservative total biomass estimate.
3) AIBT survey estimates of biomass are uncertain with an average CV of 0.36 . The 2002, 2004, and 2006 estimates are especially uncertain with CVs of $0.38,0.78$, and 0.48 respectively. This results in considerable uncertainty in the projections.

## ABC Recommendations

The pollock spawning stock biomass in the NRA appears to be increasing, even in light of the latest low values for the AIBT survey and the total biomass appears to be stable. The estimated female spawning biomass projected for 2007 is expected to be $82,210 \mathrm{t}$. The total age $3+$ biomass for 2007 is expected to be $167,581 \mathrm{t}$. The year 2007 maximum permissible ABC based on $\mathrm{F}_{40 \%}$, is $44,470 \mathrm{t}(\mathrm{F}=0.29)$.

## Standard Harvest Scenarios and Projection Methodology

A standard set of projections is required for each stock managed under Tiers 1, 2, or 3, of Amendment 56. This set of projections encompasses seven harvest scenarios designed to satisfy the requirements of Amendment 56, the National Environmental Policy Act, and the Magnuson-Stevens Fishery Conservation and Management Act (MSFCMA).
For each scenario, the projections begin with the vector of 2006 numbers at age estimated in the assessment. This vector is then projected forward to the beginning of 2007 using the schedules of natural mortality and selectivity described in the assessment and the best available estimate of total (year-end) catch for 2006. In each subsequent year, the fishing mortality rate is prescribed on the basis of the spawning biomass in that year and the respective harvest scenario. In each year, recruitment is drawn from an inverse Gaussian distribution whose parameters consist of maximum likelihood estimates determined from recruitments estimated in the assessment. Spawning biomass is computed in each year based on the time of peak spawning and the maturity and weight schedules described in the assessment. Total catch is assumed to equal the catch associated with the respective harvest scenario in all years. This
projection scheme is run 1000 times to obtain distributions of possible future stock sizes, fishing mortality rates, and catches.
Five of the seven standard scenarios will be used in an Environmental Assessment prepared in conjunction with the final SAFE. These five scenarios, which are designed to provide a range of harvest alternatives that are likely to bracket the final TAC for 2007, are as follows (a " $m a x F_{A B C}$ " refers to the maximum permissible value of $F_{A B C}$ under Amendment 56):

Scenario 1: In all future years, $F$ is set equal to $\max F_{A B C}$. (Rationale: Historically, TAC has been constrained by ABC, so this scenario provides a likely upper limit on future TACs.)
Scenario 2: In all future years, $F$ is set equal to a constant fraction of $\max F_{A B C}$, where this fraction is equal to the ratio of the $F_{A B C}$ value for 2007 recommended in the assessment to the max $F_{A B C}$ for 2007. (Rationale: When $F_{A B C}$ is set at a value below $\max F_{A B C}$, it is often set at the value recommended in the stock assessment.)

Scenario 3: In all future years, $F$ is set equal to $50 \%$ of $\max F_{A B C}$. (Rationale: This scenario provides a likely lower bound on $F_{A B C}$ that still allows future harvest rates to be adjusted downward when stocks fall below reference levels.)
Scenario 4: In all future years, $F$ is set equal to the 2002-2006 average $F$. (Rationale: For some stocks, TAC can be well below ABC, and recent average $F$ may provide a better indicator of $F_{T A C}$ than $F_{A B C}$.)
Scenario 5: In all future years, $F$ is set equal to zero. (Rationale: In extreme cases, TAC may be set at a level close to zero.)

Two other scenarios are needed to satisfy the MSFCMA's requirement to determine whether a stock is currently in an overfished condition or is approaching an overfished condition. These two scenarios are as follow (for Tier 3 stocks, the MSY level is defined as $B_{35 \%}$ ):
Scenario 6: In all future years, F is set equal to FOFL. (Rationale: This scenario determines whether a stock is overfished. If the stock is expected to be 1) above its MSY level in 2007 or 2) above $1 / 2$ of its MSY level in 2008 and above its MSY level in 2007 under this scenario, then the stock is not overfished.)
Scenario 7: In 2007 and 2008, $F$ is set equal to $\max F_{A B C}$, and in all subsequent years, $F$ is set equal to $F_{\text {OFL }}$. (Rationale: This scenario determines whether a stock is approaching an overfished condition. If the stock is expected to be above its MSY level in 2019 under this scenario, then the stock is not approaching an overfished condition.)

The author included one more scenario in order to take into consideration of the Exempted fishing permit studies currently being carried out.

Scenario 8: In 2007 the TAC is set at 4,000 t , increased to $7,600 \mathrm{t}$ for 2008 and 2009 then increased to $19,000 \mathrm{t}$ from 2010 through 2019. (Rationale: This scenario seems like a plausible outcome given the current proposal for conducting the Aleutian Islands Cooperative Acoustic Survey Study under an exempted fishing permit, 7,600 t is $40 \%$ of the 19,000 cap which is the A-season allocation, and 19,000 is the AI pollock cap by law).

## Projections and status determination

The projected yields, female spawning biomass, and the associated fishing mortality rates for the seven harvest strategies for the reference model are shown in Table 1A.27. Under a harvest strategy of $F_{40 \%}$ (Scenario 1), female spawning biomass is projected to be above $B_{40 \%}$ for all 13 years of the projection Fig. 1A. 28 and Fig. 1A.29. Female spawning biomass is projected to fall below $B_{40 \%}$ when fishing at $F_{\text {OFL }}$ (Scenarios $6 \& 7$, Table 1A.27) in 2009 and remain below for the remainder of the projection. Please note
again that the fishing mortality rates are prescribed on the basis of the harvest scenario and the spawning biomass in each year. Thus, fishing mortality rates may not be constant within the projection if spawning biomass drops below $B_{40 \%}$ in any run.

The associated long-term average female spawner biomass that would be expected under average estimated recruitment from 1978-2003 (67.49 million age 2 fish) and $F=F_{35 \%}$, denoted $B_{35 \%}$ is estimated to be $35,331 \mathrm{t}$. This value ( $B_{35 \%}$ ), is used in the status determination criteria. Female spawning biomass for $2007(82,210 \mathrm{t})$ is projected to be above $B_{35 \%}$ thus, the NRA pollock stock would be determined to be above its minimum stock size threshold (MSST) and is not overfished. Female spawning biomass for 2009 is projected to be above $B_{35 \%}$ in scenario 7, thus the NRA pollock stock is not expected to fall below its MSST in two years and is not approaching an overfished condition.

Scenario 8 (Fig. 1A.28, Fig. 1A.29, and Table 1A.27) shows that a harvest of $4,000 \mathrm{t}$ for 2007 an increase to $7,600 \mathrm{t}$ for 2008 and 2009, and an increase to $19,000 \mathrm{t}$ through 2019 would be a conservative harvest strategy under Model 2B, resulting in $\mathrm{F}_{07}=0.002$ and a slowly declining female spawning stock biomass trajectory approaching $\mathrm{B}_{40}$ by 2019.

## Ecosystem Considerations

Pollock is a commercially important species which is also important as prey to other fish, birds, and marine mammals, and has been the focus of substantial research in Alaskan ecosystems, especially in the Gulf of Alaska (GOA; e.g. Hollowed et al 2000). To determine the ecosystem relationships of juvenile and adult pollock in the Aleutian Islands (AI), we first examine the diet data collected for pollock. Diet data are collected aboard NMFS bottom trawl surveys in the AI ecosytem during the summer (May August). In the AI, a total of 1458 pollock stomachs were collected between the 1991 and 1994 bottom trawl surveys ( $\mathrm{n}=688$ and 770 , respectively) and used in this analysis. The diet compositions reported here reflect the size and spatial distribution of pollock in each survey (see Appendix A, "Diet calculations" for detailed methods). Juvenile pollock were defined as fish less than 20 cm in length, which roughly corresponds to 0 and 1 year old fish in the stock assessment, and adult pollock were defined as fish 20 cm in length or greater, roughly corresponding to age $2+$ fish.
In the AI, pollock diet data reflects a closer connection with open oceanic environments than in either the Eastern Bering Sea (EBS) or the GOA. Similar to the other ecosystems, euphausiids and copepods together make up the largest proportion of AI adult pollock diet ( $29 \%$ and $19 \%$, respectively); however, it is only in the AI that adult pollock rely on mesopelagic forage fish in the family Myctophidae for $24 \%$ of their diet, and AI juvenile pollock have a lower proportion of euphausiids and a higher proportion of gelatinous filter feeders than in the GOA or EBS (Fig. 1A.30, left panels). We can take this diet composition information and convert it to broad ranges of tons consumed annually by pollock in the AI using the Sense routine (Aydin et al. in review), which incorporates information on pollock consumption derived from the stock assessment (see Appendix A, "ration calculations" for detailed methods), as well as uncertainty in all other food web model parameters. As estimated by the Sense routine, AI adult pollock consumed between 100 and 900 thousand metric tons of euphausiids annually during the early 1990s, with similar ranges of myctophid and copepod consumption. Juvenile AI pollock consumed an additional estimated 100 to 900 thousand tons of copepods per year (Fig. 1A.30, right panels).

Using diet data for all predators of pollock and consumption estimates for those predators, as well as fishery catch data, we next estimate the sources of pollock mortality in the AI (see detailed methods in Appendix A). Sources of mortality are compared against the total production of pollock as estimated in the AI pollock stock assessment model. In the AI, integration of this single species information with predation within the food web model suggests that most adult pollock mortality was caused by the pollock trawl fishery during the early 1990s (48\%; Fig. 1A.31, left panels). (Fishery catch of pollock in the AI has subsequently declined to less than half the early 1990s catch by the late 1990s, and the directed
fishery was closed in 1999 (Ianelli et al 2005). Therefore, AI pollock likely now experience predation mortality exceeding fishing mortality as in the EBS and GOA ecosystems.) The major predators of AI adult pollock are Pacific cod, Steller sea lions, pollock themselves, halibut, and skates. In the AI, juvenile pollock have a very different set of predators from adult pollock; Atka mackerel cause most juvenile pollock mortality ( $71 \%$ ). Estimates of the tonnage of adult pollock consumed by predators from the Sense routines (Aydin et al in review) ranged from 8 to 27 thousand tons consumed by cod annually during the early 1990s, while Atka mackerel were estimated to consume between 75 and 410 thousand tons of juvenile pollock annually in the AI ecosystem (Fig. 1A.31, right panels).

After reviewing the diet compositions and mortality sources of pollock in the AI, we shift focus slightly to view pollock and the pollock fishery within the context of the larger AI food web. When viewed within the AI food web, the pollock trawl fishery (in red; Fig. 1A.32) is a relatively high trophic level (TL) predator which interacts mostly with adult pollock, but also with many other species (in green; Fig. 1A.32). The diverse pollock fishery bycatch ranges from high TL predators such as salmon sharks, sleeper sharks, and arrowtooth flounder, to mid TL pelagic forage fish and squid, to low TL benthic invertebrates such as crabs and shrimp, but all of these catches represent extremely small flows. Because the pollock trawl fishery contributes significant fishery offal and discards back into each ecosystem, these flows to fishery detritus groups are represented as the only "predator consumption" flows from the fishery; the biomass of retained catch represents a permanent removal from the system.

In the AI food web model, we included detailed information on bycatch for each fishery. This data was collected in the early 1990s when the AI pollock fishery was much larger than it is at present. During the early 1990's, the pollock trawl fishery was extremely species-specific in the AI ecosystem, with pollock representing over $90 \%$ of its total catch by weight (Fig. 1A.33). No single bycatch species accounted for more than $1 \%$ of the catch. Although these catches are small in terms of percentage, the high volume pollock fisheries still account for the majority of bycatch of pelagic species in the BSAI management areas, including smelts, salmon sharks, and squids (Gaichas et al 2004).
The intended target of the pollock trawl fishery is also a very important prey species in the wider AI food web. When both adult and juvenile pollock food web relationships are included, over two thirds of all species groups turn out to be directly linked to pollock either as predators or prey in the food web model (Fig. 1A.34). In the AI, the significant predators of pollock (blue boxes joined by blue lines) include halibut, cod, Alaska skates, Steller sea lions, and the pollock trawl fishery. Significant prey of pollock (green boxes joined by green lines) are myctophids, euphausiids, copepods, benthic shrimps, and amphipods, with juveniles preying on the euphausiids and copepods.
We can investigate whether these differences in pollock diet, mortality, and relationships between the EBS and AI might suggest different ecosystem roles for pollock in these areas. We use the diet and mortality results integrated with information on uncertainty in the food web using the Sense routines (Aydin et al in review) and a perturbation analysis with each model food web to explore the ecosystem relationships of pollock further. Two questions are important in determining the ecosystem role of pollock: which species groups are pollock important to, and which species groups are important to pollock?
First, the importance of pollock to other groups within the AI ecosystem was assessed using a model simulation analysis where pollock survival was decreased (mortality was increased) by a small amount, $10 \%$, over 30 years to determine the potential effects on other living groups. This analysis also incorporated the uncertainty in model parameters using the Sense routines, resulting in ranges of possible outcomes which are portrayed as $50 \%$ confidence intervals (boxes in Fig. 1A.35) and $95 \%$ confidence intervals (error bars in Figure 1A. 6). Species showing the largest median changes from baseline conditions are presented in descending order from left to right. Therefore, the largest change resulting from a $10 \%$ decrease in pollock survival in both ecosystems is a decrease in adult pollock biomass, as might have been expected from such a perturbation. However, the decrease in pollock biomass resulting
from the $10 \%$ survival reduction is uncertain in AI : the $50 \%$ intervals range from a $5-37 \%$ decrease in the AI (Fig. 1A.35, upper panel). Along with the decrease in pollock biomass predicted in this simulation is a decrease in pollock fishery catch. The next largest median effect is on juvenile pollock, which are predicted to decrease in $50 \%$ of feasible ecosystems, but he $95 \%$ interval includes zero, suggesting that the decrease is uncertain. The simulation further suggests the possibility that herring, Atka mackerel, and other miscellaneous deepwater fish might increase slightly as a result of a decrease in pollock survival; however, for all of these species groups the $95 \%$ intervals cross zero, so the direction of change is uncertain. Therefore, this analysis suggests that in the AI ecosystem during the early 1990's, pollock were most important to themselves, and to the pollock fishery.

To determine which groups were most important to pollock in each ecosystem, we conducted the inverse of the analysis presented above. In this simulation, each species group in the ecosystem had survival reduced by $10 \%$ and the system was allowed to adjust over 30 years. The strongest median effects on AI adult pollock are presented in Fig. 1A. 35 (lower panel). The largest effect on adult pollock was the reduction in biomass resulting from the reduced survival of juvenile pollock, although the $95 \%$ intervals include zero change, indicating considerable uncertainty in this result. (The same caution applies to the interpretation of all of the results of this simulation as all of the $95 \%$ intervals contain zero). It is interesting, however, that reduced survival of juvenile Atka mackerel had a larger median effect on adult pollock biomass than the direct effect of reduced adult pollock survival itself (Fig. 1A.35, lower panel), and that the effect is positive. Adult Atka mackerel show the same pattern, which is likely explained by the amount of mortality caused by Atka mackerel on juvenile pollock in the AI food web model (see Fig. 1A.31, lower panels). Reduced survival of Atka mackerel adults or juveniles apparently relieves considerable mortality on juvenile pollock in this model, accounting for the increases in pollock biomass predicted (which is similar in magnitude to the increase predicted from reducing the pollock fishery catch by $10 \%$ ). Although this result is uncertain, it does indicate an important interaction between two commercially important species in the AI ecosystem which might be further investigated.

## Ecosystem effects on Aleutian Islands Walleye Pollock

The following ecosystem considerations are summarized in Table 1A. 28.

## Prey availability/abundance trends

Adult walleye pollock in the Aleutian Islands consume a variety of prey, primarily large zooplankton, copepods, and myctophids. Figure 36 highlights the trophic level of pollock in relation to its prey and predators. No time series of information is available on Aleutian Islands for large zooplankton, copepod, or myctophid abundance.

## Predator population trends

The abundance trend of Aleutian Islands Pacific cod is decreasing, and the trend for Aleutian Islands arrowtooth flounder is relatively stable. Northern fur seals are showing declines, and Steller sea lions have shown some slight increases. Declining trends in predator abundance could lead to possible decreases in walleye pollock mortality. The population trends of seabirds are mixed, some increases, some decreases, and others stable. Seabird population trends could affect young-of-the-year mortality.

## Changes in habitat quality

The 2006 Aleutian Islands summer bottom temperatures indicated that water temperatures were slightly cooler at shallower depths than 2004, but was otherwise an average year. Bottom temperatures could possibly affect fish distribution, but there have been no directed studies, and there is no time series of data which demonstrates the effects on Aleutian Islands walleye pollock.

## Al pollock fishery effects on the ecosystem

## AI pollock fishery contribution to bycatch

The 2006 AI pollock fishery was conducted in conjunction with the 2006 AICASS, Pacific Ocean perch (POP) was the most substantial bycatch species and made up $3 \%$ of the catch. The AI pollock fishery opening in 2005 was limited to only four hauls, within these four hauls the bycatch level of POP was very high ( $\sim 50 \%$ ). Besides the lack of commercially harvestable levels of pollock, the high levels of POP bycatch convinced fishers to discontinue the fishery in 2005. Prior to 1998, levels of bycatch in the pollock fishery of prohibited species, forage, HAPC biota, marine mammals and birds, and other sensitive non-target species was very low compared to other fisheries in the region.

## Concentration of AI pollock catches in time and space

Since the AI pollock fishery is expected to be conducted in conjunction with the 2007 AICASS, catch is expected to be limited to $3,000 \mathrm{t}$ and distributed evenly between $173^{\circ} \mathrm{W}$ and $179^{\circ} \mathrm{W}$ longitude. The impacts of this fishery due to temporal and spatial concentration are not expected be substantial due to the relatively low fishing mortality expected.

## Al pollock fishery effects on amount of large size walleye pollock

The AI pollock fishery in the Aleutian Islands was closed between 1999 and 2005. There was only a very limited fishery in $2005(<200 t)$ and $2006(932 \mathrm{t})$. Year to year differences observed in the previous seven years can not be attributed to the fishery and must be attributed to natural fluctuations in recruitment. Fishers have indicated that the larger pollock in the Aleutian Islands will be targeted. But the low level of fishing mortality is not expected to greatly affect the size distribution of pollock in the AI.

## Al pollock fishery contribution to discards and offal production

The 2007 Aleutian Islands pollock fishery, if pursued, is expected to be conducted by catcher vessels delivering unsorted catch to the Adak Fisheries LLC. processing plant, and therefore very little discard or offal production is expected from this fishery.

## AI Pollock fishery effects on AI pollock age-at-maturity and fecundity

The effects of the fishery on the age-at-maturity and fecundity of AI pollock are unknown. No studies on AI pollock age-at-maturity or fecundity have been conducted. Studies are needed to determine if there have been changes over time and whether changes could be attributed to the fishery.

## Data gaps and research priorities

Very little is known about the AI pollock stock structure and their relation to Western Bering Sea, Eastern Bering Sea, Gulf of Alaska, Bogoslof and Central Bering Sea pollock. Genetic work on the relationship of NRA pollock to other stocks in the North Pacific is essential for further assessment work. Tissue samples were collected during the 2006 AICASS for this analysis but genetic analysis of these samples are waiting on funding. In addition, studies on the migration of pollock in the North Pacific should be explored in order to obtain an understanding of how the stocks relate spatially and temporally and how neighboring fisheries affect local abundances. Time series data sets on prey species abundance in the Aleutian Islands would be useful for a more clear understanding of ecosystem affects. Studies to determine the impacts of environmental indicators such as temperature regime on AI Aleutian pollock are needed. Currently, we rely on studies from the eastern Bering Sea for our estimates of life history parameters (e.g. maturity-atage, fecundity, and natural mortality) for the NRA pollock. Studies specific to the NRA to determine whether there are any differences from the eastern Bering Sea stock and whether there have been any changes in life history parameters over time would be informative.

## Summary

Model 2B Parameters
Natural Morality: $\quad \mathrm{M}=0.235$
Initial Biomass (1978): $B_{0}=280,040 t$

2007

| Maximum permissible ABC : | Tier 3a Model 2B F $\mathrm{F}_{40 \%}=0.29$ Tier 5 ( $\mathrm{M}=0.235$ ) | $\begin{aligned} & \text { yield }=44,470 \mathrm{t} \\ & \text { yield }=16,800 \mathrm{t} \end{aligned}$ |
| :---: | :---: | :---: |
|  | Tier $5(\mathrm{M}=0.3)$ | yield $=21,370 \mathrm{t}$ |
| Overfishing (OFL): | Tier 3a Model 2B $\mathrm{F}_{35 \%}=0.38$ Tier $5(\mathrm{M}=0.235)$ | $\begin{aligned} & \text { yield }=54,540 \mathrm{t} \\ & \text { yield }=22,350 \mathrm{t} \end{aligned}$ |
|  | Tier 5 ( $\mathrm{M}=0.3$ ) | yield $=28,500 \mathrm{t}$ |
| 2008 |  |  |
| Maximum permissible ABC : | Tier 3a Model 2B $\mathrm{F}_{40 \%}=0.29$ | yield $=33,090 \mathrm{t}$ |
|  | Tier 5 ( $\mathrm{M}=0.235$ ) | yield $=16,800 \mathrm{t}$ |
|  | Tier $5(\mathrm{M}=0.3)$ | yield $=21,370 \mathrm{t}$ |
| Overfishing (OFL): | Tier 3a Model 2B F $35 \%$ \% 0.38 | yield $=29,630 \mathrm{t}$ |
|  | Tier 5( $\mathrm{M}=0.235$ ) | yield $=22,350 \mathrm{t}$ |
|  | Tier 5 ( $\mathrm{M}=0.3$ ) | yield $=28,500 \mathrm{t}$ |

Model 2B Equilibrium female spawning biomass

$$
\begin{aligned}
& \mathrm{B}_{100 \%}=100,945 \mathrm{t} \\
& \mathrm{~B}_{40 \%}=40,378 \mathrm{t} \\
& \mathrm{~B}_{35 \%}=35,331 \mathrm{t}
\end{aligned}
$$

Model 2B Projected 2007 biomass
Age 3+ biomass $=167,581 \mathrm{t}$
Female spawning biomass $=82,210 \mathrm{t}$

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## Tables

Table 1A.1. Time series of $\mathrm{ABC}, \mathrm{TAC}$, and total catch for Aleutian Islands Region walleye pollock fisheries 1991-2006. Units are in metric tons. Note: There was no OFL level set in 1991 and the 1993 harvest specifications were not available

| YEAR | ABC | TAC | OFL | CATCH | CATCH/TAC |
| :--- | ---: | ---: | ---: | ---: | ---: |
| 1991 | 101,460 | 72,250 | NA | 98,604 | $136 \%$ |
| 1992 | 51,600 | 47,730 | 62,400 | 52,352 | $110 \%$ |
| 1993 |  |  |  | 57,132 |  |
| 1994 | 56,600 | 56,600 | 60,400 | 58,659 | $104 \%$ |
| 1995 | 56,600 | 56,600 | 60,400 | 64,925 | $115 \%$ |
| 1996 | 35,600 | 35,600 | 47,000 | 29,062 | $82 \%$ |
| 1997 | 28,000 | 28,000 | 38,000 | 25,940 | $93 \%$ |
| 1998 | 23,800 | 23,800 | 31,700 | 23,822 | $100 \%$ |
| 1999 | 23,800 | 2,000 | 31,700 | 1,010 | $51 \%$ |
| 2000 | 23,800 | 2,000 | 31,700 | 1,244 | $62 \%$ |
| 2001 | 23,800 | 2,000 | 31,700 | 824 | $41 \%$ |
| 2002 | 23,800 | 1,000 | 31,700 | 1,156 | $116 \%$ |
| 2003 | 39,400 | 1,000 | 52,600 | 1,653 | $165 \%$ |
| 2004 | 39,400 | 1,000 | 52,600 | 1,150 | $115 \%$ |
| 2005 | 29,400 | 19,000 | 39,100 | 1,556 | $8 \%$ |
| 2006 | 29,400 | 19,000 | 39,100 | 1,829 | $10 \%$ |

Table 1A.2. Estimates of walleye pollock catches from the entire Aleutian Islands Region by source, 1977-2003. Units are in metric tons.

| Year | Official <br>  <br> JV Blend | Domestic <br> Blend | Foreign <br> Reported | NMFS <br> Observer <br> Data | Current <br> estimates |
| ---: | ---: | ---: | ---: | ---: | ---: |
| 1977 | 7,367 |  | 7,827 | 5 | 7,367 |
| 1978 | 6,283 |  | 6,283 | 234 | 6,283 |
| 1979 | 9,446 |  | 9,505 | 58 | 9,446 |
| 1980 | 58,157 |  | 58,477 | 883 | 58,157 |
| 1981 | 55,517 |  | 57,056 | 2,679 | 55,517 |
| 1982 | 57,753 |  | 62,624 | 11,847 | 57,753 |
| 1983 | 59,021 |  | 44,544 | 12,429 | 59,021 |
| 1984 | 77,595 |  | 67,103 | 48,383 | 77,595 |
| 1985 | 58,147 |  | 48,733 | 43,844 | 58,147 |
| 1986 | 45,439 |  | 14,392 | 29,464 | 45,439 |
| 1987 | 28,471 |  |  | 17,944 | 28,471 |
| 1988 | 41,203 |  |  | 21,987 | 41,203 |
| 1989 | 10,569 |  |  | 5,316 | 10,569 |
| 1990 |  | 79,025 |  | 51,137 | 79,025 |
| 1991 |  | 98,604 |  | 20,493 | 98,064 |
| 1992 |  | 52,352 |  | 20,853 | 52,352 |
| 1993 |  | 57,132 |  | 22,804 | 57,132 |
| 1994 |  | 58,659 |  | 37,707 | 58,659 |
| 1995 |  | 64,925 |  | 18,023 | 64,925 |
| 1996 |  | 29,062 |  | 5,982 | 29,062 |
| 1997 |  | 25,940 |  | 5,580 | 25,940 |
| 1998 |  | 23,822 |  | 1,882 | 23,822 |
| 1999 |  | 1,010 |  | 24 | 1,010 |
| 2000 |  | 1,244 |  | 75 | 1,244 |
| 2001 |  | 824 |  | 88 | 824 |
| 2002 |  | 1,156 |  | 144 | 1,156 |
| 2003 |  | 1,653 |  |  | 1,653 |
| 2004 |  | 1,150 |  |  | 1,150 |
| 2005 |  | 1,610 |  |  | 1,610 |
| 2006 |  | 1,829 |  |  | 1,829 |

Table 1A.3. Estimated walleye pollock catch discarded and retained for the Aleutian Islands Region based on NMFS blend data, 1991-2001.

|  | Catch <br> Retained | Discard | Total | Discard <br> Percentage |
| ---: | ---: | ---: | ---: | ---: |
| 1990 | 69,682 | 9,343 | 79,025 | $12 \%$ |
| 1991 | 93,059 | 5,441 | 98,500 | $6 \%$ |
| 1992 | 49,375 | 2,986 | 52,361 | $6 \%$ |
| 1993 | 55,399 | 1,740 | 57,138 | $3 \%$ |
| 1994 | 57,308 | 1,373 | 58,681 | $2 \%$ |
| 1995 | 63,545 | 1,380 | 64,925 | $2 \%$ |
| 1996 | 28,067 | 994 | 29,062 | $3 \%$ |
| 1997 | 25,323 | 617 | 25,940 | $2 \%$ |
| 1998 | 23,657 | 164 | 23,822 | $1 \%$ |
| 1999 | 361 | 446 | 807 | $55 \%$ |
| 2000 | 455 | 790 | 1,244 | $64 \%$ |
| 2001 | 445 | 380 | 824 | $46 \%$ |
| 2002 | 398 | 758 | 1,156 | $66 \%$ |
| 2003 | 1184 | 468 | 1,653 | $28 \%$ |
| 2004 | 871 | 278 | 1,150 | $24 \%$ |
| 2005 | 200 | 1,410 | 1,610 | $88 \%$ |
| 2006 | 1,082 | 747 | 1,829 | $41 \%$ |

Table 1A.4. Estimates of Aleutian Islands Region walleye pollock catch by the three management subareas. Foreign reported data were used from 1977-1984, from 1985-1998 observer data were used to partition catches among the areas. Units are in metric tons.

|  | East <br> $(541)$ | Central <br> $(542)$ | West <br> $(543)$ | Total |
| :---: | :---: | :---: | :---: | :---: |
| Year |  | 0 | 2,965 | 7,367 |
| 1977 | 4,402 | 0, | 305 | 6,283 |
| 1978 | 5,267 | 712 | 6,203 | 9,446 |
| 1979 | 1,488 | 1,756 | 22,775 | 58,157 |
| 1980 | 28,284 | 7,097 | 1,982 | 55,517 |
| 1981 | 43,461 | 10,074 | 2,376 | 57,753 |
| 1982 | 54,173 | 1,205 | 1,194 | 59,021 |
| 1983 | 56,577 | 1,250 | 7,663 | 77,595 |
| 1984 | 64,172 | 5,760 | 100 | 58,147 |
| 1985 | 19,885 | 38,163 | 0 | 45,439 |
| 1986 | 38,361 | 7,078 | 0 | 28,471 |
| 1987 | 28,086 | 386 | 0 | 41,203 |
| 1988 | 40,685 | 517 | 0 | 10,569 |
| 1989 | 10,569 | 0 | 430 | 79,025 |
| 1990 | 69,170 | 9,425 | 11 | 98,604 |
| 1991 | 98,032 | 561 | 6 | 52,352 |
| 1992 | 52,140 | 206 | 83 | 57,132 |
| 1993 | 54,512 | 2,536 | 15 | 58,659 |
| 1994 | 58,091 | 554 | 102 | 64,925 |
| 1995 | 28,109 | 36,714 | 261 | 29,062 |
| 1996 | 9,226 | 19,574 | 1,031 | 25,940 |
| 1997 | 8,110 | 16,799 | 18,127 | 23,822 |
| 1998 | 1,837 | 3,858 |  |  |

Table 1A.5. Estimates of pollock catch (metric tons) by new area definitions. "NRA" stands for Near, Rat, and Andreanof island groups, "NRA w/o E" signifies the NRA region without the area east of $174^{\circ} \mathrm{W}$, "Basin" represents the northern portions of areas 541 and 542. See Fig. 1A. 3 for locations on a map. (Note: 1977-1984 area assignments are based on foreign reported data, 1985-2006 are based on observer data).

| Year | NRA | NRA w/o E | Basin | Basin + E |
| :---: | :---: | :---: | :---: | :---: |
| 1977 | 7,367 | 2,965 | 0 | 4,402 |
| 1978 | 6,283 | 1,016 | 0 | 5,267 |
| 1979 | 9,446 | 7,959 | 0 | 1,488 |
| 1980 | 58,157 | 29,873 | 0 | 28,284 |
| 1981 | 31,258 | 14,811 | 24,259 | 40,706 |
| 1982 | 50,322 | 3,149 | 7,863 | 54,605 |
| 1983 | 44,442 | 1,669 | 15,354 | 57,352 |
| 1984 | 42,901 | 9,171 | 39,140 | 68,424 |
| 1985 | 47,070 | 870 | 48,472 | 57,278 |
| 1986 | 23,810 | 704 | 28,003 | 44,735 |
| 1987 | 26,257 | 2,720 | 2,251 | 25,752 |
| 1988 | 36,864 | 574 | 4,339 | 40,628 |
| 1989 | 10,569 | 0 | 0 | 10,569 |
| 1990 | 79,025 | 10,477 | 0 | 68,548 |
| 1991 | 98,604 | 561 | 230 | 98,043 |
| 1992 | 52,352 | 8,519 | 29,455 | 43,833 |
| 1993 | 57,132 | 16,162 | 22,404 | 40,970 |
| 1994 | 58,659 | 5,965 | 26,288 | 52,694 |
| 1995 | 64,925 | 58,203 | 3,015 | 6,723 |
| 1996 | 29,062 | 23,187 | 899 | 5,875 |
| 1997 | 25,940 | 25,774 | 0 | 166 |
| 1998 | 23,822 | 23,335 | 67 | 486 |
| 1999 | 1,010 | 631 | 0 | 378 |
| 2000 | 1,244 | 891 | 0 | 354 |
| 2001 | 824 | 575 | 0 | 249 |
| 2002 | 1,156 | 351 | 1 | 805 |
| 2003 | 1,653 | 1,430 | 0 | 222 |
| 2004 | 1,150 | 962 | 0 | 188 |
| 2005 | 1,610 | 1,330 | 0 | 280 |
| 2006 | 1,829 | 1,657 | 0 | 172 |
|  |  |  |  |  |

Table 1A.6. Sampling levels in Aleutian Islands Region sub-regions based on foreign, J.V., and domestic walleye pollock observer data 1978 - 1998.

| Year | NRA West of $174^{\circ}$ Longitude |  |  | NRA East of $174^{\circ}$ Longitude |  |  | Aleutian Islands Area Basin |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Fish Measured | Hauls Sampled | Vessels Sampled | Fish Measured | Hauls Sampled | Vessels Sampled | Fish <br> Measured | Hauls Sampled | Vessels Sampled |
| 1978 | 1,503 | 64 | 4 | 4,831 | 135 | 11 | 0 | 0 | 0 |
| 1979 | 1,317 | 16 | 4 | 977 | 33 | 6 | 0 | 0 | 0 |
| 1980 | 2,154 | 53 | 4 | 4,753 | 119 | 10 | 0 | 0 | 0 |
| 1981 | 4,782 | 37 | 7 | 6,617 | 96 | 14 | 1,913 | 15 | 3 |
| 1982 | 7,713 | 102 | 13 | 29,549 | 331 | 30 | 11,151 | 84 | 7 |
| 1983 | 2,977 | 35 | 12 | 24,793 | 242 | 27 | 20,744 | 174 | 21 |
| 1984 | 10,844 | 111 | 22 | 46,037 | 541 | 49 | 157,388 | 1,223 | 81 |
| 1985 | 780 | 9 | 2 | 33,471 | 259 | 37 | 68,923 | 460 | 58 |
| 1986 | 0 | 0 | 0 | 22,939 | 195 | 18 | 39,875 | 268 | 48 |
| 1987 | 4,045 | 26 | 5 | 43,093 | 352 | 29 | 2,665 | 26 | 8 |
| 1988 | 378 | 3 | 2 | 28,423 | 249 | 24 | 4,528 | 37 | 14 |
| 1989 | 0 | 0 | 0 | 7,424 | 57 | 8 | 0 | 0 | 0 |
| 1990 | 12,303 | 131 | 14 | 55,837 | 587 | 47 | 55 | 1 | 1 |
| 1991 | 0 | 1 | 1 | 26,035 | 211 | 32 | 24,025 | 194 | 26 |
| 1992 | 7,405 | 59 | 15 | 18,771 | 178 | 50 | 20,769 | 179 | 27 |
| 1993 | 13,471 | 130 | 15 | 13,264 | 137 | 34 | 22,022 | 185 | 30 |
| 1994 | 5,025 | 47 | 18 | 29,805 | 305 | 64 | 5,314 | 56 | 16 |
| 1995 | 29,070 | 324 | 34 | 2,963 | 212 | 31 | 1,922 | 19 | 7 |
| 1996 | 15,307 | 160 | 35 | 3,462 | 179 | 41 | 0 | 0 | 0 |
| 1997 | 17,239 | 189 | 33 | 64 | 122 | 26 | 77 | 1 | 1 |
| 1998 | 10,439 | 122 | 15 | 148 | 107 | 12 | 0 | 0 | 0 |
| Total | 146,752 | 1,619 | 255 | 403,256 | 4,647 | 600 | 381,371 | 2,922 | 348 |

Table 1A.7. NRA pollock fishery average weight-at-age in kilograms. Shaded cells had missing observations and were filled with their mean values

| Year | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 5+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1978 | 0.3318 | 0.3933 | 0.7603 | 0.6877 | 0.8097 | 0. | 0.9065 | 0.9722 | 0.9281 | 1.0613 | 74 | 87 | 49 | 1.0729 |
| 1979 | 0.2314 | 0.3476 | 0.5293 | 0.7306 | 0.6727 | 825 | 0.9435 | 0.9532 | 1.0381 | 1.1638 | 1.0598 | 1.5186 | 5788 | 06 |
| 1980 | 0.2392 | 0.5526 | 0.7651 | 0.8412 | 0.8629 | 0.9129 | 1.0002 | 1.089 | 1.0628 | 1.0204 | 1.1568 | 1.1019 | 0.8521 | . 5242 |
| 1981 | 0.3392 | 0.4778 | 0.5521 | 0.7286 | 0.7637 | 0.7817 | 0.8096 | 0.8953 | 0.90 | 0.8598 | 1.0199 | 1.0259 | 29 | 79 |
| 1982 | 0.3392 | 0.4179 | 0.5414 | 0.6436 | 0.7838 | . 82 | 0.8417 | 0.8921 | 0.984 | 1.0011 | 0.9575 | 0.9546 | 0.9058 | . 966 |
| 1983 | 0.3392 | 0.4736 | 0.6609 | 0.7333 | 0.7796 | 0.7954 | 0.9264 | 0.9574 | 1.0146 | 0.9024 | 1.1892 | 1.1496 | 0.974 | 1.14 |
| 19 | 0.426 | 0.4459 | 0.6609 | 0.7419 | 0.8099 | 0.8721 | 0.968 | 0.9963 | 1.2704 | 1.6431 | 1.1351 | 1.2212 | 1.1943 | 1.14 |
| 1985 | 0.4675 | 0.5656 | 0.6705 | 0.6896 | 0.8028 | 0.8536 | 0.8567 | 1.0909 | 1.23 | 1.5996 | 1.664 | 1.1496 | 1.6448 | 14 |
| 1986 | 0.3392 | 0.5114 | 0.6019 | 0.7 | 0.8266 | 0.8698 | 0.9506 | 0.9266 | 1.0 | 0.9428 | 1.07 | 0.8963 | 943 | 14 |
| 1987 | 0.3392 | 0.4736 | 0.6852 | 0.7562 | 0.8335 | 0.8504 | 0.8715 | 0.9809 | 1.0725 | 0.9915 | 1.3379 | 1.1546 | 1.0065 | 1.0935 |
| 1988 | 0.3392 | 0.4736 | 0.6609 | 0.8013 | 0.7905 | 0.8208 | 0.9279 | 0.8883 | 0.9839 | 0.8933 | 0.7843 | 0.7223 | 0.8976 | . 0621 |
| 19 | 0.3 | 0.4736 | 0.6609 | 0.7536 | 0.851 | 0.926 | 0.9927 | 1.0611 | 1.110 | 1.1501 | 1.1892 | 1.1496 | 1.1943 | 1.14 |
| 1990 | 0.3392 | 0.4778 | 0.5521 | 0.7286 | 0.7637 | 0.7817 | 0.8096 | 0.8953 | 0.9021 | 0.8598 | 1.0199 | 1.0259 | 0.8929 | 0.9079 |
| 19 | 0.3392 | 0.4736 | 0.6668 | 0.6551 | 0.7989 | 0.962 | 1.0755 | 1.1731 | 1.099 | 1.2177 | 1.1573 | 1.0955 | 1.2898 | . 0856 |
| 1992 | 0.3392 | 0.4736 | 0.6401 | 0.7418 | 0.7254 | 0.797 | 0.9356 | 1.2457 | 1.0267 | 1.0034 | 1.2501 | 1.1451 | 1.0514 | 1.0976 |
| 1993 | 0.3392 | 0.4736 | 0.8862 | 0.8237 | 1.0335 | 1.0315 | 1.1399 | 1.0808 | 1.1638 | 1.1905 | 1.2027 | 1.3256 | 1.1373 | 1.1352 |
| 1994 | 0.3392 | 0.4736 | 0.6373 | 0.8437 | 0.9743 | 1.1 | 1.14 | 1.1 | 1.190 | 1.2437 | 1.2659 | 1.0591 | 1.09 | 17 |
| 1995 | 0.3392 | 0.5512 | 0.8471 | 0.7536 | 1.1264 | 1.3303 | 1.3972 | 1.3551 | 1.4333 | 1.4197 | 1.501 | 1.4466 | 1.6582 | 1.3206 |
| 1996 | 0.3392 | 0.5391 | 0.4753 | 0.9301 | 1.0287 | 1.1796 | 1.2751 | 1.3945 | 1.4682 | 1.3548 | 1.3777 | 1.3619 | 1.4562 | 1.3013 |
| 1997 | 0.3 | 0.4 | 0. | 0.7536 | 0.851 | 0.926 | 0.992 | 1.0 | 1.11 | 1.1501 | 1.18 | 1.1 | 1.1943 | 1.14 |
| 19 | 0.3392 | 0.403 | 0.7631 | 0.7398 | 0.9826 | 1.0575 | 1.085 | 1.2532 | 1.3137 | 1.4826 | 1.2785 | 1.3012 | 1.3597 | . 4522 |
| 199 | 0.3392 | 0.4736 | 0.6609 | 0.7536 | 0.851 | 0.926 | 0.9927 | 1.0611 | 1.1106 | 1.1501 | 1.1892 | 1.1496 | 1.1943 | 1.1 |
| 00 | 0.3392 | 0.4736 | 0.6609 | 0.7536 | 851 | 92 | 0.9927 | 1.0611 | 1.1106 | 1.1501 | 1.1892 | 1.1496 | 1.1943 | 1.14 |
| 2001 | 0.3392 | 0.4736 | 0.6609 | 0.7536 | 0.851 | 0.926 | 0.9927 | 1.0611 | 1.1106 | 1.1501 | 1.1892 | 1.1496 | 1.1943 | 1.14 |
| 2002 | 0.3392 | 0.4736 | 0.6609 | 0.7536 | 0.851 | 0.926 | 0.9927 | 1.0611 | 1.1106 | 1.1501 | 1.1892 | 1.1496 | 1.1943 | 1.1 |
| 2003 | 0.3318 | 0.3933 | 0.7603 | 0.6877 | 0.8097 | 0.9151 | 0.9065 | 0.9722 | 0.9281 | 1.0613 | 1.1674 | 1.187 | 1.6149 | 1.0729 |
| 2004 | 0.3318 | 0.3933 | 0.7603 | 0.6877 | 0.8097 | 0.9151 | 0.9065 | 0.9722 | 0.9281 | 1.0613 | 1.1674 | 1.187 | 1.6149 | 1.0729 |
| 2005 | 0.3318 | 0.3933 | 0.7603 | 0.6877 | 0.8097 | 0.9151 | 0.9065 | 0.9722 | 0.9281 | 1.0613 | 1.1674 | 1.187 | 1.6149 | 1.0729 |
| 2006 | 0.3318 | 0.3933 | 0.7603 | 0.6877 | 0.8097 | 0.9151 | 0.9065 | 0.9722 | 0.9281 | 1.0613 | 1.1674 | 1.187 | 1.6149 | 1.0729 |

Table 1A.8. Number of aged and measured fish in the NRA pollock fishery used to estimate fishery age composition. Shaded values were not used in assessment. Data for 2006 from 2006 AICASS.

|  | Number Aged |  |  | Number Measured |  |  |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| Year | Males | Females | Total | Males | Females | Total |
| 1978 | 209 | 322 | 531 | 490 | 1,013 | 1,503 |
| 1979 | 124 | 178 | 302 | 611 | 706 | 1,317 |
| 1980 | 93 | 167 | 260 | 971 | 1,183 | 2,154 |
| 1981 | 124 | 152 | 276 | 2,226 | 2,556 | 4,782 |
| 1982 | 564 | 640 | 1,204 | 3,655 | 4,058 | 7,713 |
| 1983 | 132 | 145 | 277 | 1,493 | 1,484 | 2,977 |
| 1984 | 294 | 312 | 606 | 5,273 | 5,571 | 10,844 |
| 1985 | 210 | 265 | 475 | 349 | 431 | 780 |
| 1986 | 77 | 113 | 190 | 0 | 0 | 0 |
| 1987 | 131 | 142 | 273 | 1,670 | 2,375 | 4,045 |
| 1988 | 34 | 33 | 67 | 188 | 190 | 378 |
| 1989 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1990 | 46 | 49 | 95 | 5,209 | 7,094 | 12,303 |
| 1991 | 36 | 47 | 83 | 0 | 0 | 80 |
| 1992 | 110 | 121 | 231 | 3,755 | 3,650 | 7,405 |
| 1993 | 81 | 82 | 163 | 7,701 | 5,770 | 13,471 |
| 1994 | 157 | 151 | 308 | 2,644 | 2,381 | 5,025 |
| 1995 | 74 | 106 | 180 | 16,518 | 12,552 | 29,070 |
| 1996 | 95 | 84 | 179 | 8,933 | 6,374 | 15,307 |
| 1997 | 15 | 15 | 30 | 9,232 | 8,007 | 17,239 |
| 1998 | 144 | 170 | 314 | 5,992 | 4,447 | 10,439 |
| 1999 | 0 | 0 | 0 | 75 | 60 | 135 |
| 2000 | 0 | 1 | 1 | 70 | 114 | 184 |
| 2001 | 0 | 1 | 1 | 52 | 106 | 158 |
| 2002 | 0 | 0 | 0 | 46 | 61 | 107 |
| 2003 | 0 | 0 | 0 | 0 | 0 | 0 |
| 2004 | 0 | 0 | 0 | 153 | 212 | 365 |
| 205 | 0 | 0 | 0 | 309 | 260 | 569 |
| 2006 | 74 | 87 | 161 | 1,315 | 1,630 | 2,945 |

Table 1A.9. Number of individual vessels and hauls sampled by observers in the NRA pollock fishery west of $174^{\circ} \mathrm{W}$ longitude, 1990-1998.

| Year | NRA Area 541 West of 174W |  |  |  | NRA Area 542 |  |  |  | NRA Area 543 |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | CatcherProcessorVessel |  | Catcher Only |  | Catcher Processor |  | Catcher Only |  | Catcher Processor |  | Catcher Only |  |
|  | S | Hauls | S | S | Vessels | Hauls | S | S | Vessels | Hauls | S | S |
| 1990 | 12 | 50 | 0 | 0 | 16 | 132 | 0 | 0 | 2 | 4 | 0 | 0 |
| 1991 | 2 | 3 | 0 | 0 | 2 | 2 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1992 | 18 | 126 | 0 | 0 | 4 | 5 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1993 | 18 | 195 | 0 | 0 | 6 | 25 | 0 | 0 | 3 | 5 | 0 | 0 |
| 1994 | 18 | 76 | 0 | 0 | 3 | 6 | 0 | 0 | 0 | 0 | 0 | 0 |
| 1995 | 22 | 200 | 8 | 39 | 15 | 272 | 11 | 77 | 0 | 0 | 0 | 0 |
| 1996 | 5 | 12 | 7 | 15 | 25 | 198 | 10 | 38 | 0 | 0 | 0 | 0 |
| 1997 | 13 | 66 | 11 | 30 | 14 | 93 | 10 | 60 | 1 | 6 | 0 | 0 |
| 1998 | 4 | 6 | 5 | 16 | 3 | 24 | 5 | 19 | 2 | 97 | 4 | 24 |

Table 1A.10. Estimated NRA region pollock catch at age (millions). Highest mode for each year is shaded.

| Year | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | Total |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 1978 | 0.01 | 0.14 | 0.12 | 0.07 | 0.36 | 0.10 | 0.14 | 0.13 | 0.13 | 0.06 | 0.02 | 0.01 |  | 0.00 | 1.27 |
| 1979 | 0.01 | 2.18 | 2.22 | 2.02 | 2.43 | 1.73 | 0.65 | 0.63 | 0.37 | 0.03 | 0.22 |  |  | 0.05 | 12.53 |
| 1980 | 8.20 | 3.24 | 2.64 | 3.71 | 6.94 | 4.05 | 2.47 | 0.73 | 1.07 | 0.53 | 0.16 | 0.01 | 0.14 | 0.01 | 33.91 |
| 1981 |  | 5.72 | 3.36 | 2.19 | 1.65 | 2.55 | 2.54 | 1.93 | 1.37 | 0.73 | 0.20 | 0.15 | 0.20 | 0.04 | 22.64 |
| 1982 |  | 0.01 | 3.00 | 0.51 | 0.23 | 0.31 | 0.38 | 0.35 | 0.15 | 0.07 | 0.04 | 0.03 | 0.01 | 0.01 | 5.10 |
| 1983 |  |  |  | 0.74 | 0.44 | 0.17 | 0.11 | 0.24 | 0.23 | 0.05 | 0.04 | 0.01 | 0.00 | 0.00 | 2.04 |
| 1984 | 0.14 | 3.97 |  | 4.12 | 4.12 | 1.46 | 1.10 | 0.74 | 0.51 | 0.34 | 0.09 | 0.06 | 0.03 | 0.01 | 16.68 |
| 1985 | 0.01 | 0.01 | 0.17 | 0.06 | 0.17 | 0.46 | 0.20 | 0.08 | 0.08 | 0.04 | 0.01 | 0.01 | 0.00 | 0.00 | 1.30 |
| 1986 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1987 |  |  | 1.40 | 0.31 | 0.23 | 0.04 | 0.09 | 1.01 | 0.09 | 0.12 | 0.00 | 0.03 | 0.01 | 0.04 | 3.36 |
| 1988 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1989 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1990 |  | 0.95 | 0.26 | 0.96 | 0.78 | 0.78 | 0.93 | 0.17 | 1.10 | 0.34 | 0.56 | 0.28 | 0.13 | 0.21 | 7.45 |
| 1991 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1992 |  |  | 0.03 | 0.33 | 0.60 | 0.30 | 0.60 | 0.12 | 0.69 | 0.39 | 0.52 | 0.36 | 1.71 | 1.91 | 7.55 |
| 1993 |  |  | 0.18 | 0.47 | 1.12 | 1.34 | 0.54 | 1.46 | 0.81 | 0.88 | 0.83 | 0.38 | 0.70 | 4.34 | 13.05 |
| 1994 |  | 0.07 | 1.00 | 0.31 | 0.42 | 0.60 | 0.43 | 0.33 | 0.17 | 0.39 | 0.10 | 0.08 | 1.30 | 5.20 |  |
| 1995 | 0.22 | 0.38 | 0.00 | 10.22 | 1.19 | 5.10 | 4.84 | 1.42 | 2.36 | 2.08 | 3.82 | 0.77 | 8.32 | 40.71 |  |
| 1996 | 0.17 | 0.15 | 0.56 | 1.42 | 5.15 | 1.53 | 2.09 | 1.21 | 0.92 | 0.64 | 0.20 | 0.77 | 2.00 | 16.79 |  |
| 1997 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1998 | 0.05 | 0.08 | 5.66 | 1.65 | 1.05 | 0.96 | 1.71 | 1.20 | 1.00 | 2.40 | 1.30 | 1.17 | 1.49 | 19.73 |  |
| 2006 |  |  |  | 0.01 | 0.33 | 0.13 | 0.04 | 0.02 | 0.08 | 0.06 | 0.06 | 0.05 | 0.12 | 0.12 | 1.02 |

Table 1A.11. Pollock biomass estimates from the Aleutian Islands Groundfish Survey, 1980-2002.

|  | Aleutian Islands Region <br> NRA West <br> $(174 \mathrm{~W}-170 \mathrm{E})$ | NRA East <br> $(170 \mathrm{~W}-174 \mathrm{~W})$ | NRA <br> total | Unalaska-Umnak <br> area $(\sim 165 \mathrm{~W}-170 \mathrm{~W})$ | Combined |
| :--- | :---: | :---: | :---: | :---: | :---: |
| $\mathbf{1 9 8 0}$ |  |  | 243,695 | 56,732 | 300,427 |
| $\mathbf{1 9 8 3}$ |  |  | 495,775 | 282,648 | 778,423 |
| $\mathbf{1 9 8 6}$ |  |  | 439,461 | 102,379 | 541,840 |
|  |  |  |  |  |  |
| $\mathbf{1 9 9 1}$ | 83,337 | 53,865 | 137,202 | 51,644 | 188,846 |
| $\mathbf{1 9 9 4}$ | 47,623 | 29,879 | 77,502 | 39,696 | 117,199 |
| $\mathbf{1 9 9 7}$ | 57,577 | 39,935 | 97,512 | 65,400 | 158,912 |
| $\mathbf{2 0 0 0}$ | 76,613 | 28,985 | 105,598 | 22,462 | 128,060 |
| $\mathbf{2 0 0 2}$ | 121,915 | 53,368 | 175,283 | 181,334 | 356,617 |
| $\mathbf{2 0 0 4}$ | 19,201 | 111,250 | 130,451 | 235,658 | 366,110 |
| $\mathbf{2 0 0 6}$ | 25,471 | 69,522 | 94,993 | 18,006 | 112,999 |

Table 1A.12. Results of the 2002 Aleutian Islands echo integration-trawl survey conducted by the R/V Kaiyo Maru.

|  | Leg 2-1 | Leg 2-2 | Leg 2-3 | Leg 2-4 |
| ---: | ---: | ---: | ---: | ---: |
| Area $\left(\mathrm{km}^{2}\right)$ | 27,902 | 10,433 | 4,045 | 1,413 |
| Density $\left(\mathrm{t} / \mathrm{km}^{2}\right)$ | 2.18 | 1.82 | 2.46 | 1.79 |
| Population $\left(10^{6}\right)$ | 37 | 12 | 6 | 2 |
| Biomass $\left(10^{3} \mathrm{t}\right)$ | 61 | 19 | 10 | 3 |
| CV | 0.31 | 0.33 | 0.21 | 0.76 |

Table 1A.13. Results from the 2006 Aleutian Islands Cooperative Acoustic Survey.

| Survey | Area <br> $\left(\mathrm{n} . \mathrm{mi.}^{2}\right)$ | Deadzone <br> $(\mathrm{Y} / \mathrm{N})$ | Biomass <br> $(\mathrm{t})$ | Relative <br> Precision <br> $\left(\mathrm{E}_{\mathrm{i}}\right)$ | High Biom. <br> $(\mathrm{t})$ | Low Biom. <br> $(\mathrm{t})$ | Density $(\mathrm{t}$ <br> /n.mi. $\left.{ }^{2}\right)$ |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 2 | 180 | N | 8233.8 | $8.67 \%$ | 9632.5 | 6835.1 | 45.7 |
| 2 | 180 | Y | 8809.9 | $8.04 \%$ | 10198.4 | 7421.4 | 48.9 |
| 2 | 72 | N | 6484.5 | $12.29 \%$ | 8046.1 | 4922.9 | 90.1 |
| 2 | 72 | Y | 6706.6 | $14.32 \%$ | 8589.2 | 4824.0 | 93.1 |
| 4 | 180 | N | 6600.4 | $7.96 \%$ | 7630.1 | 5570.7 | 36.7 |
| 4 | 180 | Y | 7980.2 | $7.87 \%$ | 9210.6 | 6749.8 | 44.3 |
| 4 | 72 | N | 5246.4 | $12.31 \%$ | 6512.6 | 3980.2 | 72.9 |
| 4 | 72 | Y | 6149.8 | $11.89 \%$ | 7582.5 | 4717.1 | 85.4 |
| 5 | 9 | N | 890.8 | $5.29 \%$ | 983.2 | 798.4 | 99.0 |
| 5 | 9 | Y | 1036.6 | $4.75 \%$ | 1133.1 | 940.1 | 115.2 |
| 6 | 72 | N | 3015.0 | $6.64 \%$ | 3407.4 | 2622.6 | 41.9 |
| 6 | 72 | Y | 3458.5 | $6.44 \%$ | 3894.9 | 3022.1 | 48.0 |
| 7 | 72 | N | 1159.0 | $6.83 \%$ | 1314.2 | 1003.8 | 16.1 |
| 7 | 72 | Y | 2179.7 | $5.05 \%$ | 2395.4 | 1964.0 | 30.3 |
| 8 | 180 | N | 2313.6 | $14.51 \%$ | 2971.6 | 1655.6 | 12.9 |
| 8 | 180 | Y | 2845.2 | $14.24 \%$ | 3639.0 | 2051.4 | 15.8 |
| 8 | 72 | N | 559.2 | $14.32 \%$ | 716.1 | 402.3 | 7.8 |
| 8 | 72 | Y | 677.0 | $12.96 \%$ | 848.9 | 505.1 | 9.4 |

Table 1A.14. Estimated instantaneous natural mortality rates (M) by age from Wespestad and Terry (1984).

| Age | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| M | 0.85 | 0.45 | 0.3 | 0.3 | 0.3 | 0.3 | 0.3 | 0.3 | 0.3 | 0.3 | 0.4 | 0.4 | 0.4 | 0.5 | 0.5 |

Table 1A.15. Estimated von Bertalanffy growth curve parameters and length-weight regression parameters for walleye pollock sampled during the U.S.-Japan 1980, 1983, and 1986 groundfish surveys and the 1991, 1994, 1997, 2000, 2002, and 2004 RACE groundfish surveys.

|  | $\mathbf{L}_{\text {inf }}$ | $\mathbf{K}$ | $\mathbf{t}_{\mathbf{0}}$ | $\mathbf{A}$ | $\mathbf{b}$ |
| ---: | ---: | ---: | ---: | ---: | ---: |
| 1980 | 51.92 | 0.414 | -0.525 | 0.0132 | 2.858 |
| 1983 | 53.26 | 0.383 | 0.002 | 0.0178 | 2.768 |
| 1986 | 51.02 | 0.443 | -0.084 | 0.0142 | 2.831 |
| 1991 | 54.55 | 0.392 | -0.361 | 0.0104 | 2.912 |
| 1994 | 61.58 | 0.330 | -0.102 | 0.0069 | 3.022 |
| 1997 | 61.41 | 0.286 | -0.397 | 0.0081 | 2.983 |
| 2000 | 62.58 | 0.306 | -0.048 | 0.0064 | 3.019 |
| 2002 | 64.36 | 0.289 | -0.127 | 0.0066 | 3.018 |
| 2004 | 61.76 | 0.332 | -0.189 | 0.0065 | 3.022 |

Table 1A.16. Average weight-at-age for Aleutian Islands pollock as estimated from NMFS summer bottom trawl survey estimates. Values between survey years (shaded) were set to the mean of the nearest two surveys (or single year for 1978-79, 2003-04).

|  | Age |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Year | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15+ |
| 1978 | 0.31 | 0.50 | 0.65 | 0.78 | 0.87 | 0.93 | 0.97 | 1.00 | 1.02 | 1.03 | 1.04 | 1.04 | 1.05 | 1.05 |
| 1979 | 0.31 | 0.50 | 0.65 | 0.78 | 0.87 | 0.93 | 0.97 | 1.00 | 1.02 | 1.03 | 1.04 | 1.04 | 1.05 | 1.05 |
| 1980 | 0.31 | 0.50 | 0.65 | 0.78 | 0.87 | 0.93 | 0.97 | 1.00 | 1.02 | 1.03 | 1.04 | 1.04 | 1.05 | 1.05 |
| 1981 | 0.25 | 0.43 | 0.60 | 0.73 | 0.83 | 0.90 | 0.95 | 0.99 | 1.01 | 1.03 | 1.04 | 1.05 | 1.05 | 1.05 |
| 1982 | 0.25 | 0.43 | 0.60 | 0.73 | 0.83 | 0.90 | 0.95 | 0.99 | 1.01 | 1.03 | 1.04 | 1.05 | 1.05 | 1.05 |
| 1983 | 0.19 | 0.37 | 0.54 | 0.69 | 0.80 | 0.88 | 0.94 | 0.98 | 1.01 | 1.02 | 1.04 | 1.05 | 1.05 | 1.06 |
| 1984 | 0.21 | 0.40 | 0.56 | 0.70 | 0.80 | 0.87 | 0.92 | 0.95 | 0.97 | 0.99 | 1.00 | 1.01 | 1.01 | 1.01 |
| 1985 | 0.21 | 0.40 | 0.56 | 0.70 | 0.80 | 0.87 | 0.92 | 0.95 | 0.97 | 0.99 | 1.00 | 1.01 | 1.01 | 1.01 |
| 1986 | 0.23 | 0.42 | 0.59 | 0.71 | 0.80 | 0.86 | 0.90 | 0.92 | 0.94 | 0.95 | 0.96 | 0.96 | 0.97 | 0.97 |
| 1987 | 0.23 | 0.46 | 0.64 | 0.75 | 0.91 | 1.01 | 1.08 | 1.06 | 1.10 | 1.08 | 1.06 | 1.04 | 1.06 | 1.03 |
| 1988 | 0.23 | 0.46 | 0.64 | 0.75 | 0.91 | 1.01 | 1.08 | 1.06 | 1.10 | 1.08 | 1.06 | 1.04 | 1.06 | 1.03 |
| 1989 | 0.23 | 0.46 | 0.64 | 0.75 | 0.91 | 1.01 | 1.08 | 1.06 | 1.10 | 1.08 | 1.06 | 1.04 | 1.06 | 1.03 |
| 1990 | 0.23 | 0.46 | 0.64 | 0.75 | 0.91 | 1.01 | 1.08 | 1.06 | 1.10 | 1.08 | 1.06 | 1.04 | 1.06 | 1.03 |
| 1991 | 0.22 | 0.51 | 0.69 | 0.79 | 1.01 | 1.15 | 1.26 | 1.21 | 1.27 | 1.21 | 1.16 | 1.12 | 1.16 | 1.10 |
| 1992 | 0.21 | 0.51 | 0.78 | 0.89 | 1.08 | 1.22 | 1.25 | 1.33 | 1.36 | 1.32 | 1.35 | 1.33 | 1.35 | 1.22 |
| 1993 | 0.21 | 0.51 | 0.78 | 0.89 | 1.08 | 1.22 | 1.25 | 1.33 | 1.36 | 1.32 | 1.35 | 1.33 | 1.35 | 1.22 |
| 1994 | 0.20 | 0.52 | 0.87 | 1.00 | 1.14 | 1.29 | 1.24 | 1.45 | 1.44 | 1.43 | 1.54 | 1.54 | 1.54 | 1.35 |
| 1995 | 0.22 | 0.48 | 0.82 | 0.97 | 1.07 | 1.24 | 1.26 | 1.38 | 1.44 | 1.45 | 1.53 | 1.52 | 1.57 | 1.47 |
| 1996 | 0.22 | 0.48 | 0.82 | 0.97 | 1.07 | 1.24 | 1.26 | 1.38 | 1.44 | 1.45 | 1.53 | 1.52 | 1.57 | 1.47 |
| 1997 | 0.25 | 0.43 | 0.78 | 0.95 | 1.00 | 1.19 | 1.29 | 1.31 | 1.44 | 1.47 | 1.52 | 1.51 | 1.60 | 1.60 |
| 1998 | 0.21 | 0.47 | 0.77 | 0.92 | 0.95 | 1.17 | 1.28 | 1.31 | 1.43 | 1.50 | 1.62 | 1.59 | 1.53 | 1.65 |
| 1999 | 0.21 | 0.47 | 0.77 | 0.92 | 0.95 | 1.17 | 1.28 | 1.31 | 1.43 | 1.50 | 1.62 | 1.59 | 1.53 | 1.65 |
| 2000 | 0.17 | 0.51 | 0.77 | 0.89 | 0.90 | 1.15 | 1.26 | 1.32 | 1.41 | 1.52 | 1.71 | 1.67 | 1.47 | 1.70 |
| 2001 | 0.19 | 0.49 | 0.74 | 1.02 | 1.03 | 1.23 | 1.29 | 1.43 | 1.53 | 1.56 | 1.74 | 1.68 | 1.58 | 1.67 |
| 2002 | 0.21 | 0.47 | 0.70 | 1.15 | 1.16 | 1.32 | 1.32 | 1.53 | 1.65 | 1.61 | 1.76 | 1.69 | 1.68 | 1.64 |
| 2003 | 0.21 | 0.47 | 0.70 | 1.15 | 1.16 | 1.32 | 1.32 | 1.53 | 1.65 | 1.61 | 1.76 | 1.69 | 1.68 | 1.64 |
| 2004 | 0.22 | 0.46 | 0.70 | 0.83 | 0.96 | 1.21 | 1.15 | 1.38 | 1.48 | 1.55 | 1.63 | 1.65 | 1.55 | 1.68 |
| 2005 | 0.22 | 0.46 | 0.70 | 0.83 | 0.96 | 1.21 | 1.15 | 1.38 | 1.48 | 1.55 | 1.63 | 1.65 | 1.55 | 1.68 |
| 2006 | 0.22 | 0.46 | 0.70 | 0.83 | 0.96 | 1.21 | 1.15 | 1.38 | 1.48 | 1.55 | 1.63 | 1.65 | 1.55 | 1.68 |

Table 1A.17. Average weight-at-age for Aleutian Islands pollock as estimated from fishery data.

|  | Age |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Year | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15+ |
| 1978 | 0.33 | 0.39 | 0.76 | 0.69 | 0.81 | 0.92 | 0.91 | 0.97 | 0.93 | 1.06 | 1.17 | 1.19 | 1.61 | 1.07 |
| 1979 | 0.23 | 0.35 | 0.53 | 0.73 | 0.67 | 0.83 | 0.94 | 0.95 | 1.04 | 1.16 | 1.06 | 1.52 | 1.58 | 1.02 |
| 1980 | 0.24 | 0.55 | 0.77 | 0.84 | 0.86 | 0.91 | 1.00 | 1.09 | 1.06 | 1.02 | 1.16 | 1.10 | 0.85 | 1.52 |
| 1981 | 0.34 | 0.48 | 0.55 | 0.73 | 0.76 | 0.78 | 0.81 | 0.90 | 0.90 | 0.86 | 1.02 | 1.03 | 0.89 | 0.91 |
| 1982 | 0.34 | 0.42 | 0.54 | 0.64 | 0.78 | 0.82 | 0.84 | 0.89 | 0.98 | 1.00 | 0.96 | 0.95 | 0.91 | 0.97 |
| 1983 | 0.34 | 0.47 | 0.66 | 0.73 | 0.78 | 0.80 | 0.93 | 0.96 | 1.01 | 0.90 | 1.19 | 1.15 | 0.97 | 1.14 |
| 1984 | 0.43 | 0.45 | 0.66 | 0.74 | 0.81 | 0.87 | 0.97 | 1.00 | 1.27 | 1.64 | 1.14 | 1.22 | 1.19 | 1.14 |
| 1985 | 0.47 | 0.57 | 0.67 | 0.69 | 0.80 | 0.85 | 0.86 | 1.09 | 1.23 | 1.60 | 1.66 | 1.15 | 1.64 | 1.14 |
| 1986 | 0.34 | 0.51 | 0.60 | 0.75 | 0.83 | 0.87 | 0.95 | 0.93 | 1.01 | 0.94 | 1.07 | 0.90 | 1.19 | 1.14 |
| 1987 | 0.34 | 0.47 | 0.69 | 0.76 | 0.83 | 0.85 | 0.87 | 0.98 | 1.07 | 0.99 | 1.34 | 1.15 | 1.01 | 1.09 |
| 1988 | 0.34 | 0.47 | 0.66 | 0.80 | 0.79 | 0.82 | 0.93 | 0.89 | 0.98 | 0.89 | 0.78 | 0.72 | 0.90 | 1.06 |
| 1989 | 0.34 | 0.47 | 0.66 | 0.75 | 0.85 | 0.93 | 0.99 | 1.06 | 1.11 | 1.15 | 1.19 | 1.15 | 1.19 | 1.14 |
| 1990 | 0.34 | 0.48 | 0.55 | 0.73 | 0.76 | 0.78 | 0.81 | 0.90 | 0.90 | 0.86 | 1.02 | 1.03 | 0.89 | 0.91 |
| 1991 | 0.34 | 0.47 | 0.67 | 0.66 | 0.80 | 0.96 | 1.08 | 1.17 | 1.10 | 1.22 | 1.16 | 1.10 | 1.29 | 1.09 |
| 1992 | 0.34 | 0.47 | 0.64 | 0.74 | 0.73 | 0.80 | 0.94 | 1.25 | 1.03 | 1.00 | 1.25 | 1.15 | 1.05 | 1.10 |
| 1993 | 0.34 | 0.47 | 0.89 | 0.82 | 1.03 | 1.03 | 1.14 | 1.08 | 1.16 | 1.19 | 1.20 | 1.33 | 1.14 | 1.14 |
| 1994 | 0.34 | 0.47 | 0.64 | 0.84 | 0.97 | 1.14 | 1.14 | 1.12 | 1.19 | 1.24 | 1.27 | 1.06 | 1.09 | 1.15 |
| 1995 | 0.34 | 0.55 | 0.85 | 0.75 | 1.13 | 1.33 | 1.40 | 1.36 | 1.43 | 1.42 | 1.50 | 1.45 | 1.66 | 1.32 |
| 1996 | 0.34 | 0.54 | 0.48 | 0.93 | 1.03 | 1.18 | 1.28 | 1.39 | 1.47 | 1.35 | 1.38 | 1.36 | 1.46 | 1.30 |
| 1997 | 0.34 | 0.47 | 0.66 | 0.75 | 0.85 | 0.93 | 0.99 | 1.06 | 1.11 | 1.15 | 1.19 | 1.15 | 1.19 | 1.14 |
| 1998 | 0.34 | 0.40 | 0.76 | 0.74 | 0.98 | 1.06 | 1.09 | 1.25 | 1.31 | 1.48 | 1.28 | 1.30 | 1.36 | 1.45 |
| 1999 | 0.34 | 0.47 | 0.66 | 0.75 | 0.85 | 0.93 | 0.99 | 1.06 | 1.11 | 1.15 | 1.19 | 1.15 | 1.19 | 1.14 |
| 2000 | 0.34 | 0.47 | 0.66 | 0.75 | 0.85 | 0.93 | 0.99 | 1.06 | 1.11 | 1.15 | 1.19 | 1.15 | 1.19 | 1.14 |
| 2001 | 0.34 | 0.47 | 0.66 | 0.75 | 0.85 | 0.93 | 0.99 | 1.06 | 1.11 | 1.15 | 1.19 | 1.15 | 1.19 | 1.14 |
| 2002 | 0.34 | 0.47 | 0.66 | 0.75 | 0.85 | 0.93 | 0.99 | 1.06 | 1.11 | 1.15 | 1.19 | 1.15 | 1.19 | 1.14 |
| 2003 | 0.33 | 0.39 | 0.76 | 0.69 | 0.81 | 0.92 | 0.91 | 0.97 | 0.93 | 1.06 | 1.17 | 1.19 | 1.61 | 1.07 |
| 2004 | 0.33 | 0.39 | 0.76 | 0.69 | 0.81 | 0.92 | 0.91 | 0.97 | 0.93 | 1.06 | 1.17 | 1.19 | 1.61 | 1.07 |
| 2005 | 0.33 | 0.39 | 0.76 | 0.69 | 0.81 | 0.92 | 0.91 | 0.97 | 0.93 | 1.06 | 1.17 | 1.19 | 1.61 | 1.07 |
| 2006 | 0.33 | 0.39 | 0.76 | 0.69 | 0.81 | 0.92 | 0.91 | 0.97 | 0.93 | 1.06 | 1.17 | 1.19 | 1.61 | 1.07 |

Table 1A.18. Percentage mature females at age from Wespestad and Terry (1984).

| Age | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| Percent | 0.0 | 0.8 | 28.9 | 64.1 | 84.2 | 90.1 | 94.7 | 96.3 | 97.0 | 97.8 | 98.4 |

Table 1A.19. Comparisons of fits for the evaluations of Aleutian Islands pollock Model 1 and Model 2.

|  | Model 1 | Model 2A | Model 2B |
| :--- | ---: | ---: | ---: |
| Number of Parameters | 194 | 278 | 279 |
| Survey catchability | 1.00 | 1.00 | 1.00 |
| Fishery Average Effective N | 37 | 43 | 44 |
| Survey Average Effective N | 115 | 190 | 205 |
| RMSE Survey | 0.667 | 0.363 | 0.356 |
| -log Likelihoods |  |  |  |
| Survey index | 28.17 | 8.97 | 8.20 |
| Fishery age comp | 89.26 | 76.59 | 72.83 |
| Survey age comp | 27.91 | 20.34 | 19.42 |
| Sub total | 145.34 | 105.91 | 100.45 |
|  |  |  |  |
| Recruitment | -4.86 | -14.41 | -14.85 |
| Selectivity constraint | 15.53 | 15.42 | 17.08 |
| Prior | 1.20 | 0.00 | 0.08 |
| Total | 162.27 | 115.89 | 11.02 |

Table 1A.20. Key results for the evaluations of Aleutian Islands pollock Model 1 and Model 2.

|  | Model 1 | Model 2A | Model 2B |
| :---: | :---: | :---: | :---: |
| Model conditions |  |  |  |
| Survey catchability | 1.00 | 1.00 | 1.00 |
| Natural mortality | 0.30 | 0.30 | 0.22 |
| Fishing mortalities |  |  |  |
| Max F 1978-2006 | 0.909 | 0.742 | 0.699 |
| F 2006 | 0.030 | 0.013 | 0.029 |
| Stock abundance |  |  |  |
| Initial Biomass (1978; thousands of tons) | 327 | 467 | 280 |
| CV | 16\% | 17\% | 19\% |
| 2006 total biomass (thousands of tons) | 140 | 354 | 225 |
| CV | 22\% | 18\% | 20\% |
| 2007 Age 3+ biomass (thousands of tons) | 141 | 363 | 229 |
| 1978 year class (at age 2) | 140 | 354 | 225 |
| CV | 22\% | 18\% | 20\% |
| Recruitment Variability | 0.64 | 0.44 | 0.42 |
| Specified Sigma R | 0.60 | 0.60 | 0.60 |
| Steepness (h) | 0.7 | 0.7 | 0.7 |
| Projected catch (unadjusted) |  |  |  |
| F50\% 2006 catch | 21.2 | 66.4 | 34.5 |
| CV | 21\% | 20\% | 27\% |
| F40\% 2006 catch | 31.6 | 96.7 | 50.9 |
| CV | 21\% | 20\% | 27\% |
| F35\% 2006 catch | 38.3 | 114.9 | 61.8 |
| CV | 21\% | 20\% | 27\% |

Table 1A.21. Model 2B estimates of pollock biomass with approximate lower (LCI) and upper (UCI) $95 \%$ confidence bounds for age $2+$ biomass. Also included are the age $3+$ biomass and female spawning stock biomass (SSB) estimates.

| $\begin{aligned} & \hline \text { M 2B } \\ & \text { Year } \end{aligned}$ | Total Biomass (age 2+) |  |  | Biomass <br> Age 3+ | $\begin{aligned} & \text { Female } \\ & \text { SSB } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | LCI | UCI |  |  |
| 1978 | 280,040 | 173,988 | 386,092 | 229,574 | 98,023 |
| 1979 | 286,860 | 180,416 | 393,304 | 212,469 | 102,161 |
| 1980 | 330,910 | 200,418 | 461,402 | 257,290 | 105,797 |
| 1981 | 343,790 | 192,886 | 494,694 | 279,021 | 107,579 |
| 1982 | 346,550 | 190,000 | 503,100 | 257,546 | 103,767 |
| 1983 | 342,060 | 193,970 | 490,150 | 274,820 | 106,445 |
| 1984 | 335,280 | 198,728 | 471,832 | 279,987 | 124,051 |
| 1985 | 329,870 | 200,668 | 459,072 | 275,938 | 133,284 |
| 1986 | 325,790 | 208,136 | 443,444 | 259,501 | 130,771 |
| 1987 | 321,530 | 214,894 | 428,166 | 258,733 | 129,452 |
| 1988 | 314,320 | 218,188 | 410,452 | 234,958 | 127,921 |
| 1989 | 309,980 | 223,008 | 396,952 | 256,822 | 127,718 |
| 1990 | 308,690 | 229,096 | 388,284 | 224,199 | 124,334 |
| 1991 | 310,680 | 232,162 | 389,198 | 241,176 | 120,865 |
| 1992 | 316,670 | 241,098 | 392,242 | 257,961 | 118,736 |
| 1993 | 302,600 | 231,996 | 373,204 | 297,755 | 115,371 |
| 1994 | 282,190 | 217,486 | 346,894 | 259,711 | 115,861 |
| 1995 | 272,210 | 211,834 | 332,586 | 289,642 | 113,767 |
| 1996 | 218,720 | 160,800 | 276,640 | 214,014 | 110,218 |
| 1997 | 198,570 | 140,972 | 256,168 | 163,659 | 102,222 |
| 1998 | 175,220 | 115,374 | 235,066 | 154,908 | 82,340 |
| 1999 | 158,800 | 96,656 | 220,944 | 132,515 | 72,929 |
| 2000 | 167,890 | 102,650 | 233,130 | 136,434 | 62,024 |
| 2001 | 178,240 | 108,294 | 248,186 | 143,819 | 57,292 |
| 2002 | 201,260 | 119,648 | 282,872 | 156,677 | 61,644 |
| 2003 | 217,390 | 127,544 | 307,236 | 185,704 | 64,425 |
| 2004 | 221,000 | 129,030 | 312,970 | 185,179 | 68,821 |
| 2005 | 220,130 | 129,564 | 310,696 | 173,778 | 76,233 |
| 2006 | 219,820 | 130,172 | 309,468 | 173,463 | 83,038 |

Table 1A.22. Results from MCMC simulations with 1 million iterations sampled every $200^{\text {th }}$ iteration for reference Model 2B.

| Parameter | Mean | CV |
| ---: | ---: | ---: |
| Natural Mortality | 0.235 | $10 \%$ |
| Steepness | 0.65 | $27 \%$ |
| Depletion | 0.78 | $20 \%$ |
| 2006 Total Biomass | 248.45 | $22 \%$ |
| $\mathrm{~F}_{35 \%}$ | 0.90 | $32 \%$ |
| $\mathrm{~F}_{40 \%}$ | 0.68 | $30 \%$ |
| $\mathrm{~F}_{50 \%}$ | 0.40 | $27 \%$ |

Table 1A.23. Estimates of full-selection fishing mortality and exploitation rates for pollock based on the reference model (Model 2B).

| Year | $\boldsymbol{F}^{\boldsymbol{a}}$ | Catch/Biomass <br> Rate $^{\mathbf{b}}$ |
| :---: | :---: | :---: |
| 1978 | 0.008 | 0.004 |
| 1979 | 0.063 | 0.037 |
| 1980 | 0.211 | 0.116 |
| 1981 | 0.125 | 0.053 |
| 1982 | 0.022 | 0.012 |
| 1983 | 0.010 | 0.006 |
| 1984 | 0.055 | 0.033 |
| 1985 | 0.005 | 0.003 |
| 1986 | 0.004 | 0.003 |
| 1987 | 0.017 | 0.011 |
| 1988 | 0.004 | 0.002 |
| 1989 | 0.000 | 0.000 |
| 1990 | 0.085 | 0.047 |
| 1991 | 0.006 | 0.002 |
| 1992 | 0.093 | 0.033 |
| 1993 | 0.177 | 0.054 |
| 1994 | 0.067 | 0.023 |
| 1995 | 0.699 | 0.201 |
| 1996 | 0.359 | 0.108 |
| 1997 | 0.653 | 0.157 |
| 1998 | 0.642 | 0.151 |
| 1999 | 0.018 | 0.005 |
| 2000 | 0.024 | 0.007 |
| 2001 | 0.014 | 0.004 |
| 2002 | 0.008 | 0.002 |
| 2003 | 0.030 | 0.008 |
| 2004 | 0.032 | 0.009 |
| 2005 | 0.029 | 0.009 |
| 2006 | 0.034 | 0.009 |

${ }^{a}$ Full selection fishing mortality rates.
${ }^{\mathrm{b}}$ Catch/biomass rate is the ratio of catch to beginning year age 3+ biomass.

Table 1A.24. Estimated pollock numbers at age in millions, 1978-2006 for reference Model 2B.

| Model 2B | $\mathbf{2}$ | $\mathbf{3}$ | $\mathbf{4}$ | $\mathbf{5}$ | $\mathbf{6}$ | $\mathbf{7}$ | $\mathbf{8}$ | $\mathbf{9}$ | $\mathbf{1 0}$ | $\mathbf{1 1}$ | $\mathbf{1 2}$ | $\mathbf{1 3}$ | $\mathbf{1 4}$ | $\mathbf{1 5 +}$ | Total | $\mathbf{\%} \mathbf{\text { of } \mathbf { 1 5 + }}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1978 | 70 | 62 | 55 | 42 | 39 | 18 | 16 | 13 | 11 | 8 | 6 | 5 | 4 | 21 | 368 | $5.66 \%$ |
| 1979 | 92 | 56 | 49 | 44 | 33 | 31 | 14 | 13 | 10 | 9 | 7 | 5 | 4 | 20 | 387 | $5.10 \%$ |
| 1980 | 315 | 73 | 44 | 39 | 34 | 25 | 23 | 11 | 10 | 8 | 7 | 5 | 4 | 18 | 616 | $2.97 \%$ |
| 1981 | 71 | 249 | 56 | 32 | 27 | 23 | 17 | 15 | 7 | 7 | 5 | 5 | 4 | 16 | 533 | $2.99 \%$ |
| 1982 | 46 | 57 | 195 | 42 | 24 | 20 | 16 | 12 | 11 | 5 | 5 | 4 | 4 | 15 | 455 | $3.28 \%$ |
| 1983 | 75 | 37 | 45 | 154 | 33 | 19 | 16 | 13 | 9 | 8 | 4 | 4 | 3 | 15 | 436 | $3.38 \%$ |
| 1984 | 77 | 60 | 30 | 36 | 123 | 27 | 15 | 12 | 10 | 7 | 7 | 3 | 3 | 14 | 425 | $3.38 \%$ |
| 1985 | 78 | 62 | 48 | 23 | 28 | 95 | 20 | 11 | 9 | 8 | 6 | 5 | 2 | 14 | 410 | $3.32 \%$ |
| 1986 | 71 | 62 | 50 | 38 | 19 | 22 | 76 | 16 | 9 | 8 | 6 | 5 | 4 | 13 | 399 | $3.22 \%$ |
| 1987 | 57 | 57 | 50 | 40 | 31 | 15 | 18 | 61 | 13 | 7 | 6 | 5 | 4 | 14 | 377 | $3.63 \%$ |
| 1988 | 85 | 45 | 46 | 40 | 32 | 24 | 12 | 14 | 48 | 10 | 6 | 5 | 4 | 14 | 385 | $3.58 \%$ |
| 1989 | 78 | 68 | 36 | 37 | 32 | 25 | 19 | 9 | 11 | 38 | 8 | 5 | 4 | 14 | 387 | $3.67 \%$ |
| 1990 | 73 | 63 | 55 | 29 | 30 | 26 | 20 | 16 | 8 | 9 | 31 | 7 | 4 | 14 | 384 | $3.78 \%$ |
| 1991 | 140 | 58 | 50 | 43 | 22 | 22 | 19 | 15 | 12 | 6 | 7 | 23 | 5 | 14 | 437 | $3.13 \%$ |
| 1992 | 69 | 113 | 47 | 40 | 34 | 18 | 18 | 15 | 12 | 9 | 4 | 5 | 19 | 15 | 419 | $3.55 \%$ |
| 1993 | 54 | 55 | 90 | 37 | 32 | 27 | 14 | 14 | 12 | 9 | 7 | 3 | 4 | 24 | 383 | $6.38 \%$ |
| 1994 | 56 | 44 | 44 | 71 | 29 | 24 | 20 | 10 | 10 | 9 | 7 | 5 | 2 | 19 | 350 | $5.43 \%$ |
| 1995 | 61 | 45 | 35 | 35 | 57 | 23 | 19 | 16 | 8 | 8 | 7 | 5 | 4 | 16 | 337 | $4.73 \%$ |
| 1996 | 45 | 49 | 36 | 27 | 25 | 37 | 14 | 11 | 9 | 5 | 4 | 3 | 2 | 7 | 274 | $2.63 \%$ |
| 1997 | 55 | 36 | 39 | 28 | 20 | 18 | 26 | 10 | 8 | 6 | 3 | 3 | 2 | 5 | 257 | $1.94 \%$ |
| 1998 | 61 | 44 | 28 | 30 | 20 | 14 | 12 | 16 | 6 | 5 | 4 | 1 | 1 | 3 | 243 | $1.07 \%$ |
| 1999 | 42 | 48 | 34 | 22 | 21 | 14 | 9 | 7 | 10 | 4 | 3 | 2 | 1 | 1 | 218 | $0.65 \%$ |
| 2000 | 57 | 34 | 39 | 28 | 17 | 17 | 11 | 7 | 6 | 8 | 3 | 2 | 1 | 2 | 232 | $0.68 \%$ |
| 2001 | 71 | 46 | 27 | 31 | 22 | 14 | 14 | 9 | 6 | 5 | 6 | 2 | 2 | 2 | 256 | $0.90 \%$ |
| 2002 | 114 | 57 | 37 | 22 | 25 | 18 | 11 | 11 | 7 | 4 | 4 | 5 | 2 | 3 | 319 | $0.98 \%$ |
| 2003 | 45 | 91 | 46 | 30 | 17 | 20 | 14 | 9 | 9 | 5 | 4 | 3 | 4 | 4 | 302 | $1.31 \%$ |
| 2004 | 41 | 36 | 73 | 37 | 24 | 14 | 16 | 11 | 7 | 7 | 4 | 3 | 2 | 6 | 281 | $2.23 \%$ |
| 2005 | 55 | 33 | 29 | 59 | 29 | 19 | 11 | 13 | 9 | 6 | 5 | 3 | 2 | 7 | 280 | $2.39 \%$ |
| 2006 | 59 | 45 | 26 | 23 | 47 | 23 | 15 | 9 | 10 | 7 | 4 | 4 | 3 | 7 | 283 | $2.44 \%$ |

Table 1A.25. Estimates of age-2 pollock recruitment (in millions) based on reference model.

| Year | Index at age 2 |
| :---: | ---: |
| 1978 | 69.8 |
| 1979 | 91.9 |
| 1980 | 314.5 |
| 1981 | 71.4 |
| 1982 | 46.0 |
| 1983 | 74.9 |
| 1984 | 77.1 |
| 1985 | 77.6 |
| 1986 | 71.3 |
| 1987 | 56.6 |
| 1988 | 84.8 |
| 1989 | 78.5 |
| 1990 | 73.0 |
| 1991 | 140.5 |
| 1992 | 68.9 |
| 1993 | 54.3 |
| 1994 | 56.4 |
| 1995 | 61.1 |
| 1996 | 44.6 |
| 1997 | 54.5 |
| 1998 | 60.5 |
| 1999 | 41.8 |
| 2000 | 57.2 |
| 2001 | 71.1 |
| 2002 | 114.0 |
| 2003 | 45.4 |
| 2004 | 40.5 |
| 2005 | 55.4 |
| 2006 | 59.4 |
| Ave $83-06$ | 66.2 |
| Med $83-06$ | 60.0 |
|  |  |

Table 1A. 26 Estimates of 2005 pollock fishery, survey, and projected fishery selectivity-at-age for Model 2B.

|  | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| M2B Fishery | 0.025 | 0.060 | 0.153 | 0.376 | 0.716 | 0.814 | 0.801 | 0.843 | 0.987 | 1.216 | 1.647 |
| Projected* | 0.016 | 0.110 | 0.438 | 0.949 | 1.470 | 1.639 | 1.621 | 1.470 | 1.290 | 1.198 | 1.198 |
| M2B Survey | 0.065 | 0.183 | 0.404 | 0.565 | 0.656 | 0.714 | 0.783 | 0.861 | 0.982 | 1.157 | 1.282 |

* From the 2005 EBS pollock stock assessment (Ianelli et al. 2005).

Table 1A.27. Projections of Model 2B (with adjusted selectivity) female spawning biomass (in thousands of t ), $F$, and catch (in thousands of t ) for NRA pollock for the 8 scenarios.
Fishing mortality rates given are based on the average fishing mortality over all ages.

| Sp.Biomass | Scenario 1 | Scenario 2 | Scenario 3 | Scenario 4 | Scenario 5 | Scenario 6 | Scenario 7 | Scenario 8 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2006 | 90.45 | 90.45 | 90.45 | 90.45 | 90.45 | 90.45 | 90.45 | 90.45 |
| 2007 | 82.21 | 82.21 | 84.86 | 87.04 | 87.61 | 80.70 | 82.21 | 87.19 |
| 2008 | 58.41 | 58.41 | 70.37 | 82.14 | 85.56 | 52.63 | 58.41 | 82.60 |
| 2009 | 48.47 | 48.47 | 63.75 | 81.48 | 87.13 | 42.11 | 47.77 | 80.32 |
| 2010 | 44.48 | 44.48 | 60.40 | 81.46 | 88.69 | 38.79 | 41.43 | 77.53 |
| 2011 | 42.97 | 42.97 | 58.61 | 81.65 | 90.06 | 38.08 | 39.13 | 70.52 |
| 2012 | 42.42 | 42.42 | 57.78 | 82.52 | 92.00 | 37.91 | 38.31 | 65.41 |
| 2013 | 42.04 | 42.04 | 57.35 | 83.82 | 94.42 | 37.71 | 37.86 | 61.54 |
| 2014 | 41.70 | 41.70 | 56.69 | 83.82 | 95.02 | 37.50 | 37.54 | 57.76 |
| 2015 | 41.55 | 41.55 | 56.29 | 83.80 | 95.41 | 37.42 | 37.43 | 54.78 |
| 2016 | 41.63 | 41.63 | 56.40 | 84.84 | 97.14 | 37.53 | 37.53 | 52.75 |
| 2017 | 41.77 | 41.77 | 56.55 | 85.65 | 98.49 | 37.67 | 37.67 | 51.09 |
| 2018 | 41.72 | 41.72 | 56.56 | 86.25 | 99.55 | 37.62 | 37.62 | 49.58 |
| 2019 | 41.46 | 41.46 | 56.32 | 86.33 | 99.95 | 37.35 | 37.35 | 48.06 |
| $F$ | Scenario 1 | Scenario 2 | Scenario 3 | Scenario 4 | Scenario 5 | Scenario 6 | Scenario 7 | Scenario 8 |
| 2006 | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 | 0.01 |
| 2007 | 0.29 | 0.29 | 0.15 | 0.03 | 0.00 | 0.38 | 0.29 | 0.02 |
| 2008 | 0.29 | 0.29 | 0.15 | 0.03 | 0.00 | 0.38 | 0.29 | 0.05 |
| 2009 | 0.29 | 0.29 | 0.15 | 0.03 | 0.00 | 0.37 | 0.38 | 0.05 |
| 2010 | 0.29 | 0.29 | 0.15 | 0.03 | 0.00 | 0.34 | 0.36 | 0.14 |
| 2011 | 0.28 | 0.28 | 0.15 | 0.03 | 0.00 | 0.34 | 0.34 | 0.15 |
| 2012 | 0.28 | 0.28 | 0.15 | 0.03 | 0.00 | 0.34 | 0.34 | 0.17 |
| 2013 | 0.28 | 0.28 | 0.15 | 0.03 | 0.00 | 0.34 | 0.34 | 0.18 |
| 2014 | 0.28 | 0.28 | 0.15 | 0.03 | 0.00 | 0.34 | 0.34 | 0.20 |
| 2015 | 0.28 | 0.28 | 0.15 | 0.03 | 0.00 | 0.33 | 0.33 | 0.21 |
| 2016 | 0.28 | 0.28 | 0.15 | 0.03 | 0.00 | 0.34 | 0.34 | 0.23 |
| 2017 | 0.28 | 0.28 | 0.15 | 0.03 | 0.00 | 0.34 | 0.34 | 0.24 |
| 2018 | 0.28 | 0.28 | 0.15 | 0.03 | 0.00 | 0.34 | 0.34 | 0.25 |
| 2019 | 0.28 | 0.28 | 0.15 | 0.03 | 0.00 | 0.33 | 0.33 | 0.26 |
| Catch | Scenario 1 | Scenario 2 | Scenario 3 | Scenario 4 | Scenario 5 | Scenario 6 | Scenario 7 | Scenario 8 |
| 2006 | 1.90 | 1.90 | 1.90 | 1.90 | 1.90 | 1.90 | 1.90 | 1.90 |
| 2007 | 44.47 | 44.47 | 24.40 | 5.39 | 0.00 | 54.54 | 44.47 | 4.00 |
| 2008 | 29.63 | 29.63 | 19.23 | 4.89 | 0.00 | 33.09 | 29.63 | 7.60 |
| 2009 | 22.86 | 22.86 | 16.33 | 4.57 | 0.00 | 23.98 | 28.29 | 7.60 |
| 2010 | 20.12 | 20.12 | 15.17 | 4.48 | 0.00 | 20.45 | 22.77 | 19.00 |
| 2011 | 19.56 | 19.56 | 15.15 | 4.65 | 0.00 | 20.26 | 21.20 | 19.00 |
| 2012 | 19.47 | 19.47 | 15.21 | 4.80 | 0.00 | 20.40 | 20.74 | 19.00 |
| 2013 | 19.33 | 19.33 | 15.17 | 4.91 | 0.00 | 20.33 | 20.45 | 19.00 |
| 2014 | 19.18 | 19.18 | 15.09 | 4.99 | 0.00 | 20.20 | 20.24 | 19.00 |
| 2015 | 18.99 | 18.99 | 14.85 | 4.93 | 0.00 | 20.05 | 20.06 | 19.00 |
| 2016 | 18.95 | 18.95 | 14.80 | 4.96 | 0.00 | 20.04 | 20.05 | 19.00 |
| 2017 | 19.02 | 19.02 | 14.81 | 4.98 | 0.00 | 20.15 | 20.15 | 19.00 |
| 2018 | 19.04 | 19.04 | 14.80 | 5.01 | 0.00 | 20.17 | 20.17 | 19.00 |
| 2019 | 18.95 | 18.95 | 14.79 | 5.03 | 0.00 | 20.04 | 20.04 | 19.00 |


| Table 1A.28. Ecosystem effects on AI walleye pollock |  |  |  |
| :---: | :---: | :---: | :---: |
| Indicator | Observation | Interpretation | Evaluation |
| Prey availability or abundance trends |  |  |  |
| Zooplankton | Stomach contents, ichthyoplankton surveys | None | Unknown |
| Predator population Marine mammals | trends <br> Fur seals declining, Steller sea lions increasing slightly | Possibly lower mortality on walleye pollock | No concern |
| Birds | Stable, some increasing some decreasing | May affect young-of-year mortality | Unknown |
| Fish (Pacific cod, arrowtooth flounder) | Pacific cod-decreasing, arrowtooth--stable | Possible decreases to walleye pollock mortality | No concern |
| Changes in habitat quality |  |  |  |
| Temperature regime | The 2004 and 2006AI summer bottom temperature was near average. A warming since 2000 and 2002 were coldest and second coldest survey years respectively. | Warming from 2002 could affect apparent distribution. | Unknown |
| The AI walleye pollock effects on ecosystem |  |  |  |
| Indicator | Observation | Interpretation | Evaluation |
| Fishery contribution to bycatch |  |  |  |
| Prohibited species | Expected to be heavily monitored | Likely to be a minor contribution to mortality | No concern |
| Forage (including herring, Atka mackerel, cod, and pollock) | Expected to be heavily monitored. | Bycatch levels should be low. | Unknown |
| HAPC biota (seapens/whips, corals, sponges, anemones) | Very low bycatch levels of seapens/whips, sponge and coral catches expected in the pelagic fishery | Bycatch levels and destruction of benthic habitat expected to be minor given the pelagic fishery. | No concern |
| Marine mammals and birds | Very minor direct-take expected | Likely to be very minor contribution to mortality | No concern |
| Sensitive nontarget species | Expected to be heavily monitored | Unknown given that this fishery was closed between 1999 and 2005. The 2006 AICASS had 3\% POP bycatch, the only significant bycatch. The 2005 fishery had a high bycatch of POP, but bycatch of other species was very low in fishery prior to 1999. | No concern |
| Other non-target species | Very little bycatch. | Unknown | No concern |
| Fishery concentration in space and time | Steller sea lion protection measures may concentrate fishery spatially to very small areas between 20 nm closures | Depending on concentration of pollock outside of critical habitat could possibly have an effect. | Possible concern |
| Fishery effects on amount of large size target fish | Depends on highly variable year-class strength | Natural fluctuation | Possible Concern |
| Fishery contribution to discards and offal production | Offal production-unknown. Fishery in 2005 expected to be conducted by CPs which may have fish meal production capabilities | Unknown | Unknown |
| Fishery effects on age-at-maturity and fecundity | Unknown | Unknown | Unknown |

## Figures



Figure 1A.1. Estimated pollock catch by sub-area of the Aleutian Islands Region, 1977-2003. Units in metric tons.


Figure 1A.2. Observed foreign and J.V. (1978-1989), and domestic (1989-2002) pollock catch in the Aleutian Islands Area summed over all years and 10 minute latitude and longitude blocks. Both maps use the same scale (maximum observed catch per 10 minute block: foreign and J.V. 8,000 t and Domestic 19,000 t). Catches of less than 1 t were excluded from cumulative totals.


Figure 1A.3. Regions defined for consideration of alternative data partitions for Aleutian Islands Region pollock. The abbreviation "NRA" represents the Near, Rat, and Andreanof Island groups.


Figure 1A.4. Mean longitude of observed targeted domestic (1990-1998) pollock catch in the NRA west of 174 W longitude. Error bars indicate one standard deviation from the mean.


Figure 1A.5. Catch per unit effort ( kg per $\mathrm{m}^{3}$ ) for surveys of pollock in the Aleutian Islands Region, 2002-2006. The shaded area is the region surveyed.


Figure 1A.6. Pollock CPUE (KG per $\mathrm{m}^{3}$ ) by depth and temperature from the 2004 Aleutian Islands and Bering Sea and 2005 Gulf of Alaska bottom trawl surveys. Circle area is proportional to CPUE.


Figure 1A.7. Pollock CPUE (KG per m ${ }^{3}$ ) by depth and temperature from the 2004 (red) and 2006 (blue) Aleutian Islands bottom trawl surveys. Circle area is proportion to CPUE.


Figure 1A.8. Pollock CPUE (KG per $\mathrm{m}^{3}$ ) by depth and temperature from the 1999 (blue) and 2004 (red) Bering Sea bottom trawl surveys. Circle area is proportional to CPUE.


Figure 1A.9. Length at age for Aleutian Islands (red), Gulf of Alaska (blue), and Bering Sea (grey) pollock from the 2004 Aleutian Islands, 2004 Bering Sea, and 2005 Gulf of Alaska bottom trawl surveys.



Figure 1A.11. R/V Kaiyo Maru 2002 echo integration-trawl survey (above) strata for leg2 and below observed $\mathrm{S}_{\mathrm{A}}$ in both legs. Please note that in the bottom picture the encircled area is leg 2.


Figure 1A.12. 2006 AICASS Survey Area.


Figure 1A.13. Proportion of total weight of pollock at age from the 2006 AICASS. The age 15 group represents all fish age 15 and older.


Figure 1A.14. Pollock abundance estimation and cumulative catch for large (top) and small (bottom) survey areas. Note error bars are $\pm 1.96 \times E_{i} \times B_{i}$. Method proposed by Kloser 1996 used to estimate biomass in the "Deadzone."


Figure 1A.15. 2006 AICASS distributions of pollock. Figures from left to right correspond to Surveys $2,4,6,7$, and 8 .


Figure 1A.16. Change in abundance for All, High Density, and Low Density areas. The High Density area corresponds with the small survey area while All corresponds with the large survey area, and Low Density corresponds with the large survey area outside of the small survey area. Arrows indicate a significant change in abundance from the first survey. Note: 935 t of the total 965 t caught during the 2006 AICASS were removed from the High Density area.


Figure 1A.17. Female pollock maturity over the duration of the 2006 AICASS, $\mathrm{m} 1=$ immature, $\mathrm{m} 2=$ developing, $\mathrm{m} 3=$ pre-spawning, $\mathrm{m} 4=$ spawning, and $\mathrm{m} 5=$ spent. There were no spent fish observed during this survey.


Figure 1A.18. Biomass trajectories under the three evaluated models.


Figure 1A. 192006 total biomass (right) and natural mortality (left) distributions from MCMC runs of Model 2B. Distributions were generated through 1,000,000 MCMC simulations sampled every 200 simulation.


Figure 1A. 20 Trace of natural mortality from MCMC simulations generated through 1,000,000 simulations sampled every $200^{\text {th }}$ iteration for the two 6 models. The purple line is a running mean for every $200^{\text {th }}$ sampled iteration and the red line is a linear fit to the data showing a flat (slope of $-9 \times 10^{-7}$ ) trajectory over 5000 iterations.


Figure 1A.21. Fit (solid line) to NMFS summer trawl survey (dots) for Model 2B. Dashed lines represent upper and lower confidence bounds of survey estimates.
Model_2B Survey Age Composition Data

AGE

+ Observed - Predicted

Figure 1A.22. Fits to NMFS summer trawl survey age composition data for Model 2B for Aleutian Islands pollock.


Figure 1A.23. Fit to fishery age composition data for Model 2B for Aleutian Islands (NRA) pollock.


Figure 1A.24. Selectivity estimates for Aleutian Islands pollock for the bottom trawl survey (left) and the fishery (right) Model 2B.


Figure 1A.25. Model 2B estimates of Aleutian Islands pollock age $2+$ total biomass (in tons); dashed lines represent approximate upper and lower confidence bounds.


Figure 1A. 26 Spawning biomass relative to F40\% values and fishing mortality rates for Model 2B AI pollock over time (top) and plotted jointly (bottom) for 1978-2006. Fishing mortality rates are based on the average over ages 2-15.


Figure 1A.27. Model 2B estimates of Aleutian Islands (NRA assessment area) pollock year-class estimates; vertical bars represent approximate upper and lower confidence bounds.


Figure 1A. 28 Projected spawning biomass for $\mathrm{F}_{40 \%}$ and Alternative 8 ABC scenarios from Model 2B with adjusted selectivity-at-age.


Figure 1A. 29 Projected catch for $\mathrm{F}_{40 \%}$ and Alternative 8 ABC scenarios from Model 2B with adjusted selectivity-at-age.


Figure 1A.30. Diet composition (left) and estimated consumption of prey (right) by AI adult (top) and juvenile (bottom) pollock. Diets are estimated from stomach collections taken aboard NMFS bottom trawl surveys in 1991-1994. See Appendix A for detailed methods.


Figure 1A.31. Mortality sources (left) and estimated consumption by predators (right) of AI adult (top) and juvenile (bottom) pollock. Mortality sources reflect pollock predator diets estimated from stomach collections taken aboard NMFS bottom trawl surveys in 1991-1994, pollock predator consumption rates estimated from stock assessments and other studies, and catch of pollock by all fisheries in the same time periods. Annual consumption ranges incorporating uncertainty in food web model parameters were estimated by the Sense routines (Aydin et al in review). See Appendix A for detailed methods.


Figure 1A.32. The pollock trawl fishery in the AI food web. Species taken by the pollock fishery (in red) are highlighted in green, with the most significant flow to pollock indicated with a green line. Box size is proportional to biomass and lines between boxes represent the most significant energy flows. From Aydin et al (in review).


Figure 1A.33. Catch composition of the AI pollock trawl fishery during the early 1990 's, as used in the food web model (Aydin et al Tech Memo).


Figure 1A.34. Adult and juvenile pollock (highlighted in red) in the AI food web (Aydin et al Tech Memo). Predators of pollock are dark blue, prey of pollock are green, and species that are both predators and prey of pollock are light blue. Box size is proportional to biomass and lines between boxes represent the most significant energy flows.


Figure 1A.35. (upper panel) Effect of changing pollock survival on fishery catch (yellow) and biomass of other species (dark red), from a simulation analysis where pollock survival was decreased by $10 \%$ and the rest of the ecosystem adjusted to this decrease for 30 years. (lower panel) Effect of reducing fisheries catch (yellow) and other species survival (dark red) on pollock biomass, from a simulation analysis where survival of each X axis species group was decreased by $10 \%$ and the rest of the ecosystem adjusted to this decrease for 30 years. In both panels, boxes show resulting percent change in the biomass of each species on the x axis after 30 years for $50 \%$ of feasible ecosystems, error bars show results for $95 \%$ of feasible ecosystems (see Aydin et al in review for detailed Sense methods).

## Appendix A

## Diet composition calculations

Notation:

$$
\begin{array}{lll}
\text { DC = diet composition } & \mathrm{s}=\text { predator size class } & \mathrm{v}=\text { survey } \\
\mathrm{W}=\text { weight in stomach } & \mathrm{h}=\text { survey haul } & \mathrm{a}=\text { assessment } \\
\mathrm{n}=\text { prey } & \mathrm{r}=\text { survey stratum } & \mathrm{R}=\text { ration estimate } \\
\mathrm{p}=\text { predator } & \mathrm{B}=\text { biomass estimate } &
\end{array}
$$

The diet composition for a species is calculated from stomach sampling beginning at the level of the individual survey haul (1), combining across hauls within a survey stratum (2), weighting stratum diet compositions by stratum biomass (3), and finally combining across predator size classes by weighting according to size-specific ration estimates and biomass from stock assessment estimated age structure (4). Ration calculations are described in detail below.

Diet composition (DC) of prey $n$ in predator $p$ of size $s$ in haul $h$ is the total weight of prey $n$ in all of the stomachs of predator $p$ of size $s$ in the haul divided by the sum over all prey in all of the stomachs for that predator size class in that haul:

$$
\begin{equation*}
D C_{n, p, s, h}=W_{n, p, s, h} / \sum_{n} W_{n, p, s, h} \tag{1}
\end{equation*}
$$

Diet composition of prey n in predator p of size s in survey stratum r is the average of the diet compositions across hauls within that stratum:
$D C_{n, p, s, r}=\sum_{h} D C_{n, p, s, h} / h$
Diet composition of prey $n$ in predator $p$ of size $s$ for the entire area $t$ is the sum over all strata of the diet composition in stratum $r$ weighted by the survey biomass proportion of predator $p$ of size $s$ in stratum $r$ :
$D C_{n, p, s, t}=\sum_{r} D C_{n, p, s, r} * B_{p, s, r}^{v} / \sum_{r} B_{p, s, r}^{v}$
Diet composition of prey n in predator p for the entire area t is the sum over all predator sizes of the diet composition for predator $p$ of size $s$ as weighted by the relative stock assessment biomass of predator size $s$ times the ration of predator p of size s :

$$
\begin{equation*}
D C_{n, p, t}=\sum_{s} D C_{n, p, s, t} * B_{p, s}^{a} * R_{p, s} / \sum_{s} B_{p, s}^{a} * R_{p, s} \tag{4}
\end{equation*}
$$

## Ration Calculations

Size specific ration (consumption rate) for each predator was determined by the method of fitting the generalized Von Bertalanffy growth equations (Essington et al. 2001) to weight-at-age data collected aboard NMFS bottom trawl surveys.

The generalized Von Bertalanffy growth equation assumes that both consumption and respiration scale allometrically with body weight, and change in body weight over time (dW/dT) is calculated as follows (Paloheimo and Dickie 1965):

$$
\begin{equation*}
\frac{d W_{t}}{d t}=H \cdot W_{t}^{d}-k \cdot W_{t}^{n} \tag{5}
\end{equation*}
$$

Here, $W_{t}$ is body mass, $t$ is the age of the fish (in years), and $H, d, k$, and $n$ are allometric parameters. The term $H \cdot W_{t}^{d}$ is an allometric term for "useable" consumption over a year, in other words, the consumption (in wet weight) by the predator after indigestible portions of the prey have been removed and assuming constant caloric density between predator and prey. Total consumption $(Q)$ is calculated as $(1 / A) \cdot H \cdot W_{t}^{d}$, where $A$ is a scaling fraction between predator and prey wet weights that accounts for indigestible portions of the prey and differences in caloric density. The term $k \cdot W_{t}^{n}$ is an allometric term for the amount of biomass lost yearly as respiration.
Based on an analysis performed across a range of fish species, Essington et al. (2001) suggested that it is reasonable to assume that the respiration exponent $n$ is equal to 1 (respiration linearly proportional to body weight). In this case, the differential equation above can be integrated to give the following solution for weight-at-age:

$$
\begin{equation*}
W_{t}=W_{\infty} \cdot\left(1-e^{-k(1-d)\left(t-t_{0}\right)}\right)^{\frac{1}{1-d}} \tag{6}
\end{equation*}
$$

Where $W_{\infty}$ (asymptotic body mass) is equal to $(H / k)^{\frac{1}{1-d}}$, and $t_{0}$ is the weight of the organism at time $=0$. If the consumption exponent $d$ is set equal to $2 / 3$, this equation simplifies into the "specialized" von Bertalanffy length-at-age equation most used in fisheries management, with the "traditional" von Bertalanffy K parameter being equal to the $k$ parameter from the above equations divided by 3 .
From measurements of body weight and age, equation 2 can be used to fit four parameters ( $W_{\infty}, d, k$, and $t_{0}$ ) and the relationship between $W_{\infty}$ and the $H, k$, and $d$ parameters can then be used to determine the consumption rate $H \cdot W_{t}^{d}$ for any given age class of fish. For these calculations, weight-at-age data available and specific to the modeled regions were fit by minimizing the difference between $\log$ (observed) and $\log$ (predicted) body weights as calculated by minimizing negative log likelihood: observation error was assumed to be in weight but not aging. A process-error model was also examined but did not give significantly different results.
Initial fitting of 4-parameter models showed, in many cases, poor convergence to unique minima and shallow sum-of-squares surfaces: the fits suffered especially from lack of data at the younger age classes that would allow fitting to body weights near $t=0$ or during juvenile, rapidly growing life stages. To counter this, the following multiple models were tested for goodness-of-fit:

1. All four parameters estimated by minimization;
2. $d$ fixed at $2 / 3$ (specialized von Bertalanffy assumption)
3. $d$ fixed at 0.8 (median value based on metaanalysis by Essington et al. 2001).
4. $t_{0}$ fixed at 0 .
5. $d$ fixed at $2 / 3$ with $t_{0}$ fixed at 0 , and $d$ fixed at 0.8 with $t_{0}$ fixed at 0 .

The multiple models were evaluated using Aikeike's Information Criterion, AIC (spreadsheet review). In general, the different methods resulted in a twofold range of consumption rate estimates; consistently, model \#3, $d$ fixed at 0.8 while the other three parameters were free, gave the most consistently good results using the AIC. In some cases model \#1 was marginally better, but in some cases, model \#1 failed to converge. The poorest fits were almost always obtained by assuming that d was fixed at $2 / 3$.
To obtain absolute consumption $(Q)$ for a given age class, the additional parameter $A$ is required to account for indigestible and otherwise unassimilated portions of prey. We noted that the range of indigestible percentage for
a wide range of North Pacific zooplankton and fish summarized in Davis (2003) was between 5-30\%, with major zooplankton (copepods and euphasiids), as well as many forage fish, having a narrower range of indigestible percentages, generally between 10-20\%. Further, bioenergetics models, for example for walleye pollock (Buckley and Livingston), indicate that nitrogenous waste (excretion) and egestion resulted in an additional 20-30\% loss of consumed biomass. As specific bioenergetics models were not available for most species, we made a uniform assumption of a total non-respirative loss of $40 \%$ (from a range of $25-60 \%$ ) for all fish species, with a corresponding $A$ value of 0.6 .

Finally, consumption for a given age class was scaled to population-level consumption using the available numbers-at-age data from stock assessments, or using mortality rates from stock assessments and the assumption of an equilibrium age structure in cases where numbers-at-age reconstructions were not available.

## Production rates

Production per unit biomass $(\mathrm{P} / \mathrm{B})$ and consumption per unit biomass $(\mathrm{Q} / \mathrm{B}=\mathrm{R}$, ration above) for a given population depend heavily on the age structure, and thus mortality rate of that population. For a population with an equilibrium age structure, assuming exponential mortality and Von Bertalanffy growth, $\mathrm{P} / \mathrm{B}$ is in fact equal to total mortality Z (Allen 1971) and $\mathrm{Q} / \mathrm{B}$ is equal to $(\mathrm{Z}+3 \mathrm{~K}) / \mathrm{A}$, where K is Von Bertalanffy's K , and A is a scaling factor for indigestible proportions of prey (Aydin 2004). If a population is not in equilibrium, $\mathrm{P} / \mathrm{B}$ may differ substantially from Z although it will still be a function of mortality.

For the Bering Sea, Aleutian Islands, and Gulf of Alaska ECOPATH models, $\mathrm{P} / \mathrm{B}$ and $\mathrm{Q} / \mathrm{B}$ values depend on available mortality rates, which were taken from estimates or literature values used in single-species models of the region. It is noted that the single-species model assumptions of constant natural mortality are violated by definition in multispecies modeling; therefore, these estimates should be seen as "priors" to be input into the ECOPATH balancing procedures or other parameter-fitting (e.g. Bayesian) techniques.

Several methods were used to calculate $\mathrm{P} / \mathrm{B}$, depending on the level of data available. Proceeding from most data to least data, the following methods were used:

1. If a population is not in equilibrium, total production $P$ for a given age class over the course of a year can be approximated as $\left(\mathrm{N}_{\mathrm{at}} \cdot \Delta \mathrm{W}_{\mathrm{at}}\right)$, where $\mathrm{N}_{\mathrm{at}}$ is the number of fish of a given age class in a given year, exponentially averaged to account for mortality throughout the year, and $\Delta \mathrm{W}_{\mathrm{at}}$ is the change in body weight of that age class over that year. For a particular stock, if weight-at-age data existed for multiple years, and stock-assessment reconstructed numbers-at-age were also available, production was calculated by summing this equation over all assessed age classes. Walleye pollock $\mathrm{P} / \mathrm{B}$ for both the EBS and GOA were calculated using this method: examining the components of this sum over the years showed that numbers-at-age variation was responsible for considerably more variability in overall P/B than was weight-at-age variation.
2. If stock assessment numbers-at-age were available, but a time series of weight-at-age was not available and some weight-at-age data was available, the equation in (1), above, was used, however, the change in body weight over time was estimated using fits to the generalized Von Bertalanffy equations described in the consumption section, above.
3. If no stock assessment of numbers-at-age was available, the population was assumed to be in equilibrium, so that $\mathrm{P} / \mathrm{B}$ was taken to equal Z . In cases for many nontarget species, estimates of Z were not available so estimates of M were taken from conspecifics with little assumed fishing mortality for this particular calculation.

[^0]:    ${ }^{1}$ The likelihood is quasi because model penalties (e.g., non-parametric smoothers) are included.

