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Jack mackerel status in the South Eastern Pacific and exploitation management strategies
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## Technical Report

Jack mackerel (Trachurus murphyi) status in the South Eastern Pacific and exploitation management strategies

## OCEANA-CAPES PROJECT

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## SUMMARY

A stock assessment of the South Eastern Pacific (SEP) jack mackerel stock is conducted considering two hypotheses about stock structure and two scenarios for the steepness of the stock-recruitment function. An integrated statistical catch-at-age model was implemented considering the following fishery-dependent data: i) catches in the SEP from 1970 to 2014 grouped in four main fleets (North Chile, South Central Chile, Far North and Offshore trawl), ii) CPUE indices for South-central Chile, Far North, China, European Union and Russia, and iii) age- structures from north Chile, south central Chile, Offshores Trawl and length-structures for Far North. Independent-fishery data corresponded to biomass acoustic surveys for the North and South central Chile and Far North, spawning biomass estimates from the daily egg production method for centralsouth of Chile and age structures from Chilean surveys. Life history parameters such as somatic growth were taken from the most recent and updated parameters available from literature. Catches and age-structures of the Chilean fleets were updated with complete information up to 2014. We used the model developed by the Scientific Committee of the South Pacific Regional Fishery Management Organization (SPRFMO), an adopted in 2010 as an assessment method of the jack mackerel stock. The two hypotheses evaluated were: i) Jack mackerel is one single population in the South Eastern Pacific area (Model1) and ii) Jack mackerel constitute two discrete populations one in the Peruvian and Ecuadorian waters, and one off Chile extending onto the high seas, (Model-2). We assessed Model-1 by optimizing the model keeping all the information available. Model2 was evaluated by removing all data from the Far North fleet, CPUE and survey from Peru and related underpinning processes in the model. Testing both hypotheses lead to the conclusion that the biomass from Peru and Ecuador did not make a significant contribution to the spawning biomass of the jack mackerel stock in the South Pacific. All combinations of stock hypotheses and scenarios of $h(h=0.80$ and $h=0.65)$ lead to the same diagnosis of stock status. Jack mackerel stock in South Eastern Pacific is overexploited and with some probability of being depleted (close to Blim). We explore robustness of the model using a retrospective analysis concluding that he model proposed is robust to estimate abundance in jack mackerel.
Projections of the abundance included the combination of two level of recruitments (1970-2012 and 2000-2012), two level of steepness ( $h=0.80$ and $h=0.65$ ) and five level of fishing mortality (F) obtained from different multiplier of F in 2014. Constant fishing mortality was assumed as an exploitation strategy. We assessed the probabilities to reach the maximum sustainable yield (MSY) or the $80 \%$ of MSY under different combinations of recruitment, steepness and F. Results show that the selection of recruitment period for projections is the factor that mostly influences the recovery of abundance. Steepness plays a secondary role, by influencing the time or the probability required to accomplish the management strategies based on MSY. Using long periods of recruitment (1970-
2012) is misleading and overestimates the rebuilding capacity of the stock; thus the last period of recruitment (2000-2012) is recommended when simulating exploitation strategies. Harvesting at a $50 \%$ of fishing mortality estimated in 2014 has a $30 \%$ chance of reaching Bmsy at the end of the projection period (year 2034). Harvesting with any higher fishing mortality ( $>50 \% \mathrm{~F}$ ), will reduce the chances below an $8 \%$ to reach $B m s y$ in 2034 and thus it is not recommendable as a sustainable management strategy. Finally, we present preliminary results of a population dynamic model for understanding the interaction between jack mackerel (Trachurus murphyi) and the fishery. This analysis suggests that jack mackerel and fishery dynamics are linked in a predator/prey like system of mutual causal second-order loop. In addition, recruitment dynamics appears to interact with density-dependent process and El Niño variability. This dynamics model shed light about population processes usually not considered in integrated stock assessments models and thus, it provides a promising and complementary tool in the analysis of fish dynamics. Thus, further work in this project involves the integration of these results of population dynamic models in the context of stock assessment and management framework.

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## 1. INTRODUCTION

Jack mackerel (Trachurus murphyi, Nichols 1920) is a transboundary fishery resource and widespread specie throughout the South Pacific (SP), along the coastal and oceanic waters adjacent to Ecuador, Peru, and Chile, and along the Subtropical Convergence Zone. This has been described as the "Jack mackerel belt" that goes from the coast of Chile to New Zealand within a $35^{\circ} \mathrm{S}$ to $50^{\circ} \mathrm{S}$ variable band across the South Pacific (JMSWG-Report, 2014).

Five population structure hypotheses have been proposed for jack mackerel in the South Pacific, however only two of them are considered in the recent stock assessment (JMSWG-Report, 2014). First, jack mackerel conforms only one single population in the South Eastern Pacific, and the second proposes that jack mackerel constitute two discrete populations in the South Pacific: one in the Peruvian waters and, one off Chile extending onto the high seas (Hintzen et al. 2014; JMSWG-Report, 2014), although only the single population hypothesis is considered for management purposes.

The biological aspects indicate that jack mackerel has a low individual growth rate. A recent age validation study for jack mackerel caught off the Chilean coast was conducted by Cerna et al. (2011). The result of the study confirms that the by counting microincrements of the first annual annulus, also supported by previous studies conducted by Castillo and Arrizaga (1987) and Gili et al (1995), von Bertalanffy growth parameters estimated by Cerna et al. (2011) were: $L_{\infty}=72,7 \mathrm{~cm}, k=0,074\left(\right.$ year $\left.^{-1}\right)$ and $t_{0}=-$ $1,97\left(\right.$ year $\left.^{-1}\right)$. Differences in somatic growth have also been identified for the jack mackerel caught off the coast of Peru, with higher somatic growth rate compared to the jack mackerel caught off the coast of Chile. Growth parameters of jack mackerel caught off Peru corresponded to $L_{\infty}=80,77 \mathrm{~cm}, k=0,16\left(\right.$ year $\left.^{-1}\right)$ and $t_{0}=-0,356\left(\right.$ year $\left.^{-1}\right)$ (Dioses et al. 2015). Natural mortality for jack mackerel estimated using the empirical method in Pauly (1980) is $M=0.23$ ( year $^{-1}$ ) for Chile and $M=0.33$ ( year $^{-1}$ ), for Peru (JMSWG-Report, 2014).

Maturity at age also differs between Peru and Chile. The maturity at $L_{50 \%}$ off Chile occurs at 22.7 cm (between 2-3 years old) meanwhile for Peru $L_{50 \%}$ is estimated as 26.5 cm (near 4 years old) (Leal et al. 2012, Pereas et al. 2013 respectively). The main spawning period takes place between October and December, with a small spawning period reported between July and March (JMSWG-Report, 2014, Leal et al 2012).

The jack mackerel fishery in the South Eastern Pacific is shared by the fleets of Chile, Peru, Ecuador and by the distant water fleets from China, European Union, Faroe Islands, Korea, Japan, Russian Federation, Ukraine and Vanuatu operating outsize of the Exclusive Economic Zone (EEZ) (JMSWG-Report, 2014) off south central Chile.

Fisheries management of jack mackerel in the coastal states (Ecuador, Peru and Chile) started in the middle of 1990s. The total allowed catch was introduced in 1999 in Chile and in 1995 in Peru. In this last country there is a ban to use jack mackerel catches for fishmeal production. The jack mackerel fishery outsize of EEZ has been banned since 2010 to the entry of new fishing vessels, and since 2011 total allowable catch has been defined in the South Pacific Regional Fisheries Management Organization (SPRFMO) as tenure to control fishing mortality.

The international fishery management of the jack mackerel started in 2010, when the first stock assessment of the jack mackerel at the South Pacific scale was developed. These assessments have been developed in the context of SPRFMO where the management unit takes into account the jack mackerel fisheries from coastal states and distant water fleets.

The SPRFMO has established that non-government organization such as OCEANA can attend the Annual Commission and the Scientific Committee (SC) meetings (www.sprfmo.int/participation/). Latest catch recommendations for jack mackerel have been established with a strategy of constant fishing mortality at high risk of not recovering the spawning biomass by 2034. According to Canales (2015) jack mackerel stock is close to $50 \%$ of the maximum sustainable yield, and thus close to a status of collapse or depletion according to the criteria used by the New Chilean Fisheries Law. In order to assess the current state of the jack mackerel stock and its exploitation strategies, Oceana requested (CAPES) to elaborate a report to examine key parameters that guarantee the sustainability of the stock and provide recommendation for precautionary quota decisions in the future. This document contains the status of jack mackerel with complete information until 2014 and the analysis of its exploitation strategies.

## 2. GENERAL OBJECTIVE

The aim of this project is to assess the status and exploitation strategies of jack mackerel considering the two main hypotheses about its population structure in the South Eastern Pacific.

## 3. SPECIFIC OBJECTIVES

3.1 Abundance estimation of jack mackerel considering two stock hypotheses in the SEP.
3.2 Estimates the biological references points of jack mackerel in the SEP following the method proposed for Chilean fisheries.
3.3 Define the status of jack mackerel stock (s) in the SEP and analysis its equilibrium points.

### 3.4 Evaluate exploitation strategies and risk analyses of jack mackerel in the SEP

## 2. METHODS

### 2.1. Data

The data used in this study correspond to the last jack mackerel stock assessment conducted in October of 2014 by the Scientific Committee (SC) of the SPRFMO, as well as from Canales (2014). Some modifications were introduced according to the criteria of the Working Team that carried out the present study. The stock assessment considered information updated until 2014.

## Fishery data

The catch data used in this assessment is summarized in Fig. 1 and Table 1 (Annex 1 and 2 respectively). The records include reported catches by the fishing fleets from coastal countries and distant water fleets in the SEP, which are grouped into four main fleets according to the criteria of gear and fishing areas as defined in JMSWG-Report (2014) (Northern Chile, South Chile, Far North and Offshore Trawl fleets). The catches by fleet used in the year 2014 were updated with the available information at July 2015 for the Chilean fisheries. For other fleets, the data are the same as in First Report of the Project. The total catch of jack mackerel in the year 2014 was 406.942 t in the SEP and within the range of the maximum of 460.000 t , proposed as TAC for the year 2014 by the Commission at SPMFRO.

CPUE time series (Table 2, Annex 2) used in this model include: i) Fleet 2: SouthCentral Chile CPUE from the purse seiner fleet (1983-2014), ii) Fleet 3: Far North CPUE (2002-2014), iii) Fleet 4: three different CPUE time series are used: China (2001-2013), European Union (EU) (2006-2011; 2013) and Russia (1987-1991; 2008-2009; 2011). This information is the same as in JMSWG-Report (2014).

Catch at age in Tables 3, 4 and 5 (Annex 2) are available for the fleets of north and south central Chile and for the offshore trawl fleet. Length structures are only available for far north fleet (Table 6, Annex 2). Mean weights at age by fleet are presented on Table 7 to 10 (Annex 2) and are available for all fleets. In the present report age structures for the fleets of north and south central Chile encompassed the whole year 2014.

## Survey data

Biomass indices from acoustic surveys are available for north Chile (1984-1988; 1991; 2006-2014), south central Chile (1997-2009), far north Peru (1986-2009; 2011-2013) (Table 11, Annex 2). Spawning biomass estimated (Table 11, Annex 2) by the Daily Egg Production Method (DEPM) is available for south central of Chile (1999-2001; 20032008) (Table 11, Annex 2). Age structures from acoustic surveys are only available for the Chilean data and they are summarized in Tables 12 to 14. Note that although an acoustic survey was carried in early 2015 off Northern Chile, the information was not yet available and therefore not included in this assessment.

## Biological parameters

In jack mackerel, age at first reproduction is estimated between 2-3 years (Table 15, Annex 2). Simulated age-at-length key used to fit length structures from Peruvian data, was computed using growth parameters in Cerna et al. (2011) with $L_{\infty}=72,7 \mathrm{~cm}, k=0,074$ (year ${ }^{-1}$ ) and $t_{0}=-1,97\left(\right.$ year $\left.^{-1}\right)$. Natural mortality is considered $\mathrm{M}=0.23\left(\mathrm{year}^{-1}\right)$ following JMSWG-Report (2014).

### 2.2. Stock assessment model

The model used to assess the jack mackerel in the SEP was requested to the Subsecretaría de Pesca de Chile (www.subpesca.cl), we also obtained information from SPRFMO reports (www.sprfmo.int) and from de Instituto de Fomento Pesquero reports (www.ifop.cl). The current model was developed by the SC at the SPRFMO, and was adopted as an assessment method for the Jack mackerel in the South Eastern Pacific in 2010 (JMSWG-Report, 2014).

The stock assessment model corresponded to an integrated statistical catch-at-age approach, implemented in AD Model Builder (ADMB) with different sources of information from the jack mackerel fisheries in the SEP area as well as fisheryindependent information (surveys) covering the period from 1970 to 2014 (Canales 2014, JMSWG-Report, 2014).

The model consists of four main components: 1) the dynamics of the fish population (Table 1, Annex 3); 2) the fishery dynamics (Table 1, Annex 3), 3) observations models (Table 2, Annex 3), and 4) parameter estimation (Table 3, Annex 3).

## Population and fishery dynamics

Population dynamic considers ages from 1 to $12+$ years old, the recruitment is assumed to take place on the first of January of each year, and the spawning occurs instantaneously in the middle of November. The initial population is based on equilibrium condition in 1958, 12 years before the information start in 1970 (landings are available from 1970). Cohort dynamic considers exponential decay of the abundance affected only by natural mortality up to 1970 and thereafter from natural mortality and fishing. Natural mortality is assumed to be time and age invariant (Table 1, Annex 3).

Fishing mortality is modeled using the separability hypotheses based on an annual component, and the selectivity that describes the proportion of individuals at age removed by fishing. Selectivity is non-parametric, fishery-specific and time-variant. The catchability is assumed variable (changes in the fishing effort) modeled by a random walk (Table 1, Annex 3). Stochastic density-dependence relationship between recruitments and spawning biomass is modeled using the Beverton and Holt stock recruitment function (Table 1, Annex 3).

## Observations model

Observations come from four datasets as follows. (1) The abundance indices (CPUE, acoustic survey and spawning biomass from DEPM), (2) total catches, (3) ages and (4) length structures from fishing operations, acoustic surveys and DEPM. The observation models for each dataset are described in Table 2 (Annex 3). The observation models of the indices are obtained according to the fraction of year that they take place. Catches by year, age and fleet (Table 2, Annex 3) are all weighted by ageing error matrix based on Chilean age data (Table 16, Annex 2). Table 2 (Annex 3) shows the form of transition matrix to convert length distribution into age distribution, based on the von Bertalanffy growth parameters $L_{\infty}$ and $k$.

## Parameters estimation

The parameters of the model were estimated maximizing the log-likelihood of each dataset together with $\log$ probability density functions of the priors and smoothing penalties of selectivity. The list of parameters is summarized in Table 3 (Annex 3).

The log-likelihood (Table 3, Annex 3) of indices (CPUE and surveys) and catches are assumed as lognormal distributed. Likewise, multinomial distribution is considered for the age and length frequencies. Constrain for the fishing mortality are assumed for the last year of the assessment due to effect of the presence of incomplete cohorts. Estimates of recruitment are conditioned to a stock-recruitment curve over the period 1977-2011.

The weight of each dataset (coefficients of variations and sample sizes) of each likelihood function was taken from JMSWG-Report, (2014) to maintain consistency. Thus, CV of the catches was $c v=0.05$ for the four fleets keeping the assumption of high precision of the catch data. Smoothness for selectivity (indexes) were all were kept as $\lambda=$ 100 , the $\mathrm{S}-\mathrm{R}$ function fit was kept with a $\mathrm{cv}=0.7$ and the recruitment regularity $\lambda=1.4$. Sample size for the proportions at age of the Chilean acoustic survey were $\mathrm{n}=30$, and DEPM $n=20$. Table 4 (Annex 3) summarizes the weights used for abundance indices, smoothness for selectivity of the fleets and samples size of the proportions at age/length of each fleet.

## Hypotheses - Stock Structure

We explored the trends and levels of the spawning biomasses for two hypotheses about the stock structure of jack mackerel in the SEP. Two hypotheses were evaluated as follows. H1: Jack mackerel is one single population in the South Eastern Pacific area, this hypothesis, hereafter Model-1, and H2: Jack mackerel constitute two discrete populations one in the Peruvian and Ecuadorian waters, and one off Chile extending onto the high seas, hereafter Model-2. We assessed both hypotheses by optimizing the stock assessment models, first keeping all the information (Model-1). Hypothesis H2 all the data and related underpinning processes in the model belonging to the Far North fleet (CPUE and survey from Peru) were removed from the assessment (Model-2). This means that only the portion of the jack mackerel population inhabiting off Chile and distributed in coastal waters and high sea is assessed. Results showed that when information from Peru were removed from the assessment, no significant differences in the trend and level of the spawning biomass were observed, and we concluded that the biomass from Peru did not make a significant addition to the spawning biomass of the jack mackerel stock in the SEP. Thus, the following analyses is based only in the H1 (Model-1), however we take into account two scenarios regarding the assumptions of the steepness of the stockrecruitment relationship of jack mackerel, this means $h=0.65$ (low resilience) y $h=0.8$ (high resilience).

## Biological references points (BRP)

To estimate the biological reference points of jack mackerel we reviewed the methodological approach proposed by Payá et al. (2014) for Chilean Fisheries, Canales (2014) and JMSWG-Report (2014).

Payá et al. (2014) classified the jack mackerel fishery as a Tier 1a, which means there is enough information to apply an age- or length-structured assessment model which will provide usable estimates of the current abundance of the stock. This tier also implies that species-specific MSY can be estimated and therefore the steepness parameter ( $h$ ) of Beverton and Holt stock-recruitment relationship can be estimated within the assessment model. Therefore, Fmsy (fishing mortality at maximum sustainable yield), Bmsy (spawning biomass at maximum sustainable yield), and Blim (spawning biomass at collapse or depletion limit) can be estimated with reliability. Canales (2015) discussed the status of jack mackerel based in two values of steepness implies high $(h=0.8)$ and low ( $h=0.65$ ) resilience to fishing exploitation. However, both assumptions lead to a similar status of jack mackerel.

Here, we used the stock assessment model to estimate BRPs and define the status of jack mackerel population. We analyzed two scenarios for steepness parameter, considered fixed $h=0.8$ and 0.65 following Canales 2015; JMSWG-Report, 2014. Although $h$ could have been estimated within the stock assessment models, as proposed in Payá et al (2014), we decided to fix it in order to maintain consistency with previous assessments. Note that, when the steepness parameter ( $h$ ) is known, BPRs as MSY, Fmsy and Bmsy can be directly derived from yield per recruit analysis. We computed these BRPs using a routine already implemented in the jack mackerel stock assessment model (JMSWGReport, 2014). Notice that Fmsy and Bmsy are target management values according to the Chilean Fishing Law, but they are also used as management values at the SPRMFO. We also estimated Blim (spawning biomass limit) a half of Bmsy (Blim=Bmsy/2) following Payá et al. (2014). This reference point indicates a non-desirable state for jack mackerel spawning biomass, because beyond this point the risk of collapse is high.

To define population status we used the Kobe o Phase Diagram plot. This is a scatter plot of the ratio between annual estimates of spawning biomass (1970-2014) and the target spawning biomass ( $x$-axis) and, the ratio of annual estimates of fishing mortality (19702014) and target fishing mortality ( $y$-axis). In the diagram, we distinguish different areas following recommendations from Payá et al. (2014) for Chilean fisheries resources. Thus, we identified different areas: i) under-exploitation, where the actual point of the spawning biomass of jack mackerel is higher than the target biomass $(B / B m s y>1)$, or
lower if the criteria is fishing mortality $(F / F m s y<1)$, ii) full-exploitation, a level in which the biological point has been reached or is close to Bmsy $(B / B m s y \approx 1)$, iii) overexploitation, a level where the current or actual biomass is below the target biomass $(B / B m s y<1)$, or higher if the fishing mortality is considered $(F / F m s y>1)$, and iv) depleted or collapsed, where biomass is below the biomass of the biological limit point (Blim).

## Harvest control rule (HCR)

During 2008 and 2012, the Science Group (SG) of SPFRMO made an important progress in estimating the status of jack mackerel stock. As a result of this work, the spawning biomass was diagnosed as depleted where its lowest level was reached in 2010. Thus SPRMFO proposed a rebuilding plan for the jack mackerel stock over the whole southeast Pacific Ocean (http://www.southpacificrfmo.org/2nd-commission-meeting/). The first management action was reduced the jack mackerel catches in 2011. Afterwards, a substantial decrease of the fishing mortality was estimated, however, the spawning biomass was still at $51 \%$ of Bmsy during 2013. The nearest objective of the management procedure was to ensure that the spawning biomass of jack mackerel will increase up to $80 \%$ of Bmsy. Notice that $80 \%$ Bmsy was set as a management objective for the SPRMFO which is different than the one established in the Chilean Fishing Law, which uses Bmsy.

Projections of population abundance of jack mackerel population were conducted following Canales (2015) and JMSWG-Report (2014). The starting point of the projection was the year 2015, and the population was projected forward until 2034. Recruitments were modeled using the Beverton \& Holt function with two assumptions about steepness $h=0.8$ and 0.65 and two value of long-term recruitment or maximum recruitment (Rmed) as 1970-2012 and 2000-2012. The fishing strategy considered was fishing at a constant rate, and using five levels of F . The fishing mortality estimated in 2014 was multiplied by the following factors (multipliers): $\mathrm{mf}=\{0,0.25,0.5,0.75,1,1.25\}$ creating five levels of fishing mortality for the projections of the spawning biomass. Thus, four scenarios for the projection of the jack mackerel stock and catches were proposed as follows:

- Scenario 1: Model-1, $h=0.8$, and mean recruitment (Rmed) computed from the period 1970-2012.
- Scenario 2: Model-1, $h=0.8$, and mean recruitment (Rmed) computed from the period 2000-2012.
- Scenario 3: Model-1, $h=0.65$, and mean recruitment (Rmed) computed from the
period 1970-2012.
- Scenario 4: Model-1, $h=0.65$, and mean recruitment (Rmed) computed from the period 2000-2012.

Note that each scenario simulated the jack mackerel abundance (stock) using the five level of fishing mortality described before.

## 3. ASSESSMENT RESULTS

### 3.1. Fitting model to data

A comparison of the fit in Model-1 in all indices used and for the two scenarios of steepness ( $\mathrm{h}=0.80$ and $\mathrm{h}=0.65$ ) is shown in Fig. 2 (Annex 1). Almost no differences were observed between the fits of both models. In general, the fits are better for CPUE series than the surveys biomass indices. Within the time series of CPUE, the model predictions were better for SC-Chile, China and EU. The fit of age structures for fleets and survey data for the two scenarios of the steepness of Model-1, as the same as the indices show little differences between the two scenarios of $h$ (Fig. 3 to Fig. 9, Annex 1).

### 3.2. Stock assessment

Abundance estimates and the associated uncertainty for each scenario of $h$ are summarized in Fig. 10, Fig. 11 (Annex 1) and Table 17 (Annex 2). The comparison between scenarios (Fig. 12, Annex 1) showed almost no differences between $h$ scenarios on the mean value of the spawning biomass time series. Between 1974 and 2002 the highest recruitment period is observed in both models. Since 2003, jack mackerel recruitments fluctuated with values below mean recruitment estimates over the period 1974 to 2002 in both models. The highest spawning biomass (SSB) in both models is estimated in 1988, as well as the second maximum in the year 2003. Since 2010, the spawning biomass had increased, although estimates are still on the range of its lowest values. This slight recovery is explained by higher recruitments observed between 2007 to 2009 and an important decrease of the fishing mortality experienced since 2011. This increase of recruitments is mainly caused by an increase in the last few years of the CPUE in Chile CS. Spawning biomass estimates in 2014 for the Model-1 with two scenarios of steepness reached almost 3.6 million tones (Model-1, $h=0.8$ ), and 3.4 million t Model-1, $h=0.65$. The fishing mortality of both scenarios followed the same trend. A
significant decrease of the fishing mortality takes place from 2009 to 2011 as a consequence of a decrease of total landings in almost a half.

### 3.3. Retrospective analysis

Retrospective analysis in stock assessment, refers to the examination of the consistency among successive estimates of the same parameter obtained as new data are gathered. This analysis constitutes a useful tool to examine systematic patterns or inconsistency in state variables derived from integrated stock assessment models (Mohn, 1999). There are two types of retrospective analysis: historical and within-model. Historical retrospective analysis compare results of each final assessment with those conducted in previous years, and it is usually implemented to evaluate the effects of changing stock assessment methodologies. In contrast, within-model retrospective analysis uses the same data and assessment model to trim the most recent year's data in successive models runs. Thus, this analysis reproduces what would have been obtained annually if the current method had been used for past assessments. The within-model retrospective analysis is the most useful for determining internal inconsistency in the data, because the only change in different runs is the number of years in the model (Clark et al. 2012). We conducted a within-model retrospective analysis to evaluate consistency in the stock assessment proposed for jack mackerel. We used spawning biomass time series as a control variable dropping five years of the most recent data.

In Fig. 13, evaluations with previous data are labeled as "R-1" to "R-5" indicating the number of years that has been dropped from the last assessment. We can see that for the retrospective analysis in $\mathrm{R}-1$ to $\mathrm{R}-4$, an overestimated spawning biomass respect to the mean value of the most recent assessment (actual) is found. In addition, assessment considering the least data (R-5) showed an inverse behavior respect to the rest of retrospective analyses and provided estimates of spawning biomass smaller of what was predicted using the most recent assessment. Higher differences are shown only in the last few years of each assessment (R1 to R5). Spawning biomass estimates from R-1 to R-5 fall onto the $95 \%$ confidence intervals computed for the last year assessment (Fig. 13).

In Fig. 14, retrospective analysis is showed in terms of percent differences between the spawning biomass in each year to the terminal year estimates. Here, we see that the main difference is detected when dropping two years from the last assessment (R-2), which in percent differences reached about $30 \%$ of the spawning biomass estimated in 2010 with the last assessment. On the other hand, for the same year, retrospective analysis in R-5 produces estimates of spawning biomass that are $20 \%$ lower of what is estimated when using the last year assessment.

## 4. EXPLOITATION STATUS

Biological references points computed for Model-1, $h=0.8$ and Model-1, $h=0.65$ are shown in Table 18 (Annex 2). Depletion of the unexploited biomass is slightly greater in the case when $h=0.65$. Spawning biomass in the year 2014 in Model $-1, h=0.8$ and Model$1, h=0.65$ are near to Blim rather than Bmsy in both cases; however F in 2014 is below Fmsy (Table 17, Annex 2).

The status of jack mackerel stock in the SEP under the scenarios of Model-1, $h=0.8$ and Model-1, $h=0.65$ are shown in the Fig. 15 and Fig. 16 (Annex 1). Differences between trajectories of spawning biomass under both escenarios of $h$ are small. Both diagrams show that during the 1970s, jack mackerel was underexploited, and during 1980s and 1990s the stock reached overexploitation with spawning biomass below maximum sustainable yield ( $\mathrm{B}<B m s y$ ) and $\mathrm{F}>F m s y$ ). For a few years, during the year 2000 the stock seems to be near Bmsy, although always above Fmsy. From 2008 to 2012 the stock status is in depletion. Steepness scenarios change the perceived status in 2013. When using $h=0.8$ the status is overexploited, whereas with $h=0.65$ the population is in depletion. However, in 2014 both scenarios of steepness lead to overexploited condition (Fig. 15 and Fig. 16).

Confidence intervals over the fishing mortality and spawning biomass estimates for the year 2014 in the scenarios $h=0.8$ suggest a 0.5 probability of being below $B_{\text {lim }}$, and thus, in a depleted condition (Fig. 15, Annex 1). In addition, confidence intervals of fishing mortality show there is a $10 \%$ chance of exceeding target fishing mortality level (F $>$ Fmsy) (Fig. 15, Annex 1). Kobe plot for $h=0.65$ (Fig. 16, Annex 1) shows that spawning biomass (B) has a probability higher than 0.75 of being depleted where $\mathrm{B}<\mathrm{B} m s y$ and low a probability of $\mathrm{F}>F m s y$.

## 5. PROJECTIONS AND EXPLOITATION STRATEGIES

The scenario 1 (Model $1, h=0.80$, Rmed $=1970-2012$ ) of spawning biomass projection (Fig. 17) showed that under all levels of constant fishing mortality the spawning biomass of jack mackerel would be higher than the spawning biomass in 2015 in the short (2016), medium (2024) and long term (2034). In addition, all these projections show that spawning biomass was higher than the Bmsy (Table 18) in short, medium and long term. Scenario 2 (Model 1, $h=0.80$, Rmed=2000-2012) (Fig. 20) shows different results than
the scenario 1 . At all levels of $\mathrm{F}>0$, the spawning biomass tends to decrease in the long term (2034). The biggest decrease in biomass is observed when projection are set with F values $25 \%$ higher than F 2014. In this case, biomass in 2015 is below the level defined by Bmsy and thus, in an overexploited condition.

None of the F scenarios analyzed recovers the spawning biomass over Bmsy. The scenarios of Rmed used to project the abundance explains the differences observed in these two scenarios. The first scenario used Rmed, which was based in the years 1970 to 2012 (see recruitment time series, Fig. 10) and the second from 2000-2012. In the first case, maximum recruitment for the population growth is higher than in the second case. Thus, the first case has higher compensation as fishing mortality increases across different F scenarios projected.

In Fig. 18 and 21 show the probability distribution to fall below $80 \%$ Bmsy (yellow area) and Bmsy (blue line) for scenarios 1 and 2, respectively. For scenario 2 (Fig. 21) in the short term (2016), the probability to be below the $80 \%$ Bmsy or Bmsy is higher than in scenario 1 (Fig. 18). In the medium term, scenario 1 has a lower chance to be below $80 \%$ Bmsy or Bmsy than scenario 2, and the same results are shown for long-term for all levels of F. On the other hand, Fig. 19 and 22 show the probability of recovery of the spawning biomass of jack mackerel over Bmsy. In the short term (2016), the probability to recover the spawning biomass is lower in the scenario 2 (Fig. 22) than scenario 1 (Fig. 19). In the scenario 2 , spawning biomass has a higher probability to recover the spawning biomass over Bmsy only when $\mathrm{F}=0$. In the medium (2024) and long-term (2034) scenario 2 is worse, with the highest chances to recover the spawning biomass over Bmsy only if the F is half of F in 2014 ( $35 \%$ and $27 \%$ respectively) and above $>90 \%$ if $\mathrm{F}=0$.

Scenarios 3 and 4 (Fig. 23 and 26) correspond to the projections of spawning biomass of jack mackerel with value of steepness of $h=0.65$. Scenario 3 (Model $1, h=0.65$, Rmed=1970-2012) showed that at almost all levels of constant fishing mortality produced in the short, medium and long term spawning biomasses higher than the value in 2015 (Fig. 23). In all these projections (except at $125 \% \mathrm{~F}$ ) spawning biomass is higher than Bmsy (Table 18) in the medium and long term. In cases where fishing mortality is equal to $25 \%$ or above the F estimates in 2014 ( $>25 \% \mathrm{~F}$ ), the spawning biomass does not recover over the Bmsy in the short, medium and long term. Scenario 4 (Model 1, $h=0.65$, Rmed=2000-2012) (Fig. 26) shows different results than scenario 3. At all levels of F, except when $\mathrm{F}=0$, spawning biomass decreases below Bmsy (Table 18) in the medium and long term (2034). The highest decline takes place with the highest values of $F, 125 \%$ of the estimates F in $2014(125 \% \mathrm{~F})$. Different Rmed used in the simulation explains the differences observed between scenarios 3-4, and 1-2. The first scenario used an Rmed based in the years 1970 to 2012 (see recruitment time series, Fig. 11), and the second
from 2000-2012. Thus, scenario 3 sets a higher Rmed than the scenario 4. In addition, scenarios 3-4 show a slightly deeper decline than scenarios 1-2, due to the lower resilience set by the steepness value of $h=0.65$.

Probability distribution to fall below $80 \%$ of Bmsy (yellow area) and Bmsy (blue line) for the scenario 3 and 4, respectively are shown in Fig. 24 and 27. Scenario 4, short term (2016), shows that the probability to be below to $80 \%$ Bmsy or Bmsy is higher than in scenario 3. In the medium term, scenario 3 has a lower chance for the spawning biomass to fall below $80 \%$ Bmsy or Bmsy than scenario 4, in the medium and long term. Fig. 25 and 28 show the probability to recover the spawning biomass of jack mackerel over Bmsy, in scenarios 3 and 4 respectively. In the short term (2016) the probability is lower for scenario 4 than scenario 3. Scenario 3 has the highest probability (0.12) to recover spawning biomass over Bmsy but only if $\mathrm{F}=0$. In the medium term (2024) and long-term (2034) scenario 4 is worse than scenario 3. Scenario 3 in general has better chances to recover the spawning biomass over Bmsy than scenario 4. In scenario 4, the highest probability to recover the spawning biomass over Bmsy, in the medium and long term takes place only when $\mathrm{F}=0$. Likewise $50 \% \mathrm{~F}$ has a probability to recover the spawning biomass above Bmsy of 0.07 in the short term (2016), 0.28 in a medium term (2024) and 0.22 in long term (2034).

Probabilities of the jack mackerel spawning biomass to recover over Bmsy or to fall below $80 \%$ Bmsy for each scenario and F level are summarized in Table 19. The table also contains the predicted catches of jack mackerel in the year 2016 and 2017. It is clear from the results in Table 19 that the recovery of the spawning biomass of jack mackerel over Bmsy in the short term (2016), has the highest probability ( $>0.2$ ) only if $h=0.8$ and Rmed $=1970-2012$ and $F=0$. For all levels of $F$, the probability to recover biomass above Bmsy in the four scenarios is $<0.2$. In the medium term projection, the highest probability ( $>0.5$ ) to recover the spawning biomass above Bmsy occurs when Rmed is computed from the period 1970-2012, however the probability decreases $(<0.35)$ if Rmed is computed from the period 2000-2012 and $F \neq 0$. In the medium term projection, $50 \% \mathrm{~F}$ increases the chances to recover the biomass above Bmsy between $28 \%$ and $35 \%$ depending on $h$ value used on the projection. Long-term results are similar to medium term although, chances to recover the spawning biomass over Bmsy drop slightly more ( $<$ $27 \%$ ) if fishing mortality is $50 \%$ F. Using Rmed for the period 2000-2012 and $h=0.8$ (scenario 2), TAC of jack mackerel 2016 takes a range between 357 to 768 thousands $t$ depending on the level of F and risk of not accomplish the management rule proposed. When using $h=0.65$ this TAC range changes slightly between 353 to 759 thousand tones also depending on F and risk (Table 19).

Although, there are no strong differences between the range of TAC 2016 depending on the scenario of $h$, the higher impact of this parameter is reflected in the time and probability that the stock of jack mackerel will leave the overexploited area. As we can see in Table 19, the probability to recover the jack mackerel stock to the sustainable area (over the Bmsy) in the long term (2034) taking a $h=0.8$ varied between 0.002 to 0.266 when $\mathrm{F}>0$. With a value of steepness of $h=0.65$, these probabilities became smaller, for instance increasing F over the value of F in 2014 will not recover the stock by 2034 (Table 19), and if $F$ is kept equal to the value of $F$ in 2014 the probability is 0.006 ( $h=0.65$ ) and $0.016(h=0.80)$ under the same scenario of Rmed.

## 6. DISCUSSION

Main results of this work indicate that jack mackerel stock in the South Eastern Pacific is overexploited and with some probability of being depleted (close to $B_{\text {lim }}$ ). We conducted a stock assessment considering two hypotheses regarding the stock structure of the jack mackerel in the South Pacific (Canales et al. 2015), and two scenarios of steepness (h). All these combinations between stock hypotheses and scenarios of $h$ lead to the same diagnosis of stock status. Main differences between hypotheses and scenarios of $h$ are related with the probability of depletion, which varies between 0.25 (one single stock) and 0.35 (two stock). In addition, based on the hypotheses of one single stock (Model-1), probability of depletion varies between $0.5(h=0.80)$ and 0.75 when $h=0.65$. A similar conclusion is found in Canales (2015) who reported that jack mackerel is overexploited with spawning biomass near Blim, and thus, with probability of depletion.

The stock assessment presented in this report was conducted using updated information of the catches of Chilean fleets in 2014, based in the landings available up to July 2015. The complete catch at age structures of the North and Central south of Chile fleet observed in 2014 was also updated. The new information included in the assessment did not produce significant variation in the trend, level and status of jack mackerel. This robustness of the state variables to the incorporation of new information if expected in lengthy-data and age-structured stock assessment models like the one used here, because historical information has a high relative weight in the analysis. In addition, note that here, we use a different set of growth parameters than those reported in previous assessments in JMSWG-Report (2014) ( $k=0.16$ year $^{-1}, L_{\infty}=74.4 \mathrm{~cm}$ ). These parameters showed inconsistencies, because $k$ estimates seem to be close to what is reported to the far north stock, while $L_{\infty}$, is similar of what is reported in Chilean waters. $k$ and $L_{\infty}$, are highly correlated parameters and thus, consistency between their estimates is expected.

Growth parameters in this assessment are used to model a simulated age at length key to account for length structures observed on the far north fleet. Misspecification of growth parameters in age-structured stock assessment models that fit length structures usually result in biased estimations of fleet-specific selectivity-at-age, which are then propagated in a cascade effect to, biased abundance estimates, biological reference points and population status. Given the importance of growth parameters in this assessment, we used the latest revision of growth parameters available in this species from Cerna et al (2014). These parameters are as follows $k=0.074$ year $^{-1}$ and $L_{\infty}=72.7 \mathrm{~cm}$. Changes in growth parameters usually trigger a revision of natural mortality. However, in order to maintain consistency with previous assessments we kept $M=0.23$ year $^{-1}$. Nevertheless, using
growth parameters in Cerna et al (2014), and using the M estimator in Pauly (1980) and temperatures from $15-20^{\circ} \mathrm{C}$ show a variation of $M$ from 0.19 to 0.22 , a value close to what is currently used in the assessment.

Comparing stock assessment outputs presented here with previous assessments in JMSWG-Report (2014) (comparison not shown here) we note that observed differences are mainly related with modifications of Chilean catches and age-structures and, secondarily with changes in growth parameters described above. In addition, results presented here for both hypotheses about stock structure were not significant (for details, see Canales et al. 2015). Thus, we conclude that magnitude of the Peruvian catches is small leading to a relative small estimate of abundance off shore Peru and thus making almost negligible to the contribution of this abundance to the mega scale stock of jack mackerel in the SEP.

We conducted a retrospective analysis based on the spawning biomass of jack mackerel to assess if there is a systematic over- or under-estimation of the last year of the assessment. Spawning biomass was overestimated in four of the retrospective analyses (R-1 to R-4), representing the most recent years. In the last analysis R-5 this pattern is inverted and biomass observed is underestimated. We conclude that abundance estimates in jack mackerel are highly influenced by the CPUE index from Center-South (CS) off Chile. The retrospective pattern found in jack mackerel appears to be random and thus, only caused by the addition of new data, because changes in historical spawning biomass estimates from R-1 to R-5 are small and falling into the confidence intervals computed in the last assessment. The amount of percent differences reported here for jack mackerel is small in comparison with other species (see Legault 2009) and this kind of patterns in retrospective analysis are probably of little concern (Clark et al. 2012). Retrospective analyses should be considered for further investigations where "one-way" pattern is observed. This means that as data is added, the recent estimates of the spawning biomass changed for each of the years considered in the analysis. Nevertheless, this is not the case for the assessment presented here for jack mackerel and thus, we conclude that the present assessment provides a robust estimation of abundance.

Scenarios of projections included the combination of two levels of $h$ and two levels of Rmed. Here, we conclude that Rmed has an important impact in the projections and predicted recovery of the spawning biomass over Bmsy or $80 \%$ Bmsy. Instead, the level of h chosen has less relevance on the projections, which may seem a bit confusing giving the importance of in MSY-based management approaches (Mangel et al 2013). Here, projections of abundance are based on the Beverton-Holt function parameterized in terms of Rmed and $h$. Rmed is used to compute the unexploited recruitment $\left(R_{0}\right)$, which defines the asymptotic level of the Beverton-Holt function. Thus, the period over Rmed
computed has a direct effect on defining the upper limit of "ceiling" that simulated recruitments can reach on abundance projections. On the other hand, $h$ defines the slope of the Beverton-Holt function and thus, this gives an idea on how fast the asymptotic recruitment is reached when simulating projections. Here, Rmed is computed over two periods, 1970-2012 and 2000-2012. It is important to notice that the period 2000-2012 was selected based on JMSWG-Report (2014). When using the longest period of recruitment (1970-2012), Rmed gives higher values of what is observed in the last two decades period (2000-2012). Higher value of Rmed using the longest period is highly influenced by extreme high values of recruitments estimated in 1984 and 1985. Last period of recruitments are lower and thus more precautionary and will give a better idea of the current level of recruitments, which may be more likely to occur during the horizon of projections. On the other hand, the impact of the two different scenarios of steepness ( $h=0.8$ and $h=0.65$ ) is negligible in comparison with changes in Rmed, although it affects the probability of reaching Bmsy in a certain period of time. For the same Rmed, the uses of higher values of resilience ( $\mathrm{h}=0.8$ ) increase the probability to reach Bmsy, in comparison with low resilience scenarios ( $h=0.65$ ). Thus, $h$ has a secondary importance on the projections and it is mainly associated with the probability of accomplished management decision based on MSY. In the current stock assessment, steepness value is treated as fixed parameter and sensitivity analysis is conducted using two levels of $h$. Levels of $h$ chosen here seem to be according of what we would expected in species with life histories such as jack mackerel (Payá et al 2014). Projections using multipliers of the fishing mortality estimated for 2014 have high probability of accomplish the management rule proposed only in case of high Rmed is considered in the longest period. However, the Rmed for the period 1970-2012 does not represent the recruitment level for the most recent years and is likely to overestimate recruitment in the short and medium term projections. Therefore, the use of Rmed for the longest period may be misleading and overestimate the rebuilding capacity of the jack mackerel stock. Thus using Rmed from the period 2000-2012 is more appropriate to represent the current productive capacity of the stock. The choice of this scenario will imply that harvesting at $50 \%$ of fishing mortality estimates in 2014 has a chance of $30 \%$ to accomplish Bmsy at the end of the projection period (year 2034). Harvesting with any higher F ( $>50 \%$ F), will reduce the chances below an $8 \%$ to reach Bmsy in 2034 and thus it is not recommendable as a sustainable management strategy. At the current exploitation status presented here for jack mackerel, according to the precautionary approach FAO guidelines state, it is fundamental to take this recommendation into account in order to ensure the sustainability of the stock. This means when a course of action has to be chosen (e.g catch limits), negative impacts need to be avoided or minimized and management decisions should ensure a low risk in order drive the stock to desirable levels of abundance. The acceptance of the precautionary approach usually involve the adoption of harvest rules that only allows $10 \%$ of risk of not accomplish the management objective.

Thus, even reducing fishing mortality to be applied at only half of what was estimated for 2015, the chances of reaching Bmsy by 2034 are only $30 \%$, a relative large risk in the light of a precautionary approach. In order to reach $10 \%$ of risk, fishing mortality and related catches need to be reduced even more of what is presented in Table 19, for those scenarios considering Rmed (2000-2012) and a multiplier of $50 \%$ of the fishing mortality in 2014.

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## ANNEX 1: FIGURES



Fig 1. Catches ( $\mathrm{x} 10^{3}$ metric ton) of jack mackerel at the Southern-eastern Pacific 19702014. (Fleet 1: Northern Chile, Fleet 2: Central-south Chile, Fleet 3: Far North - Peru, Fleet 4: Offshore Trawl).


Fig. 2. Comparison of the Model-1 of two scenarios of steepness, $h=0.8$ and $h=0.65$. Black dots: observed values, red solid line: predicted values Model-1, and $h=0.8$; blue dotted line: predicted values Model-1, $h=0.65$ (CS: central south, EU: European Union, DEMP: Daily Egg Production Method).


Fig. 3. Comparison fit catch at age of the North Chilean fleet (Fleet 1). Two scenarios of steepness, $h=0.8$ and $h=0.65$ (Model-1). Grey bars: observed values, red solid line: predicted values Model-1, and $h=0.8$; blue dotted line: predicted values Model-1, $h=0.65$.


Fig. 4. Comparison fit catch at age of the central-south Chilean fleet (Fleet 2). Two scenarios of steepness, $h=0.8$ and $h=0.65$ (Model-1). Grey bars: observed values, red solid line: predicted values Model-1, and $h=0.8$; blue dotted line: predicted values Model-1, $h=0.65$.


Fig. 5. Comparison fit catch at age of the Far North (Fleet 3). Two scenarios of steepness, $h=0.8$ and $h=0.65$ (Model-1). Grey bars: observed values, red solid line: predicted values Model-1, and $h=0.8$; blue dotted line: predicted values Model-1, $h=0.65$.


Fig. 6. Comparison fit catch at age of the Offshore Trawl (Fleet 4). Two scenarios of steepness, $h=0.8$ and $h=0.65$ (Model-1). Grey bars: observed values, red solid line: predicted values Model-1, and $h=0.8$; blue dotted line: predicted values Model-1, $h=0.65$.


Fig. 7. Comparison fit catch at age of the Acoustic survey - central South Chile. Two scenarios of steepness, $h=0.8$ and $h=0.65$ (Model-1). Grey bars: observed values, red solid line: predicted values Model-1, and $h=0.8$; blue dotted line: predicted values Model-1, $h=0.65$.


Fig. 8. Comparison fit catch at age of the Acoustic survey - North - Chile. Two scenarios of steepness, $h=0.8$ and $h=0.65$ (Model-1). Grey bars: observed values, red solid line: predicted values Model-1, and $h=0.8$; blue dotted line: predicted values Model-1, $h=0.65$.


Fig. 9. Comparison fit catch at age of the DEPM - Central south Chile. Two scenarios of steepness, $h=0.8$ and $h=0.65$ (Model-1). Grey bars: observed values, red solid line: predicted values Model-1, and $h=0.8$; blue dotted line: predicted values Model-1, $h=0.65$.


Fig. 10. Summary estimates of the population of jack mackerel 1970-2014. Model-1, $h=0.8$. Spawning biomass (SSB): 1000 t , Fishing mortality: year ${ }^{-1}$, Recruitments: $10^{9}$ number and Landings: 1000 t .


Fig. 11. Summary estimates of the population of jack mackerel 1970-2014. Model-1, $h=0.65$. Spawning biomass (SSB): 1000 t , Fishing mortality: year ${ }^{-1}$, Recruitments: $10^{9}$ number and Landings: 1000 t .


Fig. 12. Comparison of population estimates of Jack mackerel from Model 1, $h=0.8$ (dark green line) and Model-1, $h=0.65$ (black line) (1970 to 2014). Total Biomass: 1000 t , Spawning biomass (SSB): 1000 t , Total Abundance: $10^{9}$ number, Recruitments: $10^{9}$ number, Fishing mortality: year ${ }^{-1}$, and Landings: 1000 t .


Fig. 13. Retrospective analysis in Jack mackerel stock assessment. Absolute changes in spawning biomass. Vertical lines represent the $95 \%$ confidence internals for the last year assessment.


Fig. 14. Retrospective analysis in Jack mackerel stock assessment. Relative differences of spawning biomass in each year to the terminal year estimates.


Fig. 15. Kobe diagram of the Jack mackerel Status (Model 1, $h=0.8$ ). Lines show the confidence region (90\%) for spawning biomass and fishing mortality on year 2014. $\mathrm{SB}=$ spawning biomass (Dark grey area: depleted condition, light grey area: overexploited condition, light green area: under-exploited condition of the stock).


Fig. 16. Kobe diagram of Jack mackerel Status (Model 1, $h=0.65$ ). Lines show the confidence region (90\%) for spawning biomass and fishing mortality on year 2014. $\mathrm{SB}=$ spawning biomass. (Dark grey area: depleted condition, light grey area: overexploited condition, light green area: under-exploited condition of the stock).


Fig. 17. Scenario 1: Projection of spawning biomass of jack mackerel 2015-2034 (Model $1, h=0.80$, Rmed $=1970-2012$ ). F constant $=$ fishing mortality $(\mathrm{F})$ is equal to the value in 2014; $125 \% * \mathrm{~F}=$ fishing mortality is increased in a $25 \% ; 50 \% * \mathrm{~F}=$ fishing mortality is reduced in a $50 \% ; 75 \% * \mathrm{~F}=$ fishing mortality $(\mathrm{F})$ is reduced in a $75 \%$, and $\mathrm{M}=$ natural mortality operates and $\mathrm{F}=0$.


Fig. 18. Scenario 1: Probability distribution of the spawning biomass (Model 1, $h=0.80$, Rmed=1970-2012) in the years 2016, 2024 and 2034 (Vertical). The yellow color indicates the probability of the spawning biomass to fall below the $80 \% \mathrm{Bmsy}$ and the blue line below the Bmsy . Horizontal: $0 \% * \mathrm{~F}=\mathrm{F}=0 ; 50 \% * \mathrm{~F}=$ fishing mortality ( F ) value in 2014 is reduced in a $50 \% ; 75 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $75 \% ; 100 \% * \mathrm{~F}=\mathrm{F}$ was kept equal to the value estimated in 2014. $125 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is increased in a $25 \%$. Xaxis corresponds to the spawning biomass $\left(\mathrm{x} 10^{3} \mathrm{t}\right)$. Notice that each title in each graph contains the risk (\%) to be below $80 \% \mathrm{Bmsy}$ for each year and level of F .


Fig. 19. Scenario 1: Probability distribution of the spawning biomass (Model 1, $h=0.80$, Rmed=1970-2012) in the years 2016, 2024 and 2034 (Vertical). The red color indicates the probability of the spawning biomass to recover over $\mathrm{B} m s y$. Horizontal: $0 \% * \mathrm{~F}=\mathrm{F}=0$; $50 \% * \mathrm{~F}=$ fishing mortality $(\mathrm{F})$ value in 2014 is reduced in a $50 \% ; 75 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $75 \% ; 100 \% * \mathrm{~F}=\mathrm{F}$ was kept equal to the value estimated in $2014.125 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is increased in a $25 \%$. X-axis corresponds to the spawning biomass ( $\times 10^{3} \mathrm{t}$ ). Notice that each title in each graph contains the probability (\%) to be located over Bmsy for each year and level of F.


Fig. 20. Scenario 2: Projection of spawning biomass of jack mackerel 2015-2034 (Model $1, h=0.80$, Rmed $=2000-2012$ ). F constant $=$ fishing mortality $(\mathrm{F})$ is equal to the value in $2014 ; 125 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is increased in a $25 \% ; 50 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $50 \%$; $75 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $75 \%$, and $\mathrm{M}=$ natural mortality operates and $\mathrm{F}=0$.


Fig. 21. Scenario 2: Probability distribution of the spawning biomass (Model 1, $h=0.80$, Rmed=2000-2012) in the years 2016, 2024 and 2034 (Vertical). The yellow color indicates the probability of the spawning biomass to fall below the $80 \% \mathrm{Bmsy}$ and the blue line to fall below the $\mathrm{B} m s y$. Horizontal: $0 \% * \mathrm{~F}=\mathrm{F}=0 ; 50 \% * \mathrm{~F}=$ fishing mortality ( F ) value in 2014 is reduced in a $50 \% ; 75 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $75 \% ; 100 \% * \mathrm{~F}=\mathrm{F}$ was kept equal to the value estimated in 2014. $125 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is increased in a $25 \%$. X -axis corresponds to the spawning biomass ( $\mathrm{x} 10^{3} \mathrm{t}$ ). Notice that each title in each graph contains the risk (\%) to be below $80 \%$ Bmsy in each year and level of F .


Fig. 22. Scenario 2: Probability distribution of the spawning biomass (Model 1, $h=0.80$, Rmed=2000-2012) in the years 2016, 2024 and 2034 (Vertical). The red color indicates the probability of the spawning biomass to recover over Bmsy. Horizontal: $0 \% * \mathrm{~F}=\mathrm{F}=0$; $50 \% * \mathrm{~F}=$ fishing mortality $(\mathrm{F})$ value in 2014 is reduced in a $50 \% ; 75 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $75 \% ; 100 \% * \mathrm{~F}=\mathrm{F}$ was kept equal to the value estimated in $2014.125 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is increased in a $25 \%$. X-axis corresponds to the spawning biomass ( $\times 10^{3} \mathrm{t}$ ). Notice that each title in each graph contains the risk (\%) to be located over Bmsy in each year and level of $F$.


Fig. 23. Scenario 3: Projection of spawning biomass of jack mackerel 2015-2034 (Model $1, h=0.65$, Rmed $=1970-2012$ ). F constant $=$ fishing mortality $(\mathrm{F})$ is equal to the value in $2014 ; 125 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is amplified in a $25 \% ; 50 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $50 \%$; $75 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $75 \%$, and $\mathrm{M}=$ natural mortality operates and $\mathrm{F}=0$.


Fig. 24. Scenario 3: Probability distribution of the spawning biomass (Model $1, h=0.65$, Rmed=1970-2012) in the years 2016, 2024 and 2034 (Vertical). The yellow color indicates the probability of the spawning biomass to be located below the $80 \% \mathrm{Bmsy}$ and the blue line to be located below the Bmsy. Horizontal: $0 \% * \mathrm{~F}=\mathrm{F}=0 ; 50 \% * \mathrm{~F}=$ fishing mortality ( F ) value in 2014 is reduced in a $50 \% ; 75 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $75 \%$; $100 \% * \mathrm{~F}=\mathrm{F}$ was kept equal to the value estimated in 2014. $125 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is increased in a $25 \%$. X-axis corresponds to the spawning biomass (x10 t ). Notice that each title in each graph contains the risk (\%) to be below $80 \% \mathrm{Bmsy}$ in each year and level of F.


Fig. 25. Scenario 3: Probability distribution of the spawning biomass (Model 1, $h=0.65$, Rmed=1970-2012) in the years 2016, 2024 and 2034 (Vertical). The red color indicates the probability of the spawning biomass to be above of Bmsy. Horizontal: $0 \% * \mathrm{~F}=\mathrm{F}=0$; $50 \% * \mathrm{~F}=$ fishing mortality $(\mathrm{F})$ value in 2014 is reduced in a $50 \% ; 75 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $75 \% ; 100 \% * \mathrm{~F}=\mathrm{F}$ was kept equal to the value estimated in $2014.125 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is increased in a $25 \%$. X-axis corresponds to the spawning biomass ( $\times 10^{3} \mathrm{t}$ ). Notice that each title in each graph contains the risk (\%) to be located over Bmsy in each year and level of $F$.


Fig. 26. Scenario 4: Projection of spawning biomass of jack mackerel 2015-2034 (Model $1, h=0.65$, Rmed $=2000-2012$ ). F constant $=$ fishing mortality $(\mathrm{F})$ is equal to the value in $2014 ; 125 \% * \mathrm{~F}=\mathrm{F}$ in 2014 was amplified in a $25 \% ; 50 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in $50 \%$; $75 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in $75 \%$, and $\mathrm{M}=$ natural mortality operates and $\mathrm{F}=0$.


Fig. 27. Scenario 4: Probability distribution of the spawning biomass (Model 1, $h=0.65$, Rmed=2000-2012) in the years 2016, 2024 and 2034 (Vertical). The yellow color indicates the probability of the spawning biomass to be located below the $80 \% \mathrm{Bmsy}$ and the blue line to be located below the Bmsy. Horizontal: $0 \% * \mathrm{~F}=\mathrm{F}=0 ; 50 \% * \mathrm{~F}=$ fishing mortality ( F ) value in 2014 is reduced in a $50 \% ; 75 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $75 \%$; $100 \% * \mathrm{~F}=\mathrm{F}$ was kept equal to the value estimated in 2014. $125 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is increased in a $25 \%$. X-axis corresponds to the spawning biomass (x10 t ). Notice that each title in each graph contains the risk (\%) to be below $80 \% \mathrm{~B} m s y$ in each year and level of $F$.


Fig. 28. Scenario 4: Probability distribution of the spawning biomass (Model $1, h=0.65$, Rmed=2000-2012) in the years 2016, 2024 and 2034 (Vertical). The red color indicates the probability of the spawning biomass to be above of Bmsy . Horizontal: $0 \% * \mathrm{~F}=\mathrm{F}=0$; $50 \% * \mathrm{~F}=$ fishing mortality $(\mathrm{F})$ value in 2014 is reduced in a $50 \% ; 75 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $75 \% ; 100 \% * \mathrm{~F}=\mathrm{F}$ was kept equal to the value estimated in $2014.125 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is increased in a $25 \%$. X-axis corresponds to the spawning biomass ( $\times 10^{3} \mathrm{t}$ ). Notice that each title in each graph contains the risk (\%) to be located over Bmsy in each year and level of F.

## ANNEX 2: TABLES

Table 1. Input catch data ( 1000 t ) of Jack mackerel from the South-Eastern Pacific. (Fleet 1: Northern Chile, Fleet 2: Central-south Chile, Fleet 3: Far North - Peru, Fleet 4: Offshore Trawl).

| Year | Fleet 1 | Fleet 2 | Fleet 3 | Fleet 4 |
| :---: | :---: | :---: | :---: | :---: |
| 1970 | 101.685 | 10.309 | 4.711 | 0.000 |
| 1971 | 143.454 | 14.988 | 9.189 | 0.000 |
| 1972 | 64.457 | 22.546 | 18.782 | 5.500 |
| 1973 | 83.204 | 38.391 | 42.781 | 0.000 |
| 1974 | 164.762 | 28.750 | 129.211 | 0.000 |
| 1975 | 207.327 | 53.878 | 37.899 | 0.000 |
| 1976 | 257.698 | 84.571 | 54.154 | 0.035 |
| 1977 | 226.234 | 114.572 | 504.992 | 2.273 |
| 1978 | 398.414 | 188.267 | 386.793 | 51.290 |
| 1979 | 344.051 | 253.460 | 333.810 | 370.290 |
| 1980 | 288.809 | 273.453 | 414.299 | 339.802 |
| 1981 | 474.817 | 586.092 | 445.638 | 438.123 |
| 1982 | 789.912 | 704.771 | 143.724 | 733.204 |
| 1983 | 301.934 | 563.338 | 110.690 | 894.300 |
| 1984 | 727.000 | 699.301 | 200.674 | 1,059.927 |
| 1985 | 511.150 | 945.839 | 114.622 | 799.323 |
| 1986 | 55.210 | 1,129.107 | 51.029 | 837.502 |
| 1987 | 313.310 | 1,456.727 | 46.304 | 863.423 |
| 1988 | 325.462 | 1,812.793 | 244.229 | 863.215 |
| 1989 | 338.600 | 2,051.517 | 316.247 | 875.821 |
| 1990 | 323.089 | 2,148.786 | 370.823 | 872.059 |
| 1991 | 346.245 | 2,674.267 | 213.447 | 543.659 |
| 1992 | 304.243 | 2,907.817 | 111.682 | 37.932 |
| 1993 | 379.467 | 2,856.777 | 133.354 | 0.000 |
| 1994 | 222.254 | 3,819.193 | 233.346 | 0.000 |
| 1995 | 230.177 | 4,174.016 | 550.993 | 0.000 |
| 1996 | 278.439 | 3,604.887 | 495.518 | 0.000 |
| 1997 | 104.198 | 2,812.866 | 680.053 | 0.000 |
| 1998 | 30.273 | 1,582.639 | 412.846 | 0.000 |
| 1999 | 55.654 | 1,164.035 | 203.751 | 0.007 |
| 2000 | 118.734 | 1,115.565 | 303.700 | 2.318 |
| 2001 | 248.097 | 1,401.836 | 857.744 | 20.090 |
| 2002 | 108.727 | 1,410.266 | 154.823 | 76.261 |
| 2003 | 143.277 | 1,278.019 | 217.734 | 158.199 |
| 2004 | 158.656 | 1,292.943 | 187.369 | 295.443 |
| 2005 | 165.626 | 1,264.808 | 80.663 | 243.576 |
| 2006 | 155.256 | 1,224.685 | 277.568 | 362.627 |
| 2007 | 172.701 | 1,130.083 | 255.360 | 438.831 |
| 2008 | 167.258 | 728.850 | 169.537 | 406.986 |
| 2009 | 134.022 | 700.905 | 76.629 | 371.918 |
| 2010 | 169.012 | 295.796 | 22.172 | 239.593 |
| 2011 | 30.825 | 216.470 | 326.394 | 60.891 |
| 2012 | 13.256 | 214.204 | 187.396 | 39.918 |
| 2013 | 16.361 | 214.999 | 80.586 | 41.177 |
| 2014 | 18.230 | 254.280 | 74.530 | 62.720 |

Table 2. CPUE time series used in the stock assessment of jack mackerel. (Fleet 2: Central-south Chile, Fleet 3: Far North - Peru, Fleet 4: Offshore Trawl).

| Year | Fleet 2 - Chile | Fleet 3 - Peru | Fleet 4 - China | Fleet 4-EU | Fleet 4 - RUSSIA |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1983 | 0.797 |  |  |  |  |
| 1984 | 0.700 |  |  |  |  |
| 1985 | 0.568 |  |  |  |  |
| 1986 | 0.491 |  |  |  |  |
| 1987 | 0.590 |  |  |  | 55.020 |
| 1988 | 0.493 |  |  |  | 58.240 |
| 1989 | 0.506 |  |  |  | 51.060 |
| 1990 | 0.401 |  |  |  | 52.570 |
| 1991 | 0.497 |  |  |  | 60.990 |
| 1992 | 0.419 |  |  |  |  |
| 1993 | 0.368 |  |  |  |  |
| 1994 | 0.441 |  |  |  |  |
| 1995 | 0.392 |  |  |  |  |
| 1996 | 0.408 |  |  |  |  |
| 1997 | 0.362 |  |  |  |  |
| 1998 | 0.347 |  |  |  |  |
| 1999 | 0.401 |  |  |  |  |
| 2000 | 0.382 |  |  |  |  |
| 2001 | 0.473 |  | 1.400 |  |  |
| 2002 | 0.416 | 212.7 | 1.97 |  |  |
| 2003 | 0.365 | 244.1 | 1.740 |  |  |
| 2004 | 0.397 | 276.6 | 1.44 |  |  |
| 2005 | 0.363 | 193.2 | 1.44 |  |  |
| 2006 | 0.398 | 245.9 | 1.02 | 310 |  |
| 2007 | 0.302 | 231.0 | 1.13 | 308 |  |
| 2008 | 0.204 | 222.6 | 0.86 | 256 | 77.419 |
| 2009 | 0.167 | 184.1 | 0.81 | 209 | 59.563 |
| 2010 | 0.120 | 255.4 | 0.57 | 124 |  |
| 2011 | 0.069 | 264.9 | 0.33 | 57 | 45.213 |
| 2012 | 0.217 | 264.7 | 0.37 |  |  |
| 2013 | 0.162 | 139.3 | 0.58 | 81 |  |
| 2014 | 0.135 | 240.4 |  |  |  |

Table 3. Catch age-structure Fleet 1 (North Chile) 1975 - 2014 (numbers $10^{9}$ ).

| Year | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1975 | 0 | 4,073 | 13,776 | 44,143 | 60,659 | 166,146 | 171,002 | 81,053 | 29,641 | 8,253 | 4,552 | 1,070 |
| 1976 | 0 | 27 | 1,676 | 10,342 | 62,211 | 190,616 | 229,530 | 109,553 | 17,356 | 4,612 | 104 | 0 |
| 1977 | 122 | 12,767 | 20,036 | 48,155 | 149,760 | 239,459 | 184,136 | 68,069 | 10,018 | 155 | 0 | 0 |
| 1978 | 0 | 5,910 | 92,821 | 171,908 | 149,947 | 99,992 | 274,829 | 227,407 | 75,025 | 11,649 | 99 | 0 |
| 1979 | 0 | 0 | 40,320 | 104,406 | 202,313 | 247,236 | 261,704 | 212,251 | 72,124 | 7,503 | 87 | 0 |
| 1980 | 0 | 5,663 | 19,209 | 39,611 | 120,365 | 158,767 | 188,736 | 133,955 | 25,201 | 792 | 0 | 0 |
| 1981 | 0 | 0 | 17,592 | 107,026 | 226,852 | 272,757 | 332,572 | 167,387 | 32,014 | 4,135 | 31 | 0 |
| 1982 | 0 | 1,674 | 29,021 | 332,623 | 362,536 | 484,944 | 639,872 | 367,034 | 127,157 | 21,795 | 225 | 0 |
| 1983 | 0 | 102 | 1,966 | 49,812 | 151,839 | 222,327 | 205,704 | 103,468 | 21,178 | 556 | 27 | 0 |
| 1984 | 0 | 4,148 | 232,259 | 599,923 | 284,517 | 284,809 | 377,052 | 318,705 | 67,881 | 4,009 | 0 | 0 |
| 1985 | 0 | 886 | 53,397 | 255,301 | 400,091 | 427,340 | 252,577 | 73,876 | 12,239 | 1,256 | 0 | 0 |
| 1986 | 4,646 | 15,143 | 14,153 | 8,069 | 9,791 | 17,324 | 26,790 | 15,727 | 6,358 | 1,078 | 0 | 0 |
| 1987 | 940 | 69,685 | 612,169 | 638,980 | 149,771 | 36,285 | 27,361 | 8,978 | 428 | 431 | 16 | 0 |
| 1988 | 0 | 3,652 | 130,324 | 489,772 | 452,240 | 105,585 | 5,445 | 647 | 14 | 0 | 0 | 0 |
| 1989 | 0 | 7,901 | 4,867 | 43,820 | 326,596 | 271,953 | 55,735 | 9,161 | 2,786 | 295 | 47 | 0 |
| 1990 | 6,262 | 77,422 | 5,668 | 27,866 | 236,685 | 412,335 | 84,292 | 8,079 | 388 | 94 | 3 | 0 |
| 1991 | 21 | 17,154 | 218,253 | 217,953 | 121,225 | 180,648 | 259,236 | 64,615 | 5,402 | 5,551 | 1,113 | 542 |
| 1992 | 1,787 | 30,253 | 252,037 | 142,917 | 268,721 | 274,311 | 149,879 | 60,020 | 8,877 | 790 | 39 | 0 |
| 1993 | 2,413 | 66,476 | 1,485,903 | 597,010 | 115,335 | 99,352 | 19,235 | 7,088 | 2,161 | 730 | 123 | 77 |
| 1994 | 431 | 139,526 | 339,253 | 101,904 | 266,129 | 131,505 | 23,152 | 3,211 | 1,618 | 68 | 0 | 0 |
| 1995 | 2,315 | 170,501 | 345,192 | 296,965 | 146,207 | 84,328 | 17,213 | 801 | 82 | 0 | 0 | 0 |
| 1996 | 131,844 | 269,877 | 533,140 | 572,880 | 155,446 | 30,636 | 7,068 | 63 | 68 | 0 | 0 | 0 |
| 1997 | 191 | 26,244 | 307,207 | 204,870 | 23,575 | 694 | 72 | 0 | 0 | 0 | 0 | 0 |
| 1998 | 0 | 3,406 | 89,297 | 37,875 | 18,933 | 5,847 | 285 | 1 | 0 | 0 | 0 | 0 |
| 1999 | 38 | 24,289 | 319,634 | 75,711 | 15,598 | 5,394 | 392 | 7 | 0 | 0 | 0 | 0 |
| 2000 | 57,122 | 235,887 | 136,283 | 236,690 | 110,317 | 15,424 | 39 | 0 | 0 | 0 | 0 | 0 |
| 2001 | 1,568 | 256,795 | 1,326,138 | 491,732 | 25,070 | 1,848 | 0 | 0 | 0 | 0 | 0 | 0 |
| 2002 | 48,483 | 98,920 | 391,463 | 176,981 | 92,150 | 18,789 | 4,497 | 276 | 26 | 0 | 0 | 0 |
| 2003 | 7,504 | 158,188 | 604,518 | 242,543 | 53,916 | 21,616 | 9,440 | 1,894 | 295 | 0 | 0 | 0 |
| 2004 | 747 | 17,164 | 103,070 | 464,915 | 191,312 | 7,389 | 275 | 0 | 0 | 0 | 0 | 0 |
| 2005 | 43,804 | 324,087 | 476,065 | 193,396 | 151,443 | 43,843 | 5,150 | 0 | 0 | 0 | 0 | 0 |
| 2006 | 27,386 | 38,392 | 390,068 | 607,711 | 68,098 | 25,256 | 8,161 | 1,256 | 49 | 0 | 0 | 0 |
| 2007 | 681 | 93,552 | 346,671 | 475,258 | 113,550 | 27,460 | 17,712 | 7,169 | 1,191 | 165 | 165 | 66 |
| 2008 | 14,499 | 712,726 | 359,139 | 117,862 | 138,886 | 110,359 | 12,931 | 844 | 191 | 41 | 0 | 0 |
| 2009 | 569 | 58,921 | 250,894 | 432,581 | 34,649 | 70,639 | 3,089 | 78 | 0 | 0 | 0 | 0 |
| 2010 | 4 | 524,298 | 57,658 | 360,130 | 140,820 | 36,448 | 9,825 | 1,044 | 506 | 146 | 0 | 0 |
| 2011 | 20,901 | 199,846 | 94,185 | 11,650 | 3,387 | 5,921 | 383 | 0 | 0 | 0 | 0 | 0 |
| 2012 | 2,080,455 | 6,700,598 | 8,465,934 | 34,863,481 | 4,648,381 | 106,332 | 0 | 0 | 0 | 0 | 0 | 0 |
| 2013 | 5,232,401 | 86,320,606 | 33,345,959 | 861,630 | 357,301 | 415,201 | 245,386 | 34,652 | 14,421 | 0 | 0 | 0 |
| 2014 | 3,726,116 | 61,365,991 | 27,398,847 | 8,823,215 | 4,763,121 | 790,779 | 35,958 | 6,085 | 0 | 0 | 0 | 0 |

Table 4. Catch age-structure of the Fleet 2 (Central South Chile) 1975-2014 (numbers $10^{9}$ ).

| Year | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1975 | 0 | 0 | 355 | 633 | 2,259 | 7,108 | 11,305 | 10,032 | 5,551 | 1,867 | 999 | 190 |
| 1976 | 0 | 1,272 | 364 | 353 | 1,550 | 17,545 | 45,885 | 38,920 | 18,306 | 2,707 | 37 | 0 |
| 1977 | 1 | 223 | 462 | 5,493 | 19,717 | 34,341 | 61,594 | 47,567 | 7,128 | 684 | 70 | 4 |
| 1978 | 0 | 595 | 4,226 | 20,574 | 70,250 | 115,605 | 96,857 | 45,404 | 12,171 | 1,330 | 0 | 0 |
| 1979 | 0 | 20 | 13,439 | 147,552 | 206,176 | 210,012 | 208,192 | 180,490 | 120,831 | 48,163 | 4,206 | 904 |
| 1980 | 2,152 | 3,996 | 8,314 | 128,820 | 322,829 | 356,395 | 311,998 | 166,713 | 78,325 | 17,706 | 686 | 995 |
| 1981 | 0 | 2,419 | 8,973 | 83,655 | 392,031 | 696,981 | 627,159 | 326,988 | 99,193 | 23,220 | 3,581 | 800 |
| 1982 | 1,324 | 4,221 | 8,889 | 118,067 | 618,244 | 826,235 | 877,968 | 626,789 | 224,844 | 57,471 | 10,391 | 13,914 |
| 1983 | 1,624 | 98,937 | 191,076 | 314,984 | 749,070 | 1,084,641 | 1,113,032 | 548,218 | 207,964 | 30,697 | 3,760 | 1,259 |
| 1984 | 98 | 7,977 | 190,091 | 357,918 | 447,250 | 985,396 | 1,175,198 | 851,961 | 292,315 | 38,931 | 7,977 | 394 |
| 1985 | 53 | 889 | 39,763 | 372,792 | 621,888 | 1,131,123 | 1,405,139 | 725,890 | 182,303 | 21,926 | 2,839 | 2,111 |
| 1986 | 7,703 | 32,892 | 50,408 | 254,334 | 720,436 | 1,125,301 | 1,563,727 | 833,125 | 141,461 | 12,640 | 1,048 | 651 |
| 1987 | 8,538 | 240,224 | 509,673 | 459,902 | 311,739 | 907,439 | 1,929,691 | 1,290,613 | 257,500 | 39,754 | 3,890 | 922 |
| 1988 | 442 | 23,756 | 228,313 | 1,415,721 | 1,662,909 | 665,913 | 1,203,766 | 1,215,469 | 405,628 | 50,539 | 6,623 | 114 |
| 1989 | 0 | 5,570 | 34,610 | 283,575 | 1,634,407 | 2,293,278 | 1,376,907 | 1,070,507 | 406,813 | 64,106 | 1,490 | 0 |
| 1990 | 248 | 5,228 | 1,826 | 31,608 | 506,751 | 1,598,666 | 2,003,162 | 1,148,443 | 668,436 | 128,313 | 8,666 | 130 |
| 1991 | 54 | 30,081 | 134,146 | 122,717 | 56,100 | 419,889 | 1,682,863 | 1,831,750 | 982,207 | 504,943 | 158,725 | 46,297 |
| 1992 | 0 | 0 | 71,389 | 186,534 | 321,697 | 367,160 | 405,356 | 1,258,212 | 1,072,392 | 952,609 | 406,683 | 151,907 |
| 1993 | 0 | 11,391 | 231,606 | 759,545 | 940,331 | 854,977 | 790,767 | 758,606 | 893,719 | 721,295 | 259,075 | 41,588 |
| 1994 | 0 | 21,702 | 87,328 | 808,441 | 1,200,387 | 1,266,242 | 802,876 | 692,317 | 1,102,792 | 853,702 | 284,938 | 26,970 |
| 1995 | 760 | 9,375 | 365,691 | 1,727,987 | 1,350,915 | 2,318,999 | 1,687,597 | 807,698 | 562,790 | 385,464 | 170,813 | 31,602 |
| 1996 | 3,354 | 48,674 | 835,128 | 1,041,899 | 1,421,658 | 1,327,055 | 1,172,716 | 792,734 | 374,612 | 171,145 | 70,209 | 20,378 |
| 1997 | 7,893 | 191,472 | 1,428,562 | 2,627,586 | 1,898,630 | 906,334 | 488,229 | 377,039 | 302,621 | 132,121 | 75,881 | 42,083 |
| 1998 | 25,251 | 242,671 | 1,516,538 | 1,864,189 | 763,106 | 345,371 | 165,969 | 178,113 | 173,183 | 79,161 | 32,019 | 13,170 |
| 1999 | 2,407 | 190,107 | 1,825,090 | 1,676,166 | 718,361 | 267,024 | 76,733 | 34,947 | 59,069 | 54,516 | 34,827 | 28,903 |
| 2000 | 780 | 46,357 | 597,816 | 1,633,359 | 1,015,296 | 412,689 | 115,221 | 42,928 | 47,268 | 58,944 | 37,371 | 31,373 |
| 2001 | 3 | 32,966 | 361,663 | 970,229 | 1,270,037 | 594,890 | 184,069 | 82,580 | 61,986 | 57,150 | 47,079 | 76,019 |
| 2002 | 3,553 | 44,875 | 395,001 | 846,794 | 853,666 | 522,140 | 191,428 | 97,160 | 80,106 | 76,745 | 63,094 | 99,435 |
| 2003 | 5 | 16,651 | 232,038 | 908,724 | 1,101,223 | 740,572 | 303,108 | 100,321 | 77,824 | 61,641 | 37,835 | 38,325 |
| 2004 | 0 | 1,617 | 128,611 | 449,306 | 920,244 | 917,951 | 422,033 | 156,434 | 98,973 | 58,716 | 27,757 | 30,207 |
| 2005 | 14,398 | 15,576 | 14,953 | 145,195 | 460,828 | 1,047,649 | 518,471 | 208,577 | 141,084 | 66,743 | 28,283 | 32,878 |
| 2006 | 401 | 4,986 | 11,959 | 82,227 | 150,385 | 390,459 | 490,794 | 255,963 | 191,215 | 127,571 | 67,970 | 60,057 |
| 2007 | 0 | 260 | 26,043 | 250,100 | 293,211 | 206,428 | 282,545 | 280,026 | 242,885 | 165,923 | 92,414 | 86,474 |
| 2008 | 6,542 | 23,768 | 1,806 | 6,768 | 74,663 | 237,017 | 215,953 | 168,781 | 124,776 | 104,046 | 55,045 | 100,625 |
| 2009 | 0 | 1,581 | 43,860 | 108,580 | 22,418 | 222,409 | 250,865 | 193,736 | 120,657 | 77,999 | 55,350 | 47,561 |
| 2010 | 0 | 1,799 | 24,389 | 176,203 | 122,711 | 59,484 | 63,728 | 38,970 | 55,112 | 40,186 | 12,155 | 13,967 |
| 2011 | 0 | 29 | 3,626 | 55,109 | 45,036 | 121,204 | 33,101 | 20,726 | 45,281 | 16,068 | 3,724 | 4,334 |
| 2012 | 0 | 0 | 0 | 33,833 | 151,629 | 151,544 | 85,613 | 34,624 | 20,360 | 3,061 | 842 | 834 |
| 2013 | 2,000 | 40,778 | 12,467,295 | 101,991,694 | 162,856,221 | 181,274,927 | 78,563,279 | 14,567,793 | 2,215,252 | 980,360 | 171,018 | 459,068 |
| 2014 | 149,950 | 1,068,084 | 12,787,928 | 46,238,563 | 109,660,285 | 111,079,109 | 99,304,853 | 48,000,998 | 13,232,481 | 4,157,327 | 1,653,121 | 1,232,270 |

Table 5. Catch age-structure of the Fleet 4 (Offshore Trawl) (numbers $10^{9}$ ).

| Year | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1975 |  |  |  |  |  |  |  |  |  |  |  |  |
| 1976 |  |  |  |  |  |  |  |  |  |  |  |  |
| 1977 |  |  |  |  |  |  |  |  |  |  |  |  |
| 1978 |  |  |  |  |  |  |  |  |  |  |  |  |
| 1979 | 0 | 0 | 49 | 1905 | 20679 | 69124 | 1E+05 | $2 \mathrm{E}+05$ | 97437 | 40291 | 3780 | 904 |
| 1980 | 0 | 3842 | 4615 | 38803 | 1E+05 | 2E+05 | 2E+05 | 1E+05 | 64819 | 17129 | 686 | 995 |
| 1981 | 0 | 239 | 2919 | 18545 | 99141 | 2E+05 | 3E+05 | $2 \mathrm{E}+05$ | 75791 | 19940 | 3221 | 800 |
| 1982 | 0 | 2756 | 1078 | 9883 | 1E+05 | $3 \mathrm{E}+05$ | 5E+05 | 4E+05 | 2E+05 | 45982 | 10106 | 13829 |
| 1983 | 0 | 55448 | 99493 | 73750 | 3E+05 | 6E+05 | 8E+05 | 5E+05 | 2E+05 | 28049 | 3760 | 1259 |
| 1984 | 0 | 2428 | 53471 | 2E+05 | 2E+05 | 4E+05 | 6E+05 | 5E+05 | 2E+05 | 26780 | 7176 | 394 |
| 1985 | 0 | 539 | 20116 | 2E+05 | 3E+05 | 5E+05 | 6E+05 | 3E+05 | 96589 | 16896 | 2661 | 1840 |
| 1986 | 0 | 27483 | 33424 | 1E+05 | $4 \mathrm{E}+05$ | 5E+05 | 5E+05 | $3 \mathrm{E}+05$ | 62152 | 6636 | 748 | 651 |
| 1987 | 0 | 93104 | 1E+05 | 73668 | $2 \mathrm{E}+05$ | 5E+05 | 7E+05 | 4E+05 | 79536 | 18316 | 2974 | 878 |
| 1988 | 0 | 12902 | 89058 | $3 \mathrm{E}+05$ | $3 \mathrm{E}+05$ | 4E+05 | 5E+05 | 4E+05 | 1E+05 | 23035 | 5183 | 114 |
| 1989 | 0 | 265 | 12357 | 1E+05 | 5E+05 | 6E+05 | 4E+05 | 3E+05 | 2E+05 | 35858 | 1293 | 0 |
| 1990 | 0 | 536 | 316 | 10917 | 2E+05 | 6E+05 | 5E+05 | $3 \mathrm{E}+05$ | 2E+05 | 56477 | 5452 | 130 |
| 1991 | 0 | 3175 | 4982 | 9073 | 21237 | $2 \mathrm{E}+05$ | 4E+05 | $2 \mathrm{E}+05$ | 1E+05 | 57856 | 22617 | 9811 |
| 1992 ( 192617 |  |  |  |  |  |  |  |  |  |  |  |  |
| 1993 |  |  |  |  |  |  |  |  |  |  |  |  |
| 1994 |  |  |  |  |  |  |  |  |  |  |  |  |
| 1995 |  |  |  |  |  |  |  |  |  |  |  |  |
| 1996 |  |  |  |  |  |  |  |  |  |  |  |  |
| 1997 |  |  |  |  |  |  |  |  |  |  |  |  |
| 1998 |  |  |  |  |  |  |  |  |  |  |  |  |
| 1999 |  |  |  |  |  |  |  |  |  |  |  |  |
| 2000 | 6306 | 137881 | 9E+05 | 1E+06 | 9E+05 | 6E+05 | 4E+05 | $3 \mathrm{E}+05$ | 3E+05 | $2 \mathrm{E}+05$ | 1E+05 | 0 |
| 2001 | 0 | 934657 | 9E+06 | $2 \mathrm{E}+07$ | $2 \mathrm{E}+07$ | 9E+06 | 3E+06 | 1E+06 | 1E+06 | 6E+05 | $3 \mathrm{E}+05$ | 0 |
| 2002 | 4000081 | 5531716 | 7E+07 | 1E+08 | 7E+07 | 3E+07 | 9E+06 | 4E+06 | 3E+06 | 1E+06 | $4 \mathrm{E}+05$ | 0 |
| 2003 | 0 | 6341275 | $9 \mathrm{E}+07$ | $2 \mathrm{E}+08$ | 1E+08 | 3E+07 | $3 \mathrm{E}+06$ | $3 \mathrm{E}+05$ | 12549 | 0 | 0 | 0 |
| 2004 |  |  |  |  |  |  |  |  |  |  |  |  |
| 2005 |  |  |  |  |  |  |  |  |  |  |  |  |
| 2006 | 0 | 0 | 16612 | 4E+06 | 5E+07 | 1E+08 | 1E+08 | 4E+07 | $2 \mathrm{E}+07$ | 1E+07 | 5E+06 | 0 |
| 2007 | 0 | 0 | 78923 | 4E+06 | 3E+07 | 1E+08 | 1E+08 | 8E+07 | 7E+07 | $5 \mathrm{E}+07$ | 3E+07 | 0 |
| 2008 | 0 | 26553 | 1E+05 | 1E+05 | 4E+06 | 4E+07 | 1E+08 | 1E+08 | 6E+07 | 4E+07 | 2E+07 | 0 |
| 2009 | 0 | 1333996 | 2E+06 | $2 \mathrm{E}+06$ | 7E+05 | 6E+06 | 5E+07 | 1E+08 | 9E+07 | $5 \mathrm{E}+07$ | $3 \mathrm{E}+07$ | 0 |
| 2010 | 0 | 3782200 | 8E+07 | 4E+07 | 1E+06 | 1E+06 | 1E+07 | $3 \mathrm{E}+07$ | 5E+07 | 4E+07 | 1E+07 | 0 |
| 2011 | 0 | 858 | 2E+05 | $2 \mathrm{E}+06$ | 1E+06 | 3E+06 | 2E+06 | 2E+06 | 4E+06 | 2E+06 | 9E+05 | 0 |
| 2012 | 0 | 0 | 0 | 0 | 0 | 40 | 598 | 1601 | 18477 | 10000 | 3052 | 2932 |
| 2013 | 0 | 1 | 1044 | 18799 | 21113 | 25626 | 16848 | 8282 | 3278 | 4366 | 1374 | 1040 |
| 2014 | 0 | 0 | 0 | 1326 | 18534 | 38572 | 28875 | 8395 | 2064 | 1421 | 570 | 159 |

Table 6．Catch length－structure of the Fleet 3 （Far North Peru）1980－2014（numbers $10^{9}$ ）．

| Year | 10 | ${ }_{11}$ | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20 | 21 | 22 | ${ }^{23}$ | ${ }^{24}$ | 25 | 26 | 27 | 28 | 29 | 30 | 31 | 32 | 33 | ${ }^{34}$ | 35 | 36 | 37 | 38 | 39 | 40 | ${ }^{41}$ | 42 | 43 | 44 | 45 | 46 | 47 | 48 | 49 | 50 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1975 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1977 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1978 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1980 | 44 | 59 | 71 | 59 | 126 | 67 | 184 | 105 | 67 | 50 | 0 | 0 | 0 | 33 | ${ }^{46}$ | 25 | 9 | 9 | 50 | 101 | 134 | 209 | 318 | 471 | 419 | ${ }^{34}$ | 251 | 184 | 134 | ${ }^{84}$ | 50 | ${ }^{33}$ | ${ }^{33}$ | 29 | 17 | 9 | 9 | 0 | 0 | 0 | 0 |
| 1981 | 。 | 0 | 0 | 0 | 0 | ， | 0 | 0 | ， | 0 | 。 | 0 | 0 | 0 | 0 | 0 | 6 | 14 | 28 | 101 | 129 | 105 | 109 | 112 | 105 | 87 | 78 | 62 | 50 | 36 | 34 | 22 | 20 | 17 | 8 | 3 | 1 | 0 | 。 | － | \％ |
| 1982 | － | 23 | 91 | 72 | 545 | 545 | 523 | 477 | 409 | 432 | 545 | 364 | 68 | 23 | 0 | 0 | 0 | 0 | 45 | 91 | ${ }^{136}$ | 409 | 795 | 1136 | 840 | 545 | 296 | 227 | 181 | 159 | 136 | 113 | 23 | 0 |  | 0 |  | $\bigcirc$ | 0 | 0 |  |
| 1983 | － | 0 | 0 | 0 |  | 0 |  | ， | 0 |  | 0 | 0 | 0 | 0 | 0 | 22 | 114 | 137 | 182 | 206 | 297 | 707 | 1426 | 1711 | 1426 | 1254 | 707 | 502 | 297 | 228 | 114 | 69 | 137 | 69 | 22 | 46 | 22 | $\bigcirc$ | 0 | 0 | － |
| 1984 1985 | $\bigcirc$ | 0 | $\bigcirc$ | $\bigcirc$ | $\bigcirc$ | ${ }_{8}^{8}$ | ${ }^{16}$ | ${ }_{3}^{8}$ | ${ }_{6}^{4}$ | 8 | 4 16 | 8 16 | ${ }_{19}^{41}$ | 19 | 160 22 | ${ }_{31}^{340}$ | ${ }_{62}^{872}$ | 1743 <br> 136 | ${ }_{215}^{2472}$ | ${ }_{399}^{3355}$ | 3441 535 | 3270 615 | 4325 818 | 4657 <br> 848 | ${ }_{949}^{4153}$ | 3245 <br> 987 | 2545 <br> 888 | 1731 <br> 906 | ${ }^{1166}$ | ${ }_{821}^{771}$ | 585 725 | （401 | ${ }_{439}^{274}$ | ${ }_{353}^{258}$ | ${ }_{27}^{209}$ | ${ }_{259}^{147}$ | ${ }_{242}^{217}$ | 147 179 | 123 198 | 45 150 | 0 |
| 1986 | 3 | 8 | ${ }^{18}$ | ${ }_{13}$ | ${ }_{13}$ | ${ }_{27}$ | 39 | 29 | 14 | ${ }_{17}$ | 24 | ${ }_{36}$ | 51 | 63 | ${ }_{64}^{22}$ | 67 | 57 | 37 | ${ }_{23}$ | 17 | 30 | 53 | 165 | 397 | 691 | 1066 | 1337 | 1260 | ${ }_{1255}$ | 896 | 637 | 478 | 369 | 287 | ${ }_{257}$ | 207 | 195 | 191 | 190 | 150 109 | 0 |
| 1987 | 2 | 6 | 8 | 10 | 6 | 21 | 52 | 61 | ${ }^{43}$ | 44 | 74 | 95 | 106 | 169 | 254 | 395 | 533 | 830 | 1110 | 1215 | 1071 | 861 | 537 | 337 | 249 | 308 | ${ }^{376}$ | 408 | 335 | 213 | 200 | 160 | 130 | 92 | 61 | 44 | 19 | 10 | 7 | 8 | 。 |
| 1988 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 4 | 22 | 60 | ${ }^{73}$ | 95 | 186 | 354 | 739 | 1663 | 2985 | 3961 | 4250 | 3615 | 2985 | 2134 | 1559 | 1378 | 1399 | 1304 | 1266 | ${ }^{1136}$ | 981 | 1019 | 972 | 1149 | 1058 | 1045 | 1088 | 1145 | 1045 | 747 | 596 | ${ }^{354}$ | 0 |
| 1989 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 7 | 179 | 365 | 3204 | 4837 | 2294 | 2723 | 1988 | 1212 | 877 | 1009 | 1157 | 1309 | 2110 | 2874 | 3570 | 3979 | 3959 | 2848 | 2080 | 1295 | 705 | 498 | 386 | 297 | 260 | 251 | 260 | 237 | 191 | 185 | 0 |
| 1990 | 0 | 0 | $\bigcirc$ | 0 | 0 | 0 | 0 | ， | 8 | 0 | ${ }^{22}$ | 45 | 90 | 90 | ${ }^{127}$ | 187 | 247 | 284 | ${ }^{486}$ | 860 | 1533 | ${ }_{1236}$ | ${ }^{3373}$ | ${ }^{45488}$ | 5221 | 6769 | ${ }^{8579}$ | ${ }^{9410}$ | ${ }^{7353}$ | 6119 | 4473 | 3194 | 2409 | 1855 | ${ }_{1279}^{127}$ | ${ }^{808}$ | 554 | ${ }^{292}$ | ${ }_{687}^{202}$ | ${ }^{90}$ | 0 |
| 1991 | 0 | 0 | $\bigcirc$ | ${ }^{3}$ | 0 | ${ }^{13}$ | 62 | 112 | 85 | 102 | ${ }^{131}$ | 99 | 118 | 927 | 2308 | 1107 | 686 | 628 | 658 | ${ }^{813}$ | 1134 | 1749 | ${ }^{2249}$ | ${ }_{1}^{2785}$ | ${ }_{2851}$ | ${ }_{341}^{341}$ | 4354 | ${ }^{4262}$ | ${ }^{3939}$ | 3496 | 293 | 1877 | 1405 | 1469 | ${ }_{1}^{1237}$ | ${ }_{1167}$ | 1130 | ${ }^{864}$ | 687 | ${ }^{616}$ | 504 |
| 1992 | 0 | 0 | 0 |  | 0 | 0 | 0 | 0 |  | 19 | 103 | ${ }^{306}$ | 166 | 124 | ${ }_{1}^{162}$ | ${ }^{131}$ | 116 | ${ }^{136}$ | 164 | ${ }^{273}$ | ${ }^{471}$ | ${ }^{698}$ | 1108 | 1555 | 1948 | ${ }^{1862}$ | ${ }^{1393}$ | ${ }^{1054}$ | 979 | 866 224 | ${ }_{164}^{812}$ | ${ }^{597}$ | ${ }^{532}$ | ${ }^{401}$ | 168 43 | $\begin{array}{r}104 \\ \hline\end{array}$ |  | 64 | ${ }^{63}$ | 76 | $\bigcirc$ |
| 1993 | 0 | 3 | ${ }^{20}$ | 16 | 3 | 0 | ${ }^{3}$ | ${ }^{26}$ | 66 | 201 | 497 | 691 | 1036 | 1266 | 1963 | 2943 | 3798 | 2821 | 2430 | 1934 | ${ }_{2175}^{1726}$ | 1605 | ${ }_{1809}$ | 1670 | 1684 | 1506 | 1154 | ${ }^{658}$ | 480 | ${ }^{224}$ | 164 | ${ }^{155}$ | 122 | ${ }_{9}^{128}$ | ${ }_{9}^{43}$ | 30 815 | 510 | 45 | ${ }^{2} 8$ | 3 <br> 3 <br> 3 | 0 |
| 1994 1995 | ${ }_{0}^{4}$ | ${ }_{0}^{4}$ | \％ | ${ }_{22}^{4}$ | 73 | 11 103 | 125 | ${ }_{95}$ | ${ }_{51}^{4}$ | ${ }_{44}^{29}$ | 72 103 | 133 396 | ${ }_{792}^{176}$ | ${ }_{1041}^{240}$ | 341 1459 | 452 <br> 2236 | 625 <br> 3043 | ${ }_{3}^{958}$ | ${ }_{4326}^{1253}$ | ${ }_{5635}^{1450}$ | ${ }_{7159}^{2175}$ | ${ }_{917}^{4824}$ | ${ }_{\text {c }}^{6041} 1$ | 4810 10768 | ${ }_{\text {c }}^{3331}$ | ${ }_{19496}^{2495}$ | 1920 12580 | ${ }_{1}^{1407} 688$ | ${ }_{4115}^{1472}$ | ${ }_{2602}^{1088}$ | ${ }_{2003}^{1141}$ | 962 1675 | ${ }_{1201}^{1267}$ | ${ }_{789}^{908}$ | ${ }_{761} 965$ | 815 59 | ${ }_{501}^{510}$ | 4458 | ${ }_{232}^{298}$ | 334 119 |  |
| 1996 | － | 0 | 0 | 0 | 6 | 44 | 200 | 194 | 237 | 712 | 1437 | 1469 | 1462 | 2550 | 3834 | 4445 | 4353 | 4211 | 4457 | 5153 | 6419 | 6045 | 4674 | ${ }^{4311}$ | 3460 | 2887 | 2815 | ${ }^{2487}$ | ${ }^{2268}$ | 1352 | ${ }_{84} 8$ | 549 | 1319 | 211 | ${ }_{76}$ | 4 | 12 | 12 | ${ }_{6}$ | 0 |  |
| 1997 1998 | $\bigcirc$ | $\bigcirc$ | \％ | ${ }_{5}$ | ${ }_{65}{ }^{0}$ | 207 | 109 | 71 | 283 | 14 1301 | 1927 | 109 160 | 166 139 | $\stackrel{\substack{332 \\ 1937}}{1}$ | 3940 | ${ }_{8593}^{1832}$ | ${ }_{\substack{3506 \\ 1027}}$ | 5749 7402 | ${ }_{4272} 5$ | ${ }_{244}^{2855}$ | ${ }_{1253}^{2347}$ | 3963 1012 | 7246 936 | 825 1176 1 | 4460 980 | 2297 <br> 555 <br> 1 | 1380 425 | 737 343 | 506 261 | 371 <br> 234 <br> 1 | 234 174 | 163 147 | 103 109 | ${ }_{98}^{65}$ | 38 87 | 24 60 | ${ }_{49}^{10}$ | 27 |  | 5 16 |  |
| 1999 | 。 | － | － | 0 | 21 | 80 | 80 | 101 | 155 | 174 | 140 | 186 | 270 | 457 | 762 | 1035 | 1838 | 2102 | 2532 | 2300 | 1279 | 755 | 464 | ${ }^{284}$ | 183 | 127 | 120 | 57 | 64 | 106 | ${ }_{43}^{174}$ | ${ }_{27}$ | ${ }_{27}$ | ${ }_{21}^{98}$ | 15 | 17 | 49 | 17 | ${ }_{3}$ | 16 |  |
| 2000 |  | 0 | 0 | $\stackrel{4}{4}$ | 7 | 7 | 7 | 18 188 | ${ }_{178}^{178}$ | ${ }^{1995}$ | 2766 | ${ }^{2473}$ | 1856 | 1341 | ${ }^{1339}$ | ${ }_{3922}^{3492}$ | 2839 | 2483 | 3039 1509 | ${ }_{4523}^{45238}$ | 4097 | ${ }_{803}^{273}$ | 1151 | ${ }^{387}$ | 119 | ${ }_{3}^{31}$ | ${ }_{28}^{11}$ | 4 | 4 | 4 | 7 | $\bigcirc$ | 4 | $\bigcirc$ | 0 | 0 | $\bigcirc$ | 0 | 0 | 0 |  |
| ${ }_{2001}^{2002}$ | ${ }_{8}^{0}$ |  | ${ }_{33}^{16}$ | 33 71 | ${ }_{81}^{33}$ | 16 101 | ${ }_{71}^{41}$ | 188 47 | ${ }_{35}^{459}$ | 942 45 | 1500 40 | 1344 57 | 1304 93 | $\begin{aligned} & 2041 \\ & 123 \end{aligned}$ | 4038 140 | ${ }_{68}^{6242}$ | 9220 61 | 11965 120 | 15404 449 | 15088 1426 | 13405 2689 | ${ }_{3181}^{839}$ | ${ }_{3146}^{2959}$ | 850 2300 | 239 1102 | ${ }_{462}^{97}$ | 28 337 | 183 | 93 | 28 | ${ }_{8}^{0}$ | 0 | $\bigcirc$ | ： | $\bigcirc$ | ： | ： | $\bigcirc$ | 0 | ： | $\bigcirc$ |
| 2003 | 0 | 0 | 0 | 0 | 2 | 2 | 8 | 31 | 61 | 159 | 278 | 453 | 905 | 1419 | 1555 | 1173 | 1063 | 1254 | 1732 | 1911 | 1399 | 961 | 795 | 687 | 697 | 937 | 972 | 640 | 348 | 154 | 63 | 18 | 4 | 2 | 2 | 0 | 0 | 0 | 0 | 0 |  |
| 2004 | 8 | 15 | 47 | 51 | 45 | ${ }^{40}$ | 55 | 48 | 19 |  |  |  | 1 |  | 1 | 1 |  | 38 | 196 | 577 | ${ }^{879}$ | 1015 | 1269 | 1665 | 1779 | 1478 | 1086 | 1040 | 946 | 657 | ${ }^{350}$ | ${ }^{206}$ | ${ }^{103}$ | 45 | 22 | ${ }^{8}$ | 3 | 1 | 0 | 0 | $\bigcirc$ |
| 2005 2006 | 9 | 71 | ${ }^{220}$ | ${ }^{226}$ | 151 0 | ${ }^{93}$ | ¢ ${ }_{6}{ }_{0}$ | 950 | 581 40 | 1147 392 | 1409 1200 | $\begin{aligned} & 1553 \\ & 2435 \end{aligned}$ | 1803 5056 | （1846 | 1130 5992 | 471 6200 | ${ }_{6424}^{263}$ | ${ }_{\text {200 }}$ | 198 5784 | 123 6176 | 97 6664 | 71 6224 | （ ${ }_{547}$ | 34 3928 | 2464 | 11 1296 | ${ }_{712}^{22}$ | 78 392 | 329 216 | 942 120 | ${ }_{\substack{1676 \\ 56}}$ | 1618 24 | ${ }_{16}^{994}$ | 499 | 314 0 | 163 8 | ${ }_{8}^{84}$ | ${ }_{8}^{28}$ | ${ }_{8}$ | $\bigcirc$ | $\bigcirc$ |
| 2007 | $\bigcirc$ | $\bigcirc$ |  | 0 | ${ }_{34}$ | 52 | 52 | 155 | 258 | ${ }^{306}$ | 189 | ${ }_{2}^{245}$ | ${ }_{430}$ | 1324 | 7738 | 15184 | ${ }_{1324} 6424$ | ${ }_{8}^{6976}$ | ${ }_{9853}$ | 6895 | ${ }_{5984}^{664}$ | ${ }_{1024}^{624}$ | ${ }_{15972}^{548}$ | 3928 20704 | ${ }_{19190}^{2464}$ | ${ }_{14358}^{1296}$ | ${ }_{9853}^{712}$ | ${ }_{5399}$ | ${ }_{2528}^{216}$ | ${ }_{1135}^{120}$ | ${ }_{4}^{56}$ | ${ }_{258}^{24}$ | 103 | ${ }_{17}$ | $\bigcirc$ | ${ }_{0}^{8}$ | ${ }_{0}^{8}$ | ${ }_{8}^{8}$ | ${ }_{8}^{8}$ | 0 |  |
| 2008 | 0 |  |  |  | ${ }^{13}$ | 22 | 11 | 4 | 2 | 4 | 11 | 22 | 20 | 31 |  | 120 | 142 | 179 |  | 763 | 2120 | 3978 | 4641 | ${ }^{3688}$ | ${ }^{2233}$ | 1213 | 652 | 375 | 277 | 338 | 275 | 166 | 74 | 24 | 7 |  | 2 | 0 | 0 | 0 |  |
| 2009 2010 | ${ }^{169}$ | ${ }^{161}$ | ${ }_{0}^{227}$ | ${ }^{227}$ | 54 | ${ }_{0}^{21}$ | $\bigcirc$ | $\bigcirc$ | $\bigcirc$ | 27 | 182 17 | 729 0 | 822 17 | ${ }_{231}^{756}$ | ${ }_{451}^{467}$ | ${ }_{162}^{314}$ |  |  |  |  |  | 43 0 | ${ }_{6}^{64}$ | ${ }_{0}^{91}$ | ${ }_{8}^{89}$ | ${ }^{145}$ | 460 0 | ${ }^{1034} 0$ | ${ }^{2137}{ }_{0}$ |  |  | ${ }^{2182}$ | 915 0 | 29 | ${ }^{95}$ | 31 | $\bigcirc$ | $0$ |  | $0$ |  |
| 2011 | $\bigcirc$ | 0 |  | 53 | 53 | 53 | 53 | 0 | 0 | 0 |  | 0 | 53 |  | ${ }^{106}$ |  |  |  |  |  |  |  | 30214 | 425 | ${ }_{53}$ | － | 0 | 0 | 0 |  |  |  | 0 | 0 | 0 | 0 | 0 | 。 |  | － | $\bigcirc$ |
| 2012 | 0 | － |  | 0 | 0 | 0 | 0 | $\bigcirc$ | $\bigcirc$ | $\bigcirc$ |  |  |  |  |  | 1 |  |  |  |  |  | ${ }_{5}^{15}$ | ${ }_{2}^{32}$ | 27 | ${ }_{1}^{14}$ |  |  | 3 | 12 | 20 |  | ${ }_{4}$ | ${ }_{1}$ | 0 | $\bigcirc$ | $\bigcirc$ | $\bigcirc$ | $\bigcirc$ | 0 | 0 | 0 |
| ${ }_{2014}$ | 0 | $\bigcirc$ |  | $\bigcirc$ | 4328 | 15869 | 25667 | 25967 | 7213 | 2885 | 1443 | 2885 | 7213 | 17311 | 14226 |  | 819 | ${ }_{636}$ | ${ }^{4} 12$ | 306273 | 1589 | 2088 | 4617 | 4605 | 791082 | 4928 | 63517 | 166541 | 181363 | 695 | ${ }_{242}$ | ${ }^{4} 34$ | 524 | 12984 | $\bigcirc$ | $\bigcirc$ | $\bigcirc$ | $\bigcirc$ | 1443 | 0 | ${ }^{\circ}$ |

Table 7. Mean body mass (kg) at age over time assumed for Fleet 1 (North Chile).

| Year | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1970 | 0.05 | 0.089 | 0.129 | 0.189 | 0.248 | 0.313 | 0.396 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 1971 | 0.05 | 0.089 | 0.129 | 0.189 | 0.248 | 0.313 | 0.396 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 1972 | 0.05 | 0.089 | 0.129 | 0.189 | 0.248 | 0.313 | 0.396 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 1973 | 0.05 | 0.089 | 0.129 | 0.189 | 0.248 | 0.313 | 0.396 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 1974 | 0.05 | 0.089 | 0.129 | 0.189 | 0.248 | 0.313 | 0.396 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 1975 | 0.05 | 0.089 | 0.129 | 0.189 | 0.248 | 0.313 | 0.396 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 1976 | 0.05 | 0.089 | 0.129 | 0.189 | 0.248 | 0.313 | 0.396 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 1977 | 0.05 | 0.089 | 0.129 | 0.189 | 0.248 | 0.313 | 0.396 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 1978 | 0.05 | 0.105 | 0.124 | 0.163 | 0.204 | 0.314 | 0.369 | 0.405 | 0.434 | 0.453 | 0.59 | 1.115 |
| 1979 | 0.05 | 0.108 | 0.163 | 0.179 | 0.217 | 0.274 | 0.37 | 0.42 | 0.474 | 0.629 | 0.633 | 1.115 |
| 1980 | 0.05 | 0.069 | 0.118 | 0.21 | 0.256 | 0.324 | 0.41 | 0.451 | 0.511 | 0.998 | 0.88 | 1.115 |
| 1981 | 0.05 | 0.094 | 0.139 | 0.214 | 0.269 | 0.331 | 0.412 | 0.481 | 0.58 | 0.661 | 1.112 | 1.115 |
| 1982 | 0.071 | 0.093 | 0.168 | 0.202 | 0.248 | 0.305 | 0.356 | 0.411 | 0.446 | 0.471 | 0.719 | 1.115 |
| 1983 | 0.084 | 0.099 | 0.119 | 0.221 | 0.264 | 0.314 | 0.377 | 0.429 | 0.475 | 0.528 | 0.54 | 1.115 |
| 1984 | 0.05 | 0.164 | 0.186 | 0.217 | 0.273 | 0.345 | 0.394 | 0.437 | 0.497 | 0.568 | 0.786 | 1.115 |
| 1985 | 0.05 | 0.167 | 0.173 | 0.224 | 0.271 | 0.34 | 0.401 | 0.465 | 0.536 | 0.582 | 0.726 | 1.115 |
| 1986 | 0.096 | 0.099 | 0.143 | 0.222 | 0.289 | 0.332 | 0.418 | 0.497 | 0.55 | 0.869 | 0.88 | 1.115 |
| 1987 | 0.092 | 0.121 | 0.146 | 0.189 | 0.233 | 0.336 | 0.427 | 0.477 | 0.513 | 0.65 | 0.803 | 1.115 |
| 1988 | 0.05 | 0.11 | 0.167 | 0.197 | 0.23 | 0.298 | 0.472 | 0.545 | 0.586 | 0.6095 | 0.88 | 1.115 |
| 1989 | 0.05 | 0.123 | 0.167 | 0.23 | 0.27 | 0.31 | 0.379 | 0.491 | 0.541 | 0.569 | 0.713 | 1.115 |
| 1990 | 0.069 | 0.099 | 0.16 | 0.248 | 0.29 | 0.338 | 0.409 | 0.533 | 0.651 | 0.677 | 0.756 | 1.115 |
| 1991 | 0.049 | 0.121 | 0.143 | 0.201 | 0.277 | 0.366 | 0.408 | 0.478 | 0.637 | 0.72 | 0.794 | 0.883 |
| 1992 | 0.069 | 0.092 | 0.127 | 0.201 | 0.268 | 0.3 | 0.373 | 0.444 | 0.512 | 0.595 | 0.681 | 0.786 |
| 1993 | 0.021 | 0.116 | 0.152 | 0.205 | 0.298 | 0.364 | 0.422 | 0.489 | 0.528 | 0.596 | 0.774 | 0.889 |
| 1994 | 0.059 | 0.097 | 0.107 | 0.235 | 0.291 | 0.33 | 0.387 | 0.459 | 0.565 | 0.748 | 0.798 | 0.898 |
| 1995 | 0.069 | 0.101 | 0.137 | 0.186 | 0.263 | 0.321 | 0.357 | 0.434 | 0.561 | 0.668 | 0.88 | 1.115 |
| 1996 | 0.067 | 0 | 0.14 | 0.17 | 0.229 | 0.295 | 0.367 | 0.507 | 0.657 | 0.639 | 0.88 | 1.115 |
| 1997 | 0.029 | 0.063 | 0.125 | 0.177 | 0.246 | 0.357 | 0.503 | 0.615 | 0.584 | 0.728 | 0.88 | 1.115 |
| 1998 | 0 | 0.082 | 0.104 | 0.195 | 0.249 | 0.29 | 0.39 | 0.475 | 0.634 | 0.728 | 0.88 | 1.115 |
| 1999 | 0.071 | 0.074 | 0.089 | 0.147 | 0.27 | 0.315 | 0.446 | 0.722 | 0.584 | 0.728 | 0.88 | 1.115 |
| 2000 | 0.043 | 0.054 | 0.138 | 0.191 | 0.225 | 0.251 | 0.372 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 2001 | 0.066 | 0.093 | 0.112 | 0.133 | 0.204 | 0.286 | 0.421 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 2002 | 0.029 | 0.059 | 0.092 | 0.172 | 0.238 | 0.327 | 0.398 | 0.416 | 0.628 | 0.728 | 0.88 | 1.115 |
| 2003 | 0.036 | 0.082 | 0.102 | 0.141 | 0.227 | 0.309 | 0.416 | 0.464 | 0.534 | 0.728 | 0.88 | 1.115 |
| 2004 | 0.037 | 0.078 | 0.164 | 0.186 | 0.203 | 0.257 | 0.342 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 2005 | 0.029 | 0.076 | 0.111 | 0.175 | 0.222 | 0.268 | 0.281 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 2006 | 0.032 | 0.074 | 0.114 | 0.132 | 0.204 | 0.374 | 0.442 | 0.506 | 0.606 | 0.728 | 0.88 | 1.115 |
| 2007 | 0.087 | 0.075 | 0.122 | 0.158 | 0.222 | 0.296 | 0.404 | 0.514 | 0.614 | 0.723 | 0.723 | 1.115 |
| 2008 | 0.042 | 0.047 | 0.066 | 0.187 | 0.243 | 0.291 | 0.388 | 0.563 | 0.616 | 0.748 | 0.88 | 1.115 |
| 2009 | 0.015 | 0.047 | 0.106 | 0.138 | 0.239 | 0.285 | 0.335 | 0.526 | 0.584 | 0.728 | 0.88 | 1.115 |
| 2010 | 0.013 | 0.048 | 0.101 | 0.172 | 0.233 | 0.301 | 0.397 | 0.493 | 0.639 | 0.772 | 0.88 | 1.115 |
| 2011 | 0.019 | 0.065 | 0.095 | 0.167 | 0.276 | 0.314 | 0.398 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 2012 | 0.016 | 0.048 | 0.088 | 0.202 | 0.235 | 0.269 | 0.396 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |
| 2013 | 0.038 | 0.052 | 0.069 | 0.151 | 0.255 | 0.43 | 0.495 | 0.664 | 0.525 | 0.687 | 0.821 | 1.086 |
| 2014 | 0.018 | 0.04 | 0.082 | 0.189 | 0.248 | 0.313 | 0.396 | 0.488 | 0.584 | 0.728 | 0.88 | 1.115 |

Table 8. Mean body mass (kg) at age over time assumed for Fleet 2 (Central South Chile).

| Year | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1970 | 0.052 | 0.093 | 0.131 | 0.178 | 0.262 | 0.294 | 0.34 | 0.396 | 0.549 | 0.738 | 0.984 | 1.093 |
| 1971 | 0.052 | 0.093 | 0.131 | 0.178 | 0.262 | 0.294 | 0.34 | 0.396 | 0.549 | 0.738 | 0.984 | 1.093 |
| 1972 | 0.052 | 0.093 | 0.131 | 0.178 | 0.262 | 0.294 | 0.34 | 0.396 | 0.549 | 0.738 | 0.984 | 1.093 |
| 1973 | 0.052 | 0.093 | 0.131 | 0.178 | 0.262 | 0.294 | 0.34 | 0.396 | 0.549 | 0.738 | 0.984 | 1.093 |
| 1974 | 0.052 | 0.093 | 0.131 | 0.178 | 0.262 | 0.294 | 0.34 | 0.396 | 0.549 | 0.738 | 0.984 | 1.093 |
| 1975 | 0.052 | 0.093 | 0.131 | 0.178 | 0.262 | 0.294 | 0.34 | 0.396 | 0.549 | 0.738 | 0.984 | 1.093 |
| 1976 | 0.052 | 0.078 | 0.155 | 0.214 | 0.275 | 0.336 | 0.394 | 0.472 | 0.632 | 0.714 | 0.898 | 1.538 |
| 1977 | 0.055 | 0.092 | 0.109 | 0.236 | 0.275 | 0.314 | 0.375 | 0.456 | 0.521 | 0.732 | 0.651 | 1.137 |
| 1978 | 0.052 | 0.084 | 0.104 | 0.147 | 0.211 | 0.327 | 0.394 | 0.449 | 0.514 | 0.583 | 0.631 | 1.538 |
| 1979 | 0.052 | 0.108 | 0.16 | 0.199 | 0.241 | 0.301 | 0.388 | 0.466 | 0.588 | 0.871 | 1.265 | 1.972 |
| 1980 | 0.026 | 0.06 | 0.132 | 0.231 | 0.272 | 0.35 | 0.447 | 0.519 | 0.716 | 0.82 | 1.073 | 1.854 |
| 1981 | 0.052 | 0.095 | 0.149 | 0.242 | 0.294 | 0.34 | 0.407 | 0.503 | 0.637 | 0.765 | 1.184 | 1.9 |
| 1982 | 0.055 | 0.085 | 0.166 | 0.207 | 0.269 | 0.323 | 0.378 | 0.472 | 0.536 | 0.644 | 0.987 | 1.185 |
| 1983 | 0.07 | 0.099 | 0.122 | 0.23 | 0.273 | 0.32 | 0.374 | 0.461 | 0.596 | 0.709 | 1.196 | 1.769 |
| 1984 | 0.035 | 0.135 | 0.154 | 0.185 | 0.266 | 0.33 | 0.383 | 0.449 | 0.577 | 0.685 | 1.012 | 1.846 |
| 1985 | 0.058 | 0.148 | 0.181 | 0.223 | 0.27 | 0.339 | 0.398 | 0.473 | 0.573 | 0.796 | 1.376 | 1.647 |
| 1986 | 0.073 | 0.075 | 0.172 | 0.247 | 0.286 | 0.346 | 0.427 | 0.518 | 0.64 | 0.844 | 1.351 | 2.11 |
| 1987 | 0.076 | 0.117 | 0.14 | 0.191 | 0.27 | 0.357 | 0.434 | 0.503 | 0.577 | 0.689 | 1.089 | 1.979 |
| 1988 | 0.1 | 0.124 | 0.159 | 0.197 | 0.233 | 0.342 | 0.444 | 0.512 | 0.588 | 0.75 | 1.012 | 1.372 |
| 1989 | 0.052 | 0.103 | 0.22 | 0.241 | 0.278 | 0.339 | 0.467 | 0.585 | 0.702 | 0.779 | 0.88 | 1.538 |
| 1990 | 0.064 | 0.091 | 0.153 | 0.264 | 0.309 | 0.373 | 0.461 | 0.582 | 0.694 | 0.835 | 0.97 | 1.598 |
| 1991 | 0.037 | 0.106 | 0.132 | 0.186 | 0.271 | 0.381 | 0.451 | 0.542 | 0.667 | 0.787 | 0.901 | 1.053 |
| 1992 | 0.063 | 0.083 | 0.118 | 0.177 | 0.239 | 0.275 | 0.409 | 0.524 | 0.594 | 0.709 | 0.851 | 1.046 |
| 1993 | 0.011 | 0.089 | 0.121 | 0.181 | 0.246 | 0.32 | 0.408 | 0.579 | 0.719 | 0.853 | 0.965 | 1.174 |
| 1994 | 0.041 | 0.084 | 0.112 | 0.224 | 0.27 | 0.336 | 0.462 | 0.643 | 0.808 | 0.868 | 1.058 | 1.421 |
| 1995 | 0.07 | 0.098 | 0.145 | 0.192 | 0.27 | 0.34 | 0.429 | 0.577 | 0.807 | 0.965 | 1.115 | 1.367 |
| 1996 | 0.061 | 0.092 | 0.151 | 0.191 | 0.28 | 0.352 | 0.524 | 0.683 | 0.945 | 1.216 | 1.426 | 1.477 |
| 1997 | 0.104 | 0.106 | 0.146 | 0.201 | 0.26 | 0.355 | 0.495 | 0.683 | 0.884 | 1.088 | 1.467 | 1.647 |
| 1998 | 0.084 | 0.128 | 0.138 | 0.178 | 0.248 | 0.34 | 0.545 | 0.806 | 1.035 | 1.246 | 1.412 | 1.655 |
| 1999 | 0.09 | 0.109 | 0.134 | 0.174 | 0.25 | 0.331 | 0.465 | 0.742 | 1.021 | 1.258 | 1.376 | 1.776 |
| 2000 | 0.043 | 0.064 | 0.163 | 0.196 | 0.255 | 0.346 | 0.466 | 0.756 | 0.999 | 1.141 | 1.228 | 1.563 |
| 2001 | 0.066 | 0.098 | 0.122 | 0.179 | 0.258 | 0.325 | 0.461 | 0.614 | 0.828 | 1.074 | 1.36 | 1.671 |
| 2002 | 0.031 | 0.074 | 0.13 | 0.2 | 0.257 | 0.329 | 0.445 | 0.645 | 0.883 | 1.102 | 1.321 | 1.649 |
| 2003 | 0.036 | 0.086 | 0.117 | 0.186 | 0.245 | 0.307 | 0.4 | 0.564 | 0.768 | 1.005 | 1.209 | 1.537 |
| 2004 | 0.034 | 0.08 | 0.158 | 0.193 | 0.247 | 0.307 | 0.387 | 0.528 | 0.7 | 0.897 | 1.087 | 1.541 |
| 2005 | 0.029 | 0.075 | 0.113 | 0.196 | 0.259 | 0.318 | 0.399 | 0.517 | 0.641 | 0.767 | 0.918 | 1.296 |
| 2006 | 0.033 | 0.076 | 0.116 | 0.141 | 0.261 | 0.35 | 0.419 | 0.516 | 0.631 | 0.752 | 0.924 | 1.263 |
| 2007 | 0.086 | 0.074 | 0.121 | 0.172 | 0.226 | 0.331 | 0.431 | 0.51 | 0.621 | 0.756 | 0.903 | 1.177 |
| 2008 | 0.036 | 0.048 | 0.069 | 0.186 | 0.254 | 0.312 | 0.416 | 0.515 | 0.605 | 0.719 | 0.861 | 1.148 |
| 2009 | 0.014 | 0.045 | 0.109 | 0.142 | 0.253 | 0.33 | 0.411 | 0.532 | 0.625 | 0.764 | 0.886 | 1.144 |
| 2010 | 0.014 | 0.052 | 0.101 | 0.175 | 0.237 | 0.313 | 0.415 | 0.539 | 0.649 | 0.787 | 0.964 | 1.473 |
| 2011 | 0.019 | 0.067 | 0.101 | 0.19 | 0.287 | 0.353 | 0.466 | 0.613 | 0.774 | 0.923 | 1.173 | 1.514 |
| 2012 | 0.007 | 0.014 | 0.082 | 0.202 | 0.264 | 0.353 | 0.476 | 0.558 | 0.711 | 0.912 | 1.146 | 1.6 |
| 2013 | 0.054 | 0.158 | 0.251 | 0.26 | 0.318 | 0.385 | 0.45 | 0.553 | 0.705 | 0.829 | 1.117 | 1.977 |
| 2014 | 0.052 | 0.093 | 0.182 | 0.247 | 0.375 | 0.485 | 0.534 | 0.682 | 1.094 | 1.281 | 1.302 | 1.656 |

Table 9. Mean body mass (kg) at age over time assumed for Fleet 3 (Far North).

| Year | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1970 | 0.045 | 0.171 | 0.377 | 0.642 | 0.945 | 1.265 | 1.587 | 1.9 | 2.196 | 2.47 | 2.721 | 2.946 |
| 1971 | 0.045 | 0.171 | 0.377 | 0.643 | 0.946 | 1.266 | 1.588 | 1.902 | 2.198 | 2.472 | 2.723 | 2.949 |
| 1972 | 0.03 | 0.13 | 0.306 | 0.548 | 0.835 | 1.148 | 1.47 | 1.789 | 2.095 | 2.382 | 2.647 | 2.887 |
| 1973 | 0.037 | 0.147 | 0.33 | 0.568 | 0.842 | 1.134 | 1.43 | 1.718 | 1.991 | 2.246 | 2.478 | 2.688 |
| 1974 | 0.038 | 0.147 | 0.326 | 0.558 | 0.825 | 1.108 | 1.393 | 1.671 | 1.934 | 2.178 | 2.402 | 2.603 |
| 1975 | 0.034 | 0.136 | 0.31 | 0.54 | 0.808 | 1.095 | 1.387 | 1.674 | 1.946 | 2.201 | 2.434 | 2.645 |
| 1976 | 0.044 | 0.16 | 0.34 | 0.567 | 0.822 | 1.087 | 1.351 | 1.606 | 1.845 | 2.065 | 2.266 | 2.446 |
| 1977 | 0.032 | 0.13 | 0.294 | 0.51 | 0.76 | 1.028 | 1.3 | 1.566 | 1.818 | 2.054 | 2.27 | 2.465 |
| 1978 | 0.032 | 0.129 | 0.295 | 0.516 | 0.774 | 1.05 | 1.332 | 1.608 | 1.872 | 2.117 | 2.343 | 2.547 |
| 1979 | 0.036 | 0.138 | 0.304 | 0.518 | 0.762 | 1.02 | 1.28 | 1.532 | 1.77 | 1.991 | 2.193 | 2.375 |
| 1980 | 0.036 | 0.136 | 0.298 | 0.506 | 0.743 | 0.994 | 1.245 | 1.49 | 1.721 | 1.934 | 2.13 | 2.306 |
| 1981 | 0.041 | 0.148 | 0.314 | 0.524 | 0.758 | 1.003 | 1.247 | 1.481 | 1.702 | 1.905 | 2.089 | 2.255 |
| 1982 | 0.039 | 0.144 | 0.309 | 0.519 | 0.755 | 1.002 | 1.249 | 1.488 | 1.712 | 1.92 | 2.108 | 2.278 |
| 1983 | 0.042 | 0.138 | 0.28 | 0.451 | 0.638 | 0.828 | 1.014 | 1.191 | 1.356 | 1.507 | 1.643 | 1.764 |
| 1984 | 0.044 | 0.156 | 0.328 | 0.541 | 0.778 | 1.024 | 1.267 | 1.501 | 1.719 | 1.921 | 2.103 | 2.267 |
| 1985 | 0.04 | 0.149 | 0.322 | 0.541 | 0.789 | 1.048 | 1.308 | 1.558 | 1.794 | 2.012 | 2.211 | 2.389 |
| 1986 | 0.042 | 0.151 | 0.323 | 0.539 | 0.781 | 1.033 | 1.285 | 1.527 | 1.755 | 1.965 | 2.156 | 2.327 |
| 1987 | 0.034 | 0.132 | 0.294 | 0.504 | 0.745 | 1.001 | 1.26 | 1.512 | 1.751 | 1.973 | 2.176 | 2.359 |
| 1988 | 0.038 | 0.145 | 0.315 | 0.533 | 0.78 | 1.041 | 1.302 | 1.554 | 1.793 | 2.013 | 2.215 | 2.396 |
| 1989 | 0.044 | 0.158 | 0.337 | 0.561 | 0.812 | 1.074 | 1.334 | 1.585 | 1.821 | 2.038 | 2.236 | 2.413 |
| 1990 | 0.042 | 0.15 | 0.32 | 0.532 | 0.769 | 1.017 | 1.263 | 1.499 | 1.722 | 1.927 | 2.113 | 2.28 |
| 1991 | 0.039 | 0.142 | 0.305 | 0.511 | 0.743 | 0.985 | 1.227 | 1.461 | 1.68 | 1.883 | 2.068 | 2.234 |
| 1992 | 0.04 | 0.148 | 0.318 | 0.534 | 0.776 | 1.031 | 1.286 | 1.531 | 1.763 | 1.976 | 2.171 | 2.346 |
| 1993 | 0.039 | 0.147 | 0.323 | 0.549 | 0.807 | 1.08 | 1.354 | 1.62 | 1.871 | 2.104 | 2.317 | 2.508 |
| 1994 | 0.036 | 0.147 | 0.335 | 0.584 | 0.874 | 1.186 | 1.503 | 1.813 | 2.109 | 2.385 | 2.638 | 2.867 |
| 1995 | 0.038 | 0.146 | 0.318 | 0.54 | 0.792 | 1.058 | 1.325 | 1.583 | 1.827 | 2.053 | 2.26 | 2.446 |
| 1996 | 0.038 | 0.145 | 0.317 | 0.537 | 0.788 | 1.053 | 1.318 | 1.576 | 1.82 | 2.045 | 2.251 | 2.436 |
| 1997 | 0.045 | 0.152 | 0.312 | 0.506 | 0.72 | 0.94 | 1.155 | 1.361 | 1.553 | 1.729 | 1.889 | 2.031 |
| 1998 | 0.04 | 0.14 | 0.294 | 0.483 | 0.693 | 0.911 | 1.126 | 1.333 | 1.526 | 1.703 | 1.864 | 2.008 |
| 1999 | 0.037 | 0.146 | 0.324 | 0.557 | 0.824 | 1.107 | 1.394 | 1.673 | 1.938 | 2.183 | 2.408 | 2.611 |
| 2000 | 0.035 | 0.145 | 0.336 | 0.592 | 0.893 | 1.218 | 1.55 | 1.877 | 2.189 | 2.481 | 2.75 | 2.994 |
| 2001 | 0.033 | 0.139 | 0.324 | 0.572 | 0.864 | 1.18 | 1.504 | 1.822 | 2.127 | 2.412 | 2.674 | 2.912 |
| 2002 | 0.036 | 0.145 | 0.33 | 0.576 | 0.861 | 1.167 | 1.478 | 1.783 | 2.074 | 2.344 | 2.593 | 2.817 |
| 2003 | 0.04 | 0.154 | 0.341 | 0.584 | 0.862 | 1.157 | 1.454 | 1.743 | 2.017 | 2.272 | 2.504 | 2.714 |
| 2004 | 0.038 | 0.149 | 0.333 | 0.574 | 0.852 | 1.148 | 1.447 | 1.74 | 2.017 | 2.275 | 2.511 | 2.724 |
| 2005 | 0.037 | 0.15 | 0.341 | 0.595 | 0.89 | 1.206 | 1.527 | 1.842 | 2.142 | 2.422 | 2.678 | 2.911 |
| 2006 | 0.038 | 0.152 | 0.347 | 0.606 | 0.907 | 1.23 | 1.558 | 1.88 | 2.187 | 2.473 | 2.735 | 2.973 |
| 2007 | 0.038 | 0.149 | 0.335 | 0.579 | 0.861 | 1.161 | 1.465 | 1.762 | 2.044 | 2.306 | 2.546 | 2.763 |
| 2008 | 0.036 | 0.146 | 0.334 | 0.585 | 0.876 | 1.19 | 1.51 | 1.823 | 2.122 | 2.4 | 2.656 | 2.888 |
| 2009 | 0.038 | 0.15 | 0.337 | 0.582 | 0.865 | 1.167 | 1.474 | 1.773 | 2.057 | 2.321 | 2.563 | 2.782 |
| 2010 | 0.039 | 0.15 | 0.332 | 0.567 | 0.837 | 1.123 | 1.411 | 1.691 | 1.956 | 2.203 | 2.428 | 2.631 |
| 2011 | 0.031 | 0.143 | 0.351 | 0.644 | 1 | 1.395 | 1.806 | 2.217 | 2.614 | 2.99 | 3.337 | 3.655 |
| 2012 | 0.032 | 0.145 | 0.349 | 0.632 | 0.971 | 1.344 | 1.731 | 2.115 | 2.485 | 2.834 | 3.156 | 3.449 |
| 2013 | 0.032 | 0.145 | 0.349 | 0.632 | 0.971 | 1.344 | 1.731 | 2.115 | 2.485 | 2.834 | 3.156 | 3.449 |
| 2014 | 0.032 | 0.145 | 0.349 | 0.632 | 0.971 | 1.344 | 1.731 | 2.115 | 2.485 | 2.834 | 3.156 | 3.449 |

Table 10. Mean body mass (kg) at age over time assumed for Fleet 4 (Offshore Trawl).

| Year | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1970 | 0.052 | 0.093 | 0.131 | 0.178 | 0.262 | 0.294 | 0.34 | 0.396 | 0.549 | 0.738 | 0.984 | 1.093 |
| 1971 | 0.052 | 0.093 | 0.131 | 0.178 | 0.262 | 0.294 | 0.34 | 0.396 | 0.549 | 0.738 | 0.984 | 1.093 |
| 1972 | 0.052 | 0.093 | 0.131 | 0.178 | 0.262 | 0.294 | 0.34 | 0.396 | 0.549 | 0.738 | 0.984 | 1.093 |
| 1973 | 0.052 | 0.093 | 0.131 | 0.178 | 0.262 | 0.294 | 0.34 | 0.396 | 0.549 | 0.738 | 0.984 | 1.093 |
| 1974 | 0.052 | 0.093 | 0.131 | 0.178 | 0.262 | 0.294 | 0.34 | 0.396 | 0.549 | 0.738 | 0.984 | 1.093 |
| 1975 | 0.052 | 0.093 | 0.131 | 0.178 | 0.262 | 0.294 | 0.34 | 0.396 | 0.549 | 0.738 | 0.984 | 1.093 |
| 1976 | 0.052 | 0.078 | 0.155 | 0.214 | 0.275 | 0.336 | 0.394 | 0.472 | 0.632 | 0.714 | 0.898 | 1.538 |
| 1977 | 0.055 | 0.092 | 0.109 | 0.236 | 0.275 | 0.314 | 0.375 | 0.456 | 0.521 | 0.732 | 0.651 | 1.137 |
| 1978 | 0.052 | 0.084 | 0.104 | 0.147 | 0.211 | 0.327 | 0.394 | 0.449 | 0.514 | 0.583 | 0.631 | 1.538 |
| 1979 | 0.052 | 0.108 | 0.16 | 0.199 | 0.241 | 0.301 | 0.388 | 0.466 | 0.588 | 0.871 | 1.265 | 1.972 |
| 1980 | 0.026 | 0.06 | 0.132 | 0.231 | 0.272 | 0.35 | 0.447 | 0.519 | 0.716 | 0.82 | 1.073 | 1.854 |
| 1981 | 0.052 | 0.095 | 0.149 | 0.242 | 0.294 | 0.34 | 0.407 | 0.503 | 0.637 | 0.765 | 1.184 | 1.9 |
| 1982 | 0.055 | 0.085 | 0.166 | 0.207 | 0.269 | 0.323 | 0.378 | 0.472 | 0.536 | 0.644 | 0.987 | 1.185 |
| 1983 | 0.07 | 0.099 | 0.122 | 0.23 | 0.273 | 0.32 | 0.374 | 0.461 | 0.596 | 0.709 | 1.196 | 1.769 |
| 1984 | 0.035 | 0.135 | 0.154 | 0.185 | 0.266 | 0.33 | 0.383 | 0.449 | 0.577 | 0.685 | 1.012 | 1.846 |
| 1985 | 0.058 | 0.148 | 0.181 | 0.223 | 0.27 | 0.339 | 0.398 | 0.473 | 0.573 | 0.796 | 1.376 | 1.647 |
| 1986 | 0.073 | 0.075 | 0.172 | 0.247 | 0.286 | 0.346 | 0.427 | 0.518 | 0.64 | 0.844 | 1.351 | 2.11 |
| 1987 | 0.076 | 0.117 | 0.14 | 0.191 | 0.27 | 0.357 | 0.434 | 0.503 | 0.577 | 0.689 | 1.089 | 1.979 |
| 1988 | 0.1 | 0.124 | 0.159 | 0.197 | 0.233 | 0.342 | 0.444 | 0.512 | 0.588 | 0.75 | 1.012 | 1.372 |
| 1989 | 0.052 | 0.103 | 0.22 | 0.241 | 0.278 | 0.339 | 0.467 | 0.585 | 0.702 | 0.779 | 0.88 | 1.538 |
| 1990 | 0.064 | 0.091 | 0.153 | 0.264 | 0.309 | 0.373 | 0.461 | 0.582 | 0.694 | 0.835 | 0.97 | 1.598 |
| 1991 | 0.037 | 0.106 | 0.132 | 0.186 | 0.271 | 0.381 | 0.451 | 0.542 | 0.667 | 0.787 | 0.901 | 1.053 |
| 1992 | 0.063 | 0.083 | 0.118 | 0.177 | 0.239 | 0.275 | 0.409 | 0.524 | 0.594 | 0.709 | 0.851 | 1.046 |
| 1993 | 0.011 | 0.089 | 0.121 | 0.181 | 0.246 | 0.32 | 0.408 | 0.579 | 0.719 | 0.853 | 0.965 | 1.174 |
| 1994 | 0.041 | 0.084 | 0.112 | 0.224 | 0.27 | 0.336 | 0.462 | 0.643 | 0.808 | 0.868 | 1.058 | 1.421 |
| 1995 | 0.07 | 0.098 | 0.145 | 0.192 | 0.27 | 0.34 | 0.429 | 0.577 | 0.807 | 0.965 | 1.115 | 1.367 |
| 1996 | 0.061 | 0.092 | 0.151 | 0.191 | 0.28 | 0.352 | 0.524 | 0.683 | 0.945 | 1.216 | 1.426 | 1.477 |
| 1997 | 0.104 | 0.106 | 0.146 | 0.201 | 0.26 | 0.355 | 0.495 | 0.683 | 0.884 | 1.088 | 1.467 | 1.647 |
| 1998 | 0.084 | 0.128 | 0.138 | 0.178 | 0.248 | 0.34 | 0.545 | 0.806 | 1.035 | 1.246 | 1.412 | 1.655 |
| 1999 | 0.09 | 0.109 | 0.134 | 0.174 | 0.25 | 0.331 | 0.465 | 0.742 | 1.021 | 1.258 | 1.376 | 1.776 |
| 2000 | 0.043 | 0.064 | 0.163 | 0.196 | 0.255 | 0.346 | 0.466 | 0.756 | 0.999 | 1.141 | 1.228 | 1.563 |
| 2001 | 0.066 | 0.098 | 0.122 | 0.179 | 0.258 | 0.325 | 0.461 | 0.614 | 0.828 | 1.074 | 1.36 | 1.671 |
| 2002 | 0.031 | 0.074 | 0.13 | 0.2 | 0.257 | 0.329 | 0.445 | 0.645 | 0.883 | 1.102 | 1.321 | 1.649 |
| 2003 | 0.036 | 0.086 | 0.117 | 0.186 | 0.245 | 0.307 | 0.4 | 0.564 | 0.768 | 1.005 | 1.209 | 1.537 |
| 2004 | 0.034 | 0.08 | 0.158 | 0.193 | 0.247 | 0.307 | 0.387 | 0.528 | 0.7 | 0.897 | 1.087 | 1.541 |
| 2005 | 0.029 | 0.075 | 0.113 | 0.196 | 0.259 | 0.318 | 0.399 | 0.517 | 0.641 | 0.767 | 0.918 | 1.296 |
| 2006 | 0.033 | 0.076 | 0.116 | 0.141 | 0.261 | 0.35 | 0.419 | 0.516 | 0.631 | 0.752 | 0.924 | 1.263 |
| 2007 | 0.086 | 0.074 | 0.121 | 0.172 | 0.226 | 0.331 | 0.431 | 0.51 | 0.621 | 0.756 | 0.903 | 1.177 |
| 2008 | 0.036 | 0.048 | 0.069 | 0.186 | 0.254 | 0.312 | 0.416 | 0.515 | 0.605 | 0.719 | 0.861 | 1.148 |
| 2009 | 0.014 | 0.045 | 0.109 | 0.142 | 0.253 | 0.33 | 0.411 | 0.532 | 0.625 | 0.764 | 0.886 | 1.144 |
| 2010 | 0.014 | 0.052 | 0.101 | 0.175 | 0.237 | 0.313 | 0.415 | 0.539 | 0.649 | 0.787 | 0.964 | 1.473 |
| 2011 | 0.019 | 0.067 | 0.101 | 0.19 | 0.287 | 0.353 | 0.466 | 0.613 | 0.774 | 0.923 | 1.173 | 1.514 |
| 2012 | 0.007 | 0.014 | 0.082 | 0.202 | 0.264 | 0.353 | 0.476 | 0.558 | 0.711 | 0.912 | 1.146 | 1.6 |
| 2013 | 0.052 | 0.125 | 0.268 | 0.263 | 0.31 | 0.362 | 0.431 | 0.507 | 0.678 | 0.726 | 0.936 | 1.143 |
| 2014 | 0.026 | 0.069 | 0.150 | 0.218 | 0.287 | 0.356 | 0.458 | 0.559 | 0.721 | 0.854 | 1.085 | 1.419 |

Table 11. Survey biomass time series (1000 t).

| Year | Biomass (North Ch ile) | Biomass (Central south Chile) | Spawning Biomass DEPM (Central south Chile) | Biomass (Peru) |
| :---: | :---: | :---: | :---: | :---: |
| 1983 |  |  |  |  |
| 1984 | 99 |  |  |  |
| 1985 | 324 |  |  |  |
| 1986 | 123 |  |  | 17811 |
| 1987 | 213 |  |  | 22955 |
| 1988 | 134 |  |  | 9459 |
| 1989 |  |  |  | 15034 |
| 1990 |  |  |  | 14139 |
| 1991 | 242 |  |  | 16486 |
| 1992 |  |  |  | 6266 |
| 1993 |  |  |  | 19659 |
| 1994 |  |  |  | 10768 |
| 1995 |  |  |  | 6429 |
| 1996 |  |  |  | 7271 |
| 1997 |  | 3530 |  | 2561 |
| 1998 |  | 3200 |  | 190 |
| 1999 |  | 4100 | 5724 | 342 |
| 2000 |  | 5600 | 4688 | 2373 |
| 2001 |  | 5950 | 5627 | 2052 |
| 2002 |  | 3700 |  | 248 |
| 2003 |  | 2640 | 1388 | 1118 |
| 2004 |  | 2640 | 3287 | 864 |
| 2005 |  | 4110 | 1043 | 1025 |
| 2006 | 112 | 3192 | 3283 | 1678 |
| 2007 | 275 | 3140 | 626 | 522 |
| 2008 | 259 | 487 | 1935 | 223 |
| 2009 | 18 | 328 |  | 849 |
| 2010 | 440 |  |  |  |
| 2011 | 432 |  |  | 678 |
| 2012 | 230 |  |  | 94 |
| 2013 | 144 |  |  | 890 |
| 2014 | 87 |  |  |  |

Table 12. Acoustic biomass age-structured (numbers, $x 10^{9}$ ). (2006-2014). North Chile.

| Year | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 |
| :---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 2006 | 116822 | 403538 | 272612 | 154651 | 21715 | 0 | 0 | 0 | 0 | 0 | 0 |
| 2007 | 273 | 69043 | 241335 | 755691 | 292140 | 19746 | 2980 | 0 | 0 | 0 | 0 |
| 2008 | 14998 | $2 \mathrm{E}+06$ | $2 \mathrm{E}+06$ | 41648 | 45795 | 16174 | 145 | 0 | 0 | 0 | 0 |
| 2009 | 0 | 54510 | 55714 | 20943 | 392 | 721 | 129 | 0 | 0 | 0 | 0 |
| 2010 | 0 | 10321 | 175262 | $2 \mathrm{E}+06$ | 401742 | 79272 | 20972 | 0 | 0 | 0 | 0 |
| 2011 | 0 | 764206 | 350733 | $1 \mathrm{E}+06$ | 103679 | 144140 | 37702 | 2300 | 0 | 0 | 0 |
| 2012 | 721 | 538523 | 63430 | 214664 | 307532 | 116579 | 9419 | 779 | 0 | 0 | 0 |
| 2013 | 83292 | 269344 | 16908 | 134710 | 128512 | 114395 | 29374 | 4131 | 0 | 0 | 0 |
| 2014 | 473762 | $1 \mathrm{E}+06$ | 14319 | 20965 | 5207 | 19510 | 12090 | 0 | 0 | 0 | 0 |

Table 13. Acoustic biomass age-structure (numbers, $x 10^{9}$ ). (1997-2009). Central South Chile.

| Year | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1997 | 74 | 265 | 7857 | 8492 | 2422 | 668 | 131 | 115 | 128 | 75 | 76 | 95 |
| 1998 | 3 | 119 | 10599 | 9851 | 912 | 236 | 142 | 241 | 166 | 24 | 4 | 6 |
| 1999 | 0 | 365 | 5368 | 7891 | 1963 | 1014 | 377 | 194 | 500 | 357 | 212 | 63 |
| 2000 | 0 | 0 | 2529 | 11296 | 7864 | 2787 | 567 | 113 | 42 | 20 | 16 | 20 |
| 2001 | 0 | 280 | 6596 | 14354 | 7124 | 1792 | 428 | 241 | 177 | 103 | 44 | 7 |
| 2002 | 0 | 1 | 498 | 2217 | 1833 | 2363 | 565 | 280 | 80 | 21 | 7 | 2 |
| 2003 | 0 | 1 | 276 | 2132 | 3089 | 2343 | 680 | 200 | 156 | 115 | 54 | 41 |
| 2004 | 0 | 0 | 120 | 662 | 1833 | 2277 | 1709 | 1128 | 910 | 893 | 320 | 62 |
| 2005 | 0 | 0 | 0 | 1210 | 2670 | 5250 | 2377 | 701 | 320 | 64 | 82 | 188 |
| 2006 | 0 | 0 | 0 | 12 | 1799 | 4266 | 2625 | 511 | 322 | 188 | 110 | 70 |
| 2007 | 0 | 0 | 0 | 0 | 1 | 339 | 1113 | 1058 | 976 | 869 | 439 | 411 |
| 2008 | 0 | 0 | 0 | 0 | 0 | 96 | 317 | 256 | 117 | 87 | 34 | 49 |
| 2009 | 0 | 0 | 0 | 0 | 0 | 0 | 7 | 265 | 169 | 143 | 127 | 98 |

Table 14. Biomass age-structured (numbers, $x 10^{9}$ ). DEPM (Daily Egg Production Method). Central south Chile.

| Year | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 2001 | 0 | 3787 | 8944 | 9208 | 1436 | 700 | 420 | 401 | 182 | 58 | 14 |
| 2003 | 0 | 54 | 529 | 837 | 336 | 563 | 398 | 400 | 219 | 84 | 35 |
| 2004 | 0 | 122 | 1217 | 2801 | 1511 | 719 | 405 | 584 | 376 | 168 | 66 |
| 2005 | 0 | 0 | 0 | 21 | 1115 | 718 | 485 | 336 | 136 | 94 | 57 |
| 2006 | 0 | 0 | 6 | 216 | 1232 | 1892 | 1473 | 849 | 304 | 114 | 78 |
| 2008 | 0 | 0 | 0 | 146 | 419 | 778 | 982 | 716 | 323 | 194 | 9 |

Table 15. Jack mackerel sexual maturity at age used in the stock assessment. Central south Chile.

| Age | Proportion |
| :---: | :---: |
| 1 | 0.07 |
| 2 | 0.31 |
| 3 | 0.72 |
| 4 | 0.93 |
| 5 | 0.98 |
| 6 | 0.99 |
| 7 | 1 |
| 8 | 1 |
| 9 | 1 |
| 10 | 1 |
| 11 | 1 |
| 12 | 1 |

Table 16. Ageing error matrix of Jack mackerel based on Chilean age studies

| Age | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12+ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 2 | 0 | 0.76 | 0.22 | 0.02 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 3 | 0 | 0.24 | 0.51 | 0.23 | 0.02 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 4 | 0 | 0.02 | 0.23 | 0.5 | 0.23 | 0.02 | 0 | 0 | 0 | 0 | 0 | 0 |
| 5 | 0 | 0 | 0.02 | 0.23 | 0.49 | 0.23 | 0.02 | 0 | 0 | 0 | 0 | 0 |
| 6 | 0 | 0 | 0 | 0.03 | 0.23 | 0.48 | 0.23 | 0.03 | 0 | 0 | 0 | 0 |
| 7 | 0 | 0 | 0 | 0 | 0.03 | 0.24 | 0.46 | 0.24 | 0.03 | 0 | 0 | 0 |
| 8 | 0 | 0 | 0 | 0 | 0 | 0.03 | 0.24 | 0.45 | 0.24 | 0.03 | 0 | 0 |
| 9 | 0 | 0 | 0 | 0 | 0 | 0 | 0.04 | 0.24 | 0.44 | 0.24 | 0.04 | 0 |
| 10 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.04 | 0.24 | 0.43 | 0.24 | 0.04 |
| 11 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.04 | 0.24 | 0.42 | 0.29 |
| 12+ | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.05 | 0.24 | 0.71 |

Table 17. Jack mackerel population estimates. Spawning biomass (SSB, 1000 t), recruitment ( R , number $10^{9}$ ) and fishing mortality $\left(\mathrm{F}\right.$, year ${ }^{-1}$ ).

|  | Model-1, $h=0.8$ |  |  | Model-2, $h=0.65$ |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Year | SSB | R | F | SSB | R | F |
| 1970 | 9276.2 | 7428.6 | 0.02 | 9749.1 | 7535.8 | 0.02 |
| 1971 | 8391.5 | 5093.5 | 0.03 ' | 8784.7 | 5142.8 | 0.02 |
| 1972 | 7782.0 | 9630.5 | 0.02 ! | 8113.2 | 9725.6 | 0.02 |
| 1973 | 7323.8 | 8670.8 | 0.03 | 7605.5 | 8718.6 | 0.03 |
| 1974 | 6975.6 | 8555.9 | 0.04 | 7217.9 | 8569.9 | 0.04 |
| 1975 | 6785.9 | 19424.6 | 0.05 | 6995.9 | 19485.7 | 0.04 |
| 1976 | 6811.3 | 21163.4 | 0.06 | 6994.4 | 21186.4 | 0.06 |
| 1977 | 7130.7 | 19648.5 | 0.08 | 7291.3 | 19588.7 | 0.08 |
| 1978 | 7596.7 | 25119.6 | 0.13 | 7737.8 | 25101.7 | 0.13 |
| 1979 | 8022.1 | 16014.8 | 0.18 ! | 8144.8 | 15904.7 | 0.17 |
| 1980 | 8640.4 | 28858.8 | 0.14 | 8744.0 | 28779.2 | 0.14 |
| 1981 | 8879.2 | 33003.1 | 0.18 | 8963.0 | 32919.1 | 0.18 |
| 1982 | 8625.0 | 28072.6 | 0.29 | 8688.6 | 28048.6 | 0.29 |
| 1983 | 9085.6 | 8958.9 | 0.25 | 9132.7 | 8892.8 | 0.25 |
| 1984 | 9023.2 | 73993.1 | 0.37 | 9056.8 | 73962.4 | 0.36 |
| 1985 | 9455.6 | 67944.0 | 0.35 | 9477.9 | 68002.8 | 0.34 |
| 1986 | 11594.7 | 13740.7 | 0.30 | 11611.7 | 13675.1 | 0.30 |
| 1987 | 13742.6 | 14412.3 | 0.38 | 13757.5 | 14486.3 | 0.37 |
| 1988 | 14050.9 | 23201.4 | 0.49 | 14063.4 | 23179.7 | 0.48 |
| 1989 | 13012.0 | 16807.5 | 0.49 | 13024.4 | 17001.7 | 0.48 |
| 1990 | 11831.6 | 39265.4 | 0.48 | 11846.3 | 39197.9 | 0.48 |
| 1991 | 10714.0 | 26152.1 | 0.48 | 10733.5 | 26255.8 | 0.48 |
| 1992 | 9922.8 | 11647.8 | 0.53 | 9945.5 | 11654.0 | 0.53 |
| 1993 | 9308.7 | 28829.8 | 0.54 | 9333.7 | 28846.9 | 0.53 |
| 1994 | 7843.7 | 16163.1 | 0.67 | 7869.4 | 16143.4 | 0.67 |
| 1995 | 5816.0 | 19783.2 | 0.85 | 5840.9 | 19840.6 | 0.84 |
| 1996 | 4158.6 | 23518.0 | 0.82 | 4182.4 | 23406.1 | 0.82 |
| 1997 | 2962.6 | 33925.9 | 0.78 | 2985.9 | 34224.0 | 0.78 |
| 1998 | 3160.1 | 22154.8 | 0.50 | 3183.2 | 21515.1 | 0.49 |
| 1999 | 4115.8 | 36747.5 | 0.28 | 4138.9 | 37093.3 | 0.28 |
| 2000 | 5236.3 | 25201.1 | 0.23 | 5249.6 | 24863.0 | 0.23 |
| 2001 | 5979.7 | 23098.6 | 0.31 | 5987.5 | 23234.3 | 0.30 |
| 2002 | 6671.1 | 12542.2 | 0.30 | 6674.4 | 12502.5 | 0.30 |
| 2003 | 6959.9 | 5396.1 | 0.31 ! | 6962.2 | 5359.8 | 0.31 |
| 2004 | 6572.2 | 11621.0 | 0.33 ! | 6574.2 | 11653.0 | 0.32 |
| 2005 | 5806.2 | 2403.0 | 0.32 | 5808.4 | 2389.4 | 0.32 |
| 2006 | 4773.1 | 3899.3 | 0.36 | 4775.8 | 3887.4 | 0.36 |
| 2007 | 3499.0 | 10293.8 | 0.46 | 3501.9 | 10151.9 | 0.46 |
| 2008 | 2575.6 | 16245.2 | 0.48 | 2575.1 | 16046.8 | 0.47 |
| 2009 | 2007.9 | 5860.2 | 0.60 | 1998.0 | 5525.0 | 0.60 |
| 2010 | 2114.6 | 10230.3 | 0.49 | 2088.0 | 9801.2 | 0.48 |
| 2011 | 2498.6 | 4874.9 | 0.25 | 2446.6 | 4468.2 | 0.25 |
| 2012 | 2898.5 | 7233.3 | 0.14 | 2814.5 | 6776.7 | 0.14 |
| 2013 | 3255.4 | 10281.2 | 0.12 | 3135.8 | 9980.7 | 0.12 |
| 2014 | 3652.4 | 12507.7 | 0.11 | 3497.7 | 12370.9 | 0.11 |

Table 18. Biological References Points of Jack mackerel, based on Model-1, $h=0.8$ and $h=0.65$. Fmsy $=$ fishing mortality at the maximum sustainable yield (MSY), Bmsy= spawning biomass at the MSY, Blim= spawning biomass limit, Bo=virginal spawning biomass, $B m s y / B o=$ depletion level of the spawning biomass.

| BRP | Model-1, $h=0.8$ | Model $-1, h=0.65$ |
| :--- | :--- | :--- |
| Fmsy | $0.221($ year 1 - $)$ | $0.152\left(\right.$ year $\left.{ }^{-1}\right)$ |
| Bmsy | 5.8 million t | 6.9 million t |
| Blim | 2.9 million t | 3.4 million t |
| Bmsy/Bo | $32 \%$ | $35 \%$ |

Table 19. Summary of the results of the short (2016), medium (2024) and long term (2034) predictions of the spawning biomass (B) of jack mackerel and catches 2016 and 2017 for each scenario of steepness ( $h$ ) and recruitment. The risk is presented in term of the probability that the spawning biomass in 2016 (B2016), 2024 (B2024) and 2034 (B2034) is lower or equal than $80 \%$ Bmsy (or 0.8 Bmsy), and higher than Bmsy. $0 \% * \mathrm{~F}=\mathrm{F}=0 ; 50 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $50 \% ; 75 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is reduced in a $75 \%$; $100 \% * \mathrm{~F}=\mathrm{F}$ is equal to the value in 2014. $125 \% * \mathrm{~F}=\mathrm{F}$ in 2014 is amplified in a $25 \%$.


| Steepness $h=0.8$; recruitment 1970-2012 |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 0\% * F | 5,355 | 0.182 | 0.218 | 12,918 | 0.000 | 0.999 | 17,263 | 0.000 | 1.000 | 0 | 0 |
| 50\% * F | 4,981 | 0.352 | 0.091 | 9,593 | 0.006 | 0.971 | 11,314 | 0.003 | 0.989 | 367.740 | 441.922 |
| 75\% * F | 4,811 | 0.447 | 0.056 | 8,503 | 0.019 | 0.923 | 9,672 | 0.010 | 0.962 | 525.908 | 612.902 |
| 100\% * F | 4,652 | 0.540 | 0.033 | 7,645 | 0.044 | 0.846 | 8,472 | 0.026 | 0.908 | 669.107 | 757.345 |
| 125\% * F | 4,502 | 0.627 | 0.019 | 6,954 | 0.084 | 0.743 | 7,558 | 0.056 | 0.824 | 798.761 | 879.359 |
| Steepness $\boldsymbol{h}=0.8$; recruitment 2000-2012 |  |  |  |  |  |  |  |  |  |  |  |
| 0\% * F | 4,840 | 0.426 | 0.051 | 8,127 | 0.008 | 0.942 | 8,218 | 0.025 | 0.903 | 0 | 0 |
| 50\% * F | 4,474 | 0.650 | 0.012 | 5,449 | 0.264 | 0.348 | 5,101 | 0.383 | 0.266 | 357.026 | 425.002 |
| 75\% * F | 4,309 | 0.743 | 0.006 | 4,643 | 0.529 | 0.118 | 4,301 | 0.645 | 0.078 | 508.921 | 586.622 |
| 100\% * F | 4,155 | 0.817 | 0.003 | 4,041 | 0.756 | 0.029 | 3,731 | 0.836 | 0.016 | 645.436 | 721.604 |
| 125\% * F | 4,010 | 0.875 | 0.001 | 3,579 | 0.896 | 0.005 | 3,304 | 0.938 | 0.002 | 768.116 | 834.299 |


| Steepness $\boldsymbol{h}=\mathbf{0 . 6 5}$; recruitment 1970-2012 |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 0\% * F | 5,111 | 0.426 | 0.122 | 12,084 | 0.008 | 0.998 | 17,422 | 0.025 | 1.000 | 0 | 0 |
| 50\% * F | 4,743 | 0.650 | 0.041 | 8,760 | 0.264 | 0.937 | 11,025 | 0.383 | 0.981 | 360.348 | 430.353 |
| 75\% * F | 4,576 | 0.743 | 0.023 | 7,664 | 0.529 | 0.845 | 9,229 | 0.645 | 0.935 | 514.917 | 596.196 |
| 100\% * F | 4,419 | 0.817 | 0.012 | 6,799 | 0.756 | 0.710 | 7,907 | 0.836 | 0.845 | 654.591 | 735.901 |
| 125\% * F | 4,272 | 0.875 | 0.006 | 6,101 | 0.896 | 0.553 | 6,895 | 0.938 | 0.710 | 780.798 | 853.543 |
| Steepness $h=0.65$; recruitment 2000-2012 |  |  |  |  |  |  |  |  |  |  |  |
| 0\% * F | 4,731 | 0.426 | 0.033 | 7,909 | 0.008 | 0.925 | 8,207 | 0.025 | 0.894 | 0 | 0 |
| 50\% * F | 4,367 | 0.650 | 0.007 | 5,231 | 0.264 | 0.277 | 4,902 | 0.383 | 0.219 | 353.668 | 419.744 |
| 75\% * F | 4,204 | 0.743 | 0.003 | 4,420 | 0.529 | 0.076 | 4,041 | 0.645 | 0.048 | 503.893 | 578.975 |
| 100\% * F | 4,050 | 0.817 | 0.001 | 3,812 | 0.756 | 0.014 | 3,424 | 0.836 | 0.006 | 638.756 | 711.730 |
| 125\% * F | 3,907 | 0.875 | 0.001 | 3,344 | 0.896 | 0.002 | 2,961 | 0.938 | 0.000 | 759.807 | 822.356 |

## ANNEX 3: STOCK ASSESSMENT MODEL

Table 1. Population and fishery dynamics equations. Year index $i=\{1970, . ., 2014\}$, age index: $j=\left\{1,12^{+}\right\}$, length index: $l=\{10,11, \ldots, 50\}$.

| Equation | Symbol/Constrains | Description |
| :---: | :---: | :---: |
| $N_{i, j=1}=e^{\mu_{R}+\varepsilon_{i}}$ | $\varepsilon_{i}, \sum_{i=1958}^{2014} \varepsilon_{i}=0$ | Year effect and individuals at age 1 and $i=1958, \ldots, 2014$ |
|  | $\mu_{R}$ | Mean survival in recruitment effect |
| $N_{1970, j}=e^{\mu_{R}+\varepsilon_{1970}}$ | $j=1$ | Initial numbers at age |
| $N_{1970, j}=e^{\mu_{R}+\varepsilon_{1971-j}} \prod_{j=1}^{j} e^{-M}$ | $1<j<11$ |  |
| $N_{1970,12}=N_{1970,11}\left(1-e^{-M}\right)^{-1}$ | $j=12+$ |  |
| $N_{i, 1}=e^{\mu_{R}+\varepsilon_{i}}$ | $j=1$ | Years $i>1970$ |
| $N_{i, j}=N_{i-1, j-1} e^{-z_{i-1, j-1}}$ |  |  |
| $N_{i, 11^{+}}=N_{i-1,11} e^{-Z_{i-1,10}}+N_{i-1,12} e^{-Z_{i-1,11}}$ | $1<j<11$ |  |
| $Z_{i, j}=\sum_{f} F_{i, j}^{f}+M$ |  | Total mortality |
| M | Fixed | Natural mortality |
| $F_{i, j}^{f}=e^{\mu^{f}+\eta_{j}^{f}+\phi_{i}}$ |  | Instantaneous Fishing mortality |
| $\mu^{t}$ |  | Mean fishing effect |
| $\eta_{j}^{f}, \sum_{j=1958}^{2014} \eta_{j}^{f}=0$ | $\begin{aligned} & S_{i j}^{f}=e^{\eta_{j}^{f}} ; j \leq \text { maxage } \\ & S_{i j}^{f}=e^{\eta_{\text {maxage }}^{f}} ; j>\text { maxage } \end{aligned}$ | Age effect of fishing (regularized). In year time variation allowed |
| $\eta_{i, j}^{f}=\eta_{i-1, j}^{f}$ | $i \neq$ change year | Years where selectivity is constant over time |
| $\phi_{i}, \sum_{i=1970}^{2014} \phi_{i}=0$ |  | Annual effect of fishing mortality in year $i$ |
| $B_{i}=\sum_{j=2}^{12} N_{i j} e^{-\frac{10,5}{12} z_{i j}} W_{i j} p_{j}$ | $B_{i}$ | Spawning biomass (spawning occurs at mid of November) |
|  | $p_{j}$ = proportion of females mature at age $j$ $W_{i j}=$ mean weight in the year $i$ and age $j$ |  |
| $\tilde{R}_{i, j=1}=\frac{\alpha \beta}{\beta+B_{j-1}}$ | $\widetilde{R}$ | Recruitment at age $j=1$ (Beverton and Holt equation) |
| $\alpha=\frac{4 h R_{0}}{5 h-1} ; \beta=\frac{B_{0}(1-h)}{5 h-1}$ | $R_{0}$ | Unfished recruitment |
| $h$ | Fixed | Steepness |
| $\begin{aligned} & B_{0}=R_{0} \varphi \\ & \varphi=\sum_{j=1}^{12} e^{-M(j-1)} W_{j} p_{j}+\frac{e^{-12 M} W_{12} p_{12}}{1-e^{-M}} \end{aligned}$ |  | Unfished biomass |

Table 2. Observations models for the survey or CPUE indexes, total catch, and proportion of individuals caught at age or length by survey and the fleets.

| Equation | Symbol/Constrains | Description |
| :---: | :---: | :---: |
| $\hat{I}_{i}^{s}=q^{s} \sum_{j=1}^{12} N_{i, j} W_{i, j} S_{j}^{s} e^{-\Delta^{s} Z_{i, j}}$ | $\overline{\hat{I}_{i}^{s}}$ <br> $q^{s}=$ survey catchability coefficient | Abundance index ( $\hat{I}_{i}^{s}$ ) by year $i$ and survey $s . \Delta^{s}$ represents the fraction of the year when the survey takes place. $s$ represents acoustic biomass, DEPM biomass or CPUE |
| $q_{i}^{s}=e^{\mu^{s}}$ | $\mu^{s}, \mu^{f}$ | Index catchability $q$, of the survey $s$ or fleet $f$ Mean effect |
| $\begin{aligned} & S_{j}^{S}=e^{\eta_{j}^{s}} ; j \leq \text { maxage } \\ & S_{j}^{S}=e^{\eta_{\text {maxage }}^{s}} ; j>\text { maxage } \end{aligned}$ | $\eta_{j}^{s}, \sum_{j=1958}^{2014} \eta_{j}^{s}=0$ | Age effect |
| $\begin{aligned} & \hat{C}_{i, j}^{f}=T\left[N_{i, j} \frac{F_{i, j}}{Z_{i, j}}\left(1-e^{-Z_{i, j}}\right)\right] \\ & \hat{Y}_{i}^{f}=\sum_{j=1}^{12} \hat{C}_{i, j}^{f} w_{i, j}^{f} \end{aligned}$ | $\begin{aligned} & \hat{C}_{i, j,}^{f}, \hat{Y}_{i}^{f} \\ & \mathrm{~T}=\text { ageing error } \\ & \text { matrix } \end{aligned}$ | Catch at the year $i$, and age $j$ and the fleet $f$ <br> Total Catch biomass by year $i$ |
| $\hat{C}_{i, l}=\Gamma_{l, j} \hat{C}_{i, j}$ | $\hat{C}_{i, l}$ | Catch at the year $i$, and length $l$. |
| $\Gamma_{l, j}=\int_{j}^{j+1} e^{-\frac{1}{2 \sigma_{j}^{2}}\left(l-L_{j}\right)^{2}} d l$ |  | $\Gamma_{l, j}$ is the proportion of length at age to transform from age to length. |
| $L_{j}=L\left(\begin{array}{ll}1 & e^{k}\end{array}\right)+e^{k} L_{j 1}$ | $\sigma_{j}^{2}$ <br> $L_{\infty}$ asymptotic length $L_{j}$ mean length at age | Variance of the length at age $j$ von-Bertalanffy mean length at age |
| ${ }_{j}=c v L_{j}$ | $c v=$ coefficient of variation of length at age |  |
| $p_{i, j}^{f}=\hat{C}_{i, j}^{f} / \sum_{j} \hat{C}_{i, j}^{f}$ | $P_{i j}, \sum_{j=1}^{12} P_{i j}=1.0$ | Proportion of the individuals caught in year $i$, at age $j$ by the fleet $f$. |
| $p_{i, j}^{S}=N_{i, j} S_{j}^{S} e^{-\Delta^{S} Z_{i, j}} / \sum_{j} N_{i, j} S_{j}^{S} e^{-\Delta^{S} Z_{i, j}}$ |  | Proportion of the individuals caught in year $i$, at age $j$ by the survey $s$ at the time of year $\Delta^{s}$ |
| $P_{i, l}=\frac{C_{i, l}}{\sum_{l=10}^{50} C_{i, l}}$ | $P_{i, l}, \sum_{l=10}^{50} P_{i, l}=1.0$ | Proportion at length $l$, in year $i$ |

Table 3. Vector of estimated parameters, likelihood component of each data set, prior and objective function.

| Likelihood/penalty component | Equations/ parameters | Description |
| :---: | :---: | :---: |
| Estimated parameters | $\phi_{i}, R_{0}, \varepsilon_{i}, \mu^{f}, \mu^{s}, \eta_{j}^{s}, \eta$ |  |
| Surveys and CPUE indexes | $L_{1}=0.5 \sum_{s} \frac{1}{c v_{s}^{2}} \sum_{j} \log \left(\frac{I_{j}}{\hat{I}_{j}}\right)^{2}$ | Surveys and CPUE indexes. $c v_{s}$ is the coefficient of variation |
| Prior for selectivities | $L_{2}=\sum_{m} \lambda_{2}^{m} \sum_{j=1}^{12}\left(\eta_{j+2}^{m}+\eta_{j}^{m}-\eta_{j+1}^{m}\right)^{2}$ | Smoothness (second differencing), <br> Note: $m=\{s$, or $f\}$ for survey and fishery selectivity. $\lambda_{2}$ is the smoothness parameter for selectivity |
| Prior for recruitment | $L_{3}=\lambda_{3} \sum_{j=1958}^{2013} \varepsilon^{2}{ }_{j}$ | Influences estimates where data are lacking (e.g., if no signal of recruitment strength is available, then the recruitment estimate will converge to median value). $\quad \lambda_{3}$ is the smoothness parameter for recruitment |
| Catch biomass likehood | $L_{4}=0.5 \sum_{f} \frac{1}{C v_{f}^{2}} \sum_{j=1970}^{2014} \log \left(\frac{C^{f}{ }_{j}}{\hat{C}_{j}}\right)^{2}$ | Fit to catch biomass in each year. $c v_{f}$ is the coefficient of variation of catches. |
| Proportion at age/length likelihood | $L_{5}=-\sum_{v, i, j} n^{v} P_{i, j / l}^{v} \log \left(\hat{P}_{i, j / l}^{v}\right)$ | $v=\{s, f\}$ for survey and fishery age composition observations. $P_{i, j / l}$ are the catch-at-age/length proportions $n$ effective sample size |
| Fishing mortality constrain | $F$ values constrained between 0 and 5 |  |
| Recruitment constrains | $L_{6}=\frac{0.5}{c v_{r}^{2}} \sum_{j=1970}^{2011} \log \left(\frac{N_{i, j=1}}{\tilde{R}_{i}}\right)^{2}$ | Conditioning on stock-recruitment curve over period 1977-2011. $c v_{r}$ is the coefficient of variation of recruitments. |
| Priors | $R_{0}$ non-informative | (Explored alternative values of $\sigma_{R}^{2}$ ) |
| Overall objective function to be minimized. | $\dot{L}=\sum_{k} L_{k}$ | $k=\{1, . .6\}$ |

Table 4. Data weighting. Coefficients of variation and sample sizes used in the likelihood functions.

| Abundance index | Model-1, <br> $h=0.8(c v)$ | Model-1, <br> $h=0.65(c v)$ |
| :--- | :--- | :--- |
| Acoustic North Chile | 0.20 | 0.20 |
| Acoustic Central south Chile | 0.50 | 0.50 |
| CPUE - Chile | 0.15 | 0.15 |
| DEPM - Chile | 0.50 | 0.50 |
| Acoustic-Peru | 0.20 | 0.20 |
| CPUE - Peru | 0.20 | 0.20 |
| CPUE- China | 0.20 | 0.20 |
| CPUE-EU | 0.20 | 0.20 |
| CPUE- ex USSR | 0.40 | 0.40 |
| Smoothness of selectivities (fleets) | $\lambda$ | $\lambda$ |
| North Chile | 1 | 1 |
| Central south Chile | 25 | 25 |
| Far North | 12.5 | 12.5 |
| Offshore Trawl | 12.5 | 12.5 |
| Proportion at age likelihood (fleets) | n | n |
| North Chile | 20 | 20 |
| Central south Chile | 50 | 50 |
| Far North | 30 | 30 |
| Offshore Trawl | 30 | 30 |

## ANNEX 4: Using population dynamic theory for understanding the interaction between jack mackerel (Trachurus murphyi) and fishery dynamics

Two kinds of models have been traditionally used in fishery management: 1) those based on the concepts of stock productivity, surplus production, and maximum sustainable yield (MSY) (Quinn and Deriso 1999) and 2) those models based in the spawner-recruit relationships (stock-recruit models) developed from ecological assumptions affecting the reproductive processes (Ricker 1954, Beverton and Holt 1957). The basic assumption of both surplus production and stock recruitment models is the existence of an equilibrium population size (or biomass) of unexploited fishes. However, equilibrium or near equilibrium dynamics can be caused only by the presence of negative feedback processes (e.g., competition) and limiting factors (Berryman 1999). For its part, the ecosystem based fishery management (EBFM) approach assumes that exploited populations are embedded in complex ecosystems involving large numbers of interactions. From this perspective, the sustainable management of fisheries can only be achieved if models used to harvest fish populations are able to include, or account for, other ecosystems components.

Here, we used another approach to deciphering the potential interactions between the fishery fleet and the stock of jack mackerel at southeastern Pacific Ocean. First, we applied the theory of population dynamics (Royama 1992, Berryman 1999, Turchin 2003 and Ginzburg and Colyvan 2004), in order to shed light on issues of basic importance in managing the harvest of the jack mackerel stock. In particular, we want to emphasize the fundamentally important role of population theory in managing fisheries, despite the attitude of many biologists that there are no laws or reasonable theories for explaining natural systems (in particular marine ecosystems) or the idea that marine ecosystems are too complex to be described by simple models. The objective is to demonstrate that ecological theory and simple models can be useful for understanding and PREDICTING the dynamics of fish populations.

## Population Dynamic Theory

Following the work of Royama (1977, 1992), other authors have proposed that there are a few simple principles that can explain much (or even most) of the apparent complexity observed in the fluctuations of natural populations (Berryman 1999, Turchin 2003, Ginzburg and Colyvan 2004). Although the experts often differ on the importance of the basic principles of population dynamics and how they should be formulated, we used these principles (especially as formulated by Berryman 1999) to analyze how the jack mackerel dynamics and the fisheries are managed.

Classical single-species fisheries models rely, in principle, on our understanding of competition as a basic component of processes that lead to population biomass showing patterns of equilibrium or near equilibrium. Predation also contributes to these patterns, and fishing (predation by humans) will reduce population size towards lower numbers where individuals are faced with more abundant resources (per capita) resulting in increased recruitment rates. Many have criticized management based on the assumption that these principles are sufficient for realistic decision-making. One of the most common criticisms relies on empirical evidence that marine ecosystems show a great deal of variability, often explained as the effects of stochastic processes, oceanographic variability, and inter-specific interactions (Spencer and Collie 1997a).

The underlying reasoning behind these criticisms is that marine ecosystems are too complex to be managed by single-stock fishery models (Pikitich et al. 2004, Frid et al. 2006). Although it has to be recognized that single stock fishery models are a simplistic metaphor of nature, the inclusion of more parameters and variables always fails to achieve a complete understanding of the causes of fish population dynamics. In fact, however, it is highly likely that many fish stocks are governed through dynamics in which simple first-order dynamics and limited by resource availability (food, refuge, etc) count as primary factors. Some factors are clearly much more influential than others. We think that the principles behind population dynamic can never be ignored, either as they are developed to better understand the underlying causes of population dynamics, or (and perhaps more importantly) to better understand their role in implementing adequate management options to assure long term sustainability.

Fishing is, in most cases, the most important extrinsic force acting on fish populations (Jackson et al. 2001), often orders of magnitude larger than that of other predators (Fowler and McCluskey ). Therefore, to understand and manage fisheries we need to understand the dynamic of the fishery fleet. For example, if fishing effort is constant in time, and the fleet behaves as a generalist or a highly mobile predator (capable of aggregating in high prey density areas), a potential consequence is to diminish prey (fish) populations toward low levels and cause a new dynamic equilibrium point (Holling 1965, Morris 1963, Berryman 1999). Under this scenario, fish populations show a tendency to be stabilized by generalist predators at low densities and regulated by enemy free space competition (Berryman 1999).

When fishing effort increases in response to economic forces (or other factors such as highly successful previous harvests), a delayed feedback can be created which leads fish population dynamics toward regular and large amplitude cycles (predator-prey cycles). It is interesting to note that predator-prey theory has been used vary rarely in fisheries, despite the classic predator-prey model developed by Vito Volterra as an attempt to give explanation to fluctuations in the Adriatic fisheries after First World War (Kingsland 1995). Cycles in marine fish or invertebrate populations can be the consequence of the destabilized forces imposed by the economic inertia behind fisheries (Berryman 1991); regular cycles in the numerical fluctuations of several exploited species have been documented (Bostford 1986, Spencer \& Collie 1997a, Higgins et al.
1997). In these cases, the fishing effort and the fish stock are mutually connected by a feedback loop. Owing to this feedback, an integral management strategy is necessary: to reduce the amplitude of fish oscillations requires reducing the fishing effort or its variability (or both). In sum, we analyze the dynamics of the adult spawning stock of jack mackerel, recruitment, fishery effort and climate under the conceptual framework of population dynamic theory.

## Statistical Models

Population dynamics of jack mackerel is the result of the combined effects of feedback structure (ecological interactions within and between populations), limiting factors, climatic influences, and stochastic forces. To understand how these factors may determine jack mackerel population fluctuations, we model both system-intrinsic processes (both within the population and between various trophic levels) and exogenous influences, as a general model based on the $R$-function (Berryman 1999). The $R$-function represents the realized per capita population growth rates that represent the processes of individual survival and reproduction (Berryman 1999). Defining $R_{\mathrm{t}}=\log \left(\mathrm{N}_{\mathrm{t}}\right)-\log \left(\mathrm{N}_{\mathrm{t}-1}\right)$, we can express the $R$-function (sensu Berryman 1999) as:

$$
\begin{equation*}
R_{t}=\ln \left(\frac{N_{t}}{N_{t-1}}\right)=f\left(N_{t-1}, N_{t-2}, \cdots, N_{t-i}, C_{t-i}, \varepsilon_{t}\right) \tag{1}
\end{equation*}
$$

Here $N_{t-i}$ is the adult spawning biomass (recruitment and Fishing effort) at different time lags; $C_{t-i}$ is exogenous effects; and $\varepsilon_{t}$ is a random normally distributed variable. This model represents the basic feedback structure and integrates the stochastic and climatic forces that drive population dynamics in nature. Our first step was to estimate the order of the dynamical processes (Royama 1977), that is how many time lags, $N_{t-i}$, should be included in the model for representing the feedback structure. To estimate the order of the process we used the partial rate correlation $(P R C F(i))$ between $R$ and $\ln N_{t-i}=X_{t-i}$ after the effects of shorter lags have been removed. We write (1) in logarithmic form to calculate the partial correlations.

$$
\begin{equation*}
R_{t}=\ln \left(\frac{N_{t}}{N_{t-1}}\right)=A+B_{1} \cdot X_{t-1}+B_{2} \cdot X_{t-2}+\varepsilon_{t} \tag{2}
\end{equation*}
$$

Where $R$, the realized per-capita rate of change, is calculated from the data, we fitted a multiple regression between the per capita growth rates and lagged population density to estimate the $P R C F_{t-d}$ coefficients at each lag ( $B_{i}, i=1,2, . .3$ ), for statistical convenience we assumed a linear relationship between $R$ and $X_{t-i}$ (Royama 1977).
The dynamics of spawning stock was better explained by a second order component $[\operatorname{PRCF}(2)]$, which suggest a second-order feedback system dominating the adult fish dynamics (Figure 1). A first-order negative feedback [PRCF (1)] was the most important component of recruitment dynamics (Figure 1). This result suggest a simple feedback structure, and low order dynamics of the recruitment process. The fishing effort dynamics
appears to be described by a third-order component $[\mathrm{PRCF}(3)]$ suggesting a complex feedback structure (Figure 1).

## The statistical models

Our starting point in the analyses was to model jack mackerel spawning stock, recruitment and fishing effort tropical using simple statistical models. The PRCF analysis suggests that jack mackerel spawning biomass and fleet dynamics could be dynamically connected by higher order processes. Our starting point for a simple statistical model is:

$$
\begin{align*}
& R S_{t}=f\left(B_{t}, E_{t}, \varepsilon_{t}\right)  \tag{3}\\
& R E_{t}=g\left(E_{t}, B_{t}, \epsilon_{t}\right)
\end{align*}
$$

Where $R S_{t}$ is the population growth rate of spawning biomass at time $\mathrm{t}, B_{t}$ is the spawning biomass and $E_{t}$ is the fishing effort and $\varepsilon_{t}$ is a random normal variable. On the other hand, $R E_{t}$ is the rate of change of fishing effort, $\varepsilon_{t}$ is another random variable an $f$ and $g$ are simple linear functions. In addition recruitment dynamics appears to be a simple first order dynamic process; hence, we used a simple statistical representation;
$R R_{t}=h\left(R C_{t}, C_{t}, \varepsilon_{t}\right)$
(4),

Where $R R_{t}$ is the realized growth rate of recruitment, $R C_{t}$ is the recruitment abundances; $C_{t}$ represents an exogenous climatic variable (e.g. El Niño) and $\varepsilon_{t}$ a random normal variable.

## RESULTS

Population dynamics of the spawning stock of jack mackerel and fleet dynamics are characterized by large and regular oscillations (Figure 2), which are typical of secondorder processes. In fact, both systems appear to be linked, the rate of growth of spawning jack mackerels seem to be a negative linear function of the logarithm of fishing effort and a positive linear function of the logarithm of recruitment $\left(R^{2}=38 \%, F_{2,28}=8.59, p=\right.$ 0.0012 ). On the other hand, the rate of change of fishing effort appears to be a positive linear function of the logarithm of spawning biomass of jack mackerel $\left(R^{2}=30 \%, F_{1,29}=\right.$ $12.40, p=0.0014$ ). The phase diagram between spawning biomass of jack mackerel and fishing effort suggest that both are mutually influenced and are connected in a predator/prey like dynamics (Figure 3). Finally, the recruitment dynamics appears to be simpler, the recruitment growth rates are negatively influenced by the logarithm of recruitment biomass (first-order negative feedback), and positively affected by the El Niño anomalies suggesting the importance of endogenous and exogenous effects ( $R^{2}=$ $44 \%, F_{3,27}=7.09, p=0.0012$ ).

## DISCUSSION

Fishery science has been developing around a variety of basic concepts and ideas since the 1950s (Quinn \& Collie 2005). One of the most important assumptions in this history is that fish stocks are near the "equilibrium" of the system and the only effect of fishing mortality is to reduce the abundance of the harvested population. Although fisheries scientist have long been aware of the multiple effects of fishing in marine communities (Pitcher 2001), the problem of dealing with this issue, using a more general and theoretical perspective, has been absent. During this time, the models used in fisheries research incorporated varying degrees of complexity by adding explanatory variables and exploring alternative sources of variability (see, Quinn \& Collie 2005 for a review). Despite these advances in accounting for environmental factors, most of the wellaccepted general ideas and concepts derived from straight-forward population dynamics theory are ignored in applied fishery models. For example, although the concept of predator functional responses is quite well developed and used in the conventional management (control) of pest species, it has not been incorporated in the theoretical toolbox of fisheries scientists (Holling 1965, Morris 1963, Royama 1977, 1992, Berryman 1999). Therefore, instead of trying to compare single species models with ecosystem models (as currently being proposed as a basis for Ecosystem Based Fishery Management), we call for an acceptance and understanding of the magnitude of our ignorance regarding simple population dynamics, especially insofar as population dynamics is brought to bear in management.
There is an important need for simpler and theoretically based models as proper diagnostic tools for analyzing fish population fluctuations. We think that the theory behind population dynamics offers the proper conceptual background to develop simple models for understanding and predicting the dynamics of fish populations.
The present analysis suggests that jack mackerel and fishery dynamics are linked in a predator/prey like system of mutual causal second-order loop. This hypothesis represents a new view of how this fish population is responded to the fishery and put new challenges to fish management. In particular, if a cyclic dynamics is underlying the mutual connection between jack mackerels and the fleet it could be needed an integral approach similar to what is used in pest control (Berryman 1999). On the other hand, recruitment dynamics appears to be simpler, basically is the interaction among density-dependent process and El Niño variability (Figure 4).
A final comment: as substantiated elsewhere (Berryman 1991, Berryman and Lima 2006), we think that a proper analysis of empirical information is preferable to ecological modeling to inform the management of harvesting to achieve sustainability in fisheries. It is the only way to deduce the proper feedback structure to be responsive to change. It provides important insight to the importance of exogenous effects (e.g., climate), and how fishing influences fish stocks. Such an approach is essential for the analysis and $a$
posteriori modeling of observed data to estimate model parameter and establish causal connections between variables Berryman (1991).

In summary, it is fundamentally important that we avoid management that focuses on ecosystems to the exclusion of populations and their dynamics: an essential part of what is brought to management must be our understanding of population dynamics. The science of population dynamics provides crucial insight to, and understanding of, populations that must be accounted for in management of fisheries.

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## FIGURES



Fig. 1. Partial rate correlation function (PRCF) of the growth rate of spawning biomass (top-left), recruitment (top-right), and fishing effort (bottom-left).


Fig. 2. Spawning biomass, recruitment, fishing effort of jack mackerel in the South Eastern Pacific. Environmental index El Niño 1.2 sea surface temperature.


Fig. 3. Phase diagrams between spawning biomass of jack mackerel and fishing effort in the South Eastern Pacific. Both variable are mutually influenced and connected in a predator/prey like dynamics.


Fig. 4. Interactions between recruitment and spawning biomass of jack mackerel and the fishing and El Niño.

