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# Quota Options for the Red Sea Urchin Fishery in British Columbia for Fishing Season 2000/2001 

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#### Abstract

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#### Abstract

Annual landings of red sea urchin (Strongylocentrotus franciscanus) increased rapidly in the early 1980s for the south coast of British Columbia (B.C.) and in the late 1980s for the north coast, but subsequently were reduced and stabilized by arbitrary quotas. Coastwide landings were 5091 t valued at $\$ 7.4 \mathrm{M}$ (Cdn.), with 110 licenses issued during the 1998/99 fishing season. Bed areas were obtained by digitizing locations on charts indicated in harvest logbooks. Analyses of recent surveys and review of published survey reports provided estimates of mean density and weights allowing preliminary estimates of red sea urchin biomass in B.C. Based on recent published reports and preliminary estimates in B.C. natural mortality rates of red sea urchins was assumed for fishery management purposes to be between 0.05 and 0.15 . Since industry and managers requested an evaluation of the implications of reducing the size limit from present legal size limit ( $\geq 100 \mathrm{~mm}$ test diameter, TD), estimated biomass and quota options were calculated and discussed for a variety of size limits of $\geq 90, \geq 100,100-140,90-120$, and $90-130 \mathrm{~mm}$ TD for red sea urchins in B.C. Further surveys for red sea urchin density are required, especially in some areas of the south coast of B.C. where surveys are $>10$ years old, and in areas in the North Coast that have been heavily fished. Reliable biomass estimates and more accurate estimates of bed areas, natural mortality and recruitment rates for red sea urchins in most areas of B.C. are required to manage this fishery effectively.


## RÉSUMÉ

Les débarquements annuels d'oursin rouge (Strongylocentrotus franciscanus) en provenance de la côte sud de la Colombie-Britannique (C.-B.) ont augmenté rapidement au début des années 1980 et vers la fin de la décennie pour ceux en provenance de la côte nord. Ils ont ensuite été réduits et stabilisés par l'imposition de quotas. À la grandeur de la côte, les débarquements de la saison de pêche 1998-1999, pour laquelle 110 permis ont été délivrés, ont atteint 5091 t (valeur de $7,4 \mathrm{M} \$$ can.). La superficie des lits a été obtenue en numérisant sur des cartes les lieux de capture indiqués dans les registres de récolte. L'analyse des relevés récents et l'examen des rapports de relevés publiés ont servi à estimer la densité et le poids moyen et ainsi à obtenir des estimations préliminaires de la biomasse de l'oursin rouge en C.-B. Le taux de mortalité naturelle, aux fins de la gestion de la pêche, a été estimé entre 0,05 et 0,15 à partir des rapports récents publiés et d'estimations provisoires. L'industrie et les gestionnaires ont demandé une évaluation des incidences d'une réduction de la taille limite légale actuelle ( $\geq 100 \mathrm{~mm}$ de diamètre du test, DT) et des options de l'estimation de la biomasse et de quotas ont été calculées et discutées pour une variété des tailles limites de $\geq 90, \geq 100,100-140,90-120$ et 90-130 mm de DT, pour l'oursin rouge de la C.-B. Il faudra effectuer d'autres relevés de la densité de l'oursin rouge notamment dans certaines zones de la côte sud de la C.-B. où les relevés datent de plus de dix ans, de même que dans certaines zones de la côte nord où la pêche a été intense. La gestion efficace de l'oursin rouge dans la plupart des zones de la C.-B. exige d'obtenir des estimations fiables de la biomasse et plus exactes de la
superficie des fonds, de la mortalité naturelle et des taux de recrutement de l'oursin rouge dans la plupart des zones de la C.-B.

## INTRODUCTION

A commercial dive fishery for the red sea urchin (Strongylocentrotus franciscanus ) started during the 1970s in British Columbia (B.C.) (Fig. 1, Table 1). Annual landings started to increase rapidly in the early 1980s for the south coast and the late 1980s for the north coast of B.C., but subsequently were reduced and stabilized by arbitrary quotas (Table 1). Coastwide landings were 5091 t valued at $\$ 7.4$ million (Cdn.) with 110 licenses issued during the 1998/99 fishing season. The history of the management of this fishery is summarized in Campbell and Harbo (1991), Heizer et al. (1997), and Rogers and Neifer (1999). A number of papers review various aspects of red sea urchin biology (Bernard and Miller 1973; Mottet 1976; Breen 1980; Sloan et al. 1987; Tegner 1989; Campbell and Harbo 1991; Botsford et al. 1993, 1994; Lai and Bradbury 1998; Ebert 1998). A few surveys to estimate standing stock of red sea urchins in B.C. during 197694 have been published (Breen et al. 1976, 1978; Adkins et al. 1981; Sloan et al. 1987; Jamieson et al. 1998a, b, c, d). Based on the results of these surveys Campbell (1998) provided quota estimates for the 1995-96 red sea urchin fishery. Campbell et al. (2000) provided quota estimates for the 1998 red sea urchin fishery based on these data and surveys during 1995-97. The results of additional surveys in 1997-99 are summarized in the present paper and Bureau et al. (2000a, b, c, d).

The B.C. coast is divided into two main regions, the 'North Coast' and the 'South Coast'. In addition, B.C. is subdivided into Pacific Fishery Management Areas (PFMA) and (PFM) sub-areas for management and economic purposes (detailed charts are not shown in this paper).

Managers have requested additional biomass estimates based on the most recent surveys and that quotas, where possible, be estimated by PFM subarea so they can be applied to the 2000/2001 red sea urchin fishery in B.C. Industry and managers have also requested an evaluation of the present legal size limit ( $\geq 100 \mathrm{~mm}$ test diameter, TD) and implications of reducing the size limit. Industry believe that reductions in the minimum legal size limit would better meet market demands for the best quality gonad (roe) product which is extracted from red sea urchins. Any changes in size limits (including minimum and maximum sizes) would require a precautionary approach to incorporate adaptive management of this valuable resource (Campbell et al. 1999).

The purpose of this paper is to (1) summarize catch trends from the sales slips and harvest logbooks, (2) summarize the density surveys conducted to date and estimate biomass of red sea urchins, (3) determine annual quotas based on several size limits options, and (4) discuss the implications of these size limit options on the red sea urchin fishery in B.C.

## METHODS

## CATCH AND Effort

Catch and effort data were summarized from sales slips and from harvest logbooks that fishers completed each day of fishing. Information from sales slips included total weight (pounds) and value (dollars) landed, commercial fishing vessel (CFV) number, date and days fished. Information from the harvest logbooks included location of bed (with diagram), date, landed weight and minutes of diving. The harvest logbooks were not completed by each vessel so the data were used as a sample (CPUE, kilograms per hour) only where both total catch ( kg ) and effort (minutes) per region were reported per diver for each area per day. Total average annual CPUE were calculated from mean daily vessel CPUE values.

## Bed Areas

Commercial bed areas of red sea urchins were indicated on charts or diagrams provided by fishers with their harvest logbooks throughout B.C. during 1982-1996. The detailed bed areas were outlined on hydrographic charts from $1-9.1 \mathrm{~m}(0-5 \mathrm{fm})$ below chart datum and were digitized and areas estimated based on sea surface area. The digitized bed areas used in the analyses for this paper are those indicated as harvested during 1982-1996. Estimation of these red urchin bed areas must be treated with caution since the beds were not measured empirically in the field, and the proportion of the suitable substrate types (e.g., boulders or flat bedrock area more suitable than sand or mud) are unknown and may differ from one area to another.

## DENSITY AND BIOMASS

Densities of red sea urchins were generally estimated within $1 \mathrm{~m}^{2}$ quadrats along randomly chosen transects. Details of survey methodology varied between surveys (Breen et al. 1976, 1978; Adkins et al. 1981; Sloan et al. 1987; Jamieson et al. 1998a, b, c, d; Bureau et al. 2000a, b, c, d). Test diameters (TD, in mm) of urchins were measured on all the surveys except that by Adkins et al. (1981). Density estimates from Adkins et al. (1981) could be biased since counts were made only at sites where there were more than 1 red sea urchin $/ \mathrm{m}^{2}$. Surveys of sea urchin density were also conducted during 1995 to 1997, using the methodology described by Jamieson and Schwarz (1998). We reanalyzed the data from the 1994-99 surveys using the following methods of estimating mean densities (the 1993 data had a different survey method (Jamieson et al. 1998a) so were used unchanged). The present data analysis methods are different compared to those used by Campbell et al. (2000) which gave slightly different overall density and biomass estimates. Campbell et al. (2000) used overall mean sizes and proportions and applied these to the density estimates for a PFMA or subarea. In contrast, the present analysis methods consider the numbers, sizes, weights and biomass of red sea urchins within each transect length and weighs this according to the total transect lengths sampled
within each PFMA or subarea. N.B. (a) the number of quadrats sampled for urchin density was usually half the potential number of quadrats (i.e. alternate quadrats were sampled) along a transect, and (b) the number of quadrats sampled for urchin sizes could be lower or equal to the quadrats sampled for density along a transect. In some areas high urchin abundance made measuring each urchin logistically unfeasible.

The estimated mean density, $\bar{d}$ (number / $\mathrm{m}^{2}$ ) or biomass density ( $\mathrm{g} / \mathrm{m}^{2}$ ), of urchins across a number of transects surveyed in a PFMA or subarea was calculated as

$$
\begin{equation*}
\bar{d}=\frac{\sum_{t}\left(d_{t}{ }^{*} L_{t}\right)}{\sum_{t} L_{t}} \tag{1}
\end{equation*}
$$

The standard error of the mean density, $\mathrm{s}_{\mathrm{d}}$, was calculated as

$$
\begin{equation*}
s_{d}=\sqrt{1-\frac{n}{T}} * \sqrt{\frac{\sum_{t}\left(d_{t} * L_{t}-\bar{d} * L_{t}\right)^{2}}{n^{*}(n-1) * \bar{L}^{2}}} \tag{2}
\end{equation*}
$$

where n is the number of transects, $\mathrm{d}_{\mathrm{t}}=\mathrm{N}_{\mathrm{t}} / \mathrm{S}_{\mathrm{t}}$ is the density at transect t , $\mathrm{S}_{\mathrm{t}}$ is the number of quadrats surveyed for density estimates in transect $\mathrm{t}, N_{t}$ is the number of red sea urchins counted for density estimates in transect $t, L_{t}$ is the length of transect t (or area in square metres since each transect was one metre wide), $\bar{L}=\frac{\sum_{t} L_{t}}{n}$ is the mean transect length (or mean area in square metres), and $T$ is the total potential number of transects that could possibly be sampled in the surveyed PFMA or subarea. The expression $\sqrt{ }(1-\mathrm{n} / \mathrm{T})$ was close to one, because the sample size n was small compared to T.

To estimate the mean densities (number $/ \mathrm{m}^{2}$ ) or biomass density $\left(\mathrm{g} / \mathrm{m}^{2}\right)$ for a specific size group $(J)$ the value $\mathrm{d}_{\mathrm{t}}$ was substituted with densities $\left(P_{t, J}\right)$ or biomass $\left(B_{t, J}\right)$, in equations 1 and 2 .

The mean density $P_{t, J}$ (number/ $\mathrm{m}^{2}$ ) of red sea urchins of size group $J$ in transect t was calculated as

$$
\begin{equation*}
P_{t, J=} \frac{\left(N_{t} * \frac{\sum_{i \in J} m_{t, i}}{M_{t}}\right)}{S_{t}} \tag{3}
\end{equation*}
$$

The mean biomass density $B_{t, J}\left(\mathrm{~g} / \mathrm{m}^{2}\right)$ of red sea urchins of size group $J$ in transect t was calculated as

$$
\begin{equation*}
B_{t, J}=\frac{\left(\frac{N_{t}}{M_{t}} * \sum_{i \in J}\left(m_{t, i} * w_{i}\right)\right)}{S_{t}} \tag{4}
\end{equation*}
$$

where $J$ is a subset of possible i values representing a range of test diameters $\left(\mathrm{TD}_{\mathrm{i}}\right)$ (e.g. $\geq$ $90, \geq 100,100-140,90-120$, or $90-130 \mathrm{~mm} \mathrm{TD}), M_{t}=\sum_{i} m_{t, i}$ the total number of red sea urchin measured for size in transect $\mathrm{t}, m_{\mathrm{t}, \mathrm{i}}$ is the number of red sea urchins measured in size range $i$ of transect t , and $w_{i}=0.0012659 * \mathrm{TD}_{\mathrm{i}}{ }^{2.7068}$ is a relationship between mean wet weight ( g ) and size i (TD in mm ) for red sea urchins (Campbell 1998).

Where there were no mean density estimates for a PFM subarea, an overall mean density for all years surveyed within the whole PFMA was used. Where there were no data for a whole PFMA then mean values of data from the nearest or adjacent PFMA were used. The accuracy of these extrapolations is unknown and will require further comparative field surveys.

Standard errors were not directly calculated for densities that included the 1993 survey data since the survey method was different from surveys of later years. However, because an approximate lower $90 \%$ confidence intervals of mean biomass density (L90CIB) was required for each PFM subarea the following method was used. A linear regression, $\mathrm{s}_{\mathrm{d}}=$ $\mathrm{b} \bar{d} / V_{\mathrm{s}}$, was used to describe the relation between the standard error $\left(\mathrm{s}_{\mathrm{d}}\right)$ and the mean biomass density ( $\bar{d}$ ) divided by the square root of the total transect length (s) used for the biomass estimates, with $b$ being a constant estimated using the least squares method. Data analysed from each red sea urchin size limit class group in all PFM subareas and years that were surveyed after $1993(\mathrm{~N}=61)$ resulted in $b$ values of $7.00\left(\mathrm{R}^{2}=0.84\right)$ for $>90$ $\mathrm{mm} \mathrm{TD} ; 7.14\left(\mathrm{R}^{2}=0.85\right)$ for $>100 \mathrm{~mm} \mathrm{TD} ; 7.40\left(\mathrm{R}^{2}=0.83\right)$ for $100-140 \mathrm{~mm} \mathrm{TD} ; 6.91$ $\left(\mathrm{R}^{2}=0.74\right)$ for $90-120 \mathrm{~mm} \mathrm{TD}$; and $6.88\left(\mathrm{R}^{2}=0.77\right)$ for $90-130 \mathrm{~mm}$ TD. The equation $\mathrm{L} 90 \mathrm{CIB}=\bar{d}(1.0-1.64 \mathrm{~b} / \sqrt{s})$ was used to calculate the approximate lower $90 \%$ confidence interval of the mean biomass density $\left(\mathrm{g} / \mathrm{m}^{2}\right)$ for each size limit class of red sea urchin and PFM subarea.

Total current biomass of red sea urchins, for each size group, for each PFM subarea was calculated as

$$
\begin{equation*}
\mathrm{B}_{\mathrm{c}}=\mathrm{A} \bar{d} \tag{5}
\end{equation*}
$$

where $\mathrm{B}_{\mathrm{c}}$ is the current average biomass $(\mathrm{g}) ; A$ is the commercial urchin bed areas (presented as hectares, but converted to $\mathrm{m}^{2}$ for biomass calculations) estimated from digitized charts and summed for each PFM subarea; $\bar{d}$ is the estimated mean biomass density $\left(\mathrm{g} / \mathrm{m}^{2}\right)$ of red sea urchins in a size group (e.g. $\geq 100 \mathrm{~mm}$ or $100-140 \mathrm{~mm}$ TD). The values for $\mathrm{B}_{\mathrm{c}}$ were subsequently converted to tonnes for presentation. Since many of the surveys were conducted $>4$ years ago and there could be considerable uncertainty in the mean biomass density estimates in some areas we have adopted the Woodby (1992) method by also including the approximate lower $90 \%$ confidence interval (L90CIB) as a reasonable alternative for the $\bar{d}$ values in the $B_{c}$ calculations.

## Natural Mortality

There are no published estimates of instantaneous natural mortality rate (M) for red sea urchins from northern B.C. Breen (1984) estimated that M ranged from 0.016 to 0.22 for red urchins from 3 sites in southern B.C. and considered a value between 0.1-0.2 to be acceptable. Woodby (1992) estimated $\mathrm{M}=0.16$ for red sea urchins from the Sitka, Alaska area. Botsford et al. (1993) estimated $\mathrm{M}=0.14$ for a population of red sea urchins in California. Lai and Bradbury (1998) estimated $M$ to be about 0.16 for red sea urchins from Washington. Based on published values Campbell (1998) assumed M to be 0.15 in calculating quotas for the 1995 red sea urchin fishery in B.C. However, all these authors considered growth rates of red sea urchins to be faster (e.g., 4-6 years to reach 100 mm TD) than that reported by Ebert (1998) who found tagged sea urchins from Washington and Oregon to take about 10 years to reach 100 mm TD and 50 years to reach 140 mm TD. Ebert (1998) calculated the mean instantaneous total mortality rate ( Z year ${ }^{-1}$ ) of red sea urchins, from a total of twelve samples collected from six locations in Oregon and Washington, to be 0.052 (min. 0.016 , max 0.133 , lower $95 \%$ confidence interval (CI) 0.028 , upper CI 0.076 ); equivalent to a mean annual survival rate of $0.949\left(\mathrm{e}^{-\mathrm{Z}}\right)$. The average mortality values reported by Ebert (1998) were generally below those previously reported in the literature. Clearly M will vary between areas and between size classes for red sea urchins in B.C. Although a similar tagging program on red sea urchins in some areas of B.C. has been conducted, further experimental work is required to estimate growth of urchins $<2$ years ( $5-30 \mathrm{~mm} \mathrm{TD}$ ). Initial analyses of the growth and mortality data indicated Z (instantaneous natural mortality rate) varied between sites from 0.03 to 0.16 (Campbell et al. unpublished data). For the purposes of the present paper, a range of M values from 0.052 to 0.150 (with an approximate mean M of 0.10 ) was considered for red sea urchins in B.C.

## Recruitment

Sloan et al. (1987) estimated recruitment to be highly variable between areas and to average about $9.5 \%$ of the total number of sea urchins in the size frequencies per area. Little is known about the stock and recruitment relations of red sea urchins in B.C. (Campbell et al. 1999).

## Quota Estimation

A conservative management approach is used to estimate quotas $(\mathrm{Q})$ for the red sea urchin fishery in B.C. A modified surplus production model is used to estimate a maximum sustainable yield (MSY) from a stock that is in the early stages of fishing (Schaefer 1954; Gulland 1971). The model assumes that the MSY occurs when the maximum sustainable fishing mortality is equal to M .

$$
\begin{equation*}
\mathrm{Q}=\mathrm{XMB} \mathrm{~B}_{\mathrm{c}} \tag{8}
\end{equation*}
$$

where $\mathrm{B}_{\mathrm{c}}$ is the current biomass, M is the instantaneous natural mortality rate and $X=\mathrm{a}$ correction factor to insure that a sustainable fishing mortality rate is well below the calculated MSY. We chose the value of $X=0.20$ in this paper as a reasonably conservative safeguard to account for errors in estimating the lower current biomass values (Caddy 1986; Garcia et al. 1989). The correction factor should provide for a conservative harvest per year in a developing fishery where little is known about the productivity of the population. Since equation 8 is derived from a Graham-Schaefer production model, recruitment is assumed to be unaltered by these low fishing levels. Although this approximation was developed for an unexploited virgin stock $\left(B_{o}\right)$ we assumed that $\mathrm{B}_{\mathrm{c}}=\mathrm{B}_{0}$.

Caution is required in the interpretation of these calculations for the quota because there are so many assumptions in the parameters used in the oversimplified model. Since many of the surveys were conducted $>4$ years ago and there could be uncertainty in the mean biomass density estimates in some areas, we have adopted the Woodby (1992) method by also including the approximate lower $90 \%$ confidence intervals as an alternative for the mean $B_{c}$ calculations.

## RESULTS AND DISCUSSION

## Catch and Effort

The number of fishing vessels peaked at 116 in 1990 (Table 1). Coastwide landings peaked in 1992 (Tables 1, 2, 3). Quotas have generally restricted landings in the South Coast since 1985 and in the North Coast since 1993 (Table 1). There was no overall general trend in annual CPUE (kilograms per diver hour) for the red sea urchin fishery in B.C. between 1983-96 (Table 1). The general lack in CPUE trends suggests that either the fishery is at an early stage of development or CPUE data for red sea urchins may not
be used to indicate fishery trends in B.C. Fishermen have increased search time for high quality urchins in response to recent changes in market demands and the implementation of an individual quota scheme. Also, fishermen may be maintaining high CPUE values, through serial depletion, by moving to unexploited sea urchin beds within a PFMA suggesting that CPUE would not decline until most legal-sized sea urchins were removed from most of the areas in the PFMA. There is a need to re-examine the distribution of effort and variability of CPUE data on a smaller spatial scale (e.g. by bed) than the PFM subarea level to determine whether CPUE is an appropriate index of red sea urchin abundance. Additional detailed work on updating the data base on bed locations and areas for the fishery during 1997-99 is underway.

## Density and Biomass

In general, there was considerable variation in the size structure and densities of local red sea urchin populations. Estimated mean densities (number $/ \mathrm{m}^{2}$ ) and mean biomass density $\left(\mathrm{g} / \mathrm{m}^{2}\right)$ by each size group of red sea urchins varied between PFM subareas and years (Tables 4,5 ). Generally, density and biomass density were highest for the widest size limit, $\geq 90 \mathrm{~mm}$ TD, and lowest for the narrowest size limit, $90-120 \mathrm{~mm}$ TD, for the size groups examined for each PFM subarea (Fig. 2, Tables 4, 5). Although mean densities for $100-140 \mathrm{~mm}$ TD were generally lower than those for $\geq 100 \mathrm{~mm}$ TD, those for $\geq 100$ and $90-130 \mathrm{~mm}$ TD were similar. In contrast there was a general lowering trend of mean biomass density from $\geq 90, \geq 100,100-140,90-130$, to $90-120 \mathrm{~mm}$ TD (Fig. 2, Tables 4, 5). An analysis of variance of all log-transformed PFM subarea data indicated significant differences ( $\mathrm{p}<0.05$ ): post hoc Tukey pairwise comparisons showed significantly ( $\mathrm{p}<0.05$ ) lower mean densities between $90-120 \mathrm{~mm} \mathrm{TD}$ and $\geq 90 \mathrm{~mm}$ TD size limits, and significantly ( $\mathrm{p}<0.05$ ) lower mean biomass densities between $90-120 \mathrm{~mm}$ TD and two other size groups, $\geq 100$ and $\geq 90 \mathrm{~mm} \mathrm{TD}$; all other size group comparisons were not significantly different (Tukey pairwise comparisons, $\mathrm{p}>0.05$ ).

The predicted lower $90 \%$ confidence intervals for mean biomass densities for $100-140 \mathrm{~mm}$ TD for each PFM subarea were presented in Table 6, as an example, and summarized for the different size groups for each region of B.C. in Table 7.

As the fishery progresses, the average density and biomass of the size group of urchins being exploited may decrease. Fishery-independent surveys of red sea urchin populations in heavily harvested areas should be repeated. Although port sampling is not conducted at present, temporal changes and area differences in size frequency, mean weights, and gonad quality of commercial-sized individuals could be monitored by sampling harvested red sea urchins at ports and or at commercial processing plants.

## Bed Areas

The total estimated bed area was $50,205.2$ ha for the North Coast and 9,669.2 ha for the South Coast as of 1996 (Table 6a). Estimated bed areas differed for each PFM subarea and increased by an average 26.8 \% between 1994 and 1996 (Campbell et al. 1999).

Using charts to estimate bed areas is crude, especially as each location may have different substrate surface areas. The harvest logbooks provide an historical cumulative estimate of fishable sea urchin areas but may include a few areas that no longer have viable red sea urchin populations. There may be areas still unexplored, especially in the North Coast, that may contain substantial unfished "virgin" populations that have not been included in the bed area estimates. Bed area estimates probably provide the most uncertainty of all the estimates used to calculate red sea urchin biomass. The most recent bed area estimates for the 1997-99 logbooks have yet to be digitized and calculated.

In some areas there may be large numbers of red sea urchins deeper than 9.1 m (D. Bureau, unpublished data; W. Bradbury, Washington State Fisheries and Wildlife, personal communication). Estimating bed areas in the reasonably shallow depth range (1 -9.1 m ) probably provides conservative bed area values to the estimation of red sea urchin biomass. In addition, the present method of multiplying the mean density estimated over the whole PFM subarea by the total bed area is a more conservative method to estimating biomass than multiplying the density by the total coastline area in the PFM subarea.

The records of log book bed area entries should be made on an annual basis rather than a cumulative basis on charts. Clear identification of beds in relation to the amount of red sea urchins removed from each bed needs to be recorded carefully by fishers and on grounds observers to allow detailed stock analyses on a bed by bed and/or PFM subarea basis in both the North and South Coasts. Although there has been an on-ground observer in the North Coast, to date there have not been any on-grounds observers for the South Coast.

## Biomass and Quota

Red sea urchin biomass $\left(B_{c}\right)$ differed considerably between PFM subareas (Table 6). Total biomass of red sea urchins in the $100-140 \mathrm{~mm}$ TD size group was estimated at $242,952 \mathrm{t}$ for the North Coast and $43,864 \mathrm{t}$ for the South Coast; lower $90 \%$ values were $173,847 \mathrm{t}$ and $25,137 \mathrm{t}$, respectively (Table 6). These biomass and quota estimates must be treated with caution, especially when considering how inaccurate the bed estimates of viable red sea urchin populations may be.

If the size limit of $\geq 100 \mathrm{~mm} \mathrm{TD}$ is to remain the same, and considering fishermen tend to select sea urchins close to legal size for better gonad quality, the more appropriate quota estimate should be made from $B_{c}$ calculated from the $100-140 \mathrm{~mm}$ TD size group (Tables 6,7 ). Choice of a conservative quota probably should be based on M values $\leq 0.10$ which would suggest that the overall B.C. coastwide 2000/2001 quota should be between $2,982.9$ and $5,736.3 \mathrm{t}$, based on the mean densities, or between $2,069.4$ and $3,979.7 \mathrm{t}$, based on the approximate lower 90 \% confidence interval estimates (Tables 6, 7).

If a lower size limit of $\geq 90 \mathrm{~mm}$ TD is considered feasible we recommend that a precautionary approach should include (a) the institution of a maximum size no larger than 130 mm TD, (b) a quota based on a low fishing mortality of $\leq 0.02$ ( $\mathrm{M} \leq 0.10$ ) (Table 7), (c) biomass estimates be based on the lower $90 \%$ confidence limits in areas where red sea urchin abundance estimates are uncertain (e.g., surveys that are $>5$ years old), (d) an adaptive management approach be taken by identifying certain PMF subareas to allow testing the feasibility and suitability of lower size limits by fishermen, managers and biologists.

Although we examined two alternative formulations, proposed by Garcia et al (1989) (i.e., equations 7 a and 8 a , based on the Schaefer (1954) and Fox (1970) production models), to estimate Q and MSY by including current yield ( $\mathrm{Y}_{\mathrm{c}}$ ) in an exploited population, both formulae were unstable when attempting to obtain MSY estimates below $Y_{c}$. Garcia et al (1989) also indicated that these formulae are unstable under other conditions. Die and Caddy (1997) question whether any simple approximation method alone, in obtaining sustainable yield indicators from biomass estimates, can provide a safe yield target. They advocate use of low conservative targets for fishing mortality and several biological reference points (BRPs) as a precautionary approach. Lai and Bradbury (1998), through simulation of red sea urchin populations in Washington, suggested that target harvest rates should be well below biological references points such as $\mathrm{F}_{\max }=0.48$ (fishing mortality at which $\mathrm{Y} / \mathrm{R}$ is maximized) and $\mathrm{F}_{0.1}=0.19$ (at which slope of $\mathrm{Y} / \mathrm{R}$ curve is $10 \%$ of the slope at origin) calculated from a yield per recruit model. Campbell and Meynert (Table 3 in appendix B), using a yield per recruit analysis, found a more conservative BRP of $\mathrm{F}_{0.35}=0.05$ and 0.09 for a size limit of $\geq 100 \mathrm{~mm}$ TD and $90-130$ mm TD, respectively.

If stock assessment and management is needed on a bed by bed and up to date basis, landings will be required in a timely way (within one year). A 3 (or longer) year periodic rotation of fishing grounds would provide for easier monitoring of the fishing fleet and landings and allow timely analysis of up to date landings data. Three or six year rotation of fishing grounds would also allow recovery of the harvestable stock through recruitment and growth. Botsford et al. (1993) and Lai and Bradbury (1998) consider periodic harvest schedules (rotation), although not increasing cumulative yield, are biologically beneficial, reduce variability of yield, risk, and probably management and enforcement costs.

## Size Limits

The influence of different size limits and fishing mortality rates on yield and egg per recruit values, where recruitment was constant, were reported by Campbell and Meynert (Appendix B) for a red sea urchin population in B.C. The long-term effects of different harvesting rates and size restrictions on a red sea urchin population, using a stochastic yield model, were examined by Hajas et al. (Appendix A). The general results of these analyses are summarized as follows. Minimum size limits of $\geq 70$ and $\geq 90 \mathrm{~mm}$ TD are clearly less precautionary than that of $\geq 100 \mathrm{~mm}$ TD. Adding a lower maximum size to a
minimum size reduces yield but increases the reproductive potential and consequently reduces the potential for collapse of red sea urchin populations. Low fishing mortality rates (e.g., $\leq 0.02$ ) are precautionary and may reduce the influence of size limits. However, having both minimum and maximum size limits and a low fishing mortality provide for a precautionary approach to exploiting red sea urchins populations when determining quotas from uncertain natural mortality and biomass estimations for particular PFM areas.

Although the results of the size limit analyses are robust, they should be treated as preliminary and with some caution since a number of factors (e.g., changes in natural mortality and density effects on growth, mortality and reproductive rates, and roe quality) were not taken into account.

The commercial quality of red sea urchin roe is based on a combination of criteria such as colour, texture, size and taste. Generally, good quality roe from red sea urchins is considered to have a yellow/gold consistent colour, firm texture (growing or premature gonads), sweet taste and of medium size which is mainly found in the approximately 90 120 mm TD size range. Poor quality roe is generally considered to have variable yellow/brown colour, soft texture and or oozing gametes (from mature, post spawn or spent gonads). Red sea urchins $>130 \mathrm{~mm}$ TD generally have large gonads and inconsistent colour quality which are less desirable by the industry than smaller mature individuals. There is a reproductive annual cycle with the timing of the spawning season varying within the period from February to September depending on local environmental conditions such as food availability and temperature (Bernard 1977). Gonads increase in size usually from September to January (Kramer and Nordin 1975). The effect of food quality and availability to support growth and reproduction are important limiting factors on urchin stocks. Although red sea urchins are omnivorous grazers, kelps, such as Nereocystis leutkeana, provide optimal growth and gonad quality (Vadas 1977; Bureau et al. 1997; Morris and Campbell 1996). In areas of low supply or quality of food, individuals may relocate nutrients causing poor quality gonads thereby reducing reproduction potential of local red sea urchin populations. The influence of adult age, especially in large old red sea urchins, on low egg production, senescence or of poor quality and low survival of larval offspring is unknown.

Breen (1984) argued that a size limit could be used to protect sufficient reproductive potential in an attempt to insure that recruitment did not fall below replacement. A special argument for a maximum size limit can be made for red sea urchins as adult urchins may provide a spine cover for juveniles that is necessary for good survival (Tegner and Dayton 1977). Minimum and maximum size limits would leave enough large adults to protect the settlement of juveniles, such as used in Washington State (Lai and Bradbury 1998). Breen (1984) suggested that using a size limit as the only control mechanism to prevent recruitment overfishing would not be achieved with the $\geq 100 \mathrm{~mm}$ TD size limit that is presently used. A larger size limit would be required to prevent recruitment overfishing but would be larger than the industry's upper limit for market quality. Breen (1984) recommended that a better way of preventing recruitment
overfishing was "controlling effort or catch in such a way as to protect local stocks from over harvesting".

The influence of market demand for high quality small sized gonads may cause fishermen to concentrate fishing only on a small size range (e.g., $90-110 \mathrm{~mm} \mathrm{TD}$ ) of red sea urchins even though the quota may be based on a legal size limit of $\geq 90 \mathrm{~mm}$ TD or 90 130 mm TD. This would result in a higher fishing mortality on the small size range than was originally intended by the managers. Clearly setting a quota on the appropriate size range that will be harvested in an area will ensure less deviation from the intended fishing mortality on local red sea urchin populations. Monitoring harvested red sea urchins at ports and or at commercial processing plants will provide temporal changes and area differences in size frequency, mean weights, and gonad quality of commercial-sized individuals, and provide a tool to assess the appropriate size range to set quotas.

Lai and Bradbury (1998) conducted simulation models to examine the relative merits of harvest rates, size limits and periodic harvesting on red sea urchin populations in Washington. They found that there was approximately twice the risk of stock collapse without a minimum size limit than with a size limit at different fishing rates. They also found that the probability of sustainability of red sea urchin populations would increase substantially by restricting effort or catch and by increasing the period between harvests (e.g. from 1 to 5 year rotations).

There are a number of complex density dependent compensatory and depensatory mechanisms that red sea urchins may elicit in growth, mortality, increased survival of juveniles due to protection by adult spine canopies, spawning success, roe quality and egg viability. A stock at low density levels may impact on the population negatively, e.g. "the Allee affect" (Allee 1931), reducing the reproductive success of the animals. This is particularly important in organisms that are broadcast spawners. Levitan et al. (1992) found that fertilization success was a function of the number of red sea urchins, distance apart, position in the cluster, flow direction and velocity of the current, etc. For animals that show this type of fertilization, this is a factor that should be considered in the management system in providing various forms of harvest refugia (e.g. size restrictions, catch limits through quotas or rotating spatial harvests and spatial closures) (Botsford et al. 1993; Quinn et al. 1993; Pfister and Bradbury 1996; Levitan and Sewell 1998). Ebert (1998) found potentially little Allee effects on growth of red sea urchins in Oregon and Washington using simulation techniques. He recommended not to automatically assume Allee effects are important in local areas without some evaluation, such as incorporating sensitivity analyses in dynamic modelling methods.

In addition to resource monitoring through fishery independent surveys, additional biological information is clearly required on age estimation and on the variation of growth, mortality and recruitment rates for production modeling of red sea urchin populations in different areas of B.C.

## RECOMMENDATIONS

(1) Biomass and quota estimates in this paper should be considered only as a preliminary indication of the status of red sea urchin stocks in B.C.
(2) If a lower size limit of $\geq 90 \mathrm{~mm}$ TD is considered feasible we recommend that a precautionary approach could include:
(a) the institution of a maximum size no larger than 130 mm TD ;
(b) a quota based on a low fishing mortality of $\leq 0.02$ ( $\mathrm{M} \leq 0.10$ );
(c) biomass estimates be based on the lower $90 \%$ confidence limits in areas where red sea urchin abundance estimates are uncertain (e.g., surveys that are 5 years old);
(d) initially an adaptive management approach be taken by identifying certain PMF subareas to allow testing the feasibility and suitability of lower size limits by fishermen, managers and biologists;
(e) monitoring harvested red sea urchins at ports and or at commercial processing plants will provide temporal changes and area differences in size frequency, mean weights, and gonad quality of commercial-sized individuals, and provide a tool to assess the appropriate size range to set quotas in the future.
(3) Surveys for the abundance estimates of red sea urchins in alternative areas in B.C. should be conducted on an annual basis. Areas with no surveys for over 10 years and for areas that have been heavily fished should be considered a priority.
(4) More accurate estimates of bed areas holding viable populations of red sea urchins are required. Bed areas fished on an annual basis need to be recorded and an update of bed areas fished during 1997-1999 is needed.
(5) Further research is required to understand age, and growth, mortality and recruitment rates of red sea urchins in B.C. to assist with production modeling.

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## REFERENCES

Adkins, B.E., R.M. Harbo, and P.A. Breen. 1981. A survey of commercial sea urchin (Strongylocentrotus franciscanus) populations in the Gulf Islands, November 1980. Can. MS. Rep. Fish. Aquat. Sci. 1618: 41 p.

Allee, W. C. 1931. Animal aggregations. A study in general sociology. University of Chicago Press, Chicago, Ill.
Bernard, F.R. 1977. Fishery and reproductive cycle of the red sea urchin Strongylocentrotus franciscanus, in British Columbia. J. Fish. Res. Board Can. 34: 604-610.
Bernard, F.R. and D.C. Miller. 1973. Preliminary investigation of the red sea urchin (Strongylocentrotus franciscanus, Agassiz) resources of British Columbia. Fish. Res. Board Can. Tech. Rep. 400.37 p.
Botsford, L.W., B.D. Smith, S.R. Wing, and F.F. Quinn, 1994. Bi-modality in size distributions: the red sea urchin Strongylocentrotus franciscanus as an example. Ecol. Appl. 4: 42-50.
Botsford, L.W., J. F. Quinn, S.R. Wing, and J.G. Brittnacher. 1993. Rotating spatial harvest of a benthic invertebrate, the red sea urchin, Strongylocentrotus franciscanus. pp. 409-428. In: Proceedings of the international symposium on management strategies for exploited fish populations, Alaska Sea Grant College Program, AK-SG-93-02.
Breen, P.A. 1980. The ecology of red sea urchins in British Columbia. pp. 3-12. In: International symposium on coastal Pacific marine life, Western Washington University, Bellingham.
Breen, P.A. 1984. Sea urchins: suitability of the present size limit. Can. MS. Rep. Fish. Aquat. Sci. 1774: 25-51.
Breen, P.A., D.C. Miller, and B.E. Adkins. 1976. An examination of harvested sea urchin populations in the Tofino area. Fish. Res. Board Can. 1401: 23 p.
Breen, P.A., B.E. Adkins, and D.C. Miller. 1978. Recovery rate of three exploited sea urchin populations from 1972 to 1977. Can. MS. Rep. Fish. Mar. Ser. 1446: 27 p.
Bureau, D., A. Campbell and W. Hajas. 2000 a. Survey of red sea urchin populations near Comox, Denman Island and Hornby Island, British Columbia, 1999. Can. MS. Rep. Fish. Aquat. Sci. In preparation.
Bureau, D., A. Campbell and W. Hajas. 2000 b. Survey of red sea urchin populations in the Kelsey Bay area, Johnstone Strait, British Columbia, 1999. Can. MS. Rep. Fish. Aquat. Sci. In preparation.
Bureau, D., A. Campbell, C.A. Ayers and W. Hajas. 2000c. Survey of red sea urchin populations in the Gulf Islands, Strait of Georgia, British Columbia, 1998 and 1999. Can. MS. Rep. Fish. Aquat. Sci. In preparation.

Bureau, D., A. Campbell and W. Hajas. 2000 d. Survey of red sea urchin populations in the Larsen Harbour and Kinkown Inlet areas, Banks Island, British Columbia, 1997. Can. MS. Rep. Fish. Aquat. Sci. In preparation.

Bureau, D., A. Campbell, and E. B. Hartwick. 1997. Roe enhancement in red sea urchin, Strongylocentrotus franciscanus, fed the kelp, Nereocystis luetkeana. Bull.

Aquacul. Assoc. Canada. 97-1: 26-30.
Caddy, J.F. 1986. Stock assessment in data-limited situations - the experience in tropical fisheries and its possible relevance to evaluation of invertebrate resources. Can. Spec. Publ. Fish. Aquat. Sci. 92: 379-392.
Campbell, A. 1998. Catch, effort and quota estimates for the red sea urchin fishery in British Columbia. Can. Tech. Rep. Fish. Aquat. Sci. 2215: 83-109.
Campbell, A., and R.M. Harbo. 1991. The sea urchin fisheries in British Columbia, Canada. pp. 191-199. In:. Yanagisawa et al. [eds.]. Biology of Echinodermata. Balkema, Rotterdam.
Campbell, A, J. Boutillier, and J. Rogers. 1999. Discussion on a precautionary approach for management of the red sea urchin fishery in British Columbia. Can. Stock Assessment Secretariat Res. Doc. 99/094.
Campbell, A., D. Bureau, and D. Brouwer. 2000. Quota estimates for the 1998 red sea urchin fishery in British Columbia. Can. MS. Rep. Fish. Mar. Sci. 2516. In press.
Die, D.J. and J.F. Caddy. 1997. Sustainable yield indicators from biomass: are there appropriate reference points for use in tropical fisheries? Fish. Res. 32: 69-79.
Ebert, T. E. 1998. An analysis of the importance of Allee effects in management of the red sea urchin Strongylocentrotus franciscanus. p. 619-627. In R. Mooi and M. Telford (eds). Echinoderms: San Francisco. A. A. Balkema, Rotterdam.
Fox, W.W. Jr. 1970. An exponential surplus-yield model for optimizing exploited fish populations. Trans. Am. Fish. Soc. 99: 80-88.
Garcia, S., P. Sparre, and J. Csirke. 1989. Estimating surplus production and maximum sustainable yield from biomass data when catch and effort time series are not available. Fish. Res. 8: 13-23.
Gulland, J.A. 1971. The Fish Resources of the Ocean. Fishing News (Books), West Byfleet. 255 p.
Heizer, S., G. Thomas, and K. Hobbs. 1997. Red sea urchin dive fishery update. Can. MS. Rep. Fish. Aquat. Sci. 2369: 93-102.
Jamieson, G.S. and C.J. Schwarz. 1998. Survey protocol considerations for the 1995 sea urchin surveys. Can. Tech. Rep. Fish. Aquat. Sci. 2215: 69-81.
Jamieson, G.S., K. Cripps, M. Gijssen, L. Greba, R. Jones, G. Martel, W. Sandoval, C.J. Schwarz, C. Taylor, and R. Routledge. 1998a. Reanalyses of 1993 red sea urchin surveys in Haida, Heiltsuk, Kitasoo and Tsimshian traditional territories. Can. Tech. Rep. Fish. Aquat. Sci. 2215: 57-68.
Jamieson, G.S., R. Jones, G. Martel, C.J. Schwarz, C., and R. Routledge. 1998b. Analysis of 1994 red sea urchin survey conducted in Haida Pacific Fishery Management Area 1. Can. Tech. Rep. Fish. Aquat. Sci. 2215: 3-18.
Jamieson, G.S., W. Sandoval, C.J. Schwarz, C. Taylor, and R. Routledge. 1998c. Analysis of the 1994 red sea urchin surveys conducted in Heiltsuk Traditional Territory, Pacific Fishery Management Area 7, subareas 18 and 25. Can. Tech. Rep. Fish. Aquat. Sci. 2215: 19-31.
Jamieson, G.S., G. Scarf, C.J. Schwarz, C. Taylor, and R. Routledge. 1998d. Analysis of 1994 red sea urchin surveys conducted in Aweena K'ola Traditional Territory, subareas of Pacific Fishery Management Area 12. Can. Tech. Rep. Fish. Aquat. Sci. 2215: 33-56.

Kramer, D.E. and D.M.A. Nordin. 1975. Physical data from a study of size, weight and gonad quality for the red sea urchin (Strongylocentrotus franciscanus (Agassiz)) over a one-year period. Fish. Res. Board Can. MS. Rep. 1372: 91p.
Lai, H.L. and Bradbury, A. 1998. A modified catch-at-size analysis model for a red sea urchin (Strongylocentrotus franciscanus) population. Can. Spec. Publ. Fish. Aquat. Sci. 125: 85-96.
Levitan, D.R. and M. A. Sewell. 1998. Fertilization success in free-spawning marine invertebrates: review of the evidence and fisheries implications. Can. Spec. Publ. Fish. Aquat. Sci. 125: 159-164.
Levitan, D.R., M.A. Sewell, and Fu-Shiang Chia. 1992. How distribution and abundance influence fertilization success in the sea urchin Strongylocentrotus franciscanus. Ecology, 73: 248-254.
Morris, T.J and A. Campbell. 1996. Growth of juvenile red sea urchins (Strongylocentrotus franciscanus) fed Zostera marina and Nereocystis luetkeana. J. Shellfish Res. 15: 777-780.

Mottet, M.G. 1976. Fishery biology of sea urchins in the family Strongylocentrotidae. Wash. Rep. Fish. Tech. Rep. 20: 66 p.
Pfister, C.A., and A. Bradbury. 1996. Harvesting red sea urchins: recent effects and future predictions. Ecol. Appl. 6: 298-310.
Quinn, J.F., S.R Wing, and L.W. Botsford. 1993. Harvest refugia in marine invertebrate fisheries: models and applications to the red sea urchin, Strongylocentrotus franciscanus. Am. Zool. 33: 537-550.
Rogers, J. and S. Neifer. 1999. Red Sea Urchin Fishery - 1997/98. Fishery Update. (manuscript in prep.).
Schaefer, M.B. 1954. Some aspects of the dynamics of populations, important for the management of commercial fisheries. Bull. Inter-Amer. Trop. Tuna Comm., 1(2): 56 p.
Sloan, N.A., C.P. Lauridsen, and R.M. Harbo. 1987. Recruitment characteristics of the commercially harvested red sea urchin Strongylocentrotus franciscanus in southern British Columbia, Canada. Fish. Res. 5: 55-69.
Tegner, M.J. 1989. The feasibility of enhancing red sea urchin, Strongylocentrotus franciscanus, stocks in California: an analysis of the options. Mar. Fish. Rev. 51 (2): 1-22.

Tegner, M.J. and P.K. Dayton. 1977. Sea urchin recruitment patterns and implications of commercial fishing. Science. 196: 324-326.
Vadas, R.L. 1977. Preferential feeding: an optimization strategy in sea urchins. Ecol. Monog. 47: 337-371.
Woodby, D. 1992. Red sea urchins in SE Alaska: status of research and management. Unpublished MS for Sea Grant Conference on Sea Urchin, Kelp, and Abalone, March, 1992. Bodega Bay, California. 11 p.

Table 1. Annual red sea urchin landings (tonnes), value and effort for British Columbia, 1978-96, as reported on sales slips and harvest logs.

| Year | $\begin{aligned} & \hline \text { Type and } \\ & \text { Number } \\ & \text { of } \\ & \text { Licenses } \\ & \text { issued } \\ & \hline \end{aligned}$ | South Coast ${ }^{1}$ Quota (t) | North <br> Coast <br> Quota <br> (t) | Number of Vessels with Landings | Total Vessel Fishing Days | Coastwide Landings <br> (t) | $\begin{gathered} \hline \text { Landed } \\ \text { Value } \\ \left(\$ .10^{3}\right) \end{gathered}$ | Whole Landed Value (\$/t) | Mean CPUE (t/vessel day) | $\begin{gathered} \text { Mean } \\ \text { CPUE }^{3} \\ (\mathrm{~kg} / \mathrm{dive} \\ \mathrm{r} \text { hr }) \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1978 | C |  |  | 4 | 54 | 75 | 16 | 213 | 1.4 | - |
| 1979 | C |  |  | 29 | 298 | 317 | 76 | 240 | 1.1 | - |
| 1980 | C |  |  | 18 | 331 | 333 | 84 | 252 | 1.0 | - |
| 1981 | C | 136 |  | 18 | 127 | 116 | 34 | 293 | 0.9 | - |
| 1982 | C |  |  | 21 | 195 | 160 | 56 | 350 | 0.8 | - |
| 1983 | Z 64 |  |  | 36 | 825 | 986 | 358 | 354 | 1.2 | 311 |
| 1984 | Z 85 |  |  | 47 | 1150 | 1764 | 712 | 403 | 1.6 | 281 |
| 1985 | Z 86 | 1803 |  | 46 | 1086 | 1815 | 764 | 419 | 1.7 | 360 |
| 1986 | Z 103 | 1500 |  | 67 | 1534 | 2067 | 1011 | 455 | 1.4 | 363 |
| 1987 | Z 184 | 1633 |  | 97 | 1807 | 2118 | 1148 | 541 | 1.2 | 325 |
| 1988 | Z 184 | 1678 |  | 84 | 1249 | 2116 | 1241 | 587 | 1.7 | 296 |
| 1989 | Z 240 | 1644 |  | 109 | 1542 | 2658 | 1631 | 614 | 1.7 | 360 |
| 1990 | Z 188 | 1668 |  | 116 | 2651 | 3158 | 1953 | 618 | 1.2 | 298 |
| 1991 | Z 102 | 1531 |  | 89 | 3862 | 6831 | 4187 | 613 | 1.8 | 363 |
| 1992 | Z 108 | 1554 |  | $110{ }^{4}$ | 6222 | 12982 | 8662 | 616 | 2.1 | 388 |
| 1993 | Z 107 | 1401 | 5400 | 103 | 3364 | 6388 | 5373 | 841 | 1.9 | 340 |
| 1994 | Z 110 | 1543 | 5897 | 98 | 3978 | 5829 | 8066 | 1384 | 1.5 | 325 |
| 1995 | Z 108 | 1386.8 | 5455 | 108 | 4167 | 6585 | 11350 | 1724 | 1.6 | 325 |
| 1996 | Z 109 | 1264.7 | 5360 | 109 | 3536 | 5753 | 11358 | 1974 | 1.6 | 340 |
| 1997/98 ${ }^{5}$ | Z110 | 1,702 | 8150 | 109 | 5292 | 8451 | 13999 | 1656 | 1.6 |  |
| 1998/99 | Z110 | 968 | 4635 | 110 | 3282 | 5091 | 7366 | 1447 | 1.6 |  |

${ }^{1}$ South Coast quota includes exploratory areas, North Coast quota was new in 1993.
${ }^{2}$ From sales slip data. ${ }^{3}$ CPUE from harvest logbook data.
${ }^{4}$ Larger than the number of licenses issued due to license transfers.
${ }^{5}$ Fishing season changed from calendar year to market driven year (July to June), 1997/98 season January 1, 1997, to June 30, 1998.

Table 2. Red sea urchin annual landings (tonnes) by South Coast Management Area 1971 to June 1999, as reported on fish slips and harvest logs. 1994 to 1999 catch data corrected to use Validation \& Harvest logs only.

| SOUTH COAST MANAGEMENT AREAS |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Year | East Coast Vancouver Island |  |  |  |  |  |  |  |  |  |  | West Coast Vancouver Island |  |  |  |  |  |  | Total Landings |
|  | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 28 | 29 | 20 | 21 | 23 | 24 | 25 | 26 | 27 |  |
| 1971 to $1973{ }^{1}$ |  |  |  |  |  |  |  |  | 110.0 |  |  |  |  |  | 254.0 |  |  |  | 364.0 |
| 1974 to $1977{ }^{1}$ |  | 1.4 |  | * | * | 1.4 |  |  | 66.0 |  |  | * |  | 1.3 |  |  |  |  | 70.1 |
| 1978 |  |  | * |  |  |  |  |  | 46.0 |  |  | 29.0 |  |  |  |  |  |  | 75.0 |
| 1979 |  |  | * | 78.0 |  |  | 57.0 | 133.0 | 45.0 |  |  | 1.8 | 0.9 | 2.5 |  |  |  |  | 318.2 |
| 1980 |  |  |  | 18.0 |  |  | 162.0 | 54.0 | 97.0 |  |  | 1.8 |  |  |  |  |  |  | 332.8 |
| 1981 |  |  | 20.0 | 4.0 | * |  | 5.3 | 47.0 | 22.0 |  |  |  |  |  | 17.0 |  |  |  | 115.3 |
| 1982 |  | 2.5 |  | 46.0 |  |  | 0.8 | 11.0 | 94.0 |  |  |  |  |  | 5.0 |  |  |  | 158.5 |
| $1983{ }^{2}$ | 7.8 | 99.0 | 264.0 | 260.0 | * | * | 59.0 | 38.0 | 112.0 |  |  | 24.0 |  | 22.0 | 38.0 |  | 62.0 |  | 985.8 |
| 1984 | 0.3 | 437.0 | 777.3 | 172.0 |  |  | 33.0 | 67.4 | 76.3 |  | 5.7 | 69.1 |  | 17.3 | 103.0 |  | 3.9 |  | 1,762.3 |
| 1985 |  | 354.0 | 492.0 | 167.0 | 106.0 | 5.9 | 29.0 | 48.0 | 77.0 |  | 47.0 | 30.0 |  | 96.0 | 158.0 | 145.0 | 15.0 | 45.0 | 1,814.9 |
| 1986 | 27.0 | 548.0 | 376.0 | 178.0 | 56.0 | 4.4 | 57.0 | 129.0 | 105.0 |  | 2.0 | 40.0 |  | 154.0 | 285.0 |  | 2.5 | 91.0 | 2,054.9 |
| $1987{ }^{3}$ | 6.9 | 420.0 | 491.0 | 193.0 | 32.4 |  | 71.0 | 71.0 | 123.0 | 17.0 | 7.8 | 17.0 |  | 63.0 | 199.0 | 95.0 | 8.3 | 12.0 | 1,827.4 |
| 1988 | 2.6 | 534.0 | 480.0 | 78.0 | 21.0 | 2.3 | * | 22.0 | 78.0 |  |  | 74.0 |  | 13.0 | 250.0 | 66.0 |  | 58.0 | 1,678.9 |
| 1989 |  | 569.0 | 493.0 | 122.0 | 6.7 |  | 9.0 | 64.0 | 57.0 |  | 1.6 | 15.0 |  |  | 223.0 | 39.0 |  | 86.0 | 1,685.3 |
| 1990 | 84.8 | 437.6 | 428.4 | 56.6 | 1.2 | 0.6 | 43.0 | 46.5 | 58.6 | 0.3 | 1.8 | 7.9 |  | 59.7 | 215.1 | 56.8 |  | 68.1 | 1,567.0 |
| 1991 | 36.4 | 358.7 | 370.7 |  | 8.6 |  | 26.6 | 94.8 | 27.2 |  | 14.1 | 31.2 | 2.7 | 58.4 | 185.1 | 115.8 |  | 121.1 | 1,451.4 |
| 1992 | 8.0 | 531.0 | 320.0 |  |  |  | 103.0 | 36.0 | 86.0 |  | 4.0 | 56.0 | 9.0 | 31.0 | 200.0 | 10.0 |  | 65.0 | 1,459.0 |
| 1993 | 55.5 | 329.0 | 184.0 |  |  |  | 21.0 | 104.7 | 17.3 |  |  | 14.5 |  | 40.4 | 92.0 | 7.0 | 2.0 | 50.0 | 917.4 |
| 1994 | 17.0 | 348.0 | 168.0 |  |  |  | 4.0 | 59.0 | 14.0 |  | 1.0 | 3.0 |  | 54.0 | 111.0 | 50.0 |  | 49.0 | 878.0 |
| 1995 | 34.0 | 364.0 | 175.0 |  |  |  | 28.0 | 69.0 | 15.0 |  | 9.0 | 20.0 |  | 57.0 | 199.0 |  |  | 98.0 | 1,068.0 |
| 1996 | 38.0 | 344.0 | 238.0 |  |  |  | 25.0 | 112.0 | 7.0 |  | 10.0 | 33.0 |  | 46.0 | 122.0 |  |  | 70.0 | 1,045.0 |
| 1997/98 | 61.0 | 594.0 | 426.0 |  | 2.3 |  | 26.0 | 67.0 | 40.0 |  | 16.0 | 32.0 |  | 54.0 | 132.0 |  |  | 85.0 | 1,535.3 |
| 1998/99 | 49.7 | 285.2 | 196.4 |  |  |  | 26.9 | 85.3 | 21.1 |  | 2.9 | 22.5 |  | 31.5 | 107.6 |  |  | 48.2 | 877.3 |
| 1971 to |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| June 1999 | 429.0 | 6,556.4 | 5,899.8 | 1,372.6 | 234.2 | 14.6 | 785.8 | 1,358.7 | 1,394.5 | 17.3 | 122.9 | 521.8 | 12.6 | 801.1 | 2,895.8 | 584.6 | 93.7 | 946.4 | 24,041.8 |

* Less than 500 kg .
${ }^{1}$ Data for each year cannot be published separately.
${ }^{2}$ Mandatory log book under Z licence came into effect in 1983.
${ }^{3}$ Sales slips were combined for red and green sea urchins in 1987, were later separated by price criteria, but 320 t remains missing in area table.

Table 3. Red sea urchin annual landings (tonnes) by North Coast Management Area 1984 to June 1999, as reported on fish slips and harvest logs. 1994 to 1999 catch data corrected to use Validation \& Harvest logs only.

| NORTH COAST MANAGEMENT AREAS |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Year | 1 | 2E | 2W | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | Total Landings |
| 1984 | 2.2 |  |  |  |  |  |  |  |  |  |  | 2.2 |
| 1986 |  |  |  |  |  |  |  |  |  |  | 12.0 | 12.0 |
| 1987 |  |  |  |  | 23.0 |  |  | 179.0 | 91.0 |  |  | 293.0 |
| 1988 |  |  |  |  | 73.0 | 11.0 | 7.3 | 314.0 | 32.0 |  |  | 437.3 |
| 1989 | 0.2 | 223.0 |  | 1.6 | 116.0 | 1.3 | 168.0 | 217.0 | 65.0 |  | 180.0 | 972.1 |
| 1990 |  | 26.6 | 10.7 | 24.5 | 156.8 | 265.3 | 67.1 | 1,040.1 |  |  |  | 1,591.1 |
| 1991 |  | 333.1 | 2.7 | 143.3 | 1,026.7 | 2,577.3 | 77.7 | 774.7 | 114.6 | 24.5 | 304.9 | 5,379.5 |
| 1992 |  | 1,111.0 |  | 1.0 |  | 3,294.0 | 4,063.0 | 2,763.0 | 140.0 | 114.0 | 38.0 | 11,524.0 |
| 1993 | 97.0 | 189.0 | 88.9 | 127.2 | 1,008.0 | 463.0 | 2,103.0 | 1,012.0 | 43.4 |  | 215.3 | 5,346.8 |
| 1994 | 221.0 | 402.2 | 167.4 | 173.0 | 687.0 | 1,056.0 | 1,244.0 | 861.0 | 57.0 | 46.0 | 164.0 | 5,078.6 |
| $1995{ }^{1}$ | 258.0 | 440.2 | 256.3 | 48.0 | 940.0 | 1,280.0 | 1,053.0 | 1,076.0 | 111.0 | 49.0 | 224.0 | 5,735.5 |
| 1996 | 259.0 | 365.0 | 241.8 | 66.0 | 851.0 | 1,156.0 | 1,213.0 | 833.0 | 122.0 | 10.0 | 248.0 | 5,364.8 |
| 1997/98 | 582.0 | 718.7 | 311.1 | $62.0$ | 1,076.0 | 1,107.0 | 2,175.0 | 870.0 | 112.0 | 41.0 | 152.0 | 7,206.8 |
| 1998/99 | 276.2 | 438.1 |  | 38.8 | 595.1 | 741.8 | 1,373.6 | 526.2 | 87.6 | 36.8 | 191.4 | 4,305.6 |
| 1984 to |  |  |  |  |  |  |  |  |  |  |  |  |
| June 1999 | 1,695.6 | 4,246.9 | 1,078.9 | 685.4 | 6,552.6 | 11,952.7 | 13,544.7 | 10,466.0 | 975.6 | 321.3 | 1,729.6 | 53,249.3 |

[^0]Table 4. Summary of estimated mean density (number/ $\mathrm{m}^{2}$ ) and mean biomass $\left(\mathrm{g} / \mathrm{m}^{2}\right.$ ) of red sea urchins in British Columbia by PMF subarea, obtained from broad-brush surveys during 1993-1999.
Data include analyses from Jamieson et al. (1998a, b, c, d), Campbell et al. (1998), and present study. NQCI, EQCI and WOCI refer to north, east and west areas of the Queen Charlotte Island
QCStrait refers to Queen Charlotte Strait. C. Georgia St refers to Central Georgia Strait (Comox and Horrby IS. Areas). SE $=$ standard error. Standard errors were not included for the 1993 surveys or if fewer
than 5 transects were surveyed.


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## EQCI EQCI

## EQCI EOCI

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## EQCI EQCI




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Banks Is.
Banks Is.
Banks Is.

$\begin{array}{ll}\text { Banks Is. } & 26 \\ \text { Banks Is. } & 22\end{array}$

| 764 | 1.53 | 0.33 | 1.98 | 0.42 | 1.52 | 0.33 | 1.59 | 0.36 | 1.84 | 0.40 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1685 | 2.89 | 0.57 | 3.56 | 0.70 | 2.79 | 0.55 | 2.50 | 0.47 | 3.26 | 0.67 |
| 1795 | 1.26 | 0.37 | 1.34 | 0.39 | 1.05 | 0.35 | 0.49 | 0.17 | 0.82 | 0.27 |
| 2859 | 0.54 | 0.20 | 0.60 | 0.23 | 0.47 | 0.17 | 0.30 | 0.13 | 0.43 | 0.17 |
| 557 | 0.42 |  | 0.57 |  | 0.42 |  | 0.39 |  | 0.53 |  |
| 476 | 0.95 |  | 1.34 |  | 0.90 |  | 1.00 |  | 1.18 |  |
| 1070 | 0.53 |  | 0.71 |  | 0.51 |  | 0.54 |  | 0.64 |  |
| 557 | 0.42 |  | 0.44 |  | 0.37 |  | 0.15 |  | 0.25 |  |
| 589 | 0.85 |  | 1.10 |  | 0.78 |  | 0.78 |  | 0.93 |  |
| 3234 | 0.07 |  | 0.09 |  | 0.07 |  | 0.07 |  | 0.09 |  |
| 1047 | 0.47 |  | 0.69 |  | 0.47 |  | 0.54 |  | 0.61 |  |
| 230 | 0.31 |  | 0.38 |  | 0.26 |  | 0.21 |  | 0.28 |  |
| 628 | 1.13 |  | 1.54 |  | 1.08 |  | 1.11 |  | 1.38 |  |
| 949 | 0.77 | 0.24 | 1.03 | 0.33 | 0.76 | 0.23 | 0.82 | 0.25 | 0.96 | 0.30 |
| 712 | 0.80 | 0.39 | 1.07 | 0.51 | 0.77 | 0.37 | 0.81 | 0.38 | 0.98 | 0.47 |
| 532 | 1.50 | 0.79 | 1.75 | 0.82 | 1.17 | 0.63 | 0.69 | 0.20 | 0.97 | 0.30 |
| 237 | 0.41 |  | 0.69 |  | 0.41 |  | 0.55 |  | 0.65 |  |
| 375 | 1.08 | 0.42 | 1.36 | 0.52 | 1.03 | 0.44 | 0.85 | 0.45 | 1.07 | 0.55 |
| 642 | 0.06 |  | 0.09 |  | 0.06 |  | 0.07 |  | 0.09 |  |
| 70 | 1.21 |  | 1.67 |  | 1.21 |  | 1.51 |  | 1.67 |  |
| 93 | 1.35 |  | 2.42 |  | 1.35 |  | 2.23 |  | 2.39 |  |
| 26 | 0.40 |  | 0.59 |  | 0.31 |  | 0.47 |  | 0.49 |  |
| 312 | 0.09 |  | 0.15 |  | 0.09 |  | 0.14 |  | 0.15 |  |
| 386 | 0.09 |  | 0.13 |  | 0.09 |  | 0.12 |  | 0.12 |  |
| 207 | 1.13 |  | 3.11 |  | 1.13 |  | 3.11 |  | 3.11 |  |
| 52 | 0.27 |  | 0.53 |  | 0.27 |  | 0.52 |  | 0.53 |  |
| 314 | 0.81 |  | 1.57 |  | 0.81 |  | 1.46 |  | 1.57 |  |
| 71 | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  |
| 124 | 1.14 |  | 2.24 |  | 1.14 |  | 2.17 |  | 2.24 |  |
| 201 | 1.15 |  | 1.81 |  | 1.15 |  | 1.68 |  | 1.81 |  |
| 411 | 1.91 |  | 2.43 |  | 1.91 |  | 2.10 |  | 2.33 |  |
| 65 | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  |
| 321 | 0.37 |  | 0.59 |  | 0.37 |  | 0.50 |  | 0.58 |  |
| 89 | 0.57 |  | 0.86 |  | 0.57 |  | 0.77 |  | 0.86 |  |
| 1011 | 0.75 |  | 1.00 |  | 0.71 |  | 0.69 |  | 0.85 |  |
| 212 | 0.31 |  | 0.37 |  | 0.28 |  | 0.21 |  | 0.28 |  |
| 1264 | 0.59 |  | 0.71 |  | 0.53 |  | 0.41 |  | 0.56 |  |
| 809 | 0.44 |  | 0.56 |  | 0.39 |  | 0.35 |  | 0.45 |  |
| 2045 | 1.25 | 0.24 | 1.75 | 0.32 | 1.06 | 0.19 | 1.16 | 0.21 | 1.41 | 0.26 |
| 194 | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  |
| 3503 | 0.30 | 0.10 | 0.40 | 0.14 | 0.29 | 0.10 | 0.30 | 0.10 | 0.36 | 0.12 |
| 27 | 1.27 |  | 1.59 |  | 1.15 |  | 1.23 |  | 1.27 |  |
| 910 | 0.26 | 0.16 | 0.33 | 0.19 | 0.24 | 0.15 | 0.25 | 0.16 | 0.28 | 0.17 |
| 393 | 0.34 |  | 0.41 |  | 0.26 |  | 0.21 |  | 0.28 |  |
| 2596 | 1.55 | 0.23 | 2.06 | 0.33 | 1.42 | 0.23 | 1.49 | 0.29 | 1.75 | 0.32 |
| 2429 | 0.97 | 0.23 | 1.15 | 0.27 | 0.82 | 0.21 | 0.75 | 0.19 | 0.92 | 0.22 |

764
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| 744.4 | 155.8 |
| :---: | :---: |
| 1471.4 | 297.9 |
| 815.3 | 222.4 |
| 323.3 | 121.3 |
| 221.3 |  |
| 498.3 |  |
| 273.2 |  |
| 262.5 |  |
| 463.1 |  |
| 36.1 |  |
| 224.3 |  |
| 185.3 |  |
| 586.6 |  |
| 367.8 | 117.3 |
| 391.9 | 193.1 |
| 1003.6 | 577.9 |
| 184.5 |  |
| 612.3 | 212.7 |
| 29.7 |  |
| 532.7 |  |
| 568.6 |  |
| 224.1 |  |
| 38.1 |  |
| 36.4 |  |
| 421.7 |  |
| 110.8 |  |
| 344.6 |  |
| 0.0 |  |
| 466.1 |  |
| 463.9 |  |
| 853.5 |  |
| 0.0 |  |
| 282.8 |  |
| 232.8 |  |
| 408.1 |  |
| 178.8 |  |
| 337.5 |  |
| 253.2 |  |
| 790.1 | 163.8 |
| 0.0 |  |
| 152.6 | 51.5 |
| 683.9 |  |
| 134.6 | 75.8 |
| 212.0 |  |
| 819.8 | 113.3 |
| 550.0 | 129.3 |

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179.9
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73.5
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229.4167.1
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| $\begin{gathered} \hline \text { PMF } \\ \text { Subarea } \end{gathered}$ | Year | Survey | Urchin density for each size group (mm TD) |  |  |  |  |  |  |  |  |  |  |  | Mean biomass for each size group (mm TD) |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | Total Transect Number Length (m) |  | $\geq 100$ |  | $\geq 90$ |  | 100-140 |  | 90-120 |  | 90-130 |  | Total Transect |  | $\geq 100$ |  | $\geq 90$ |  | 100-140 |  | 90-120 |  | 90-130 |  |
|  |  |  |  |  | Mean | SE | Mean | SE | Mean | SE | Mean | SE | Mean | SE | Number | Length (m) | Mean | SE | Mean | SE | Mean | SE | Mean | SE | Mean | SE |
| 6.010 | 94 | Campania Is. | 30 | 5028 | 1.51 | 0.18 | 1.91 | 0.28 | 1.41 | 0.18 | 1.33 | 0.25 | 1.61 | 0.25 | 30 | 5028 | 808.4 | 94.8 | 921.5 | 116.5 | 713.4 | 85.6 | 501.1 | 85.2 | 672.6 | 87.9 |
| 6.012 | 94 | Campania Is. | 7 | 524 | 0.52 | 0.27 | 0.64 | 0.36 | 0.47 | 0.27 | 0.39 | 0.25 | 0.53 | 0.32 | 7 | 524 | 287.5 | 142.8 | 322.9 | 168.2 | 245.1 | 141.7 | 145.0 | 89.6 | 231.8 | 136.6 |
| 6.013 | 93 | Kitasoo | 32 | 2434 | 1.21 |  | 1.60 |  | 1.14 |  | 1.23 |  | 1.43 |  | 32 | 2434 | 612.8 |  | 722.9 |  | 549.0 |  | 462.1 |  | 584.5 |  |
| 6.014 | 93 | Kitasoo | 11 | 762 | 1.14 |  | 1.44 |  | 1.03 |  | 1.01 |  | 1.18 |  | 11 | 762 | 640.1 |  | 722.8 |  | 518.0 |  | 386.9 |  | 491.3 |  |
| 6.015 | 93 | Kitasoo | 8 | 426 | 0.64 |  | 0.91 |  | 0.61 |  | 0.69 |  | 0.82 |  | 8 | 426 | 327.1 |  | 402.8 |  | 298.3 |  | 252.4 |  | 327.8 |  |
| 6.016 | 93 | Kitasoo | 11 | 881 | 0.70 |  | 1.07 |  | 0.69 |  | 0.95 |  | 1.03 |  | 11 | 881 | 309.4 |  | 412.2 |  | 300.0 |  | 334.4 |  | 383.2 |  |
| 6.016 | 95 | Price Is. | 27 | 1333 | 0.81 | 0.15 | 1.32 | 0.24 | 0.79 | 0.15 | 1.11 | 0.22 | 1.24 | 0.24 | 27 | 1333 | 390.8 | 71.1 | 534.4 | 97.2 | 372.9 | 67.7 | 389.4 | 78.8 | 469.7 | 88.5 |
| 6.017 | 93 | Kitasoo | 9 | 520 | 2.94 |  | 3.54 |  | 2.84 |  | 2.84 |  | 3.24 |  | 9 | 520 | 1424.0 |  | 1595.7 |  | 1328.2 |  | 1115.9 |  | 1352.9 |  |
| 6.017 | 95 | Price Is. | 9 | 554 | 0.78 | 0.38 | 1.06 | 0.48 | 0.75 | 0.37 | 0.72 | 0.34 | 0.89 | 0.40 | 9 | 554 | 419.1 | 213.9 | 499.5 | 240.3 | 393.9 | 198.1 | 268.7 | 129.5 | 370.3 | 169.1 |
| 6.018 | 93 | Kitasoo | 3 | 316 | 0.23 |  | 0.29 |  | 0.23 |  | 0.25 |  | 0.29 |  | 3 | 316 | 102.6 |  | 120.4 |  | 102.6 |  | 95.4 |  | 120.4 |  |
| 6.019 | 93 | Kitasoo | 6 | 566 | 0.45 |  | 0.64 |  | 0.45 |  | 0.59 |  | 0.62 |  | 6 | 566 | 189.4 |  | 241.5 |  | 189.4 |  | 213.2 |  | 227.9 |  |
| 7.001 | 97 | Goose | 15 | 1241 | 0.01 | 0.00 | 0.02 | 0.01 | 0.01 | 0.00 | 0.02 | 0.01 | 0.02 | 0.01 | 15 | 1241 | 2.9 | 1.7 | 5.8 | 2.9 | 2.9 | 1.7 | 5.8 | 2.9 | 5.8 | 2.9 |
| 7.002 | 93 | Kitasoo | 4 | 268 | 1.85 |  | 2.59 |  | 1.78 |  | 2.19 |  | 2.41 |  | 4 | 268 | 887.9 |  | 1093.7 |  | 820.8 |  | 827.8 |  | 956.7 |  |
| 7.003 | 93 | Kitasoo | 14 | 2086 | 1.10 |  | 1.46 |  | 1.09 |  | 1.22 |  | 1.40 |  | 14 | 2086 | 515.4 |  | 620.2 |  | 504.9 |  | 461.3 |  | 567.1 |  |
| 7.004 | 93 | Kitasoo | 4 | 916 | 0.61 |  | 0.78 |  | 0.60 |  | 0.62 |  | 0.74 |  | 4 | 916 | 295.4 |  | 342.7 |  | 281.8 |  | 238.3 |  | 313.5 |  |
| 7.008 | 93 | Heiltsuk | 5 | 3016 | 0.11 |  | 0.16 |  | 0.11 |  | 0.15 |  | 0.16 |  | 5 | 3016 | 47.9 |  | 64.0 |  | 47.9 |  | 52.5 |  | 64.0 |  |
| 7.018 | 93 | Heiltsuk | 13 | 1055 | 0.58 |  | 1.03 |  | 0.57 |  | 0.94 |  | 0.99 |  | 13 | 1055 | 271.5 |  | 397.7 |  | 253.6 |  | 329.5 |  | 357.7 |  |
| 7.018 | 94 | Heiltsuk | 26 | 1458 | 1.02 | 0.28 | 1.69 | 0.39 | 1.01 | 0.28 | 1.51 | 0.35 | 1.65 | 0.38 | 26 | 1458 | 467.8 | 130.0 | 655.6 | 156.5 | 452.7 | 127.9 | 541.0 | 128.5 | 620.6 | 149.2 |
| 7.018 | 95 | Heiltsuk | 36 | 2028 | 1.03 | 0.10 | 1.72 | 0.17 | 1.00 | 0.09 | 1.55 | 0.17 | 1.62 | 0.17 | 36 | 2028 | 479.3 | 49.0 | 671.6 | 65.0 | 450.8 | 43.9 | 544.7 | 56.2 | 589.9 | 58.6 |
| 7.018 | 96 | Heiltsuk | 58 | 5043 | 0.97 | 0.14 | 1.34 | 0.18 | 0.95 | 0.14 | 1.13 | 0.16 | 1.28 | 0.18 | 55 | 4777 | 475.5 | 66.2 | 587.6 | 77.5 | 461.2 | 65.3 | 442.2 | 61.7 | 534.8 | 70.6 |
| 7.018 | 97 | Goose | 23 | 1199 | 1.13 | 0.17 | 1.78 | 0.25 | 1.12 | 0.17 | 1.60 | 0.23 | 1.73 | 0.25 | 23 | 1199 | 500.8 | 77.5 | 685.8 | 98.6 | 494.7 | 76.8 | 570.1 | 82.7 | 645.0 | 92.9 |
| 7.020 | 93 | Heiltsuk | 4 | 471 | 0.52 |  | 1.04 |  | 0.52 |  | 0.96 |  | 1.04 |  | 4 | 471 | 216.5 |  | 359.8 |  | 216.5 |  | 315.6 |  | 359.8 |  |
| 7.023 | 94 | Heiltsuk | 1 | 36 | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  | 1 | 36 | 0.0 |  | 0.0 |  | 0.0 |  | 0.0 |  | 0.0 |  |
| 7.025 | 94 | Heiltsuk | 26 | 1286 | 1.28 | 0.25 | 1.70 | 0.32 | 1.26 | 0.25 | 1.35 | 0.27 | 1.60 | 0.30 | 25 | 1260 | 636.0 | 120.4 | 758.2 | 139.7 | 602.2 | 119.3 | 511.5 | 101.4 | 665.3 | 123.1 |
| 7.025 | 95 | Heiltsuk | 28 | 1882 | 0.50 | 0.16 | 0.75 | 0.23 | 0.48 | 0.15 | 0.66 | 0.20 | 0.72 | 0.22 | 28 | 1882 | 232.3 | 72.3 | 302.4 | 92.7 | 213.6 | 66.4 | 240.3 | 74.0 | 273.5 | 83.8 |
| 7.025 | 97 | Goose | 29 | 2729 | 0.02 | 0.02 | 0.03 | 0.03 | 0.02 | 0.02 | 0.02 | 0.02 | 0.03 | 0.03 | 29 | 2729 | 6.6 | 6.6 | 9.5 | 9.6 | 6.6 | 6.6 | 8.7 | 8.8 | 9.5 | 9.6 |
| 7.026 | 95 | Heiltsuk | 7 | 805 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 7 | 805 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 7.026 | 97 | Goose | 7 | 441 | 0.00 | 0.00 | 0.00 | 0.00 | 0.0 | 0.00 | 0.00 | 0.00 | 0.0 | 0.00 | 6 | 372 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 7.027 | 93 | Heiltsuk | 8 | 838 | 0.44 |  | 0.70 |  | 0.43 |  | 0.65 |  | 0.69 |  | 8 | 838 | 193.3 |  | 266.8 |  | 186.1 |  | 232.0 |  | 259.6 |  |
| 7.031 | 93 | kitasoo | 10 | 1026 | 1.32 |  | 1.66 |  | 1.30 |  | 1.35 |  | 1.55 |  | 10 | 1026 | 624.4 |  | 723.6 |  | 606.7 |  | 518.8 |  | 640.1 |  |
| 7.031 | 95 | Pricels | 32 | 1305 | 1.02 | 0.27 | 1.69 | 0.41 | 1.02 | 0.27 | 1.56 | 0.38 | 1.66 | 0.40 | 31 | 1288 | 442.5 | 120.0 | 632.1 | 157.7 | 438.6 | 119.3 | 551.0 | 136.5 | 609.5 | 154.2 |
| 7.032 | 93 | Heiltsuk | 4 | 730 | 0.58 |  | 0.94 |  | 0.58 |  | 0.89 |  | 0.93 |  | 4 | 730 | 252.5 |  | 352.7 |  | 252.5 |  | 319.0 |  | 342.5 |  |
| 10.001 | 93 | Heiltsuk | 5 | 2984 | 0.09 |  | 0.16 |  | 0.09 |  | 0.15 |  | 0.16 |  | 5 | 2984 | 38.0 |  | 58.9 |  | 37.0 |  | 52.8 |  | 58.0 |  |
| 106.002 | 94 | Campania Is. | 28 | 2633 | 1.75 | 0.26 | 2.46 | 0.34 | 1.72 | 0.25 | 2.13 | 0.30 | 2.34 | 0.32 | 28 | 2633 | 821.2 | 125.1 | 1020.8 | 146.5 | 783.4 | 116.8 | 789.3 | 111.4 | 914.5 | 124.0 |
| South Coast of B.C. |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 11.002 | 96 | QCStrait | 32 | 1360 | 1.10 | 0.23 | 1.24 | 0.26 | 0.91 | 0.21 | 0.42 | 0.11 | 0.72 | 0.17 | 31 | 1339 | 753.5 | 154.9 | 795.0 | 162.4 | 573.4 | 136.8 | 170.1 | 45.3 | 355.9 | 87.4 |
| 12.001 | 99 | JohnstoneSt | 13 | 627 | 0.48 | 0.19 | 0.54 | 0.21 | 0.44 | 0.17 | 0.30 | 0.13 | 0.41 | 0.17 | 13 | 627 | 273.7 | 102.9 | 289.3 | 108.7 | 232.2 | 89.7 | 121.4 | 53.5 | 183.6 | 76.2 |
| 12.002 | 99 | JohnstoneSt | 16 | 830 | 0.34 | 0.16 | 0.38 | 0.18 | 0.28 | 0.12 | 0.20 | 0.09 | 0.25 | 0.11 | 16 | 830 | 209.9 | 107.8 | 220.8 | 111.1 | 147.4 | 69.0 | 77.1 | 35.2 | 110.5 | 50.6 |
| 12.003 | 94 | QCStrait | 9 | 309 | 1.62 | 0.57 | 1.72 | 0.55 | 1.21 | 0.40 | 0.66 | 0.17 | 0.84 | 0.20 | 9 | 309 | 1120.6 | 429.1 | 1147.4 | 424.4 | 704.1 | 262.3 | 273.9 | 72.9 | 381.8 | 101.4 |
| 12.004 | 94 | QCStrait | 1 | 25 | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  | 1 | 25 | 0.0 |  | 0.0 |  | 0.0 |  | 0.0 |  | 0.0 |  |
| 12.005 | 94 | QCStrait | 11 | 484 | 1.00 | 0.24 | 1.01 | 0.24 | 0.64 | 0.20 | 0.12 | 0.05 | 0.41 | 0.14 | 9 | 381 | 963.7 | 130.8 | 969.6 | 131.1 | 522.2 | 134.4 | 62.4 | 26.1 | 287.5 | 87.9 |
| 12.006 | 94 | QCStrait | 16 | 1356 | 0.27 | 0.09 | 0.31 | 0.12 | 0.20 | 0.08 | 0.13 | 0.08 | 0.19 | 0.09 | 14 | 1292 | 199.7 | 66.5 | 212.6 | 71.4 | 124.1 | 47.1 | 55.5 | 32.8 | 88.7 | 42.2 |
| 12.007 | 95 | QCStrait | 6 | 225 | 1.54 | 0.55 | 1.58 | 0.57 | 1.18 | 0.54 | 0.31 | 0.19 | 0.73 | 0.32 | 6 | 225 | 1088.1 | 339.4 | 1097.9 | 345.5 | 742.9 | 336.6 | 131.7 | 78.1 | 385.5 | 160.0 |
| 12.008 | 94 | QCStrait | 4 | 485 | 0.09 |  | 0.13 |  | 0.06 |  | 0.07 |  | 0.08 |  | 3 | 280 | 114.1 |  | 132.8 |  | 57.1 |  | 46.0 |  | 57.3 |  |
| 12.011 | 94 | QCStrait | 15 | 713 | 1.51 | 0.33 | 1.83 | 0.40 | 1.35 | 0.30 | 1.23 | 0.27 | 1.50 | 0.33 | 15 | 713 | 843.0 | 201.3 | 935.6 | 216.7 | 688.8 | 161.0 | 486.4 | 107.8 | 652.6 | 144.2 |
| 12.012 | 94 | QCStrait | 2 | 69 | 1.73 |  | 2.08 |  | 1.73 |  | 1.60 |  | 1.87 |  | 2 | 69 | 849.3 |  | 939.6 |  | 849.3 |  | 624.7 |  | 786.3 |  |
| 12.013 | 95 | QCStrait | 21 | 1344 | 1.28 | 0.41 | 1.39 | 0.46 | 0.97 | 0.33 | 0.52 | 0.23 | 0.76 | 0.30 | 21 | 1344 | 871.0 | 281.3 | 902.4 | 290.3 | 575.4 | 190.0 | 213.0 | 95.3 | 360.9 | 140.4 |
| 12.013 | 96 | QCStrait | 11 | 332 | 1.25 | 0.53 | 1.54 | 0.53 | 1.06 | 0.47 | 0.87 | 0.27 | 1.08 | 0.38 | 10 | 269 | 944.2 | 426.3 | 1045.0 | 407.6 | 730.5 | 343.9 | 422.8 | 123.0 | 578.3 | 215.7 |
| 12.014 | 96 | CapeSutil | 8 | 809 | 0.85 | 0.70 | 1.21 | 1.03 | 0.82 | 0.67 | 1.03 | 0.88 | 1.11 | 0.94 | 6 | 609 | 530.4 | 437.1 | 660.4 | 555.6 | 489.0 | 396.3 | 493.4 | 419.8 | 557.1 | 462.3 |
| 12.015 | 94 | QCStrait | 2 | 57 | 2.40 |  | 3.20 |  | 2.40 |  | 2.61 |  | 3.20 |  | 2 | 57 | 1051.2 |  | 1270.9 |  | 1051.2 |  | 914.4 |  | 1270.9 |  |


| PMF Subarea | Year | Survey | Urchin density for each size group (mm TD) |  |  |  |  |  |  |  |  |  |  |  | Mean biomass for each size group (mm TD) |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | Total TransectNumber Length (m) |  | $\geq 100$ |  | $\geq 90$ |  | 100-140 |  | 90-120 |  | 90-130 |  | Total TransectNumber Length (m) |  | $\geq 100$ |  | $\geq 90$ |  | 100-140 |  | 90-120 |  | 90-130 |  |
|  |  |  |  |  | Mean | SE | Mean | SE | Mean | SE | Mean | SE | Mean | SE |  |  | Mean | SE | Mean | SE | Mean | SE | Mean | SE | Mean | SE |
| 12.016 | 94 | QCStrait | 20 | 1658 | 0.88 | 0.30 | 1.07 | 0.36 | 0.73 | 0.28 | 0.46 | 0.19 | 0.73 | 0.27 | 18 | 1488 | 654.4 | 208.3 | 714.5 | 226.1 | 475.5 | 183.6 | 186.2 | 75.4 | 368.4 | 136.1 |
| 12.017 | 94 | QCStrait | 2 | 443 | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  | 2 | 443 | 0.0 |  | 0.0 |  | 0.0 |  | 0.0 |  | 0.0 |  |
| 12.018 | 94 | QCStrait | 18 | 982 | 0.95 | 0.23 | 1.17 | 0.28 | 0.89 | 0.22 | 0.74 | 0.21 | 0.93 | 0.24 | 18 | 982 | 537.4 | 128.5 | 597.5 | 141.1 | 477.2 | 115.8 | 293.5 | 81.0 | 408.9 | 104.0 |
| 12.019 | 94 | QCStrait | 3 | 258 | 0.22 |  | 0.27 |  | 0.16 |  | 0.12 |  | 0.16 |  | 3 | 258 | 144.8 |  | 157.2 |  | 88.2 |  | 39.2 |  | 65.9 |  |
| 12.020 | 94 | QCStrait | 1 | 59 | 2.80 |  | 2.80 |  | 1.77 |  | 0.59 |  | 0.89 |  | 1 | 59 | 2021.8 |  | 2021.8 |  | 1087.4 |  | 226.8 |  | 423.9 |  |
| 12.021 | 94 | QCStrait | 2 | 93 | 1.70 |  | 1.83 |  | 1.58 |  | 0.76 |  | 1.25 |  | 2 | 93 | 1055.3 |  | 1095.5 |  | 933.9 |  | 310.8 |  | 621.6 |  |
| 12.024 | 99 | QCStrait | 2 | 72 | 0.48 |  | 0.51 |  | 0.43 |  | 0.27 |  | 0.40 |  | 2 | 72 | 288.5 |  | 295.1 |  | 233.8 |  | 117.9 |  | 199.4 |  |
| 12.036 | 94 | QCStrait | 1 | 83 | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  | 1 | 83 | 0.0 |  | 0.0 |  | 0.0 |  | 0.0 |  | 0.0 |  |
| 12.039 | 94 | QCStrait | 10 | 576 | 0.10 | 0.10 | 0.10 | 0.10 | 0.10 | 0.10 | 0.06 | 0.06 | 0.08 | 0.08 | 9 | 529 | 56.4 | 57.6 | 56.4 | 57.6 | 56.4 | 57.6 | 28.5 | 29.1 | 41.0 | 41.9 |
| 12.039 | 95 | QCStrait | 16 | 1203 | 0.07 | 0.04 | 0.07 | 0.04 | 0.04 | 0.02 | 0.01 | 0.01 | 0.02 | 0.01 | 15 | 1050 | 69.0 | 40.1 | 69.6 | 40.5 | 29.9 | 18.3 | 4.2 | 2.6 | 9.3 | 5.7 |
| 12.041 | 95 | QCStrait | 7 | 194 | 1.62 | 0.89 | 1.68 | 0.95 | 0.67 | 0.31 | 0.22 | 0.13 | 0.39 | 0.20 | 7 | 194 | 1383.2 | 782.5 | 1402.4 | 800.7 | 436.8 | 204.3 | 94.3 | 53.2 | 195.1 | 94.1 |
| 13.030 | 99 | Johnstone St | 4 | 162 | 0.30 |  | 0.33 |  | 0.27 |  | 0.24 |  | 0.27 |  | 4 | 162 | 159.5 |  | 170.1 |  | 139.6 |  | 100.1 |  | 122.8 |  |
| 13.031 | 99 | Johnstone St | 5 | 229 | 1.50 | 0.41 | 1.80 | 0.41 | 1.46 | 0.40 | 1.34 | 0.31 | 1.60 | 0.35 | 5 | 229 | 746.7 | 215.7 | 833.4 | 215.3 | 715.4 | 201.5 | 522.2 | 136.4 | 677.0 | 161.2 |
| 13.032 | 99 | Johnstone St | 18 | 868 | 1.23 | 0.30 | 1.42 | 0.33 | 1.19 | 0.28 | 0.95 | 0.22 | 1.24 | 0.27 | 18 | 868 | 642.1 | 158.3 | 697.1 | 167.2 | 606.2 | 146.1 | 384.9 | 89.5 | 555.0 | 120.7 |
| 13.033 | 99 | Johnstone St | 7 | 463 | 1.48 | 0.27 | 1.83 | 0.34 | 1.45 | 0.27 | 1.37 | 0.26 | 1.63 | 0.30 | 7 | 463 | 748.7 | 140.8 | 849.1 | 158.8 | 717.0 | 135.9 | 535.7 | 100.2 | 696.3 | 126.5 |
| 13.035 | 99 | Johnstone St | 9 | 319 | 0.65 | 0.30 | 0.66 | 0.30 | 0.43 | 0.23 | 0.18 | 0.09 | 0.28 | 0.15 | 9 | 319 | 473.7 | 213.9 | 477.1 | 214.9 | 260.0 | 141.8 | 78.5 | 37.7 | 139.0 | 75.1 |
| 14.005 | 99 | C. Georgia S | 2 | 202 | 0.23 |  | 0.40 |  | 0.23 |  | 0.34 |  | 0.39 |  | 2 | 202 | 112.9 |  | 160.5 |  | 104.6 |  | 122.4 |  | 152.2 |  |
| 14.007 | 99 | C. Georgia | 9 | 1383 | 0.27 | 0.13 | 0.34 | 0.15 | 0.25 | 0.12 | 0.22 | 0.10 | 0.29 | 0.14 | 9 | 1383 | 148.5 | 67.7 | 168.2 | 75.4 | 134.4 | 64.1 | 85.9 | 41.7 | 127.6 | 61.9 |
| 14.008 | 99 | C. Georgia | 6 | 812 | 0.44 | 0.14 | 0.52 | 0.16 | 0.42 | 0.13 | 0.37 | 0.12 | 0.44 | 0.14 | 6 | 812 | 232.1 | 78.0 | 256.7 | 83.0 | 215.7 | 70.8 | 147.1 | 48.8 | 191.9 | 61.8 |
| 14.009 | 99 | C. Georgia | 22 | 3776 | 0.25 | 0.06 | 0.33 | 0.07 | 0.24 | 0.06 | 0.26 | 0.06 | 0.30 | 0.06 | 22 | 3776 | 120.4 | 30.0 | 144.9 | 32.3 | 115.4 | 28.5 | 99.4 | 22.1 | 122.4 | 26.2 |
| 14.011 | 99 | C. Georgia | 17 | 1715 | 0.08 | 0.06 | 0.09 | 0.06 | 0.07 | 0.05 | 0.03 | 0.02 | 0.06 | 0.04 | 17 | 1715 | 52.3 | 35.9 | 54.6 | 37.5 | 40.9 | 29.2 | 11.5 | 8.2 | 29.5 | 21.2 |
| 14.012 | 99 | C. Georgia | 5 | 503 | 0.17 | 0.11 | 0.19 | 0.14 | 0.14 | 0.11 | 0.12 | 0.12 | 0.13 | 0.13 | 5 | 503 | 101.4 | 62.5 | 108.1 | 67.2 | 70.4 | 52.2 | 49.0 | 47.1 | 56.3 | 54.3 |
| 17.001 | 98 | S. Gulf Is. | 4 | 232 | 0.06 |  | 0.06 |  | 0.05 |  | 0.00 |  | 0.03 |  | 4 | 232 | 39.8 |  | 39.8 |  | 32.8 |  | 0.0 |  | 14.9 |  |
| 17.002 | 98 | S. Gulf Is. | 3 | 231 | 0.02 |  | 0.02 |  | 0.01 |  | 0.00 |  | 0.00 |  | 3 | 231 | 15.0 |  | 15.0 |  | 6.4 |  | 0.0 |  | 0.0 |  |
| 18.001 | 98 | S. Gulf Is. | 6 | 366 | 0.14 | 0.08 | 0.16 | 0.10 | 0.12 | 0.06 | 0.07 | 0.05 | 0.08 | 0.06 | 6 | 366 | 91.7 | 49.9 | 99.4 | 55.8 | 73.4 | 36.7 | 27.8 | 21.7 | 36.9 | 29.1 |
| 18.002 | 98 | S. Gulf Is. | 8 | 356 | 0.20 | 0.10 | 0.21 | 0.10 | 0.14 | 0.08 | 0.08 | 0.06 | 0.11 | 0.07 | 8 | 356 | 138.1 | 62.3 | 139.8 | 63.9 | 80.7 | 40.4 | 34.9 | 25.4 | 58.1 | 31.8 |
| 18.003 | 98 | S. Gulf Is. | 11 | 677 | 0.06 | 0.03 | 0.06 | 0.03 | 0.02 | 0.01 | 0.01 | 0.00 | 0.01 | 0.00 | 11 | 677 | 54.8 | 30.5 | 54.8 | 30.5 | 11.4 | 7.2 | 4.5 | 2.4 | 4.5 | 2.4 |
| 18.004 | 99 | S. Gulf Is. | 8 | 576 | 0.24 | 0.13 | 0.25 | 0.13 | 0.09 | 0.05 | 0.02 | 0.02 | 0.03 | 0.02 | 8 | 576 | 209.3 | 108.7 | 210.1 | 109.5 | 60.7 | 31.0 | 9.2 | 9.5 | 17.7 | 11.8 |
| 18.005 | 99 | S. Gulf Is. | 6 | 460 | 0.16 | 0.09 | 0.18 | 0.10 | 0.15 | 0.09 | 0.08 | 0.05 | 0.13 | 0.08 | 6 | 460 | 102.8 | 52.9 | 108.0 | 55.7 | 86.5 | 52.8 | 33.4 | 20.8 | 62.8 | 38.9 |
| 18.006 | 99 | S. Gulf Is. | 22 | 2706 | 0.37 | 0.13 | 0.39 | 0.14 | 0.24 | 0.11 | 0.09 | 0.04 | 0.17 | 0.08 | 22 | 2706 | 273.6 | 88.8 | 278.7 | 90.1 | 150.3 | 67.7 | 38.5 | 17.8 | 86.3 | 42.9 |
| 18.009 | 98 | S. Gulf Is. | 1 | 45 | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  | 0.00 |  | 1 | 45 | 0.0 |  | 0.0 |  | 0.0 |  | 0.0 |  | 0.0 |  |
| 18.011 | 98 | S. Gulf Is. | 4 | 176 | 0.64 |  | 0.72 |  | 0.49 |  | 0.18 |  | 0.40 |  | 4 | 176 | 444.5 |  | 467.1 |  | 309.8 |  | 67.0 |  | 207.2 |  |
| 111.000 | 96 | Cox Is. | 6 | 598 | 1.99 | 0.28 | 2.56 | 0.30 | 1.96 | 0.27 | 2.25 | 0.32 | 2.47 | 0.29 | 6 | 598 | 897.1 | 122.5 | 1055.6 | 126.0 | 869.2 | 112.4 | 847.7 | 122.5 | 982.0 | 108.9 |

Table 5. Summary of estimated mean density (number $/ \mathrm{m}^{2}$ ) and mean biomass ( $\mathrm{g} / \mathrm{m}^{2}$ ) of red sea urchins in British Columbia by PMF subarea, all years combined, obtained from broad-brush surveys during 1993-1999. Data include analyses from Jamieson et al. (1998a, b, c, d), Campbell et al. (1998) and present study. 2E and 2W refer to east and west areas of PMF area 2 in the Queen Charlotte Islands.
${ }^{a}=$ PFM subarea 12.014 not included.

| $\begin{gathered} \hline \text { PMF } \\ \text { Subarea } \end{gathered}$ | Year | Urchin density for each size group (mm TD) |  |  |  |  |  |  | Mean biomass for each size group (mm TD) |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | Total Transect |  | $\geq 100$ | $\geq 90$ | 100-140 | 90-120 | 90-130 | Total Transect |  | $\geq 100$ | $\geq 90$ | 100-140 | 90-120 | 90-130 |
|  |  | Number | Length (m) |  |  |  |  |  | Number | Length (m) |  |  |  |  |  |
| North coast of B.C. |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1 | 94 | 45 | 7103 | 1.39 | 1.64 | 1.28 | 1.01 | 1.35 | 45 | 7103 | 765.3 | 835.0 | 659.8 | 391.8 | 598.2 |
| 2E | 93-95 | 78 | 10049 | 0.48 | 0.63 | 0.45 | 0.44 | 0.54 | 78 | 10049 | 245.0 | 290.3 | 226.9 | 174.6 | 231.8 |
| 2W | 93-95 | 62 | 45208 | 0.65 | 1.02 | 0.61 | 0.84 | 0.93 | 61 | 4407 | 392.3 | 500.9 | 341.8 | 324.0 | 392.2 |
| 2 | 93-95 | 140 | 14557 | 0.55 | 0.77 | 0.52 | 0.58 | 0.67 | 139 | 14456 | 289.9 | 354.5 | 261.9 | 220.2 | 280.7 |
| 3 | 93 | 35 | 1223 | 0.68 | 0.89 | 0.63 | 0.61 | 0.75 | 35 | 1223 | 368.4 | 428.2 | 326.3 | 226.1 | 313.2 |
| 4 | 93-95 | 105 | 8343 | 0.59 | 0.77 | 0.53 | 0.50 | 0.63 | 104 | 7723 | 350.4 | 409.8 | 276.5 | 203.0 | 282.0 |
| 5 | 97 | 57 | 6328 | 1.06 | 1.36 | 0.95 | 0.95 | 1.13 | 57 | 6328 | 579.9 | 662.7 | 469.1 | 357.0 | 465.5 |
| 6 | 93-95 | 153 | 13344 | 0.94 | 1.27 | 0.89 | 0.98 | 1.13 | 153 | 13344 | 617.5 | 722.5 | 553.4 | 435.9 | 560.5 |
| 7 | 93-97 | 354 | 29859 | 0.69 | 1.01 | 0.67 | 0.88 | 0.97 | 348 | 29481 | 320.9 | 414.7 | 309.6 | 323.7 | 379.6 |
| 10 | 93 | 5 | 2984 | 0.087 | 0.163 | 0.086 | 0.153 | 0.162 | 5 | 2984 | 38.0 | 58.9 | 37.0 | 52.8 | 58.0 |
| 106 | 94 | 28 | 2633 | 1.75 | 2.46 | 1.72 | 2.13 | 2.34 | 28 | 2633 | 821.2 | 1020.8 | 783.4 | 789.3 | 914.5 |
| South Coast of B.C. |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 11 | 96 | 32 | 1360 | 1.10 | 1.24 | 0.91 | 0.42 | 0.72 | 31 | 1339 | 753.5 | 795.0 | 573.4 | 170.1 | 355.9 |
| 12.014 | 96 | 8 | 809 | 0.85 | 1.21 | 0.82 | 1.03 | 1.11 | 6 | 609 | 530.4 | 660.4 | 489.0 | 493.4 | 557.1 |
| 111 | 96 | 6 | 598 | 1.99 | 2.56 | 1.96 | 2.25 | 2.47 | 6 | 598 | 897.1 | 1055.6 | 869.2 | 847.7 | 982.0 |
| 12 a | 94-96 | 180 | 11020 | 0.77 | 0.88 | 0.61 | 0.40 | 0.56 | 170 | 10215 | 544.1 | 578.5 | 378.1 | 169.6 | 275.7 |
| 12 | 99 | 31 | 1529 | 0.41 | 0.45 | 0.35 | 0.24 | 0.32 | 31 | 1529 | 239.76 | 252.36 | 186.26 | 97.19 | 144.67 |
| 13 | 99 | 43 | 2041 | 1.15 | 1.35 | 1.09 | 0.91 | 1.14 | 43 | 2041 | 613.40 | 670.69 | 552.47 | 364.04 | 501.40 |
| 14 | 99 | 61 | 8391 | 0.23 | 0.29 | 0.22 | 0.21 | 0.26 | 61 | 8391 | 120.60 | 139.29 | 110.04 | 81.35 | 107.76 |
| 17 | 98 | 7 | 463 | 0.04 | 0.04 | 0.03 | 0.00 | 0.01 | 7 | 463 | 27.43 | 27.43 | 19.63 | 0.00 | 7.47 |
| 18 | 98-99 | 74 | 3570 | 0.28 | 0.29 | 0.18 | 0.07 | 0.13 | 74 | 3570 | 206.32 | 210.83 | 111.75 | 30.24 | 64.59 |

Table 6. Quota (tonnes) options for the red sea urchin fishery by PMF subarea, estimated from various natural mortality values applied to current biomass (Bc) calculated from mean and approximate $90 \%$ lower confidence interval (CI) biomass values $\left(\mathrm{g} / \mathrm{m}^{2}\right)$ for commercial red sea urchin sizes $100-140 \mathrm{~mm}$ TD, and bed areas fished up to 1996 NB : blank = average values directly from survey data within subarea; $a=$ average values combined for all years surveyed within subarea; $b=$ average values from all surveys in PMF area (may include several subareas and years); $c=$ average values of surveys in PMF area 7 used for subareas of PMF areas $8,9,10$;
$\mathrm{d}=$ data from 11.002 copied to 11.001 ; $\mathrm{e}=$ average values of surveys in PMF area 14 used in PMF area $15 ; \mathrm{f}=$ average values of PMF area 18
used in subareas of PMF areas 17, 19; g = average values of old surveys (1974-85) extrapolated for PMF areas 19, 20, 23, 24, 25, 26, 27, 29, 124, 125, after
Campbell et al. (1998), assumed transect length was 1000 m to calculate lower $90 \%$ CI of biomass density.

| PMF <br> Subarea | Year | NB | Transect |  | Biomass (g/m ${ }^{2}$ ) |  | Bed area (ha) 1996 | Bc Biomass (t) |  | Quota 0.2 M Bc (of urchins 100-140 mm TD) |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  | $\mathrm{M}=0.052$ | $\mathrm{M}=0.075$ |  |  |  | $\mathrm{M}=0.10$ |  | $\mathrm{M}=0.15$ |  |
|  |  |  | No | Length (m) |  |  | Mean | $\begin{gathered} \hline \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \hline \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \hline \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \end{gathered}$ | Mean | $\begin{gathered} \hline \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ |
| North Coast B.C. |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 1.001 | 94 |  | 5 | 764 | 729.8 | 409.2 |  | 4837.6 | 35304.8 | 19795.3 | 367.2 | 205.9 | 529.6 | 296.9 | 706.1 | 395.9 | 1059.1 | 593.9 |
| 1.002 | 94 |  | 15 | 1685 | 1369.7 | 964.5 |  | 405.0 | 5547.1 | 3906.2 | 57.7 | 40.6 | 83.2 | 58.6 | 110.9 | 78.1 | 166.4 | 117.2 |
| 1.003 | 94 |  | 16 | 1795 | 604.2 | 431.0 | 2024.2 | 12229.5 | 8724.5 | 127.2 | 90.7 | 183.4 | 130.9 | 244.6 | 174.5 | 366.9 | 261.7 |
| 1.005 | 94 | b | 45 | 7103 | 659.8 | 564.7 | 454.7 | 2999.8 | 2567.6 | 31.2 | 26.7 | 45.0 | 38.5 | 60.0 | 51.4 | 90.0 | 77.0 |
| 1.007 | 94 |  | 9 | 2859 | 257.6 | 199.1 | 1360.4 | 3504.1 | 2708.3 | 36.4 | 28.2 | 52.6 | 40.6 | 70.1 | 54.2 | 105.1 | 81.2 |
| 2.003 | 93 |  | 5 | 557 | 221.3 | 107.4 | 339.3 | 750.9 | 364.5 | 7.8 | 3.8 | 11.3 | 5.5 | 15.0 | 7.3 | 22.5 | 10.9 |
| 2.006 | 93-95 | b | 78 | 10049 | 226.9 | 199.4 | 298.0 | 676.1 | 594.2 | 7.0 | 6.2 | 10.1 | 8.9 | 13.5 | 11.9 | 20.3 | 17.8 |
| 2.007 | 93 |  | 6 | 476 | 447.8 | 198.6 | 469.4 | 2101.7 | 932.0 | 21.9 | 9.7 | 31.5 | 14.0 | 42.0 | 18.6 | 63.1 | 28.0 |
| 2.008 | 93 |  | 7 | 1070 | 254.4 | 160.0 | 331.9 | 844.3 | 531.0 | 8.8 | 5.5 | 12.7 | 8.0 | 16.9 | 10.6 | 25.3 | 15.9 |
| 2.010 | 93 |  | 4 | 557 | 218.4 | 106.1 | 2.2 | 4.9 | 2.4 | 0.1 | 0.0 | 0.1 | 0.0 | 0.1 | 0.0 | 0.1 | 0.1 |
| 2.011 | 93 |  | 8 | 589 | 393.0 | 196.4 | 973.4 | 3825.8 | 1911.7 | 39.8 | 19.9 | 57.4 | 28.7 | 76.5 | 38.2 | 114.8 | 57.4 |
| 2.012 | 93 |  | 6 | 3234 | 36.1 | 28.4 | 212.8 | 76.8 | 60.4 | 0.8 | 0.6 | 1.2 | 0.9 | 1.5 | 1.2 | 2.3 | 1.8 |
| 2.013 | 93-95 | b | 78 | 10049 | 226.9 | 199.4 | 26.2 | 59.5 | 52.3 | 0.6 | 0.5 | 0.9 | 0.8 | 1.2 | 1.0 | 1.8 | 1.6 |
| 2.014 | 93 |  | 9 | 1047 | 224.3 | 140.1 | 284.5 | 638.0 | 398.6 | 6.6 | 4.1 | 9.6 | 6.0 | 12.8 | 8.0 | 19.1 | 12.0 |
| 2.015 | 93 |  | 6 | 230 | 142.8 | 28.4 | 128.7 | 183.8 | 36.5 | 1.9 | 0.4 | 2.8 | 0.5 | 3.7 | 0.7 | 5.5 | 1.1 |
| 2.017 | 93 |  | 11 | 628 | 532.5 | 274.6 | 341.8 | 1819.9 | 938.4 | 18.9 | 9.8 | 27.3 | 14.1 | 36.4 | 18.8 | 54.6 | 28.2 |
| 2.018 | 95 |  | 11 | 949 | 358.1 | 216.9 | 291.1 | 1042.5 | 631.6 | 10.8 | 6.6 | 15.6 | 9.5 | 20.8 | 12.6 | 31.3 | 18.9 |
| 2.019 | 95 |  | 5 | 712 | 364.6 | 198.7 | 34.4 | 125.3 | 68.3 | 1.3 | 0.7 | 1.9 | 1.0 | 2.5 | 1.4 | 3.8 | 2.0 |
| 2.031 | 95 |  | 5 | 532 | 690.0 | 326.8 | 419.6 | 2895.4 | 1371.1 | 30.1 | 14.3 | 43.4 | 20.6 | 57.9 | 27.4 | 86.9 | 41.1 |
| 2.033 | 95 |  | 4 | 237 | 184.5 | 39.0 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 2.036 | 95 |  | 5 | 375 | 558.2 | 208.2 | 95.2 | 531.2 | 198.1 | 5.5 | 2.1 | 8.0 | 3.0 | 10.6 | 4.0 | 15.9 | 5.9 |
| 2.037 | 93-95 | b | 61 | 4407 | 341.8 | 279.3 | 29.0 | 99.0 | 80.9 | 1.0 | 0.8 | 1.5 | 1.2 | 2.0 | 1.6 | 3.0 | 2.4 |
| 2.049 | 93 |  | 2 | 642 | 29.7 | 15.4 | 193.4 | 57.3 | 29.9 | 0.6 | 0.3 | 0.9 | 0.4 | 1.1 | 0.6 | 1.7 | 0.9 |
| 2.050 | 93 |  | 2 | 70 | 532.7 | 0.0 | 146.2 | 779.1 | 0.0 | 8.1 | 0.0 | 11.7 | 0.0 | 15.6 | 0.0 | 23.4 | 0.0 |
| 2.051 | 93-95 | b | 61 | 4407 | 341.8 | 279.3 | 27.2 | 93.0 | 76.0 | 1.0 | 0.8 | 1.4 | 1.1 | 1.9 | 1.5 | 2.8 | 2.3 |
| 2.053 | 93 |  | 1 | 93 | 568.6 | 0.0 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |


| PMF <br> Subarea | Year | NB | Transect |  | Biomass ( $\mathrm{g} / \mathrm{m}^{2}$ ) |  | Bed area (ha) 1996 | Bc Biomass (t) |  | Quota 0.2 M Bc (of urchins 100-140 mm TD) |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  | $\mathrm{M}=0.052$ | $\mathrm{M}=0.075$ |  |  |  | $\mathrm{M}=0.10$ |  | $\mathrm{M}=0.15$ |  |
|  |  |  | No | Length (m) |  |  | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | Lower $90 \% \mathrm{CI}$ | Mean | Lower $90 \% \mathrm{CI}$ | Mean | $\begin{gathered} \hline \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | Lower $90 \% \mathrm{CI}$ | Mean | Lower $90 \% \mathrm{CI}$ |
| 2.055 | 93 |  | 1 | 26 | 138.1 | 0.0 |  |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 2.059 | 93 |  | 5 | 312 | 38.1 | 11.9 |  |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 2.060 | 93 |  | 3 | 386 | 36.4 | 13.9 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 2.063 | 93 |  | 6 | 207 | 421.7 | 66.1 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 2.064 | 93 |  | 2 | 52 | 110.8 | 0.0 | 29.4 | 32.5 | 0.0 | 0.3 | 0.0 | 0.5 | 0.0 | 0.7 | 0.0 | 1.0 | 0.0 |
| 2.065 | 93-95 | b | 61 | 4407 | 341.8 | 279.3 | 27.0 | 92.4 | 75.5 | 1.0 | 0.8 | 1.4 | 1.1 | 1.8 | 1.5 | 2.8 | 2.3 |
| 2.066 | 93-95 | b | 61 | 4407 | 341.8 | 279.3 | 5.8 | 19.8 | 16.2 | 0.2 | 0.2 | 0.3 | 0.2 | 0.4 | 0.3 | 0.6 | 0.5 |
| 2.067 | 93-95 | b | 61 | 4407 | 341.8 | 279.3 | 61.8 | 211.4 | 172.7 | 2.2 | 1.8 | 3.2 | 2.6 | 4.2 | 3.5 | 6.3 | 5.2 |
| 2.068 | 95 |  | 4 | 314 | 344.6 | 108.5 | 561.9 | 1936.3 | 609.5 | 20.1 | 6.3 | 29.0 | 9.1 | 38.7 | 12.2 | 58.1 | 18.3 |
| 2.069 | 95 |  | 1 | 71 | 0.0 | 0.0 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 2.071 | 95 |  | 2 | 124 | 466.1 | 0.0 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 2.074 | 95 |  | 1 | 201 | 463.9 | 66.6 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 2.075 | 95 |  | 4 | 411 | 853.5 | 342.3 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 2.078 | 95 |  | 2 | 65 | 0.0 | 0.0 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 2.079 | 95 |  | 2 | 200 | 117.2 | 16.6 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 2.080 | 95 |  | 1 | 89 | 232.8 | 0.0 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 2.087 | 93-95 | b | 61 | 4407 | 341.8 | 279.3 | 32.6 | 111.4 | 91.0 | 1.2 | 0.9 | 1.7 | 1.4 | 2.2 | 1.8 | 3.3 | 2.7 |
| 2.088 | 93-95 | b | 61 | 4407 | 341.8 | 279.3 | 61.9 | 211.5 | 172.8 | 2.2 | 1.8 | 3.2 | 2.6 | 4.2 | 3.5 | 6.3 | 5.2 |
| 2.094 | 93-95 | b | 61 | 4407 | 341.8 | 279.3 | 47.6 | 162.7 | 132.9 | 1.7 | 1.4 | 2.4 | 2.0 | 3.3 | 2.7 | 4.9 | 4.0 |
| 2.095 | 93-95 | b | 61 | 4407 | 341.8 | 279.3 | 68.8 | 235.3 | 192.3 | 2.4 | 2.0 | 3.5 | 2.9 | 4.7 | 3.8 | 7.1 | 5.8 |
| 2.096 | 93-95 | b | 61 | 4407 | 341.8 | 279.3 | 96.6 | 330.0 | 269.7 | 3.4 | 2.8 | 5.0 | 4.0 | 6.6 | 5.4 | 9.9 | 8.1 |
| 2.098 | 93-95 | b | 61 | 4407 | 341.8 | 279.3 | 98.8 | 337.8 | 276.0 | 3.5 | 2.9 | 5.1 | 4.1 | 6.8 | 5.5 | 10.1 | 8.3 |
| 2.100 | 93-95 | b | 61 | 4407 | 341.8 | 279.3 | 11.8 | 40.2 | 32.8 | 0.4 | 0.3 | 0.6 | 0.5 | 0.8 | 0.7 | 1.2 | 1.0 |
| 3.001 | 93 |  | 29 | 1011 | 363.0 | 224.4 | 724.2 | 2628.9 | 1625.1 | 27.3 | 16.9 | 39.4 | 24.4 | 52.6 | 32.5 | 78.9 | 48.8 |
| 3.002 | 93 |  | 6 | 212 | 150.8 | 25.0 | 33.0 | 49.8 | 8.2 | 0.5 | 0.1 | 0.7 | 0.1 | 1.0 | 0.2 | 1.5 | 0.2 |
| 3.003 | 93 | b | 35 | 1223 | 326.3 | 213.0 | 73.7 | 240.4 | 156.9 | 2.5 | 1.6 | 3.6 | 2.4 | 4.8 | 3.1 | 7.2 | 4.7 |
| 3.004 | 93 | b | 35 | 1223 | 326.3 | 213.0 | 24.4 | 79.5 | 51.9 | 0.8 | 0.5 | 1.2 | 0.8 | 1.6 | 1.0 | 2.4 | 1.6 |
| 4.001 | 93 |  | 34 | 1264 | 280.7 | 184.8 | 1074.0 | 3014.5 | 1985.1 | 31.4 | 20.6 | 45.2 | 29.8 | 60.3 | 39.7 | 90.4 | 59.6 |
| 4.002 | 93-95 | a | 41 | 2735 | 464.6 | 356.7 | 848.8 | 3943.3 | 3027.7 | 41.0 | 31.5 | 59.1 | 45.4 | 78.9 | 60.6 | 118.3 | 90.8 |
| 4.003 | 93-95 | b | 104 | 7723 | 276.5 | 238.3 | 875.2 | 2419.7 | 2085.4 | 25.2 | 21.7 | 36.3 | 31.3 | 48.4 | 41.7 | 72.6 | 62.6 |
| 4.004 | 93-95 | b | 104 | 7723 | 276.5 | 238.3 | 55.0 | 152.2 | 131.2 | 1.6 | 1.4 | 2.3 | 2.0 | 3.0 | 2.6 | 4.6 | 3.9 |
| 4.005 | 93-95 | b | 104 | 7723 | 276.5 | 238.3 | 85.5 | 236.3 | 203.7 | 2.5 | 2.1 | 3.5 | 3.1 | 4.7 | 4.1 | 7.1 | 6.1 |
| 4.009 | 95 |  | 23 | 3503 | 141.3 | 112.3 | 659.4 | 931.5 | 740.4 | 9.7 | 7.7 | 14.0 | 11.1 | 18.6 | 14.8 | 27.9 | 22.2 |
| 4.013 | 93 |  | 3 | 27 | 565.5 | 0.0 | 738.2 | 4174.5 | 0.0 | 43.4 | 0.0 | 62.6 | 0.0 | 83.5 | 0.0 | 125.2 | 0.0 |


| PMF <br> Subarea | Year | NB | Transect |  | Biomass (g/m ${ }^{2}$ ) |  | Bed area (ha) 1996 | Bc Biomass (t) |  | Quota 0.2 M Bc (of urchins 100-140 mm TD) |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  | $\mathrm{M}=0.052$ | $\mathrm{M}=0.075$ |  |  |  | $\mathrm{M}=0.10$ |  | $\mathrm{M}=0.15$ |  |
|  |  |  | NoLength <br> $(\mathrm{m})$ |  |  |  | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \hline \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \hline \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | Lower $90 \% \mathrm{CI}$ |
| 5.009 | 97 | b | 57 | 6328 | 469.1 | 397.5 |  | 342.5 | 1606.6 | 1361.3 | 16.7 | 14.2 | 24.1 | 20.4 | 32.1 | 27.2 | 48.2 | 40.8 |
| 5.010 | 97 | b | 57 | 6328 | 469.1 | 397.5 |  | 1506.4 | 7066.3 | 5987.7 | 73.5 | 62.3 | 106.0 | 89.8 | 141.3 | 119.8 | 212.0 | 179.6 |
| 5.011 | 97 |  | 6 | 910 | 113.4 | 67.8 | 557.0 | 631.6 | 377.4 | 6.6 | 3.9 | 9.5 | 5.7 | 12.6 | 7.5 | 18.9 | 11.3 |
| 5.012 | 97 | b | 57 | 6328 | 469.1 | 397.5 | 395.3 | 1854.4 | 1571.4 | 19.3 | 16.3 | 27.8 | 23.6 | 37.1 | 31.4 | 55.6 | 47.1 |
| 5.013 | 97 |  | 3 | 393 | 134.5 | 52.1 | 835.6 | 1124.1 | 435.6 | 11.7 | 4.5 | 16.9 | 6.5 | 22.5 | 8.7 | 33.7 | 13.1 |
| 5.014 | 97 | b | 57 | 6328 | 469.1 | 397.5 | 114.6 | 537.4 | 455.4 | 5.6 | 4.7 | 8.1 | 6.8 | 10.7 | 9.1 | 16.1 | 13.7 |
| 5.016 | 97 | b | 57 | 6328 | 469.1 | 397.5 | 525.8 | 2466.7 | 2090.2 | 25.7 | 21.7 | 37.0 | 31.4 | 49.3 | 41.8 | 74.0 | 62.7 |
| 5.017 | 97 | b | 57 | 6328 | 469.1 | 397.5 | 996.2 | 4673.0 | 3959.7 | 48.6 | 41.2 | 70.1 | 59.4 | 93.5 | 79.2 | 140.2 | 118.8 |
| 5.019 | 97 | b | 57 | 6328 | 469.1 | 397.5 | 3.7 | 17.5 | 14.8 | 0.2 | 0.2 | 0.3 | 0.2 | 0.3 | 0.3 | 0.5 | 0.4 |
| 5.020 | 97 |  | 26 | 2596 | 698.5 | 532.0 | 2174.2 | 15186.1 | 11567.0 | 157.9 | 120.3 | 227.8 | 173.5 | 303.7 | 231.3 | 455.6 | 347.0 |
| 5.021 | 97 |  | 22 | 2429 | 411.3 | 310.0 | 667.1 | 2743.9 | 2067.9 | 28.5 | 21.5 | 41.2 | 31.0 | 54.9 | 41.4 | 82.3 | 62.0 |
| 5.022 | 97 | b | 57 | 6328 | 469.1 | 397.5 | 2169.9 | 10178.9 | 8625.2 | 105.9 | 89.7 | 152.7 | 129.4 | 203.6 | 172.5 | 305.4 | 258.8 |
| 6.005 | 93-95 | b | 153 | 13344 | 553.4 | 495.2 | 65.7 | 363.7 | 325.4 | 3.8 | 3.4 | 5.5 | 4.9 | 7.3 | 6.5 | 10.9 | 9.8 |
| 6.009 | 93-95 | b | 153 | 13344 | 553.4 | 495.2 | 3230.7 | 17877.4 | 15998.2 | 185.9 | 166.4 | 268.2 | 240.0 | 357.5 | 320.0 | 536.3 | 479.9 |
| 6.010 | 94 |  | 30 | 5028 | 713.4 | 591.3 | 1671.7 | 11926.6 | 9884.2 | 124.0 | 102.8 | 178.9 | 148.3 | 238.5 | 197.7 | 357.8 | 296.5 |
| 6.011 | 93-95 | b | 153 | 13344 | 553.4 | 495.2 | 99.7 | 551.6 | 493.6 | 5.7 | 5.1 | 8.3 | 7.4 | 11.0 | 9.9 | 16.5 | 14.8 |
| 6.012 | 94 |  | 7 | 524 | 245.1 | 115.1 | 203.5 | 498.7 | 234.2 | 5.2 | 2.4 | 7.5 | 3.5 | 10.0 | 4.7 | 15.0 | 7.0 |
| 6.013 | 93 |  | 32 | 2434 | 549.0 | 413.9 | 2545.4 | 13973.8 | 10534.7 | 145.3 | 109.6 | 209.6 | 158.0 | 279.5 | 210.7 | 419.2 | 316.0 |
| 6.014 | 93 |  | 11 | 762 | 518.0 | 290.1 | 299.9 | 1553.2 | 869.8 | 16.2 | 9.0 | 23.3 | 13.0 | 31.1 | 17.4 | 46.6 | 26.1 |
| 6.015 | 93 |  | 8 | 426 | 298.3 | 122.7 | 520.1 | 1551.6 | 638.3 | 16.1 | 6.6 | 23.3 | 9.6 | 31.0 | 12.8 | 46.5 | 19.1 |
| 6.016 | 93-95 | a | 38 | 2214 | 343.9 | 255.2 | 903.8 | 3108.1 | 2306.1 | 32.3 | 24.0 | 46.6 | 34.6 | 62.2 | 46.1 | 93.2 | 69.2 |
| 6.017 | 93-95 | a | 18 | 1074 | 846.8 | 533.1 | 510.3 | 4321.3 | 2720.5 | 44.9 | 28.3 | 64.8 | 40.8 | 86.4 | 54.4 | 129.6 | 81.6 |
| 6.018 | 93 |  | 3 | 316 | 102.6 | 32.5 | 115.4 | 118.4 | 37.5 | 1.2 | 0.4 | 1.8 | 0.6 | 2.4 | 0.8 | 3.6 | 1.1 |
| 6.019 | 93 |  | 6 | 566 | 189.4 | 92.8 | 109.3 | 207.1 | 101.4 | 2.2 | 1.1 | 3.1 | 1.5 | 4.1 | 2.0 | 6.2 | 3.0 |
| 6.020 | 93-95 | b | 153 | 13344 | 553.4 | 495.2 | 29.8 | 164.9 | 147.6 | 1.7 | 1.5 | 2.5 | 2.2 | 3.3 | 3.0 | 4.9 | 4.4 |
| 6.025 | 93-95 | b | 153 | 13344 | 553.4 | 495.2 | 28.8 | 159.4 | 142.6 | 1.7 | 1.5 | 2.4 | 2.1 | 3.2 | 2.9 | 4.8 | 4.3 |
| 7.001 | 97 |  | 15 | 1241 | 2.9 | 1.9 | 55.2 | 1.6 | 1.1 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 7.002 | 93 |  | 4 | 268 | 820.8 | 211.9 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 7.003 | 93 |  | 14 | 2086 | 504.9 | 370.7 | 439.2 | 2217.7 | 1628.1 | 23.1 | 16.9 | 33.3 | 24.4 | 44.4 | 32.6 | 66.5 | 48.8 |
| 7.004 | 93 |  | 4 | 916 | 281.8 | 168.8 | 208.7 | 588.2 | 352.2 | 6.1 | 3.7 | 8.8 | 5.3 | 11.8 | 7.0 | 17.6 | 10.6 |
| 7.005 | 93-97 | b | 348 | 29481 | 309.6 | 287.7 | 56.5 | 175.0 | 162.6 | 1.8 | 1.7 | 2.6 | 2.4 | 3.5 | 3.3 | 5.2 | 4.9 |
| 7.006 | 93-97 | b | 348 | 29481 | 309.6 | 287.7 | 142.4 | 440.8 | 409.6 | 4.6 | 4.3 | 6.6 | 6.1 | 8.8 | 8.2 | 13.2 | 12.3 |
| 7.008 | 93 |  | 5 | 3016 | 47.9 | 37.3 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 7.009 | 93-97 | b | 348 | 29481 | 309.6 | 287.7 | 713.1 | 2207.5 | 2051.4 | 23.0 | 21.3 | 33.1 | 30.8 | 44.2 | 41.0 | 66.2 | 61.5 |


| PMF <br> Subarea | Year | NB | Transect |  | Biomass (g/m ${ }^{2}$ ) |  | Bed area (ha) 1996 | Bc Biomass (t) |  | Quota 0.2 M Bc (of urchins 100-140 mm TD) |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  | $\mathrm{M}=0.052$ | $\mathrm{M}=0.075$ |  |  |  | $\mathrm{M}=0.10$ |  | $\mathrm{M}=0.15$ |  |
|  |  |  | No | Length (m) |  |  | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | Lower $90 \%$ CI | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \text { CI } \end{gathered}$ |
| 7.012 | 93-97 | b | 348 | 29481 | 309.6 | 287.7 |  | 148.8 | 460.7 | 428.2 | 4.8 | 4.5 | 6.9 | 6.4 | 9.2 | 8.6 | 13.8 | 12.8 |
| 7.017 | 93-97 | b | 348 | 29481 | 309.6 | 287.7 |  | 3.5 | 10.8 | 10.0 | 0.1 | 0.1 | 0.2 | 0.2 | 0.2 | 0.2 | 0.3 | 0.3 |
| 7.018 | 93-97 | a | 153 | 10517 | 441.0 | 388.8 | 1294.3 | 5708.3 | 5032.4 | 59.4 | 52.3 | 85.6 | 75.5 | 114.2 | 100.6 | 171.2 | 151.0 |
| 7.019 | 93-97 | b | 348 | 29481 | 309.6 | 287.7 | 94.6 | 292.8 | 272.1 | 3.0 | 2.8 | 4.4 | 4.1 | 5.9 | 5.4 | 8.8 | 8.2 |
| 7.020 | 93 |  | 4 | 471 | 216.5 | 95.4 | 51.8 | 112.1 | 49.4 | 1.2 | 0.5 | 1.7 | 0.7 | 2.2 | 1.0 | 3.4 | 1.5 |
| 7.021 | 93-97 | b | 348 | 29481 | 309.6 | 287.7 | 40.1 | 124.1 | 115.3 | 1.3 | 1.2 | 1.9 | 1.7 | 2.5 | 2.3 | 3.7 | 3.5 |
| 7.023 | 93-97 | b | 348 | 29481 | 309.6 | 287.7 | 51.6 | 159.6 | 148.3 | 1.7 | 1.5 | 2.4 | 2.2 | 3.2 | 3.0 | 4.8 | 4.4 |
| 7.024 | 93-97 | b | 348 | 29481 | 309.6 | 287.7 | 2.6 | 8.2 | 7.6 | 0.1 | 0.1 | 0.1 | 0.1 | 0.2 | 0.2 | 0.2 | 0.2 |
| 7.025 | 94-97 | a | 82 | 5871 | 200.8 | 168.9 | 796.0 | 1598.1 | 1344.8 | 16.6 | 14.0 | 24.0 | 20.2 | 32.0 | 26.9 | 47.9 | 40.3 |
| 7.026 | 95-97 | a | 13 | 1177 | 0.0 | 0.0 | 52.2 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 7.027 | 93 |  | 8 | 838 | 186.1 | 108.1 | 204.1 | 379.9 | 220.6 | 4.0 | 2.3 | 5.7 | 3.3 | 7.6 | 4.4 | 11.4 | 6.6 |
| 7.028 | 93-97 | b | 348 | 29481 | 309.6 | 287.7 | 19.4 | 60.2 | 55.9 | 0.6 | 0.6 | 0.9 | 0.8 | 1.2 | 1.1 | 1.8 | 1.7 |
| 7.031 | 93-95 | a | 41 | 2314 | 513.1 | 383.5 | 1107.1 | 5680.1 | 4246.3 | 59.1 | 44.2 | 85.2 | 63.7 | 113.6 | 84.9 | 170.4 | 127.4 |
| 7.032 | 93 |  | 4 | 730 | 252.5 | 139.0 | 308.2 | 778.3 | 428.4 | 8.1 | 4.5 | 11.7 | 6.4 | 15.6 | 8.6 | 23.3 | 12.9 |
| 8.001 | 93-97 | c | 348 | 29481 | 309.6 | 287.7 | 27.1 | 83.8 | 77.9 | 0.9 | 0.8 | 1.3 | 1.2 | 1.7 | 1.6 | 2.5 | 2.3 |
| 8.002 | 93-97 | c | 348 | 29481 | 309.6 | 287.7 | 87.5 | 270.8 | 251.7 | 2.8 | 2.6 | 4.1 | 3.8 | 5.4 | 5.0 | 8.1 | 7.5 |
| 8.003 | 93-97 | c | 348 | 29481 | 309.6 | 287.7 | 5.5 | 17.0 | 15.8 | 0.2 | 0.2 | 0.3 | 0.2 | 0.3 | 0.3 | 0.5 | 0.5 |
| 8.004 | 93-97 | c | 348 | 29481 | 309.6 | 287.7 | 156.2 | 483.5 | 449.3 | 5.0 | 4.7 | 7.3 | 6.7 | 9.7 | 9.0 | 14.5 | 13.5 |
| 8.016 | 93-97 | c | 348 | 29481 | 309.6 | 287.7 | 64.6 | 200.1 | 185.9 | 2.1 | 1.9 | 3.0 | 2.8 | 4.0 | 3.7 | 6.0 | 5.6 |
| 9.001 | 93-97 | c | 348 | 29481 | 309.6 | 287.7 | 138.6 | 429.0 | 398.7 | 4.5 | 4.1 | 6.4 | 6.0 | 8.6 | 8.0 | 12.9 | 12.0 |
| 9.002 | 93-97 | c | 348 | 29481 | 309.6 | 287.7 | 232.2 | 718.8 | 668.0 | 7.5 | 6.9 | 10.8 | 10.0 | 14.4 | 13.4 | 21.6 | 20.0 |
| 9.010 | 93-97 | c | 348 | 29481 | 309.6 | 287.7 | 26.3 | 81.4 | 75.6 | 0.8 | 0.8 | 1.2 | 1.1 | 1.6 | 1.5 | 2.4 | 2.3 |
| 9.011 | 93-97 | c | 348 | 29481 | 309.6 | 287.7 | 5.6 | 17.2 | 16.0 | 0.2 | 0.2 | 0.3 | 0.2 | 0.3 | 0.3 | 0.5 | 0.5 |
| 9.012 | 93-97 | c | 348 | 29481 | 309.6 | 287.7 | 103.1 | 319.1 | 296.5 | 3.3 | 3.1 | 4.8 | 4.4 | 6.4 | 5.9 | 9.6 | 8.9 |
| 10.001 | 93 |  | 5 | 2984 | 37.0 | 28.8 | 266.1 | 98.6 | 76.6 | 1.0 | 0.8 | 1.5 | 1.1 | 2.0 | 1.5 | 3.0 | 2.3 |
| 10.002 | 93-97 | c | 348 | 29481 | 309.6 | 287.7 | 485.8 | 1503.8 | 1397.5 | 15.6 | 14.5 | 22.6 | 21.0 | 30.1 | 27.9 | 45.1 | 41.9 |
| 10.003 | 93-97 | c | 348 | 29481 | 309.6 | 287.7 | 214.8 | 665.1 | 618.0 | 6.9 | 6.4 | 10.0 | 9.3 | 13.3 | 12.4 | 20.0 | 18.5 |
| 10.004 | 93-97 | c | 348 | 29481 | 309.6 | 287.7 | 74.0 | 229.2 | 213.0 | 2.4 | 2.2 | 3.4 | 3.2 | 4.6 | 4.3 | 6.9 | 6.4 |
| 105.001 | 97 | b | 57 | 6328 | 469.1 | 397.5 | 50.3 | 236.1 | 200.0 | 2.5 | 2.1 | 3.5 | 3.0 | 4.7 | 4.0 | 7.1 | 6.0 |
| 105.002 | 97 | b | 57 | 6328 | 469.1 | 397.5 | 6.6 | 30.8 | 26.1 | 0.3 | 0.3 | 0.5 | 0.4 | 0.6 | 0.5 | 0.9 | 0.8 |
| 106.002 | 94 |  | 28 | 2633 | 783.4 | 598.0 | 1827.9 | 14319.7 | 10931.1 | 148.9 | 113.7 | 214.8 | 164.0 | 286.4 | 218.6 | 429.6 | 327.9 |
|  |  |  |  |  | N. Coast | Total | 50205.2 | 242951.9 | 173846.6 | 2526.7 | 1808.0 | 3644.3 | 2607.7 | 4859.0 | 3476.9 | 7288.6 | 5215.4 |
| South Coast of British Columbia |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| 11.001 | 96 | d | 31 | 1339 | 573.4 | 383.1 | 135.9 | 779.1 | 520.6 | 8.1 | 5.4 | 11.7 | 7.8 | 15.6 | 10.4 | 23.4 | 15.6 |


| PMF <br> Subarea | Year | NB | Transect |  | $\text { Biomass }\left(\mathrm{g} / \mathrm{m}^{2}\right)$ |  | Bed area (ha) 1996 | Bc Biomass (t) |  | Quota 0.2 M Bc (of urchins 100-140 mm TD) |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  | $\mathrm{M}=0.052$ | $\mathrm{M}=0.075$ |  |  |  | $\mathrm{M}=0.10$ |  | $\mathrm{M}=0.15$ |  |
|  |  |  | NoLength <br> $(\mathrm{m})$ |  |  |  | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \hline \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \hline \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \hline \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | Lower $90 \% \mathrm{CI}$ |
| 11.002 | 96 |  | 31 | 1339 | 573.4 | 383.1 |  | 142.6 | 817.9 | 546.5 | 8.5 | 5.7 | 12.3 | 8.2 | 16.4 | 10.9 | 24.5 | 16.4 |
| 12.001 | 99 |  | 13 | 627 | 232.2 | 119.6 |  | 61.6 | 143.0 | 73.7 | 1.5 | 0.8 | 2.1 | 1.1 | 2.9 | 1.5 | 4.3 | 2.2 |
| 12.002 | 99 |  | 16 | 830 | 147.4 | 85.3 | 96.5 | 142.3 | 82.3 | 1.5 | 0.9 | 2.1 | 1.2 | 2.8 | 1.6 | 4.3 | 2.5 |
| 12.003 | 94 |  | 9 | 309 | 704.1 | 217.7 | 180.6 | 1271.6 | 393.2 | 13.2 | 4.1 | 19.1 | 5.9 | 25.4 | 7.9 | 38.1 | 11.8 |
| 12.004 | 94 |  | 1 | 25 | 0.0 | 0.0 | 2.3 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 12.005 | 94 |  | 9 | 381 | 522.2 | 197.4 | 143.2 | 747.8 | 282.6 | 7.8 | 2.9 | 11.2 | 4.2 | 15.0 | 5.7 | 22.4 | 8.5 |
| 12.006 | 94 |  | 14 | 1292 | 124.1 | 82.2 | 146.1 | 181.3 | 120.0 | 1.9 | 1.2 | 2.7 | 1.8 | 3.6 | 2.4 | 5.4 | 3.6 |
| 12.007 | 95 |  | 6 | 225 | 742.9 | 141.5 | 73.5 | 545.9 | 104.0 | 5.7 | 1.1 | 8.2 | 1.6 | 10.9 | 2.1 | 16.4 | 3.1 |
| 12.008 | 94 |  | 3 | 280 | 57.1 | 15.7 | 93.0 | 53.1 | 14.6 | 0.6 | 0.2 | 0.8 | 0.2 | 1.1 | 0.3 | 1.6 | 0.4 |
| 12.011 | 94 |  | 15 | 713 | 688.8 | 375.6 | 239.4 | 1648.9 | 899.1 | 17.1 | 9.4 | 24.7 | 13.5 | 33.0 | 18.0 | 49.5 | 27.0 |
| 12.012 | 94 |  | 2 | 69 | 849.3 | 0.0 | 172.6 | 1465.9 | 0.0 | 15.2 | 0.0 | 22.0 | 0.0 | 29.3 | 0.0 | 44.0 | 0.0 |
| 12.013 | 95-96 | a | 31 | 1613 | 601.2 | 419.4 | 117.3 | 705.4 | 492.1 | 7.3 | 5.1 | 10.6 | 7.4 | 14.1 | 9.8 | 21.2 | 14.8 |
| 12.014 | 96 |  | 6 | 609 | 489.0 | 248.4 | 425.3 | 2079.8 | 1056.5 | 21.6 | 11.0 | 31.2 | 15.8 | 41.6 | 21.1 | 62.4 | 31.7 |
| 12.015 | 94 |  | 2 | 57 | 1051.2 | 0.0 | 27.6 | 289.7 | 0.0 | 3.0 | 0.0 | 4.3 | 0.0 | 5.8 | 0.0 | 8.7 | 0.0 |
| 12.016 | 94 |  | 18 | 1488 | 475.5 | 325.8 | 303.7 | 1444.1 | 989.6 | 15.0 | 10.3 | 21.7 | 14.8 | 28.9 | 19.8 | 43.3 | 29.7 |
| 12.017 | 94 |  | 2 | 443 | 0.0 | 0.0 | 39.1 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 12.018 | 94 |  | 18 | 982 | 477.2 | 292.3 | 350.0 | 1670.3 | 1023.1 | 17.4 | 10.6 | 25.1 | 15.3 | 33.4 | 20.5 | 50.1 | 30.7 |
| 12.019 | 94 |  | 3 | 258 | 88.2 | 21.5 | 65.7 | 57.9 | 14.1 | 0.6 | 0.1 | 0.9 | 0.2 | 1.2 | 0.3 | 1.7 | 0.4 |
| 12.020 | 94 |  | 1 | 59 | 1087.4 | 0.0 | 1.4 | 15.2 | 0.0 | 0.2 | 0.0 | 0.2 | 0.0 | 0.3 | 0.0 | 0.5 | 0.0 |
| 12.021 | 94 |  | 2 | 93 | 933.9 | 0.0 | 17.6 | 164.7 | 0.0 | 1.7 | 0.0 | 2.5 | 0.0 | 3.3 | 0.0 | 4.9 | 0.0 |
| 12.024 | 99 |  | 2 | 72 | 233.8 | 0.0 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 12.026 | 94-96 | b | 170 | 10215 | 378.1 | 332.7 | 4.0 | 15.0 | 13.2 | 0.2 | 0.1 | 0.2 | 0.2 | 0.3 | 0.3 | 0.4 | 0.4 |
| 12.036 | 94 |  | 1 | 83 | 0.0 | 0.0 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 12.039 | 94-95 | a | 24 | 1579 | 38.8 | 26.9 | 59.8 | 23.2 | 16.1 | 0.2 | 0.2 | 0.3 | 0.2 | 0.5 | 0.3 | 0.7 | 0.5 |
| 12.041 | 95 |  | 7 | 194 | 436.8 | 56.0 | 86.8 | 379.0 | 48.6 | 3.9 | 0.5 | 5.7 | 0.7 | 7.6 | 1.0 | 11.4 | 1.5 |
| 12.042 | 94-96 | b | 170 | 10215 | 378.1 | 332.7 | 18.0 | 68.1 | 59.9 | 0.7 | 0.6 | 1.0 | 0.9 | 1.4 | 1.2 | 2.0 | 1.8 |
| 13.001 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 170.2 | 940.5 | 687.7 | 9.8 | 7.2 | 14.1 | 10.3 | 18.8 | 13.8 | 28.2 | 20.6 |
| 13.002 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 74.7 | 412.6 | 301.7 | 4.3 | 3.1 | 6.2 | 4.5 | 8.3 | 6.0 | 12.4 | 9.1 |
| 13.003 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 40.7 | 225.0 | 164.5 | 2.3 | 1.7 | 3.4 | 2.5 | 4.5 | 3.3 | 6.7 | 4.9 |
| 13.006 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 60.8 | 336.1 | 245.8 | 3.5 | 2.6 | 5.0 | 3.7 | 6.7 | 4.9 | 10.1 | 7.4 |
| 13.007 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 51.1 | 282.2 | 206.4 | 2.9 | 2.1 | 4.2 | 3.1 | 5.6 | 4.1 | 8.5 | 6.2 |
| 13.008 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 8.4 | 46.6 | 34.1 | 0.5 | 0.4 | 0.7 | 0.5 | 0.9 | 0.7 | 1.4 | 1.0 |
| 13.009 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 19.3 | 106.7 | 78.0 | 1.1 | 0.8 | 1.6 | 1.2 | 2.1 | 1.6 | 3.2 | 2.3 |
| 13.010 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 72.3 | 399.5 | 292.2 | 4.2 | 3.0 | 6.0 | 4.4 | 8.0 | 5.8 | 12.0 | 8.8 |


| $\begin{gathered} \hline \text { PMF } \\ \text { Subarea } \end{gathered}$ | Year | NB | Transect |  | Biomass ( $\mathrm{g} / \mathrm{m}^{2}$ ) |  | Bed area (ha) 1996 | Bc Biomass (t) |  | Quota 0.2 M Bc (of urchins 100-140 mm TD) |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  | $\mathrm{M}=0.052$ | $\mathrm{M}=0.075$ |  |  |  | $\mathrm{M}=0.10$ |  | $\mathrm{M}=0.15$ |  |
|  |  |  | No | Length (m) |  |  | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | Lower $90 \% \mathrm{CI}$ | Mean | Lower $90 \% \mathrm{CI}$ | Mean | $\begin{gathered} \hline \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | Lower $90 \% \mathrm{CI}$ | Mean | Lower $90 \% \mathrm{CI}$ |
| 13.011 | 99 | b | 43 | 2041 | 552.5 | 404.0 |  | 28.8 | 159.3 | 116.5 | 1.7 | 1.2 | 2.4 | 1.7 | 3.2 | 2.3 | 4.8 | 3.5 |
| 13.012 | 99 | b | 43 | 2041 | 552.5 | 404.0 |  | 142.1 | 785.2 | 574.1 | 8.2 | 6.0 | 11.8 | 8.6 | 15.7 | 11.5 | 23.6 | 17.2 |
| 13.016 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 14.9 | 82.2 | 60.1 | 0.9 | 0.6 | 1.2 | 0.9 | 1.6 | 1.2 | 2.5 | 1.8 |
| 13.017 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 8.9 | 49.3 | 36.0 | 0.5 | 0.4 | 0.7 | 0.5 | 1.0 | 0.7 | 1.5 | 1.1 |
| 13.023 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 66.1 | 365.3 | 267.1 | 3.8 | 2.8 | 5.5 | 4.0 | 7.3 | 5.3 | 11.0 | 8.0 |
| 13.025 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 107.6 | 594.5 | 434.7 | 6.2 | 4.5 | 8.9 | 6.5 | 11.9 | 8.7 | 17.8 | 13.0 |
| 13.026 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 30.5 | 168.4 | 123.1 | 1.8 | 1.3 | 2.5 | 1.8 | 3.4 | 2.5 | 5.1 | 3.7 |
| 13.027 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 31.0 | 171.5 | 125.4 | 1.8 | 1.3 | 2.6 | 1.9 | 3.4 | 2.5 | 5.1 | 3.8 |
| 13.028 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 152.1 | 840.2 | 614.4 | 8.7 | 6.4 | 12.6 | 9.2 | 16.8 | 12.3 | 25.2 | 18.4 |
| 13.029 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 17.6 | 97.0 | 70.9 | 1.0 | 0.7 | 1.5 | 1.1 | 1.9 | 1.4 | 2.9 | 2.1 |
| 13.030 | 99 |  | 4 | 162 | 139.6 | 6.4 | 84.1 | 117.4 | 5.4 | 1.2 | 0.1 | 1.8 | 0.1 | 2.3 | 0.1 | 3.5 | 0.2 |
| 13.031 | 99 |  | 5 | 229 | 715.4 | 141.4 | 29.1 | 208.3 | 41.2 | 2.2 | 0.4 | 3.1 | 0.6 | 4.2 | 0.8 | 6.2 | 1.2 |
| 13.032 | 99 |  | 18 | 868 | 606.2 | 356.4 | 165.0 | 1000.0 | 587.9 | 10.4 | 6.1 | 15.0 | 8.8 | 20.0 | 11.8 | 30.0 | 17.6 |
| 13.033 | 99 |  | 7 | 463 | 717.0 | 312.4 | 53.2 | 381.5 | 166.2 | 4.0 | 1.7 | 5.7 | 2.5 | 7.6 | 3.3 | 11.4 | 5.0 |
| 13.035 | 99 |  | 9 | 319 | 260.0 | 83.3 | 71.9 | 187.0 | 59.9 | 1.9 | 0.6 | 2.8 | 0.9 | 3.7 | 1.2 | 5.6 | 1.8 |
| 13.036 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 46.1 | 254.6 | 186.2 | 2.6 | 1.9 | 3.8 | 2.8 | 5.1 | 3.7 | 7.6 | 5.6 |
| 13.039 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 62.6 | 345.8 | 252.9 | 3.6 | 2.6 | 5.2 | 3.8 | 6.9 | 5.1 | 10.4 | 7.6 |
| 13.040 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 23.3 | 128.8 | 94.2 | 1.3 | 1.0 | 1.9 | 1.4 | 2.6 | 1.9 | 3.9 | 2.8 |
| 13.041 | 99 | b | 43 | 2041 | 552.5 | 404.0 | 18.7 | 103.2 | 75.5 | 1.1 | 0.8 | 1.5 | 1.1 | 2.1 | 1.5 | 3.1 | 2.3 |
| 14.005 | 99 |  | 2 | 202 | 104.6 | 15.2 |  | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 14.007 | 99 |  | 9 | 1383 | 134.4 | 90.5 | 79.9 | 107.4 | 72.3 | 1.1 | 0.8 | 1.6 | 1.1 | 2.1 | 1.4 | 3.2 | 2.2 |
| 14.008 | 99 |  | 6 | 812 | 215.7 | 123.8 | 22.0 | 47.4 | 27.2 | 0.5 | 0.3 | 0.7 | 0.4 | 0.9 | 0.5 | 1.4 | 0.8 |
| 14.009 | 99 |  | 22 | 3776 | 115.4 | 92.6 | 205.4 | 236.9 | 190.1 | 2.5 | 2.0 | 3.6 | 2.9 | 4.7 | 3.8 | 7.1 | 5.7 |
| 14.010 | 99 | b | 61 | 8391 | 110.0 | 95.5 | 226.0 | 248.7 | 215.7 | 2.6 | 2.2 | 3.7 | 3.2 | 5.0 | 4.3 | 7.5 | 6.5 |
| 14.011 | 99 |  | 17 | 1715 | 40.9 | 28.9 | 75.9 | 31.1 | 22.0 | 0.3 | 0.2 | 0.5 | 0.3 | 0.6 | 0.4 | 0.9 | 0.7 |
| 14.012 | 99 |  | 5 | 503 | 70.4 | 32.3 | 55.8 | 39.3 | 18.0 | 0.4 | 0.2 | 0.6 | 0.3 | 0.8 | 0.4 | 1.2 | 0.5 |
| 14.013 | 99 | b | 61 | 8391 | 110.0 | 95.5 | 182.3 | 200.6 | 174.0 | 2.1 | 1.8 | 3.0 | 2.6 | 4.0 | 3.5 | 6.0 | 5.2 |
| 15.001 | 99 | e | 61 | 8391 | 110.0 | 95.5 | 6.7 | 7.4 | 6.4 | 0.1 | 0.1 | 0.1 | 0.1 | 0.1 | 0.1 | 0.2 | 0.2 |
| 15.002 | 99 | e | 61 | 8391 | 110.0 | 95.5 | 25.8 | 28.4 | 24.6 | 0.3 | 0.3 | 0.4 | 0.4 | 0.6 | 0.5 | 0.9 | 0.7 |
| 15.004 | 99 | e | 61 | 8391 | 110.0 | 95.5 | 4.4 | 4.9 | 4.2 | 0.1 | 0.0 | 0.1 | 0.1 | 0.1 | 0.1 | 0.1 | 0.1 |
| 17.001 | 98 |  | 4 | 232 | 32.8 | 6.6 | 1.2 | 0.4 | 0.1 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 17.002 | 98 |  | 3 | 231 | 6.4 | 1.3 | 65.8 | 4.2 | 0.8 | 0.0 | 0.0 | 0.1 | 0.0 | 0.1 | 0.0 | 0.1 | 0.0 |
| 17.003 | 98-99 | f | 74 | 3570 | 111.8 | 89.0 | 69.7 | 77.9 | 62.1 | 0.8 | 0.6 | 1.2 | 0.9 | 1.6 | 1.2 | 2.3 | 1.9 |
| 17.008 | 98-99 | f | 74 | 3570 | 111.8 | 89.0 | 30.1 | 33.6 | 26.8 | 0.3 | 0.3 | 0.5 | 0.4 | 0.7 | 0.5 | 1.0 | 0.8 |


| $\overline{\mathrm{PMF}}$ <br> Subarea | Year | NB | Transect |  | Biomass (g/m ${ }^{2}$ ) |  | $\begin{gathered} \text { Bed } \\ \text { area (ha) } \\ 1996 \\ \hline \end{gathered}$ | Bc Biomass (t) |  | Quota 0.2 M Bc (of urchins 100-140 mm TD) |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  | $\mathrm{M}=0.052$ | $\mathrm{M}=0.075$ |  |  |  | $\mathrm{M}=0.10$ |  | $\mathrm{M}=0.15$ |  |
|  |  |  | No | Length (m) |  |  | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \hline \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \hline \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \mathrm{CI} \\ \hline \end{gathered}$ |
| 17.010 | 98-99 | f | 74 | 3570 | 111.8 | 89.0 |  | 40.0 | 44.7 | 35.7 | 0.5 | 0.4 | 0.7 | 0.5 | 0.9 | 0.7 | 1.3 | 1.1 |
| 17.012 | 98-99 | f | 74 | 3570 | 111.8 | 89.0 |  | 4.0 | 4.5 | 3.6 | 0.0 | 0.0 | 0.1 | 0.1 | 0.1 | 0.1 | 0.1 | 0.1 |
| 17.017 | 98-99 | f | 74 | 3570 | 111.8 | 89.0 | 8.1 | 9.1 | 7.2 | 0.1 | 0.1 | 0.1 | 0.1 | 0.2 | 0.1 | 0.3 | 0.2 |
| 18.001 | 98 |  | 6 | 366 | 73.4 | 26.8 | 89.8 | 66.0 | 24.1 | 0.7 | 0.3 | 1.0 | 0.4 | 1.3 | 0.5 | 2.0 | 0.7 |
| 18.002 | 98 |  | 8 | 356 | 80.7 | 28.8 | 131.6 | 106.2 | 37.8 | 1.1 | 0.4 | 1.6 | 0.6 | 2.1 | 0.8 | 3.2 | 1.1 |
| 18.003 | 98 |  | 11 | 677 | 11.4 | 6.1 | 53.8 | 6.1 | 3.3 | 0.1 | 0.0 | 0.1 | 0.0 | 0.1 | 0.1 | 0.2 | 0.1 |
| 18.004 | 99 |  | 8 | 576 | 60.7 | 30.0 | 66.8 | 40.6 | 20.1 | 0.4 | 0.2 | 0.6 | 0.3 | 0.8 | 0.4 | 1.2 | 0.6 |
| 18.005 | 99 |  | 6 | 460 | 86.5 | 37.5 | 67.5 | 58.4 | 25.3 | 0.6 | 0.3 | 0.9 | 0.4 | 1.2 | 0.5 | 1.8 | 0.8 |
| 18.006 | 99 |  | 22 | 2706 | 150.3 | 115.2 | 181.3 | 272.4 | 208.8 | 2.8 | 2.2 | 4.1 | 3.1 | 5.4 | 4.2 | 8.2 | 6.3 |
| 18.007 | 98-99 | b | 74 | 3570 | 111.8 | 89.0 | 19.4 | 21.6 | 17.2 | 0.2 | 0.2 | 0.3 | 0.3 | 0.4 | 0.3 | 0.6 | 0.5 |
| 18.009 | 98 |  | 1 | 45 | 0.0 | 0.0 | 4.2 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| 18.011 | 98 |  | 4 | 176 | 309.8 | 26.2 | 54.9 | 170.2 | 14.4 | 1.8 | 0.1 | 2.6 | 0.2 | 3.4 | 0.3 | 5.1 | 0.4 |
| 19.004 | 98-99 | f | 74 | 3570 | 111.8 | 89.0 | 29.5 | 32.9 | 26.2 | 0.3 | 0.3 | 0.5 | 0.4 | 0.7 | 0.5 | 1.0 | 0.8 |
| 19.005 | 98-99 | f | 74 | 3570 | 111.8 | 89.0 | 215.9 | 241.3 | 192.3 | 2.5 | 2.0 | 3.6 | 2.9 | 4.8 | 3.8 | 7.2 | 5.8 |
| 20.003 | 74-85 | g |  |  | 589.0 | 362.8 | 294.2 | 1732.6 | 1067.3 | 18.0 | 11.1 | 26.0 | 16.0 | 34.7 | 21.3 | 52.0 | 32.0 |
| 20.005 | 74-85 | g |  |  | 589.0 | 362.8 | 172.5 | 1016.1 | 626.0 | 10.6 | 6.5 | 15.2 | 9.4 | 20.3 | 12.5 | 30.5 | 18.8 |
| 20.006 | 74-85 | g |  |  | 589.0 | 362.8 | 32.5 | 191.5 | 118.0 | 2.0 | 1.2 | 2.9 | 1.8 | 3.8 | 2.4 | 5.7 | 3.5 |
| 23.005 | 74-85 | g |  |  | 589.0 | 362.8 | 52.9 | 311.5 | 191.9 | 3.2 | 2.0 | 4.7 | 2.9 | 6.2 | 3.8 | 9.3 | 5.8 |
| 23.007 | 74-85 | g |  |  | 589.0 | 362.8 | 64.2 | 378.4 | 233.1 | 3.9 | 2.4 | 5.7 | 3.5 | 7.6 | 4.7 | 11.4 | 7.0 |
| 23.009 | 74-85 | g |  |  | 589.0 | 362.8 | 6.9 | 40.8 | 25.1 | 0.4 | 0.3 | 0.6 | 0.4 | 0.8 | 0.5 | 1.2 | 0.8 |
| 23.011 | 74-85 | g |  |  | 589.0 | 362.8 | 132.6 | 781.0 | 481.1 | 8.1 | 5.0 | 11.7 | 7.2 | 15.6 | 9.6 | 23.4 | 14.4 |
| 24.002 | 74-85 | g |  |  | 589.0 | 362.8 | 88.4 | 520.7 | 320.7 | 5.4 | 3.3 | 7.8 | 4.8 | 10.4 | 6.4 | 15.6 | 9.6 |
| 24.006 | 74-85 | g |  |  | 589.0 | 362.8 | 452.5 | 2665.1 | 1641.8 | 27.7 | 17.1 | 40.0 | 24.6 | 53.3 | 32.8 | 80.0 | 49.3 |
| 24.007 | 74-85 | g |  |  | 589.0 | 362.8 | 97.5 | 574.4 | 353.8 | 6.0 | 3.7 | 8.6 | 5.3 | 11.5 | 7.1 | 17.2 | 10.6 |
| 24.008 | 74-85 | g |  |  | 589.0 | 362.8 | 153.9 | 906.4 | 558.3 | 9.4 | 5.8 | 13.6 | 8.4 | 18.1 | 11.2 | 27.2 | 16.7 |
| 24.009 | 74-85 | g |  |  | 589.0 | 362.8 | 7.0 | 41.2 | 25.4 | 0.4 | 0.3 | 0.6 | 0.4 | 0.8 | 0.5 | 1.2 | 0.8 |
| 25.006 | 74-85 | g |  |  | 589.0 | 362.8 | 70.3 | 414.2 | 255.1 | 4.3 | 2.7 | 6.2 | 3.8 | 8.3 | 5.1 | 12.4 | 7.7 |
| 25.007 | 74-85 | g |  |  | 589.0 | 362.8 | 193.4 | 1139.4 | 701.9 | 11.8 | 7.3 | 17.1 | 10.5 | 22.8 | 14.0 | 34.2 | 21.1 |
| 25.013 | 74-85 | g |  |  | 589.0 | 362.8 | 297.4 | 1751.7 | 1079.1 | 18.2 | 11.2 | 26.3 | 16.2 | 35.0 | 21.6 | 52.6 | 32.4 |
| 25.015 | 74-85 | g |  |  | 589.0 | 362.8 | 41.4 | 244.1 | 150.4 | 2.5 | 1.6 | 3.7 | 2.3 | 4.9 | 3.0 | 7.3 | 4.5 |
| 26.001 | 74-85 | g |  |  | 589.0 | 362.8 | 12.1 | 71.4 | 44.0 | 0.7 | 0.5 | 1.1 | 0.7 | 1.4 | 0.9 | 2.1 | 1.3 |
| 26.006 | 74-85 | g |  |  | 589.0 | 362.8 | 25.7 | 151.3 | 93.2 | 1.6 | 1.0 | 2.3 | 1.4 | 3.0 | 1.9 | 4.5 | 2.8 |
| 27.001 | 74-85 | g |  |  | 589.0 | 362.8 | 29.4 | 173.4 | 106.8 | 1.8 | 1.1 | 2.6 | 1.6 | 3.5 | 2.1 | 5.2 | 3.2 |
| 27.002 | 74-85 | g |  |  | 589.0 | 362.8 | 268.4 | 1581.1 | 974.0 | 16.4 | 10.1 | 23.7 | 14.6 | 31.6 | 19.5 | 47.4 | 29.2 |



Table 7. Total quota (tonnes) options for the red sea urchin fishery by north and south B.C., estimated from various natural mortality values applied to current biomass ( Bc ) calculated from mean and approximate $90 \%$ lower confidence interval (CI) biomassvalues for five size limits of commercial red sea urchin, and bed areas fished up to 1996.

| Size Limit (mm TD) | Region | Quota 0.2 M Bc |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | $\mathrm{M}=0.052$ |  | $\mathrm{M}=0.075$ |  | $\mathrm{M}=0.10$ |  | $\mathrm{M}=0.15$ |  |
|  |  | $90 \% \mathrm{CI}$ |  | $90 \% \mathrm{CI}$ |  | Mean | $\begin{gathered} \text { Lower } \\ 90 \% \text { CI } \end{gathered}$ | $90 \% \mathrm{CI}$ |  |
| $\geq 100$ | North Coast | 2869.4 | 2084.3 | 4138.5 | 3006.3 | 5518.0 | 4008.3 | 8277.0 | 6012.5 |
|  | South Coast | 556.5 | 322.6 | 802.7 | 465.3 | 1070.3 | 620.4 | 1605.4 | 930.6 |
|  | B.C. | 3425.9 | 2407.0 | 4941.2 | 3471.6 | 6588.3 | 4628.8 | 9882.4 | 6943.1 |
| 100-140 | North Coast | 2526.7 | 1808.0 | 3644.3 | 2607.7 | 4859.0 | 3476.9 | 7288.6 | 5215.4 |
|  | South Coast | 456.2 | 261.4 | 658.0 | 377.1 | 877.3 | 502.7 | 1315.9 | 754.1 |
|  | B.C. | 2982.9 | 2069.4 | 4302.2 | 2984.8 | 5736.3 | 3979.7 | 8604.5 | 5969.5 |
| $\geq 90$ | North Coast | 3359.5 | 2457.7 | 4845.4 | 3544.7 | 6460.6 | 4726.3 | 9690.9 | 7089.5 |
|  | South Coast | 628.7 | 370.6 | 906.8 | 534.5 | 1209.0 | 712.7 | 1813.6 | 1069.0 |
|  | B.C. | 3988.2 | 2828.3 | 5752.2 | 4079.2 | 7669.6 | 5439.0 | 11504.4 | 8158.5 |
| 90-120 | North Coast | 2018.5 | 1480.1 | 2911.4 | 2134.7 | 3881.8 | 2846.3 | 5822.7 | 4269.4 |
|  | South Coast | 311.2 | 188.0 | 448.8 | 271.2 | 598.4 | 361.6 | 897.7 | 542.4 |
|  | B.C. | 2329.7 | 1668.1 | 3360.2 | 2405.9 | 4480.3 | 3207.9 | 6720.4 | 4811.8 |
| 90-130 | North Coast | 2589.3 | 1906.8 | 3734.5 | 2750.2 | 4979.4 | 3667.0 | 7469.1 | 5500.4 |
|  | South Coast | 419.1 | 253.9 | 604.4 | 366.1 | 805.9 | 488.2 | 1208.8 | 732.3 |
|  | B.C. | 3008.3 | 2160.7 | 4338.9 | 3116.4 | 5785.2 | 4155.1 | 8677.9 | 6232.7 |



Fig. 1. Annual yield (columns) and value (diamond/line) for the red sea urchin fishery in British Columbia, 1978-99.



Fig. 2. Box plots of mean (A) density and (B) biomass for different size limits ( $\geq 100, \geq 90,100$ 140, $90-120$, and $90-130 \mathrm{~mm} \mathrm{TD}$ ) calculated from surveys of red sea urchin populations in different PFM subareas throughout B.C., during 1993-99 $(\mathrm{N}=129)$. The horizontal central line of a box plot is the median and the two horizontal outer lines are the first and third quantiles of the sample, and the asterisks are outliers.

# APPENDIX A <br> PRELIMINARY PROBABILISTIC MODELING OF THE EFFECTS OF HARVESTRATES AND SIZE-RESTRICTIONS ON A POPULATION OF RED SEA URCHINS <br> (Strongylocentrotus franciscanus) 

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## Introduction

The long-term impact of various harvesting strategies are estimated using a computer program to simulate the dynamics of a population of red sea urchins (Strongylocentrotus franciscanus). The expected behaviours of the population are expressed as mathematical equations and combined into a computer model. A set of parameter values is identified that allows the model to reach an equilibrium state that is similar to results of a size-frequency survey of an urchin bed. Random variability of environmental factors is introduced into the model to allow a probabilistic assessment of various strategies for the harvest of red sea urchins.

There are many other models of red sea urchins and other marine invertebrates (Lai and Bradbury 1998 and Breen and Kendrick 1998 for example). The analyses in this report are unique in two respects:

1. The values of independent parameters are chosen such that the model predicts that in a constant environment with no harvesting, the stock will remain stable in it's virgin state. The more common approach is to determine a history of the parameter values that could lead to the present state of the population.
2. The growth of the urchins is determined by both their biological potential for growth and by the capacity of the local environment to support growth. The more common approach is to only consider the biological capacity for growth.

Sample data is analyzed to demonstrate how the impact of harvesting can be predicted from the exploitation rate (fraction of harvestable animals taken each year) and an estimate of the virginstate of the stock.

## An Equilibrium Approach to Population Modelling

The model for a red sea urchin (RSU) population is based on the concept of an equilibrium state. In an equilibrium state, environmental conditions and harvesting strategies are held constant and consequently the characteristics of the RSU population (number of animals, biomass, etc.) also reach a constant state. A set of parameter values is identified for the model that lead to a predicted steady-state that is similar to the results of a size-frequency survey of a virgin stock. Further analysis is based on variants of this equilibrium.

From a population-modeling perspective, RSU have four significant activities:

1. Settlement
2. Growth
3. Mortality
4. Spat production

Because there is equilibrium, the rate of settlement can be determined from the survival rate and the number of animals. Estimated survival rates for RSU on the west coast of Canada typically range from 0.85 to 0.97 (A.Campbell, unpublished data). For the purpose of example, a survival rate of $S_{\text {urv }}=0.9$ will be used. In order to maintain an equilibrium population of $P_{\text {equib, }}$, the settlement rate must be $S_{\text {ettle }}=\mathrm{P}_{\text {equib }} *\left(1 / \mathrm{S}_{\text {urv }}-1\right)$ spats per year. When the survival rate is applied to the population including the new spats, the resulting number of RSU is $\mathrm{S}_{\text {urv }} *\left(\mathrm{P}_{\text {equib }}+\mathrm{S}_{\text {ettle }}\right)=\mathrm{P}_{\text {equib }}$.

Similarly, the equilibrium growth rate can be determined from the survival rate and the biomass. If an equilibrium biomass of $\mathrm{B}_{\text {equib }}$ is to be maintained, then the amount of growth that occurs every year is $\mathrm{G}_{\text {rowth }}=\mathrm{B}_{\text {equib }} *\left(1 / \mathrm{S}_{\text {urv }}-1\right)$ grams per year. When the survival rate is applied after growth has occurred, the resulting biomass is $\mathrm{S}_{\text {urv }} *\left(\mathrm{~B}_{\text {equib }}+\mathrm{G}_{\text {rowth }}\right)=\mathrm{B}_{\text {equib }}$.

If meta-population effects are ignored, then the rate at which spats are produced equals the settlement rate.

Figure A1 shows the size frequency density (SFD) taken from a survey of a virgin (or close to it) RSU stock. As often occurs with RSU surveys, the SFD is very discontinuous and therefore the precision is questionable. Even though the actual SFD may not be precisely known, the survey data does have two identifiable characteristics that can serve as useful constraints on the equilibrium state:

- The maximum test diameter of an RSU is approximately 165 mm .
- The SFD is bimodal with peaks at approximately 25 and 125 mm .

These constraints on the equilibrium state are enough to estimate parameter values for the model used in the analysis. A model with more independent parameters would require more constraints on the equilibrium state and consequently be more dependent on the accuracy of survey data.

## The Model Equations

## Survival

To account for natural mortality, a single survival rate (fraction of RSU to survive a given year), $S_{\text {urv }}$, is applied to all sizes of RSU.

Harvest is represented by a lower size limit, an upper size limit and a fishing rate, $\mathrm{f}_{\text {rate }}$. The fishing rate is expressed as the fraction of RSU in the harvestable size-range to be harvested in a year. For RSU within the size limits, the survival rate is $\mathrm{S}_{\text {urv }} *\left(1-\mathrm{f}_{\text {rate }}\right)$. If the biomass falls below $20 \%$ of the equilibrium level, the stock is considered to have collapsed and harvest stops. After a stockcollapse, the stock is considered to have recovered when the biomass reaches $50 \%$ of the equilibrium level and harvest is resumed.

## Growth

An empirical equation is used to describe the growth rate of individual RSU. The equation is based on tagging studies performed on the west coast of Canada (A. Campbell, unpublished data):

$$
G(L)=\gamma \cdot e^{-\left(\frac{\zeta-L}{\eta}\right)^{2}} \cdot(1+\varepsilon \text { grow })
$$

$\mathrm{G}(\mathrm{L})$ is the growth rate (mm/year)
L is the current test diameter $\gamma$ controls the overall growth rate
$\zeta$ and $\eta$ determine the shape of the curve
$\varepsilon_{\text {grow }}$ is a Normal variate with a mean of 0 and a standard deviation of $\sigma_{\text {grow }}$
The growth equation accounts for variability amongst individual animals. As a result, growth of individual RSU can be expressed as the probability of an RSU growing from one test-diameter to another as was done in Lai and Bradbury 1998.

Figure A2 shows a fit of the growth equation to experimental data taken from the Haida Research Area, Site 3 in 1997 (Alan Campbell, unpublished data).

The chosen growth equation has two notable features that make it amenable to calibrating the model to an equilibrium state:

- It is size-based (as opposed to age-based)
- The overall growth rate is controlled by a single parameter, $\gamma$.

The value of $\gamma$ is limited in two ways:

1. The capacity of the local environment to support the growth of RSU (food in particular).
2. The biological potential growth of RSU.

The first limit is especially relevant where growth is food-limited. Studies (Vadas 1977 for example) have suggested that the growth of RSU is often food-limited. The limit is applied by assuming the local environment has potential to support a fixed amount of growth in the RSU biomass. This potential growth is distributed amongst all the RSU in the population. Generally, as this potential growth is divided amongst more RSU, the individual growth rates become smaller. A value of $\gamma$ can be found (numerical methods) such that the RSU population uses all the potential for growth provided by the local environment.

The second limit is necessary to prevent unrealistically large growth rates when the stock of RSU becomes small. A maximum value of $\gamma$ is applied. This limit is only significant when the RSU stock becomes small.

## Spat Production

Spat production is the combination of several processes and characteristics; fecundity, fertilization success, larval survival and the migration of larvae between sub-populations of a meta-population. For simplicity, the migration of larvae between sub-populations will be ignored while defining the equilibrium state. The local population is assumed to be its own source and destination of spats. Meta-population effects are expected to contribute variability into the settlement rate and this variability will be incorporated into the probabilistic simulations.

A modification of the Ricker curve (Quinn and Deriso, 1997) is used to estimate the settlement rate.
$\mathrm{S}_{\text {ettle }}=\mathrm{R}_{\text {coef }} * \mathrm{~B}_{\text {mature }} * \exp \left(-\mathrm{B} / \mathrm{B}_{\text {equib }}\right)$
$\mathrm{R}_{\text {coef }}$ is a scaling factor and one of the independent parameters of the model
$B$ is the current biomass
$\mathrm{B}_{\text {equib }}$ is the equilibrium biomass
$B_{\text {mature }}$ is the current biomass of mature RSU
At equilibrium:

$$
\mathrm{R}_{\text {coef }}=\mathrm{P}_{\text {equib }} *\left(1 / \mathrm{S}_{\text {urv }}-1\right) / \mathrm{B}_{\text {mature,equib }} / \exp (-1)
$$

$B_{\text {mature,equib }}$ is the biomass of mature RSU at equilibrium.

## An Example of an Equilibrium State

Data from a survey (Queen Charolette Islands, Site 3, 1998, Dominique Bureau, unpublished data) is used to define an equilibrium state. For simplicity, the population is normalized to unity. The following characteristics are observed from the survey:

- The mean mass per RSU is 233 grams.
- The maximum test diameter is approximately 165 mm .
- The SFD is bimodal with peaks at approximately 25 and 125 mm .

Table A1 is a list of values of model-parameter parameters used for the equilibrium. The values of $æ$ and ç are chosen by trial and error in order to get the desired shape of the SFD. Other parameters of the model are chosen by expert opinion. Figure A3 shows the SFD from the survey and from the predicted equilibrium. Table A2 gives some resulting characteristics of the equilibrium state.

## Running the Model Probabilistically

Three model parameters are allowed to vary randomly in the model; survival, growth and settlement rate.

## Survival

For the equilibrium state, a survival rate of 0.90 per year is selected. By definition, the survival rate must have a value between zero and one. The survival rate for the probabilistic simulations is assigned a normal distribution with a mean of 0.90 and a standard deviation of 0.0333 . The survival rate is greater than one in approximately $0.001 \%$ of the years.

## Growth

For the equilibrium point, the increase in biomass due to growth (ignore mortality) is 25.889 grams per year. The ecological limit to the rate of growth in the biomass is assigned a normal distribution with a mean of 25.889 and a standard deviation of $25.889 / 2=12.994$. There is a 2.28 \% probability of negative growth.

A maximum value of $17 \mathrm{~mm} /$ year is arbitrarily applied to $\gamma$ to reflect biological limitations to the growth rate.

## Settlement

Variability is introduced into the settlement rate by assigning a lognormal distribution to the multiplicative coefficient, $\mathrm{R}_{\text {coef. }}$. A lognormal distribution with a square root of ten is used to generate a sporadic settlement rate (as may occur due to metapopulation effects). An arithmetic mean of $\mathrm{R}_{\text {coef }}=0.00139277$ spats/gram/year is chosen because it equals the value in the equilibrium state. To obtain the desired arithmetic mean and geometric standard deviation, a geometric mean of 0.000411595 is assigned to $\mathrm{R}_{\text {coef. }}$.

There is a $95 \%$ probability the value of $\mathrm{R}_{\text {coef }}$ is between 0.00004 and 0.004 .

## A Probabilistic Simulation with No Harvest

Figure A4 shows some results when the model is run probabilistically for one thousand years with no harvest. The population starts out at the predicted equilibrium. The necessary random values are chosen according to the assigned probability density functions. The resulting settlement, growth and survival rates are applied to the original SFD to predict the state of the population after one year. Similarly, the first-year results and another set of random parameter values are used to estimate the state of the population in the second year. Subsequently, each year is predicted from the previous year and a new set of random values. A simulation of one thousand years was created in this manner.

The estimated biomass is rather stable in the probabilistic simulation even though the variability of the cumulative growth is significant with respect to the mean.

The variability in the settlement rate has a significant influence on the estimated population and correspondingly the mean mass per RSU.

## Incorporating Harvest into the Model

Harvest is incorporated into the model with the goal of making informative comparisons of harvest-strategies. The first step is to generate an appropriate no-harvest case to use as a benchmark when evaluating the harvest strategies.

A no-harvest case consisting of 100 simulations of 100 years is used. Each simulation starts at the predicted equilibrium but different random values are used to predict the evolution of the RSU population over 100 years.

The 100 simulations are repeated with various harvesting strategies, but the same random values are used as in the no-harvest case. For every year or simulation in a harvest-case, there is a corresponding year or simulation in the no-harvest case. All differences between the cases are directly attributable to the differences in the harvesting strategies. As a result, a statistical approach known as pairwise comparisons (Montgomery, 1991) can be used to evaluate harvest strategies. Seventy-two different harvest strategies are considered; all combinations of nine harvest rates and eight harvestable size ranges.

Figure A5 illustrates how a case with harvesting can be compared to the no-harvest case. The mean biomass for each of the one hundred simulations is shown in both the harvest and no-harvest case; the no-harvest case on the x -axis and the harvest-case on the y -axis. A diagonal line is included to show where the points would be if the harvest had no influence on the estimated biomass. The vertical distance between the diagonal line and the points indicates the drop in biomass due to harvesting.

For the harvest strategy used for Figure A5, the long-term influence of harvesting is to cause the biomass to drop by approximately 25 grams or $25 / 233^{*} 100=11 \%$ of the equilibrium biomass. The influence of the harvest is easily discernable (visually or quantitatively) even though it is small relative to the variability of the estimated mean biomass.

## Results of the Probabilistic Cases

Tables A3-6 show how the various harvesting strategies impact various characteristics of the population. Figures A6-8 show the same information graphically. As expected:

- Smaller harvest rates have smaller impacts on the RSU population
- For the same harvest rate, narrow harvestable ranges result in smaller impacts on the RSU population.

In order to evaluate different harvest strategies, it is necessary to quantify the ecological costs of the harvest and to establish an acceptable cost. For discussion purposes only a harvest strategy will be considered acceptable if the resulting drop in biomass is less than $20 \%$.

An actual fishery is more likely to be managed on the basis of yield (mass of catch per year) than on a harvest rate (fraction of haravestable animals). Therefore it is useful to look at how yield affects the biomass. Unfortunately, constant yields are not considered explicitly in the model. To get an approximation of how a constant yield will affect biomass, the drop in biomass for a
simulation of one hundred years is plotted against the mean yield. Some results are shown in Figure A9. Interesting trends are suggested, but need to be confirmed by simulations where the yield is held constant.

For yields less than $3 \%$ of the equilibrium biomass per year, there is little difference between the size ranges considered in these analyses. As a rough approximation:

For every gram per year of RSU that is pulled out of the water, the biomass will decrease by approximately ten grams.

For yields greater than three percent of the equilibrium biomass, the linearity is lost. The damage to the biomass increases at a faster rate.

## Using These Analyses

Figure A10 shows how a specific environmental impact (a $20 \%$ drop in the long-term biomass) can be traced to a range of yield rates. When the harvest rate (fraction of harvestable animals per year) is held constant, a $20 \%$ drop in biomass is associated with a yield of approximately 1.8 to $2.3 \%$ of the equilibrium biomass per year. If the yield (mass per year) is held constant, similar results are expected (but need to be confirmed with the appropriate simulations). Given the current information for the data considered, a harvest-quota of $1.8 \%$ of the virgin biomass per year appears to be precautionary.

## Software Used for the Simulations

Probabilistic Urchin Population Simulator (PUPS) is computer program created to produce the cases for these analyses.

PUPS was developed using an object-oriented approach (Rumbaugh et al., 1991). It is written in ANSI C++. As a result, PUPS is very portable and could be easily adaptable to other population analyses involving RSU or other species.

In order to complete the cases for the analyses, PUPS was run on 12 different computers with either a Windows NT or Windows 95 operating system. The time required to complete 100 simulations of 100 years ranged from 8 to 18 hours. The majority of the computational time was spent determining the correct value of $\gamma$ to use in the growth equation.

## Discussion

The analysis demonstrates how the long-term impact of an RSU fishery can be evaluated with respect to quantitative criteria for the long-term impact on the RSU population. In particular, there appears to be a simple relationship between the mass of RSU harvested every year and the resulting drop in biomass. Unfortunately, a constant yield (mass per year) is not one of the harvest strategies considered in the analysis. Any conclusions about the impact of a constant yield should be verified by further analysis.

The analysis is also an opportunity to try a new approach to selecting values for model parameters. Parameter values are chosen so that the long-term equilibrium state predicted by the model are similar to the results of a survey. The advantages of this approach are that there is less dependence on the accuracy of the survey results and there is the assurance that at the model will make sensible predictions (at least under some conditions). A combination of quantitative and qualitative constraints are applied to the equilibrium-state of the model. The approach to selecting parametervalues is successful but it could be explored further.

The experience gained by performing the analysis leads to the following recommendations for developing the methodology:

- A constant yield (mass per year) should be one of the harvest strategies that is considered. Minor modifications to PUPS would be required.
- A rotational fishery, where harvest only occurs every second or third year should be considered. Minor modifications to PUPS would be required.
- Sensitivity analyses should be performed to compare the influences of the model parameters.
- The approach to selecting parameter-values has demonstrated potential. More investigation is warranted to determine if the approach can be further developed so that values for more model-parameters can be estimated. If more model-parameters can be estimated, then there is the potential to incorporate features such as size-specific mortality and density-dependent fertilization success into PUPS.


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## References

Breen, P.A. and T.H. Kendrick. 1998. An evaluation of surplus production analysis for assessing the fishery for New Zealand red rock lobsters (Jasus edwarsii). Can. Spec. Publ. Fish. Aquat. Sci. 125: 213-223.

Botsford, L.W., B.D. Smith, and J.F. Quinn. 1994. Bimodality in size distributions: The Red Sea Urchin (Strongylocentrotus franciscanus) as an example. Ecological Applications. 5: 4250.

Lai, H.L. and A. Bradbury. 1998. A modified catch-at-size analysis model for a red sea urchin (Strongylocentrotus franciscanus) population. Can. Spec. Publ. Fish. Aquat. Sci. 125: 8596.

Montgomery, D.C. 1991. Design and Analysis of Experiments. John Wiley and Sons.
Quinn,T.J. and R.B. Deriso. 1999. Quantitative Fish Dynamics. Oxford University Press.

Rumbaugh,J., M. Blaha, W. Premerlani, F. Eddy, and W. Lorenson. 1991. Object-Oriented Modeling and Design. Prentice Hall.

Smith, B.D. and L.W. Botsford. 1998. Interpretation of growth, mortality, and recruitment in size-at-age, growth, increment and size frequency data. Can. Spec. Publ. Fish. Aquat. Sci. 125: 125-139.

Smith, B.D., L.W. Botsford and R.W. Wing. 1998. Estimation of growth and mortality parameters from size frequency distributions lacking age patterns: the red sea urchin (Strongylocentrotus franciscanus) as an example. Can. J. Fish. Aquat. Sci. 55: 1236-1247

Vadas, R.L. 1977. Preferential feeding: an optimization strategy in sea urchins. Ecological Monographs. 47: 337-371.

Table A1. Model Parameters for an Equilibrium State.

| Parameter | Value in <br> Equilibrium |
| :--- | ---: |
| Survival rate of RSU | 0.90 |
| Mean mass in equilibrium state | 233 g |
| Coefficient used to determine settlement rate from biomass | 0.0013927 <br> spat/g |
| The size at which an RSU reaches maturity | 70 mm |
| Annual increase in biomass due to growth | 25.8889 g |
| Size at which maximum growth of test-diameter occurs | 55 mm |
| A small value means that growth is concentrated in a small <br> size-range | 60 mm |
| Standard deviation of size for RSU in the first cohort. | 3.53 mm |
| Standard deviation of growth rate for an RSU. Expressed as <br> a fraction of expected growth rate | 0.33 |
| Mean test diameter of RSU in the first cohort |  |
| Scalar multiplier to convert test-diameter to RSU mass(á) | 0.0012659 |
| Power Variable used to convert test-diameter to RSU mass <br> (â) | 2.7068 |

Table A2. Results of the Equilibrium State

| Characteristic | Value in <br> Equilibrium |
| :--- | ---: |
|  | 0.4612 |
| Mature Population | $8.750 \mathrm{~mm} / \mathrm{year}$ |
| Scaling Factor used in the growth <br> equation | $0.111 \mathrm{spat} / \mathrm{year}$ |
| Settlement Rate |  |

Table A3. Effect of Harvest on the Predicted Biomass

|  | Drop in Mean Biomass Due to Harvesting (\% of Equilibrium Biomass) |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | vestable | Range |  |  |  |  |
| Harvest Rate | 100 | 90 | 100-140 | 90-120 | 90-130 | 90-140 | 90-110 | 70 |
| 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| 0.01 | 6.86 | 7.45 | 4.88 | 2.65 | 4.00 | 5.47 | 1.55 | 8.27 |
| 0.02 | 12.86 | 13.99 | 9.38 | 5.19 | 7.76 | 10.49 | 3.07 | 15.56 |
| 0.04 | 23.01 | 25.09 | 10.28 | 9.98 | 14.66 | 19.44 | 6.00 | 28.07 |
| 0.06 | 31.41 | 34.35 | 24.51 | 14.46 | 20.90 | 27.27 | 8.81 | 38.61 |
| 0.08 | 38.59 | 42.28 | 30.82 | 18.66 | 26.61 | 34.25 | 11.52 | 47.81 |
| 0.10 | 44.88 | 49.26 | 36.53 | 22.64 | 31.87 | 40.53 | 14.14 | 54.93 |
| 0.15 | 56.17 | 59.74 | 48.65 | 31.73 | 43.55 | 53.65 | 20.35 | 63.17 |
| 0.20 | 61.29 | 63.35 | 56.62 | 39.98 | 53.25 | 60.11 | 26.17 | 64.70 |
| 0.50 | 64.95 | 64.89 | 64.16 | 61.41 | 63.77 | 65.17 | 55.78 | 65.06 |

Table A4. Effect of Harvest on the Predicted Population

|  | Drop in Mean Population Due to Harvesting (\% of Equilibrium Population) |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | stabl | Range | est Diam | - |  |  |
| Harvest Rate | 100 | 90 | 100-140 | 90-120 | 90-130 | 90-140 | 90-110 | 70 |
| 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| 0.01 | 3.21 | 3.74 | 2.46 | 1.70 | 2.35 | 2.97 | 1.11 | 4.75 |
| 0.02 | 6.50 | 7.57 | 4.96 | 3.39 | 4.69 | 5.98 | 2.20 | 9.61 |
| 0.04 | 13.11 | 15.32 | 5.72 | 6.76 | 9.39 | 12.06 | 4.38 | 19.38 |
| 0.06 | 19.66 | 23.02 | 15.10 | 10.09 | 14.07 | 18.14 | 6.54 | 28.99 |
| 0.08 | 26.01 | 30.48 | 20.16 | 13.38 | 18.69 | 24.17 | 8.67 | 38.31 |
| 0.10 | 32.14 | 37.68 | 25.18 | 16.63 | 23.27 | 30.06 | 10.78 | 45.93 |
| 0.15 | 44.23 | 49.13 | 37.19 | 24.61 | 34.41 | 43.70 | 15.97 | 55.26 |
| 0.20 | 49.90 | 53.53 | 45.84 | 32.42 | 44.64 | 50.85 | 21.07 | 57.05 |
| 0.50 | 54.63 | 55.44 | 55.01 | 54.90 | 57.09 | 57.24 | 50.27 | 57.76 |

Table A5. Effect of Harvest on the Probability of Stock Collapse

|  | Probability of Stock Collapse in 100 years |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | rvestable | Range ( 7 | est Diam | eter in m |  |  |
| Harvest Rate | 100 | 90 | 100-140 | 90-120 | 90-130 | 90-140 | 90-110 | 70 |
| 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| 0.01 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| 0.02 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| 0.04 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| 0.06 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.02 |
| 0.08 | 0.02 | 0.03 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.06 |
| 0.10 | 0.04 | 0.08 | 0.00 | 0.00 | 0.00 | 0.02 | 0.00 | 0.38 |
| 0.15 | 0.56 | 0.86 | 0.07 | 0.00 | 0.02 | 0.21 | 0.00 | 1.00 |
| 0.20 | 0.97 | 1.00 | 0.61 | 0.00 | 0.18 | 0.92 | 0.00 | 1.00 |
| 0.50 | 1.00 | 1.00 | 1.00 | 1.00 | 1.00 | 1.00 | 0.36 | 1.00 |

Table A6. Effect of Harvest on Predicted Yield

|  | Yield (\% of Equilibrium Biomass per Year) |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | Har | vestable | Range ( | ast Diam | meter in m |  |  |
| Harvest Rate | 100 | 90 | 100-140 | 90-120 | 90-130 | 90-140 | 90-110 | 70 |
| 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| 0.01 | 0.79 | 0.85 | 0.56 | 0.30 | 0.45 | 0.62 | 0.17 | 0.93 |
| 0.02 | 1.47 | 1.58 | 1.07 | 0.58 | 0.87 | 1.18 | 0.34 | 1.73 |
| 0.04 | 2.56 | 2.75 | 1.16 | 1.09 | 1.61 | 2.14 | 0.65 | 2.97 |
| 0.06 | 3.39 | 3.61 | 2.68 | 1.56 | 2.25 | 2.91 | 0.95 | 3.83 |
| 0.08 | 4.02 | 4.24 | 3.29 | 1.98 | 2.79 | 3.53 | 1.23 | 4.38 |
| 0.10 | 4.48 | 4.65 | 3.78 | 2.36 | 3.25 | 4.01 | 1.49 | 4.64 |
| 0.15 | 5.04 | 5.00 | 4.60 | 3.13 | 4.07 | 4.67 | 2.08 | 4.70 |
| 0.20 | 5.16 | 5.03 | 4.90 | 3.70 | 4.47 | 4.77 | 2.59 | 4.66 |
| 0.50 | 5.26 | 5.12 | 4.93 | 4.22 | 4.44 | 4.70 | 3.96 | 4.78 |




Figure A1. Size Frequency Densities Estimated from Survey Data

## Growth Estimates



Figure A2. Estimated and Measured Growth Rates. Most Likely Rates and 95\% Confidence Bounds are Shown

Size Frequency Densities


Figure A3. Survey Results and the Equilibrium Estimated by the Model

No Harvest


Figure A4. A Probabilistic Simulation with No Harvesting

Comparison of Harvest and No-Harvest Simulations


Figure A5. Effect of Harvesting on the Predicted Mean Biomass. A Simulation-bySimulation Comparison.

## Decrease in Biomass Due to Harvesting



Figure A6 Effect of Harvesting on the Predicted Biomass

Decrease in Population Due to Harvesting


Figure A7 Effect of Harvest on Predicted Population


Figure A8 Effect of Harvest on Probability of Stock Collapse


## Figure A9. Effect of Yield on Biomass

Harvest Size Range is $90-130 \mathrm{~mm}$



Harvest Size Range is $90-140 \mathrm{~mm}$


Minimum Harvest Size is 70 mm


Figure A9. Effect of Yield on Biomass (continued)


Figure A10. Yields Corresponding to a 20\% Drop in Biomass

## Appendix B

# Preliminary Report on Egg and Yield per Recruit Analyses for a Red Sea Urchin Population 

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## Introduction

Of interest to management are the biological implications of possible changes in various size regulations under different fishing rates to protect the valuable red sea urchin resource. Dynamic pool models, widely used in fish population dynamics studies (Quinn and Desiro 1999), require data on size at maturity, rates of growth, fecundity, natural and fishing mortality, and recruitment, some of which are difficult to obtain for red sea urchin populations. Yield per recruit and egg per recruit (YPR/EPR) analyses circumvent the requirement for detailed recruitment information by assuming recruitment is constant. Consequently, this approach was used in this paper to determine the effect of different size regulation and fishing mortality rates on a red sea urchin population. The biological input data was obtained from a productive red sea urchin population that has been studied at Kendrick Island, near Nanaimo, on the east coast of Vancouver Island, during 1990-99 (Campbell et al. unpublished data). There are few YPR/EPR analyses on red sea urchins. Botsford et al. (1993) used YPR analyses to consider periodic harvest schedules (rotation) in red sea urchins. Breen (1984), although not presenting any results of YPR analyses, discussed some implications of size limits and mortality rates on red sea urchins in B.C.

The objective of this paper is to report on the effects of two different types of size regulations on YPR/EPR under different fishing mortalities for a red sea urchin population with reasonably fast growth rates by (1) changing recruit sizes, and (2) establishing maximum sizes. The current recruit size in the fishery is $\geq 100 \mathrm{~mm}$ TD. A maximum size regulation would protect large mature red sea urchins above a certain size to allow increased potential egg production and provide additional spine canopy for small juvenile survival (Tegner and Dayton 1977; Pfister and Bradbury 1996; Lai and Bradbury 1998).

## Methods

An egg per recruit (EPR) model by Tegner et al. (1989) (equation numbers up to 15, without letters) was modified by Campbell (1997, Appendix 1) to include a maximum legal size option and a size at maturity relationship. The original equation numbers (up to 15) with "a" and " $p$ " indicate modifications to accommodate the maximum legal size option and size at maturity, respectively. In the present study, we extended these models to include a size selectivity curve and a yield per recruit model (YPR) with minimum and maximum legal size options. The EPR/YPR model was written in Java programming language by A. Meynert.

The mathematical formulae used in the models were as follows.

## Egg per Recruit Model

## Variable Definitions

$h_{j}=$ midpoint of an interval of length measurements
$\mathrm{w}=$ width of each length interval
$j=$ number of each length interval
$\mathrm{t}=$ time in years
$\mathrm{M}=$ instantaneous rate of natural mortality
$\mathrm{N}_{0}=$ original number of female animals
$\mathrm{t}_{0}=$ age of zero length for a von Bertalanffy length-at-age curve
$\mathrm{K}=$ Brody growth coefficient for a von Bertalanffy length-at-age curve
$L_{\text {infinity }}=$ asymptotic length for a von Bertalanffy length-at-age curve
$\mathrm{L}_{\mathrm{t}}=$ the average length of an animal at time t according to the von Bertalanffy curve $\mathrm{SD}_{\mathrm{t}}=$ the standard deviation of length of an animal at time t
$\mathrm{F}=$ instantaneous rate of fishing mortality

## Equations

$$
\left(h_{j}-w / 2, h_{j}+w / 2\right) ; \quad j=1, J
$$

Definition of the length intervals.

$$
\begin{equation*}
N V_{t, .}=N_{0} \exp (-M t) \tag{1}
\end{equation*}
$$

The number of animals surviving each year in a virgin population.

$$
N V_{t, .}=\sum_{j=1}^{J} N V_{t, j}
$$

Number of females in a virgin population at time t for all length intervals j .

$$
N V_{., j}=\sum_{t=1}^{\lambda} N V_{t, j}
$$

Number of females in a virgin population falling in length interval $j$ for all times t.
$L_{t}=L_{\infty}\left\{1-\exp \left[-K\left(t-t_{0}\right)\right]\right\}$
Von Bertalanffy size-at-age curve.
$\mathrm{SD}_{t}=a+b \sqrt{t}$
Standard deviation varies with the square root of age.
$Q_{t, j}=\frac{1}{\mathrm{SD}_{t} \sqrt{2 \pi}} \int_{h_{j}}^{h_{j+1}} \exp \left[-\frac{\left(h-L_{t}\right)^{2}}{2 \mathrm{SD}_{t}{ }^{2}}\right] d h$
The proportion of animals falling into length interval $j$ at time $t$.

$$
\begin{equation*}
N V_{,, j}=\sum_{t=1}^{\lambda} Q_{t, j} N V_{t, .} \tag{5}
\end{equation*}
$$

Number of females in a virgin population falling in length interval j for all times $t$, calculated a different way.
$f_{j}=c h_{j}{ }^{d}$
The average fecundity of an animal in length interval j .
$E_{\max }=\sum_{j=m}^{J} N V_{, j,} f_{j}$
The maximum number of eggs an unfished population can produce, without using the factor of a mature proportion of animals.
$E_{M A X}=\sum_{j=m}^{J} N V_{, j} f_{j} M P_{j}$
The maximum number of eggs an unfished population can produce, using the factor of a mature proportion of animals.
$N_{1, .}=N_{0} \exp (-M)$
The number of animals after one year of mortality.
$Q P R_{t}=\frac{1}{\mathrm{SD}_{t} \sqrt{2 \pi}} \int_{0}^{h_{R}} \exp -\left[\frac{\left(h-L_{t}\right)^{2}}{2 \mathrm{SD}_{t}{ }^{2}}\right] d h$
The proportion of animals smaller than minimum legal size at time $t$.
$Q M L_{t}=\frac{1}{\mathrm{SD}_{t} \sqrt{2 \pi}} \int_{M L S}^{\infty} \exp -\left[\frac{\left(h-L_{t}\right)^{2}}{2 \mathrm{SD}_{t}{ }^{2}}\right] d h$
The proportion of animals greater than the maximum legal size at time $t$.
$N P R_{t, .}=Q P R_{t} N V_{t,}$
The number of animals smaller than the minimum legal size at time t .
$N M L_{t, .}=Q M L_{t} N V_{t, .}$
The number of animals greater than the maximum legal size at time $t$.
$N R_{t, .}=N_{t, .}-N P R_{t,}$
The number of animals of recruit age at time $t$, without using the maximum legal size option.
$N R_{t, .}=N_{t,-}-N P R_{t,-}-N M L_{t, .}$
The number of animals of recruit age at time $t$, using the maximum legal size option.
$S_{t}=\left[\left(\frac{N P R_{t, .}}{N_{t, .}}\right) \exp [-M]\right]+\left[\left(\frac{N R_{t, .}}{N_{t, .}}\right) \exp [-(F+M)]\right]$
The survival rate of animals at time $t$, without using the maximum legal size option.
$S_{t}=\left[\left(\frac{N P R_{t, .}}{N_{t, .}}\right) e^{-M}\right]+\left[\left(\frac{N R_{t, .}}{N_{t, .}}\right) e^{-(M+F)}\right]+\left[\left(\frac{N M L_{t, .}}{N_{t, .}}\right) e^{-M}\right]$
The survival rate of animals at time $t$, using the maximum legal size option.
$N_{t, .}=N_{t-1, .} S_{t-1}$
The number of animals in year $t$, based on the number of animals in the previous year and the survival rate from the previous year.
$N_{t, j}=N V_{t, j} \quad j<R$
$N_{t, j}=Q_{t, j} N R_{t, .} /\left(1-Q P R_{t}\right) \quad j \geq R$
The number of animals found in length interval j at time t .
$E=\sum_{t=1}^{\lambda} \sum_{j=m}^{J} N_{t, j} f_{j}$
The total number of eggs produced by a population without using a mature proportion of animals.
$E=\sum_{t=1}^{\lambda} \sum_{j=m}^{J} N_{t, j} f_{j} M P_{j}$
The total number of eggs produced by a population using a mature proportion of animals.

$$
\begin{equation*}
M P_{j}=\frac{L}{L+e^{(A-B L)}} \tag{16}
\end{equation*}
$$

The proportion of mature animals of length interval j .

## Yield per Recruit Model

In addition to the Beverton-Holt yield per recruit model (equations from Ricker 1975), an alternate yield per recruit model was developed to allow a maximum legal size option to be used. Both models gave similar yield per recruit results when compared with similar minimum size limits. The alternate YPR/EPR model program makes use of the Baranov catch equation in combination with the other equations in Appendix B which describe the number of animals of recruit age. For brevity, only the extra equations and variables are listed below.

## Variable Definitions

$\mathrm{W}_{\mathrm{j}}=$ average weight of an animal falling in length interval j

## Equations

$$
\begin{equation*}
\mathrm{W}_{\mathrm{t}}=a L_{t}^{b} \tag{17}
\end{equation*}
$$

Length-weight power relationship.

$$
\begin{equation*}
C_{t}=N R_{t} \frac{F}{F+M}\left(1-e^{-(M+F)}\right) \tag{18}
\end{equation*}
$$

Baranov catch equation. $\mathrm{C}_{\mathrm{t}}$ is the number of animals caught each year.

$$
\begin{equation*}
W .=\sum_{t=0}^{t_{\infty}} W_{t} C_{t} \tag{19}
\end{equation*}
$$

Total weight of animals caught.

$$
\begin{equation*}
N N R_{t}=N R_{t}-N R_{t-1} \tag{20}
\end{equation*}
$$

Number of new recruits each year.
$N N R=\sum_{t=0}^{t_{\infty}} N N R_{t}$
Total number of recruits during the population's lifetime.
$Y P R=\frac{W .}{N N R .}$
Yield per recruit.

## Size-Selectivity Curve

A size-selectivity curve for the minimum legal size is approximated using five values in the YPR/EPR model program.

These equations are used for the approximation of a size-selectivity (S-shaped) curve for first entry into the fishery as an option to a knife-edge entry. Red sea urchins tests are not completely symmetrical and there can be some error (up to about 5 mm ) in measuring the test diameter of urchins at the minimum legal size. A graph of a knife-edge entry would show two lines: the first running at a level of zero exposure to exploitation until the minimum legal size, at which point the level of exploitation would jump immediately to one hundred percent.

The equations used in the program split up the interval of the integral in equation (9), the equation that defines the proportion of pre-recruits at time $t$, into six equal parts.

Define $\operatorname{pf}(\mathrm{h})$ to be the proportion of animals at size x which are fished. Let h equal the minimum legal size. Therefore, $\operatorname{pf}(\mathrm{h})=1$. The user inputs values for $\mathrm{pf}(\mathrm{h}-1), \operatorname{pf}(\mathrm{h}-2), \ldots, \mathrm{pf}(\mathrm{h}-5)$. Equation (9) then becomes the sum of six equations as follows.

$$
\begin{align*}
& l_{1}=\left(\frac{1}{S D_{t}^{2} \sqrt{2 \pi}} \int_{0}^{h_{R}-5} e^{\frac{-\left(h-L_{t}\right)^{2}}{2 S D_{t}^{2}}} d h\right)  \tag{91i}\\
& l_{2}=\left(\frac{1}{S D_{t}^{2} \sqrt{2 \pi}} \int_{h_{R}-5}^{h_{R}-4} e^{\frac{-\left(h-L_{t}\right)^{2}}{2 S D_{t}^{2}}} d h\right)(1-p f(x-5))  \tag{91ii}\\
& l_{3}=\left(\frac{1}{S D_{t}^{2} \sqrt{2 \pi}} \int_{h_{R}-4}^{h_{R}-3} e^{\frac{-\left(h-L_{t}\right)^{2}}{2 S D_{t}^{2}}} d h\right)(1-p f(x-4)) \tag{9liii}
\end{align*}
$$

$$
\begin{align*}
& l_{4}=\left(\frac{1}{S D_{t}^{2} \sqrt{2 \pi}} \int_{h_{R}-3}^{h_{R}-2} e^{\frac{-\left(h-L_{t}\right)^{2}}{2 S D_{t}^{2}}} d h\right)(1-p f(x-3))  \tag{9liv}\\
& l_{5}=\left(\frac{1}{S D_{t}^{2} \sqrt{2 \pi}} \int_{h_{R}-2}^{h_{R}-1} e^{\frac{-\left(h-L_{t}\right)^{2}}{2 S D_{t}^{2}}} d h\right)(1-p f(x-2))  \tag{91v}\\
& l_{6}=\left(\frac{1}{S D_{t}^{2} \sqrt{2 \pi}} \int_{h_{R}-1}^{h_{R}} e^{\frac{-\left(h-L_{t}\right)^{2}}{2 S D_{t}^{2}}} d h\right)(1-p f(x-1))  \tag{9lvi}\\
& Q P R_{t}=l_{1}+l_{2}+l_{3}+l_{4}+l_{5}+l_{6} \tag{91}
\end{align*}
$$

## Results and Discussion

The pattern of yield per recruit at different fishing mortalities was similar for the different size limits except for $\geq 70 \mathrm{~mm}$ TD which peaked and dropped at smaller F values (Fig. B1). The $\mathrm{F}_{\max }$ values for YPR increased with increasing minimum sizes (Table B2). $\mathrm{F}_{\text {max }}$ values increased with the lowering of maximum sizes because with a smaller maximum size a greater F would be required to catch available urchins within a smaller size range (Table B2).

The relative contribution of egg production (percentage of the total egg production by an unfished population at $\mathrm{M}=0.10$ ) by female red sea urchins at various size limits and fishing mortalities is shown in Fig. B1. As F increases, fewer mature females reproduced and EPR was reduced. EPR is highest for $\geq 100 \mathrm{~mm}$ TD and lowest for $\geq 70 \mathrm{~mm}$ TD for only the minimum size limits at all F values (Fig. B1). Lowering the maximum size had a dramatic effect on protecting mature females and increasing egg production (Fig. B1).

At $\mathrm{F}=0.025$ (the approximate current fishing mortality in the B.C. red sea urchin fishery) YPR was decreased from $\geq 80 \mathrm{~g}$ to $<60 \mathrm{~g}$, but EPR increased from 73-79 $\%$ to $87-99 \%$ when different maximum sizes were included (Table B3).

The YPR/EPR used in this paper contains, as with many dynamic pool models, oversimplifications of reality, such as constant instantaneous natural mortality for all size groups and constant recuitment. Complicating the understanding of stock recruitment of red sea urchins is the sporadic recruitment and uncertainty of the source of recruitment for a given area. Further research is required to provide more information on stock recruitment relationships and effects of density on growth, mortality and reproduction potential of red sea urchins.

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## References

Breen, P.A. 1984. Sea urchins: suitability of the present size limit. Can. MS. Rep. Fish. Aquat. Sci. 1774: 2551.

Botsford, L.W., J. F. Quinn, S.R. Wing, and J.G. Brittnacher. 1993. Rotating spatial harvest of a benthic invertebrate, the red sea urchin, Strongylocentrotus franciscanus. pp. 409-428. In: Proceedings of the international symposium on management strategies for exploited fish populations, Alaska Sea Grant College Program, AK-SG-93-02.
Campbell, A. 1997. Possible criteria for reopening the northern abalone (Haliotiskamtschatkana) fishery in British Columbia. Can. Stock Assessment Secretariat Res. Doc. 97/64.
Lai, H.L. and Bradbury, A. 1998. A modified catch-at-size analysis model for a red sea urchin (Strongylocentrotus franciscanus) population. Can. Spec. Publ. Fish. Aquat. Sci. 125: 85-96.
Pfister, C.A., and A. Bradbury. 1996. Harvesting red sea urchins: recent effects and future predictions. Ecol. Appl. 6: 298-310.
Tegner, M.J. and P.K. Dayton. 1977. Sea urchin recruitment patterns and implications of commercial fishing. Science 196: 324-326.
Tegner, M.J., P.A. Breen, and C.E. Lennert. 1989. Population biology of red abalone, Haliotis rufescens, in southern California, and management of the red and pink, H. corrugata, abalone fisheries. Fish. Bull., U.S. 87: 313 - 339 .

Quinn II, T.J. and R.B. Desiro. 1999. Quantitative fish dynamics. Oxford Univ. Press, N.Y.

Table B1. Input parameters for yield per recruit and egg per recruit analysis using biological data collected from red sea urchins at Kendrick Island (after A. Campbell et al. unpublished data).

| Parameter | Values |
| :---: | :---: |
| Minimum minimum legal size | 70 |
| Maximum minimum legal size | 120 |
| Step size for minimum legal size | 5 |
| Minimum fishing mortality | 0.00 |
| Maximum fishing mortality | 0.50 |
| Step size for fishing mortality | 0.025 |
| Maximum legal size | $\begin{gathered} \text { None, } 110,120, \\ 130,140 \end{gathered}$ |
| Asymptotic length L-infinity | 145.634 |
| Brody growth coefficient | 0.157 |
| Hypothetical age of zero weight t-zero | 0.690 |
| Gonad length-weight curve coefficient a | $8.6424 \times 10^{-8}$ |
| Gonad length-weight curve coefficient b | 4.310 |
| Total length-weight curve coefficient a | $8.0715 \times 10^{-4}$ |
| Total length-weight curve coefficient b | 2.835 |
| Oldest age | 50 |
| Natural mortality | . 10 |
| Original number of females | 100 |
| Standard deviation of length-at-age curve coefficient a | 6.8319856 |
| Standard deviation of length-at-age curve coefficient b | 0 |
| Length-fecundity curve coefficient c | 16.8234 |
| Length-fecundity curve coefficient d | 3.2659 |
| Length-maturity curve coefficient a | 12.9571 |
| Length-maturity curve coefficient b | 0.1594 |
| Length interval width | 10 |
| Proportion of animals exposed to fishing at minimum legal size - 1 unit | 0.8887 |
| Proportion of animals exposed to fishing at minimum legal size - 2 units | 0.5151 |
| Proportion of animals exposed to fishing at minimum legal size - 3 units | 0.1237 |
| Proportion of animals exposed to fishing at minimum legal size - 4 units | 0.0184 |
| Proportion of animals exposed to fishing at minimum legal size - 5 units | 0.0025 |

Table B 2 . Biological reference points $\mathrm{F}_{\text {max }}, \mathrm{F}_{0.1}$, and $\mathrm{F}_{0.35}$ from the yield per recruit analyses when $\mathrm{M}=0.10$ for red sea urchins from Kendrick Island at different size limits. Methods for calculating these values are explained in Quinn and Desiro (1999).

| Size limit (mm TD) |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Min | Max | $\mathrm{F}_{\max }$ | $\mathrm{F}_{0.1}$ | $\mathrm{~F}_{0.35}$ |
| 70 | none | 0.17 | 0.08 | 0.04 |
| 90 | none | 0.30 | 0.11 | 0.05 |
| 100 | none | 0.47 | 0.13 | 0.05 |
| 100 | 110 | 1.00 | 0.56 | 0.18 |
| 100 | 120 | 1.00 | 0.45 | 0.15 |
| 100 | 130 | 1.00 | 0.31 | 0.10 |
| 100 | 140 | 0.95 | 0.19 | 0.06 |
| 90 | 110 | 1.00 | 0.84 | 0.24 |
| 90 | 120 | 1.00 | 0.33 | 0.12 |
| 90 | 130 | 0.86 | 0.25 | 0.09 |
| 90 | 140 | 0.59 | 0.16 | 0.06 |

Table B3. Yield per recruit and egg per recruit (\% of unfished population, $5.61 \mathrm{E}+12$ eggs) results when $\mathrm{M}=0.10$ and $\mathrm{F}=0.025$ for red sea urchins from Kendrick Island at different size limits.

| Size limit (mm TD) |  | EPR | YPR |
| :---: | :---: | :---: | :---: |
| Min | Max | (\%) | (g) |
| 70 | none | 73.63 | 87.4 |
| 90 | none | 77.12 | 84.8 |
| 100 | none | 79.31 | 81.5 |
| 100 | 110 | 99.78 | 9.09 |
| 100 | 120 | 98.89 | 20.0 |
| 100 | 130 | 96.29 | 34.6 |
| 100 | 140 | 88.90 | 54.5 |
| 90 | 110 | 99.44 | 14.2 |
| 90 | 120 | 98.21 | 24.8 |
| 90 | 130 | 95.15 | 38.9 |
| 90 | 140 | 87.13 | 58.1 |



Fig. B1. (A) Yield per recruit (g) and (B) egg per recruit (percentage of total eggs of unfished population) analyses using $\mathrm{M}=0.10$ and red sea urchin data from Kendrick Island. Numbers next to line in graphs indicate the different legal size limit options (e.g., 100140 indicates 100-140 mm TD).


[^0]:    ${ }^{1}$ Area $106=115.3 \mathrm{t}$

