# Dynamics of Marine Fish Assemblages in Hecate Strait, British Columbia 

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## Canadian Technical Report of Fisheries and Aquatic Sciences

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by

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#### Abstract

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Data was analysed from the Hecate Strait assemblage surveys conducted between 1984 and 2003. The results supported earlier analyses and indicated that three distinct fish assemblages were present over the depth range of 20-220 metres. The assemblages persisted over time with little bathymetric variation. The dominant species components of the assemblage included regular species, present year round, seasonal species, present seasonally and transient species, moving among assemblages. Diversity, species richness and evenness changed very little over the study period. Depth was a prominent feature of all of the Assemblages. The shallowest assemblage, Reef Island, had a median depth of 45 meters with the biomass dominated by spotted ratfish, rock sole, spiny dogfish, Pacific halibut and big skate. The intermediate assemblage, Bonilla, had a median depth of 74 meters and English sole, spotted ratfish, spiny dogfish, arrowtooth flounder and Pacific sanddab were dominant proportions of the biomass. The deepest assemblage, Butterworth, had a median depth of 101 meters and arrowtooth flounder, English sole, spotted ratfish, Dover sole and rex sole were the dominant components. Diversity, species richness and evenness metrics changed very little over the study period although species relative abundance changed over time. The assemblages defined here appear to be useful units for multispecies stock assessments for the region.


## RÉSUMÉ

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Les données des relevés des assemblages dans le détroit d'Hécate effectués entre 1984 et 2003 ont été analysées. Les résultats ont corroboré les analyses précédentes et ont indiqué que trois assemblages de poissons distincts étaient présents à des profondeurs de 20 à 220 m . Les assemblages ont résisté au cours du temps, ne présentant qu'une faible variation dans les données bathymétriques. Les espèces dominantes de l'assemblage étaient les espèces habituelles, les espèces présentes à longueur d'année, les espèces saisonnières, les espèces présentes de manière saisonnière et les espèces de passage se déplaçant parmi les assemblages. La diversité ainsi que la richesse et l'uniformité des espèces ont très peu changé au cours de la période à l'étude. La profondeur était une caractéristique importante de tous les assemblages. L'assemblage le moins profond, celui de l'île Reef, était situé à une profondeur moyenne de 45 m , où la biomasse était dominée par la chimère d'Amérique, la fausse limande, l'aiguillat commun, le flétan du Pacifique et la raie biocellée. L'assemblage intermédiaire de la pointe Bonilla, lui, était situé à une profondeur moyenne de 74 m où la sole anglaise, la chimère d'Amérique, l'aiguillat commun, la plie à grande bouche et la limande sordide étaient les espèces dominantes dans la biomasse. L'assemblage le plus profond, celui de Butterworth, était quant à lui situé à une profondeur moyenne de 101 m et la plie à grande bouche, la sole anglaise, la chimère d'Amérique, la limande-sole et la plie royale étaient les espèces dominantes de la biomasse. Les paramètres relatifs à la diversité ainsi qu'à la richesse et à l'uniformité des espèces ont très peu changé au cours de l'étude même si l'abondance relative des espèces a changé au cours du temps. Les assemblages décrits ici semblent constituer des unités utiles pour l'évaluation des stocks de bon nombre d'espèces dans la région.

## INTRODUCTION

In recent years scientists have increasingly focused on species interactions and environmental influence with primary consideration of the structure and function of a marine ecosystem in addition to sustainable fishing yields (Pikitch et al. 2004) (Table 1). Pauly and Christensen (2002) predict that if fisheries management does not progress towards incorporating ecosystem consideration the structure of the ecosystem can be altered irreversibly as in the case of the trophic cascades observed after top predators are eliminated by commercial fisheries.

A study of the Hecate Strait ecosystem began in April of 1984 with the primary objective of conducting research into the ecological basis for multispecies stock assessment for that region. The research was jointly carried out by the Pacific Biological Station in Nanaimo, B.C. and the Institute of Ocean Sciences in Sidney, B.C. on Vancouver Island. The fishery in Hecate Strait targets approximately 45 species taken by various gear types. The fishery is small on a world scale but this lends itself to modelling as many of the physical processes occur on a geographically tractable scale and the groundfish fleet size over time has included less than 50 vessels (Tyler 1989).

One of the sub projects was related to analysis and distribution of demersal fish assemblages using research survey data. Commensurate with this the multispecies groundfish Species Assemblage Research Survey was initiated in 1984 and continued through 2003. The catch rate data from those surveys are the focus of this report. Information on diet of fish species in the Strait was also collected on the surveys and those data were investigated using a mass balance approach (Pearsall and Fargo 2007).

A total of 12 systematic bottom trawl surveys of Hecate Strait were conducted by the Pacific Biological Station (Fargo et al. 1990, Tyler 1989) between 1984 and 2003. Survey haul locations are summarized in Figure 1. The fishing gear used for all surveys was a Yankee 36 trawl net. The survey design has been summarised by Fargo et al. (1990). Briefly, haul locations were arrayed over a grid of $19 \mathrm{~km}^{2}\left(10 \mathrm{~nm}^{2}\right)$ blocks, with an attempt made to establish one trawl site within each 18 m depth interval within each block. The 1984 survey involved one chartered and one research vessel, while all other surveys were single-vessel surveys. A survey in winter was conducted in 1986. The summer surveys provide synoptic data that can be used to map fish assemblages available to bottom trawlers, and to provide an index of relative abundance indices for species inhabiting the region. To date only data from the first three surveys has been analysed (Fargo and Tyler 1991).

This report summarises the results of analysis for all summer Species Assemblage Surveys from 1984-2003. It focuses on the distribution of fish
assemblages, the physical and biological characteristics associated with them and their change in structure over time. It includes a summary of the ecosystem metrics associated with the region along with an evaluation of their utility.

## METHODS

## DATA

Species catch composition data were the basis for a cluster analysis to characterize demersal fish assemblages in the Strait. Data for each survey were analysed independently in the following way: 1 . Species composition data for usable stations (trawl sites with no mishaps) were converted to species percentage composition by weight. For the two-vessel survey (1984), the data sets were combined. 2. The data was then analysed using an agglomerative clustering technique, where individual stations represented collections (entities) and species catch composition represented attributes. Cluster analysis was used to group sites into geographic areas with relatively homogeneous species composition. We rejected the reverse clustering of species that would group those species that occurred together in relatively equal proportions among sites since this technique ignores the fact that ubiquitous species should be classified as part of more than one assemblage. We preferred the alternative of examining and comparing the composition of the dominant species within clusters of sites, where sites are grouped by relatively similar percentage species composition. Previous studies have shown that fairly homogeneous habitat areas can be delineated by clustering the sites, and that a list of the species tending to cohabit will result (Gabriel \& Tyler 1980, Gomes et al. 1989, Overholtz \& Tyler 1985). We define the dominant species as collectively, making up at least $90 \%$ of the biomass of any assemblage.

## CLUSTER ANALYSIS

A general question facing researchers in many areas of inquiry is how to organize observed data into meaningful structures. Cluster analysis (first used by Tyron 1939) is an exploratory data analysis tool which aims at sorting different objects into groups in a way that the degree of association between two objects is maximal if they belong to the same group and minimal otherwise. Cluster analysis simply discovers structures in data without explaining why they exist. Discussions of the results refer to clustering algorithms and do not mention statistical significance testing. In fact, cluster analysis is a collection of different algorithms that put objects into clusters according to well defined similarity rules. Unlike many other statistical procedures, cluster analysis methods are mainly used when we do not have any a priori hypotheses. In that sense, cluster analysis finds the most significant solution possible. Therefore, statistical significance testing is really not appropriate.

Cluster analysis in this study was carried out according to the following procedure. First, a Bray-Curtis dissimilarity coefficient was calculated for each
pair of sites then clustering was based on a group average fusion strategy. A dendrogram was produced for each analysis showing the dissimilarity level among sites. The software used for the cluster analysis was an R language function calling the agglomerative nesting routine AGNES. The starting point for this analysis was a data matrix consisting of $n$ rows of samples (stations) and $p$ columns of variables (species) with individual values representing the proportion of the catch of each haul represented for each species caught, called an $n \times p$ ( $n$ by $p$ ) matrix for each survey. A matrix of dissimilarity was then calculated among sites.

The Bray-Curtis index is a modified Manhattan measurement where the summed differences between the variables are standardised by the summed variables of the objects. The general equation of the Bray-Curtis dissimilarity is:

$$
d^{B C D}(i, j)=\frac{\sum_{k=0}^{n-1}\left|y_{i, k}-y_{j, k}\right|}{\frac{\sum_{k=0}^{n-1}\left(y_{i, k}+y_{j, k}\right)}{}}
$$

In the equation above, $d^{B C D}$ is the Bray-Curtis dissimilarity between the objects $i$ and $j, k$ is the index catch proportion and $n$ is the total number of samples (stations) $y$. The Bray-Curtis dissimilarity index is bounded between 0 and 1, where 0 means the two sites have the same composition (that is they share all the species), and 1 means the two sites do not share any species.

Using this approach samples (stations) all start out as individuals, and the two samples most similar (or least dissimilar) are fused to form the first cluster. Subsequently, samples are continually fused one-by-one in order of highest similarity (or equivalently lowest dissimilarity) to the sample or cluster to which they are most similar. The hierarchy is determined by the cluster at a height characterized by the similarity at which the samples fused to form the cluster. Eventually, all samples are contained in the final cluster at similarity 0.0 or dissimilarity of 1.0.

After the dissimilarity matrices were computed for each survey, clustering was done via the group-average method (UPGMA (Unweighted Pair Group arithiMetric Average) (Kaufman, L. and Rousseeuw, P.J. (1990). For the group average version of hierarchical clustering, the proximity of two clusters is defined to be the average of the pairwise proximities between all pairs of points in the different clusters. This is expressed by the following equation:

$$
\operatorname{proximity}(\text { cluster } 1, \text { cluster } 2)=\sum_{\substack{p 1 \operatorname{ccluster} 1 \\ p 2 \in \text { cluster } 2}} \operatorname{proximity}(p 1 p 2)
$$

$\overline{\text { (size(cluster1))(size(cluster2)) }}$
(Steinbach, M. 2000)

## DENDROGRAMS

Fargo \& Tyler (1991) used a previously defined rockfish assemblage adjacent to the survey area to assist in interpretation of the dendrograms. The deep-water rockfish complex in Moresby Gully, in southern Hecate Strait is considered to be a discrete group based on biological and geographic characteristics (Leaman \& Nagtegaal 1987). The cluster for this group of species joined with other major clusters in the dendrograms at dissimilarity levels ranging from 0.65 to 0.77 , and values within this range were subsequently used to interpret the dendrograms for the period 1984-2003.

Dendrograms were also compared to Non-metric multidimensional scaling (nMDS) ordination plots to determine if assemblages were geographically distinct and persistent through time (Tyler and Gabriel 1982, Overholtz and Tyler 1985, Gabriel 1992; Gomes et al. 2001, Jørgensen et al. 2005; Duffy-Anderson et al. 2006).

## CPUE ANALYSIS

Bootstrapped mean catch rates were computed for the species components of each assemblage as indicated by the following formula (Fargo et al. 1991).

$$
U_{i j}=\left(\sum_{k=1}^{n} C_{i k} / E_{i k}\right)
$$

where:
$U_{i j}$ is the mean CPUE in kg•hr-1 for species $i$ in assemblage $j$ and $C_{j k}$ is the weight of species $i(\mathrm{~kg})$ in haul $k$ for assemblage $j, E_{i k}$ is the effort expended to catch species $i$ in haul $k$ for assemblage $j, k$ is a vector representing the individual stations in assemblage $j$ and $n$ is the total number of stations in assemblage $j$.

## ECOLOGICAL METRICS

Diversity, species richness and evenness were estimated within and among assemblages. Shannon's diversity index (Pielou 1977) was calculated as follows:

$$
\text { diversity }=1-\sum_{i=1}^{S} P_{i} \ln \left(P_{i}\right)
$$

where:
$S$ is the number of species and $P_{i}$ is the proportion of the sample comprised of species $i$. The index takes on values from 0 (the entire sample consists of only one species) to a maximum of $\ln S$ for a given $S$. Diversity increases as species number increases and dominance decreases. The index is not directly sensitive to density, since it is based on proportionalized data. Diversity is considered here as a single statistic and the number of species and the evenness of the histogram of species' biomass proportions are confounded. Thus, an assemblage with few species and high evenness could have the same diversity as another assemblage with many species and low evenness. We therefore examined species richness and evenness for each assemblage in addition to diversity.

Species 'richness', $S^{*}$, for each assemblage was equal to the number of fish species present in the data. The 'evenness', of the distribution of individuals among the assemblages was estimated using the following equation (Pielou 1977):

$$
V^{\prime}=H^{\prime} / \ln S^{*}
$$

where:
$V$ ís a measure of evenness, $H^{\prime}$ is Shannon's diversity index for an assemblage and $S^{*}$ is the number of species present in an assemblage.

It is useful to assess the total species richness of the study area as a measure of the completeness of the species inventory and this was done as well. One method to estimate total species richness or maximum number of species present in an ecosystem, $S_{\text {max }}$, is derived from a species accumulation curve (Simberloff 1972). This is a cumulative curve of the number of new species encountered in relation to the sampling effort. $S_{\max }$ is equivalent to the asymptote of the curve assuming that the sampling effort was adequate. An alternative to these computations is a simple but reliable non-parametric method that has been shown to be effective on fish communities; the Chao 2 index (Chao 1984) which can be calculated from the following formula:

$$
S_{\max }=S_{o b s}+\left(a^{2} / 2 b\right)
$$

where:
$S_{\text {obs }}$ is the number of species caught, $a$ is the number of species only found in one sample and $b$ is the number of species only found in two samples (Chao 1984).

## RESULTS

## CLUSTER ANALYSIS

Dendrograms for each survey are depicted in Figures (2-12). Species assemblages classified from the dendrograms were named after prominent landmarks in their geographic areas. The different colours for the legs of the dendrograms represent corresponding species percent composition for clustered sites. The labels at the bottom of each dendrogram leg indicate the specific geographical trawling site. The height of the dendrogram legs represents the level of dissimilarity between sites. Thus sites that cluster out at a height of 0.2 are less dissimilar than sites clustering out at a height of 0.6.

Dendrogram legs in black represent sites that could not be classified at the chosen dissimilarity level. Many of these represent species composition that is intermediate to two assemblages. This relates to the gradual transition from one assemblage to another near boundaries where species composition blurs and components of more than one assemblage occur. It is also characteristic of the bathymetry in Hecate Strait which does not follow even lines but can vary somewhat during a trawling operation. In some cases, 1984, 1989, 2000, for instance clusters for one assemblage are not adjacent to one another in the dendrogram. This was due to the fact that although the species composition was similar the relative percentage of species components differed. Fargo and Tyler (1991) noted this in their analysis as well.

Four clusters appear in the dendrogram for 1984, one each for Butterworth and Bonilla and two for Reef Island. The two for reef Island are not adjacent. This may partially reflect different catchability coefficients for the two vessels. Reef Island and Bonilla are more closely related than other combinations. Butterworth assemblage contains the fewest haul stations while the rest are evenly split between Reef Island and Bonilla.

In the dendrogram for 1987, there are three clusters, one for each assemblage. There are also several long legs in the dendrogram that lie between the Bonilla assemblage and the Reef Island assemblage. The species composition here reflected both assemblages. Reef Island and Bonilla appear to be more closely related than other combinations. Trawling sites are fairly evenly split among assemblages.

The dendrogram for 1989 shows three distinct clusters at the 0.75 dissimilarity level. Each of these represents a different assemblage. The long legs in black adjacent to the Reef Island assemblage contain catches with a low number of species that were difficult to classify. Butterworth and Bonilla are more closely related than other combinations. This was due to the presence of turbot in the catches of both assemblages. Bonilla and Butterworth contain the most stations while Reef Island contains the fewest.

The clusters in the dendrogram for 1991 represent all three assemblages at the 0.7 dissimilarity level. The Butterworth assemblage represents more stations than either of the other two assemblages and Butterworth and Bonilla are more closely related as in 1989. The Butterworth assemblage is represented by the highest number of stations while Bonilla contains the fewest.

For 1993 and 1995 three distinct groups are clustered at the 0.7 dissimilarity level. Reef Island and Bonilla are more closely related than other combinations in 1993. Butterworth contains the most stations in 1993 and Reef Island the most in 1995. Bonilla contains the fewest stations in both surveys and is smaller than in previous years.

In 1996, Butterworth and Bonilla are most closely related while Reef Island represents the most stations and Bonilla the least. In 1998 Butterworth is clustered at the lowest dissimilarity level while Reef Island and Bonilla do no fall out until a dissimilarity level of 0.78 . Reef Island Bonilla are most closely related. Butterworth contains the most stations.

In 2000 the Bonilla assemblage is represented by split clusters. Bonilla and Reef Island are most closely related. Butterworth contains the highest number of stations. In 2002 Bonilla and Reef Island assemblages are most closely related and haul stations are more evenly split between assemblages than for any other year's survey. In 2003 Bonilla and Butterworth assemblages are most closely related. Bonilla assemblage is composed of the fewest stations while the rest of the stations are evenly split between Reef Island and Butterworth.

## FISH ASSEMBLAGES

Fargo and Tyler (1991) defined three fish assemblage groups with characteristic depth ranges and geographical boundaries. The results of this analysis corroborated the results of their analysis (Figures 13 and 14). Geographic boundaries were stable among years and the list of dominant species remained stable as well.

In the Reef Island assemblage Pacific cod, Pacific halibut, rock sole, spiny dogfish and spotted ratfish were all rank 1 species over time (Figure 15). Rock sole was the Rank 1 species in 1989, 1993, 1995 and 2000, accounting for 29\%, $24 \%, 18 \%$ and $30 \%$ of the biomass, respectively. Pacific halibut was the Rank 1 species in 1991 and 1998, accounting for 23\% and 22\% of the biomass, respectively. Spiny dogfish was the Rank 1 species in 1984 and 2002, accounting for $43 \%$ and $31 \%$ of the biomass, respectively. Spotted ratfish was the rank 1 species in 1996 and 2003, accounting for $26 \%$ and $42 \%$ of the biomass, respectively. Pacific cod was the rank 1 species in 1987, accounting for $52 \%$ of the biomass. Spotted ratfish and Pacific cod are transient species, occurring in all three assemblages and spiny dogfish is a seasonal (summer)
species in this assemblage. Rock sole and spotted ratfish accounted for $38 \%$ of the total biomass of this assemblage for (Figure 16).

Arrowtooth flounder was the rank 1 species in the Butterworth assemblage in all years except 1993 when English sole was the rank 1 species, accounting for $32 \%$ of the biomass (figure 17). Arrowtooth flounder accounted for $43 \%, 58 \%$, $54 \%, 45 \%, 37 \%, 46 \%, 28 \%, 40 \%, 36 \%$ and $49 \%$ of the biomass in 1984-1991 and 1995-2003. Arrowtooth flounder accounted for $39 \%$ of the total biomass in this assemblage while English sole, spotted ratfish, Dover sole accounted for $10 \%$ each and rex sole for $9 \%$ of the biomass. Spiny dogfish was a seasonal component, accounting for $5 \%$ of the biomass overall (figure 18).

Rank 1 species in the Bonilla assemblage were; English sole accounting for $35 \%$, $46 \%, 45 \%$ and $32 \%$ of the biomass in 1984, 1989, 1991 and 2003(figure 19). Spiny dogfish accounted for $48 \%, 60 \%$ and $35 \%$ of the biomass in 1987, 1993 and 1998, arrowtooth flounder for $25 \%$ of the biomass in 1995, walleye pollock for $26 \%$ of the biomass in 1996 and spotted ratfish for $33 \%$ and $36 \%$ of the biomass in 2000 and 2002. English sole accounted for $24 \%$ of the total biomass, spotted ratfish for $13 \%$, spiny dogfish for $11 \%$, arrowtooth flounder for $8 \%$, Pacific sanddab, rex sole and Pacific cod for $6 \%$ each, spiny dogfish for $5 \%$ and walleye pollock for 4\% (Figure 20). Walleye pollock and spotted rattish were transient components in this assemblage while spiny dogfish was a seasonal component. Biomass proportion of by assemblage for dominant species components is presented in figure 21. Arrowtooth flounder is dominant in both the Butterworth and Bonilla assemblages but its presence in the Bonilla assemblage consisted mainly of juveniles. English sole, spiny dogfish and spotted ratfish are dominants in all three assemblages.

## CPUE

## Reef Island

CPUE was highest for spiny dogfish, Rock sole and Pacific cod although the high rate for Pacific cod was the result of one large catch in 1987 (Figure 22, Table 2). There was an increasing trend in CPUE for spotted ratfish, rock sole and sand sole and a decreasing trend for spiny dogfish, big skate, lingcod and Pacific halibut. Without the large CPUE in 1987 Pacific cod CPUE showed little trend. Rock sole showed the largest increase in CPUE from $20 \mathrm{~kg} / \mathrm{h}$ in 1984 to $108 \mathrm{~kg} / \mathrm{h}$ in 2003. Spiny dogfish showed the largest decrease in CPUE from $79 \mathrm{~kg} / \mathrm{h}$ in 1984 to $12 \mathrm{~kg} / \mathrm{h}$ in 2003. The average coefficient of variation ranged from 0.24 for rock sole to 0.56 for Pacific cod.

Boxplots of CPUE for all species (for which there was enough data) in the Reef Island assemblage provide further detail (figure 23 and 24). The lowest rates observed were those for small non-schooling species like poachers and sculpins. Catch rates for foraging species such as shiner perch and Pacific sand lance
were slightly higher. The highest rates observed were for flatfish species, specifically rock sole and Pacific halibut and elasmobranchs, big skate, spiny dogfish and spotted ratfish. Flatfish species also comprised much of the intervening distribution with the exception of Pacific herring, Pacific cod and yellowtail rockfish.

## Butterworth

For the Butterworth assemblage, CPUE were highest for arrowtooth flounder and lowest for spiny dogfish (Figure 25, Table 3). There was an increasing trend for rex sole, a decreasing trend for arrowtooth flounder and spiny dogfish and no obvious trend for spotted ratfish, Dover sole and English sole. Spotted ratfish is a transient component of this assemblage while spiny dogfish is a seasonal (summer) component. The CPUE in 1993 for English sole was nearly three times that of any other in that series. The largest increase in CPUE was that for rex sole which went from $31 \mathrm{~kg} / \mathrm{h}$ in 1984 to $200 \mathrm{~kg} / \mathrm{h}$ in 2002. Spiny dogfish CPUE showed the largest decrease from $109 \mathrm{~kg} / \mathrm{h}$ in 1984 to $13 \mathrm{~kg} / \mathrm{h}$ in 2003. Average CV ranged from 0.25 for spiny dogfish to 0.34 for Dover sole. Boxplots of CPUE by species for the Butterworth assemblage are presented in Figure 26 and 27. The lowest rates observed were those for poachers, eelpouts and small foraging species. The highest rates observed were those for arrowtooth flounder, rex sole and English sole, with arrowtooth flounder dominating. The slope of the medians for species in the Butterworth assemblage is steeper than in the Reef Island assemblage indicating fewer species contribute to the total biomass of this assemblage.

## Bonilla

CPUE for rex sole, Pacific sanddab and spotted ratfish increased over the period 1984-2003 (Figure 28, Table 4) but showed little trend for English sole, arrowtooth flounder and spiny dogfish. Rates were highest for English sole and lowest for Pacific sanddab and rex sole. The largest increase in CPUE occurred for spotted ratfish which went from $11 \mathrm{~kg} / \mathrm{h}$ in 1984 to $181 \mathrm{~kg} / \mathrm{h}$ in 2002 while the largest decrease occurred for spiny dogfish which went from $64 \mathrm{~kg} / \mathrm{h}$ in 1984 to $17 \mathrm{~kg} / \mathrm{h}$ in 2003. CPUE for spotted ratfish in 2000 and 2002 was more than twice as high as that for any other year in that time series. CPUE for rex sole in 2003 was more than twice as high as that for any other in that time series. Average CV for this assemblage ranged from 0.31 for spiny dogfish to 0.50 for arrowtooth flounder.

Boxplots of CPUE by species for the Bonilla assemblage are presented in Figures 29 and 30. The lowest rates observed were for small species such as sculpins, poachers, eelpouts and small foraging species. However, two flatfish species, speckled sanddab and butter sole were in this group as well. The highest rates observed were those for English sole, Pacific halibut, big skate, spiny dogfish and Pacific sanddab. Pacific sanddab is a relatively small species
that appears to be quite abundant in this assemblage. The slope of the medians indicates that species contribute more evenly to the biomass of this assemblage than was observed for the other assemblages.

## Depth distribution

The sequence of Boxplots in Figure 31 provides bathymetric information for 80 species including many non commercial species although it is incomplete for species at the outer limits. There is a continuous trend in median depths between 30 and 140 m . After that the trend becomes noticeably steeper indicating fewer species per unit depth and slope rockfish species begin to dominate. Flatfish species are found throughout the overall range. Shelf and inshore rockfish species are present throughout the range as well. Boxplots of depths for each assemblage are presented in Figures 32 and 33. Median depth was 45 meters for the Reef Island assemblage 74 meters for the Bonilla assemblage and 101 meters for the Butterworth assemblage.

Figure 34 contains boxplots of depth by dominant species for the combined surveys. Pacific sanddab occupied the depth range of the Bonilla assemblage almost exclusively. The shallowest depth preference was exhibited by sand sole encountered almost entirely in the Reef Island assemblage while the deepest preference was exhibited by flathead sole, split between the Butterworth and Bonilla assemblages. Sand sole, big skate, butter sole, rock sole, Pacific halibut and lingcod occupied the depth range of the Reef Island assemblage almost exclusively. English sole, spiny dogfish, Pacific cod and spotted ratfish inhabited the depths of all assemblages to some degree while Dover sole, rex sole, walleye pollock and flathead sole mainly occupied the depths of Bonilla and Butterworth assemblages. Overall, median depth overlapped for lingcod and Pacific sanddab only. The medians for all others in the Bonilla assemblage were greater than for dominant species in the Reef Island assemblage and less than for species in the Butterworth assemblage.

The relationship between mean depth and median depth vs the number of species encountered was dome-shaped (figure 35). A peak was apparent at ~60-69 meters for both. The mean depth for most species fell within a 30-140 m range. However because of the fishing gear used the depths of deepwater rockfish species were not sampled completely. Similarly, the depth range of 1 to 20 meters was not completely sampled.

## ECOLOGICAL METRICS

The diversity index ranged from 2.4 to 3.5 between 1984 and 2003 (Figure 36, Table 5). There is no significant trend apparent over this time period for any of the assemblages. However annual differences in the index were as high as 63\%. Species richness ranged from 28 to 56 over the time period analysed (Figure 37). The index fluctuates least in the Bonilla assemblage and most in the Butterworth
assemblage. Transient species that move among assemblages as well as the distribution of species that are in the margins of their depth range is also a factor. Species richness is highest for the Butterworth assemblage and lowest for the Reef Island assemblage. The estimated maximum number of fish species present in the Hecate Strait ecosystem over a depth range of 20-220 m the Chao 2 method was 153. Hart (1972) lists $\sim 150$ marine fish species present over that depth range.

Evenness showed the least fluctuation of all the ecological indices indicating that the proportions of biomass are relatively constant e.g. abundance has not changed dramatically over the time period for any of the assemblages (Figure 38). The relationship between richness and evenness was inverse for Reef Island and Bonilla and proportionate for Butterworth. However none of the relationships were significant.

## DISCUSSION

As mentioned previously scientists use cluster analysis mainly without a priori hypotheses. Results require close inspection to explain the grouping. Several phenomena affected the results of the cluster analysis in this case. Trawling stations near the assemblage boundaries resulted were unclassifiable or were classified as separate clusters. Stations where very few species were caught remained unclassifiable and appeared as long legs in the dendrograms. The timing of migration of important species such as spiny dogfish can produce misleading information regarding its role in assemblages. Careful examination of the results can help to avoid these problems.

Species composition and bathymetric boundaries of assemblages were relatively stable over the survey period. Historically, the abundance of English sole, rock sole and Pacific cod has exhibited dramatic fluctuations which will likely have an important effect on assemblage production. Similarly, transient species such as spiny dogfish and spotted ratfish exert an effect on all three assemblages. Given the significant proportion of the total biomass that arrowtooth flounder comprises and its position as a top predator will likely affect production in more than one assemblage.

The effect of the bottom trawl fishery in Hecate Strait is the selective removal of commercially valuable species such as Pacific cod, English sole, rock sole and Dover sole while avoiding or minimising the removal of non commercial species. This may create an advantage for the non target species and have a detrimental effect in the long term. Quotas for commercial species may need to be lowered to prevent this. To date multispecies surplus production models developed in many studies predict group quotas that are lower than the sum of individual species quotas. This type of model should be developed for Hecate Strait and the results compared with results from single species assessments. In addition,
long term effects such as climate change exacerbate the need for this comparison and for a holistic approach to stock assessment.

Multispecies management takes species interactions into account and may be an improvement over single species management. It does not require the enormous amount of data that ecosystem management requires. Managers of fisheries off the coasts of Alaska and California have used a multispecies approach (Witherell et al. 2000; Field and Francis 2006). Managers of fisheries in Florida and the southeastern United States have recently adopted a multi-species approach as well (Pierce and Mahmoudi 2001).

Ecosystem metrics such as species diversity, richness and evenness do not appear to provide enough information about ecosystem structure for use as a management tool. However the value for species richness this area is higher than many temperate inshore ecosystems and is even impressive when compared with much large offshore ecosystems in the temperate zone (Table 6.). There will, undoubtedly, be improvements in the utility of these metrics with more research. A recent global study by Fisher et al. (2010) indicated that fish body size may act as a factor of considerable importance in mediating the relationship between global marine fish species richness and ecosystem functioning. Over the time period of this study the relative abundance of assemblage components underwent some change while diversity was not affected. The value of this metric is limited and may occur only in the very long term. It is doubtful that management measures based on diversity alone would be appropriate. Marine ecosystems today have already been considerably perturbed by modern fisheries and long term changes in the physical environment; the latter entirely beyond our control. However, knowledge of species interactions and their effect on production is of value. In this case more information on diet is needed to examine how interactions between species change over time. This will allow us to illuminate some of the processes that operate and affect ecosystem structure and function. That insight will certainly improve our understanding of the ecosystem and by extension our management and sustainable use of this natural resource.

## LITERATURE CITED

Bosman, S.H. 2005. Northumberland Strait Fish Assemblages: Patterns and Processes. Master of Science Thesis. Graduate Academic Unit of Biology. University of New Brunswick.
Chao, A. 1984. Nonparametric estimation of the number of classes in a population. Scand. J. Stat. 11:265-270.
Duffy-Anderson, J.T., Busby, M.S., Mier K.L., Deliyanides, C.M., and Stabeno, P.J. 2006. Spatial and temporal patterns in summer ichthyoplankton assemblages on the eastern Bering Sea shelf 1996-2000. Fish. Oceanogr. 15: 80-94.

Fargo, J., and Tyler, A.V. 1991. Sustainability of flatfish dominated fish assemblages in Hecate Strait, British Columbia, Canada. Neth. J. Sea Res. 27(3/4): 237-253.

Field, J. C., and Francis, R.C. 2006. Considering ecosystem-based fisheries management in the California Current. Mar. Policy 30: 552-569.

Fisher, J.A.D., Frank, K.T., and Leggett, W.C. 2010. Global variation in marine fish body size and its role in biodiversity-ecosystem functioning. Mar. Ecol. Prog. Ser. 405: 1-13, 2010.

Gabriel, W. L. 1992. Persistence of demersal fish assemblages between Cape Hatteras and Nova Scotia, Northwest Atlantic. J. Northwest. Atl. Fish. Sci. 14: 29-46.

Gabriel, W.L., and Tyler, A.V. 1980. Analysis of Pacific coast demersal fish assemblages. Mar. Fish. Rev. 42 : 83-88.

Gaertner J.C., Bertrand J.A., de Sola, L.G., Durbec J.P., Ferrandis, E., and Souplet, A. Large spatial scale variation of demersal fish assemblage structure on the continental shelf of the NW Mediterranean Sea MEPS 297:245-257.

Gifford, D.J., Collie, J.S., and Steele, J.H. 2009. Functional diversity in a marine fish community. - ICES J. Mar. Sci. 66: 791-796.

Gomes, M.C., Haedrich, R., and Rice, J.C. 1989. Fish assemblages on the Grand Bank of Newfoundland. Northwest Atlantic Fisheries Organization. SCR Doc. 89/75. Serial No. N1656 :1-32.
Gomes, M.C., Serrão, E., and Borges, M.F. 2001. Spatial patterns of groundfish assemblages on the continental shelf of Portugal. ICES J. Mar. Sci. 58: 633-647.
Gratwicke, B., and Speight, M.R. 2005b. The relationship between fish species richness, abundance and habitat complexity in a range of shallow tropical marine habitats. J. Fish Biol. 66:650-667.

Hart, J.L. 1973. Pacific Fishes of Canada. Fish. Res. Bd. Can. Bull. 180: 740p.

Hiddink, J.G., and Ter Hofstede, R. 2008. Climate induced increases in species richness of marine fishes. Global Change Biology Volume 14, Issue 3, pages 453-460.
Jørgensen, O.A., Hvingel, C., Møller, P.R., and Treble, M.A. 2005. Identification and mapping of bottom fish assemblages in Davis Strait and southern Baffin Bay. Can. J. Fish. Aquat. Sci. 62: 1833-1852.

Kaufman, L., and Rousseeuw, P.J. 1990. Finding Groups in Data: An Introduction to Cluster Analysis. Wiley, New York.
Montserrat, D., and Sanchez, P. 2000. Demersal fish assemblages and habitat characteristics on the continental shelf and upper slope of the northwestern Mediterranean. J. Mar. Biol. Assoc. UK.

Overholtz, W.J., and Tyler, A.V. 1985. Long-term responses of the demersal fish assemblages of Georges Bank. Fishery Bull. 83: 507-520.
Pattengill-Semmens, C.V. 2002. The reef fish assemblage of Bonaire Marine Park: An analysis of REEF fish survey data. Proc. $53^{\text {rd }}$ Gulf Carib. Fish. Inst. 53: 591-605.

Pauly, D., and Christensen, V. 2002. Ecosystem Models. p. 211- 227. In Hart, P. J. B. and Reynolds, J. D. (eds). Handbook of Fish Biology and Fisheries. Vol. 2. Blackwell Science Ltd. Malden, MA. 410p.
Pearsall, I.A., and Fargo, J.J. 2007. Diet Composition and Habitat Fidelity for Groundfish Assemblages in Hecate Strait, British Columbia. Can. Tech. Rep. Fish. Aquat. Sci. 2692: vi + 141 p.
Pielou, E.C. 1977. Mathematical ecology. John Wiley \& Sons Inc. New York: 385p.
Pierce, D. J., and Mahmoudi, B. 2001. Nearshore fish assemblages along the central west coast of Florida. Bull. Mar. Sci. 68: 243-270.
Pikitch, E.K., Santora, C., Babcock, E.A., Bakun, A., Bonfil, R., Conover, D.O., Dayton, P., Doukakis, P., Fluharty, D., Heneman, B., Houde, E.D., Link, J., Livingston, P.A., Mangel, M.K., McAllister, M., Pope, J., and Sainsbury, K.J. 2004. Ecosystem-based fishery management. Science 305: 346-347.

Shackell, N.T., and Frank, K.T. 2003. Marine Diversity on the Scotian Shelf, Canada. Aquat. Conserv. 13(4): 305-321.

Simberloff, D. 1972. Properties of the Rarefaction Diversity Measurement. American Naturalist 106:414-418.
Steinbach, M. 2002. An introduction to cluster analysis for data mining. from http://www.cs.umn.edu/-han/dmclass/cluster survey 1002 00.pdf
Tyler, A.V. [Ed] 1989. Hecate Strait Project: Results From Four Years of Multispecies Fisheries Research. Can. Tech. Rep. Fish. And Aquat. Sci. No. 1675: 60p.

Tyler, A. V., and Gabriel, W. L. 1982. Adaptive management based on structure of fish assemblages of Northern Continental Shelves, pp. 149-156. In M. Mercer (ed.). Multispecies approaches to fisheries management advice. Can. Spec. Publ. Fish. Aquat. Sci. 59.
Tryon, R.C. 1939. Cluster Analysis. Edwards Brothers.
Walker, B.K., Jordan, L.K.B., and Spieler, R.E. 2009. Relationship of reef fish assemblages and topographic complexity on southeastern Florida coral reef habitats. J. Coastal Res. SI53, 39-48.

Witherell, D., Pautzke, C., and Fluharty, D. 2000. An ecosystem-based approach for Alaska groundfish fisheries. ICES J. Mar. Sci. 57: 771-777

Wraith, J.A. 2007. Assessing Reef Fish Assemblages in a Temperate Marine Park Using Baited Remote Underwater Video Master of Science Thesis, Biological Sciences, University of Wollongong.

Table 1. Number of fish species in recent studies of marine ecosystems

| Area | Number of <br> fish <br> species in <br> study | Time frame | Source <br> data |
| :--- | :--- | :--- | :--- |
| This study | 121 | $1984-2003$ | Trawl survey |
| Scotian shelf $^{1}$ | 79 | $1970-2000$ | Trawl survey |
| Virgin Islands $^{2}$ | 35 | $2000-2001$ | Visual survey (reefs) |
| North Sea $^{3}$ | 90 | $1985-2005$ | Trawl survey |
| Georges Bank $^{4}$ | $35^{\text {a }}$ | $1993-2002$ | Trawl survey |
| Florida | 184 | $1998-2002$ | Dive survey (reefs) |
| N.E. Mediterranean ${ }^{6}$ | 79 | 1991 | Trawl survey |
| Australia | R.W. Mediterranean ${ }^{8}$ | 60 | $1996-1998$ |
| Netherlands Antilles $^{9}$ | 286 | $1993-1999$ | Trawl survey |
| Northumberland Strait $^{10}$ | 23 | $2001-2006$ | Trawl survey |

${ }^{1}$ Shackell and Frank (2003)
${ }^{2}$ Gratwicke and Speight (2005)
${ }^{3}$ Hiddink and Hofstede (2004)
${ }^{4}$ Gifford, Collie and Steele (2009)
${ }^{5}$ Walker, Jordan, Spieler, R.E. (2009)
${ }^{6}$ Demestre, Sanchez and Abello (2000)
${ }^{7}$ Wraith (2007)
${ }^{8}$ Gaertner, Bertrand, Gil de Sola, Durbec, Ferrandis and Souplet (2005)
${ }^{9}$ Pattengill-Semmens (2002)
${ }^{10}$ Bosman (2005)
${ }^{\text {a }}$ including squid

Table 2. Mean catch rates and $90 \%$ confidence limits for dominant species in the Reef Island species assemblage in Hecate Strait based on surveys conducted from 1984-2003.
$\left.\begin{array}{lllllllll} & \text { Year } & \text { Species } & \text { Mean } & \begin{array}{l}\mathbf{1 0 \%} \\ \text { c.i. }\end{array} & \begin{array}{l}\mathbf{9 0 \% .} \\ \text { c.i }\end{array} & \begin{array}{l}\text { Standard } \\ \text { deviation }\end{array} & \begin{array}{l}\text { Standard } \\ \text { error }\end{array} & \text { Variance }\end{array} \begin{array}{l}\text { Coefficient } \\ \text { of variation }\end{array}\right]$

| Year | Species | Mean | $\begin{aligned} & \text { 10\% } \\ & \text { c.i. } \end{aligned}$ | $\begin{aligned} & \text { 90\%. } \\ & \text { c.i } \end{aligned}$ | Standard deviation | Standard error | Variance | Coefficient of variation |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1993 | Spotted ratfish | 92.605 | 48.507 | 145.312 | 41.286 | 1.306 | 1704.534 | 0.446 |
| 1995 | Spotted ratfish | 13.089 | 7.364 | 19.716 | 5.058 | 0.16 | 25.583 | 0.386 |
| 1996 | Spotted ratfish | 63.192 | 33.397 | 96.951 | 26.878 | 0.85 | 722.427 | 0.425 |
| 1998 | Spotted ratfish | 34.539 | 15.745 | 59.412 | 18.515 | 0.585 | 342.805 | 0.536 |
| 2000 | Spotted ratfish | 10.571 | 6.442 | 15.477 | 3.579 | 0.113 | 12.809 | 0.339 |
| 2002 | Spotted ratfish | 29.677 | 18.629 | 42.679 | 9.586 | 0.303 | 91.891 | 0.323 |
| 2003 | Spotted ratfish | 150.755 | 97.462 | 212.394 | 46.967 | 1.485 | 2205.899 | 0.312 |
| 1984 | Pacific cod | 9.438 | 6.28 | 13.091 | 2.847 | 0.09 | 8.105 | 0.302 |
| 1987 | Pacific cod | 201.39 | 64.026 | 370.035 | 140.836 | 4.454 | 19834.78 | 0.699 |
| 1989 | Pacific cod | 11.191 | 4.376 | 20.508 | 8.078 | 0.255 | 65.254 | 0.722 |
| 1991 | Pacific cod | 5.677 | 2.272 | 10.52 | 4.307 | 0.136 | 18.55 | 0.759 |
| 1993 | Pacific cod | 19.17 | 7.178 | 35.271 | 13.686 | 0.433 | 187.307 | 0.714 |
| 1995 | Pacific cod | 6.892 | 3.267 | 11.844 | 3.911 | 0.124 | 15.296 | 0.567 |
| 1996 | Pacific cod | 8.022 | 3.868 | 13.183 | 4.122 | 0.13 | 16.991 | 0.514 |
| 1998 | Pacific cod | 11.456 | 5.346 | 19.342 | 6.367 | 0.201 | 40.539 | 0.556 |
| 2000 | Pacific cod | 3.956 | 2.599 | 5.51 | 1.195 | 0.038 | 1.428 | 0.302 |
| 2002 | Pacific cod | 10.523 | 4.463 | 18.868 | 6.393 | 0.202 | 40.87 | 0.608 |
| 2003 | Pacific cod | 8.874 | 4.573 | 14.54 | 4.061 | 0.128 | 16.492 | 0.458 |
| 1984 | Lingcod | 19.055 | 10.051 | 30.078 | 8.313 | 0.263 | 69.106 | 0.436 |
| 1987 | Lingcod | 18.419 | 9.424 | 28.3 | 7.715 | 0.244 | 59.521 | 0.419 |
| 1989 | Lingcod | 51.161 | 23.854 | 86.14 | 27.218 | 0.861 | 740.82 | 0.532 |
| 1991 | Lingcod | 68.564 | 39.462 | 101.176 | 27.367 | 0.865 | 748.953 | 0.399 |
| 1993 | Lingcod | 33.05 | 7.348 | 73.851 | 33.692 | 1.065 | 1135.151 | 1.019 |
| 1995 | Lingcod | 11.971 | 7.807 | 16.582 | 3.317 | 0.105 | 11.002 | 0.277 |
| 1996 | Lingcod | 5.985 | 3.594 | 8.579 | 2.039 | 0.064 | 4.158 | 0.341 |
| 1998 | Lingcod | 17.078 | 7.724 | 29.019 | 8.078 | 0.255 | 65.254 | 0.473 |
| 2000 | Lingcod | 2.581 | 1.778 | 3.473 | 0.728 | 0.023 | 0.53 | 0.282 |
| 2002 | Lingcod | 19.859 | 11.416 | 29.937 | 7.484 | 0.237 | 56.01 | 0.377 |
| 2003 | Lingcod | 11.457 | 5.219 | 19.338 | 6.108 | 0.193 | 37.308 | 0.533 |
| 1984 | Pacific halibut | 50.075 | 39.784 | 61.287 | 8.316 | 0.263 | 69.156 | 0.166 |
| 1987 | Pacific halibut | 42.956 | 28.298 | 58.458 | 12.096 | 0.383 | 146.313 | 0.282 |


| Year | Species | Mean | $\begin{aligned} & 10 \% \\ & \text { c.i. } \end{aligned}$ | $\begin{aligned} & \text { 90\%. } \\ & \text { c.i } \end{aligned}$ | Standard deviation | Standard error | Variance | Coefficient of variation |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1989 | Pacific halibut | 34.399 | 26.785 | 42.251 | 6.436 | 0.204 | 41.422 | 0.187 |
| 1991 | Pacific halibut | 54.143 | 35.303 | 74.631 | 15.617 | 0.494 | 243.891 | 0.288 |
| 1993 | Pacific halibut | 72.669 | 56.093 | 91.325 | 14.18 | 0.448 | 201.072 | 0.195 |
| 1995 | Pacific halibut | 32.947 | 27.006 | 39.54 | 4.976 | 0.157 | 24.761 | 0.151 |
| 1996 | Pacific halibut | 38.939 | 28.578 | 49.649 | 8.365 | 0.265 | 69.973 | 0.215 |
| 1998 | Pacific halibut | 35.376 | 20.004 | 53.191 | 13.583 | 0.43 | 184.498 | 0.384 |
| 2000 | Pacific halibut | 32.379 | 22.876 | 42.906 | 7.638 | 0.242 | 58.339 | 0.236 |
| 2002 | Pacific halibut | 17.347 | 11.684 | 23.751 | 4.871 | 0.154 | 23.727 | 0.281 |
| 2003 | Pacific halibut | 26.261 | 18.859 | 34.206 | 6.14 | 0.194 | 37.7 | 0.234 |
| 1984 | Rock sole | 19.457 | 14.607 | 24.808 | 4.163 | 0.132 | 17.331 | 0.214 |
| 1987 | Rock sole | 35.692 | 23.296 | 49.629 | 10.255 | 0.324 | 105.165 | 0.287 |
| 1989 | Rock sole | 93.621 | 64.265 | 126.263 | 24.954 | 0.789 | 622.702 | 0.267 |
| 1991 | Rock sole | 51.181 | 39.299 | 64.021 | 9.549 | 0.302 | 91.183 | 0.187 |
| 1993 | Rock sole | 90.338 | 62.386 | 121.321 | 23.164 | 0.733 | 536.571 | 0.256 |
| 1995 | Rock sole | 35.841 | 26.91 | 45.387 | 7.297 | 0.231 | 53.246 | 0.204 |
| 1996 | Rock sole | 89.711 | 67.842 | 112.605 | 17.412 | 0.551 | 303.178 | 0.194 |
| 1998 | Rock sole | 44.582 | 27.764 | 63.908 | 14.424 | 0.456 | 208.052 | 0.324 |
| 2000 | Rock sole | 81.414 | 54.535 | 110.081 | 22.578 | 0.714 | 509.766 | 0.277 |
| 2002 | Rock sole | 66.331 | 50.767 | 82.689 | 12.614 | 0.399 | 159.113 | 0.19 |
| 2003 | Rock sole | 106.924 | 72.821 | 146.066 | 29.813 | 0.943 | 888.815 | 0.279 |
| 1984 | Sand sole | 5.325 | 3.288 | 7.536 | 1.787 | 0.057 | 3.193 | 0.336 |
| 1987 | Sand sole | 6.402 | 3.018 | 10.582 | 3.202 | 0.101 | 10.253 | 0.5 |
| 1989 | Sand sole | 9.667 | 5.982 | 13.991 | 3.413 | 0.108 | 11.649 | 0.353 |
| 1991 | Sand sole | 5.983 | 3.8 | 8.47 | 1.859 | 0.059 | 3.456 | 0.311 |
| 1993 | Sand sole | 10.695 | 6.356 | 16.239 | 4.148 | 0.131 | 17.206 | 0.388 |
| 1995 | Sand sole | 6.759 | 5.239 | 8.481 | 1.319 | 0.042 | 1.74 | 0.195 |
| 1996 | Sand sole | 3.609 | 2.803 | 4.48 | 0.657 | 0.021 | 0.432 | 0.182 |
| 1998 | Sand sole | 3.487 | 2.408 | 4.733 | 0.929 | 0.029 | 0.863 | 0.266 |
| 2000 | Sand sole | 5.068 | 3.417 | 7.131 | 1.441 | 0.046 | 2.076 | 0.284 |
| 2002 | Sand sole | 9.623 | 6.266 | 13.456 | 2.841 | 0.09 | 8.071 | 0.295 |
| 2003 | Sand sole | 20.232 | 14.024 | 26.934 | 5.056 | 0.16 | 25.563 | 0.25 |

Table 3. Mean catch rates and $90 \%$ confidence limits for dominant species in the Butterworth species assemblage in Hecate Strait based on surveys conducted from 1984-2003

| Year | Species | Mean | $\begin{aligned} & 10 \% \\ & \text { c.i. } \\ & \hline \end{aligned}$ | $\begin{aligned} & \hline 90 \% \\ & \text { c.i. } \\ & \hline \end{aligned}$ | Standard deviation | Standard error | Variance | Coefficient of variation |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1984 | Spiny dogfish | 108.583 | 83.748 | 134.72 | 19.277 | 0.61 | 371.603 | 0.178 |
| 1987 | Spiny dogfish | 79.534 | 50.252 | 110.635 | 24.518 | 0.775 | 601.132 | 0.308 |
| 1989 | Spiny dogfish | 38.19 | 23.965 | 54.613 | 12.41 | 0.392 | 154.008 | 0.325 |
| 1991 | Spiny dogfish | 31.75 | 23.776 | 41.215 | 6.974 | 0.221 | 48.637 | 0.22 |
| 1993 | Spiny dogfish | 26.286 | 19.32 | 33.933 | 5.86 | 0.185 | 34.34 | 0.223 |
| 1995 | Spiny dogfish | 41.219 | 28.644 | 56.376 | 11.037 | 0.349 | 121.815 | 0.268 |
| 1996 | Spiny dogfish | 20.463 | 15.041 | 26.663 | 4.695 | 0.148 | 22.043 | 0.229 |
| 1998 | Spiny dogfish | 31.958 | 21.886 | 43.974 | 9.217 | 0.291 | 84.953 | 0.288 |
| 2000 | Spiny dogfish | 50.495 | 36.247 | 67.421 | 12.431 | 0.393 | 154.53 | 0.246 |
| 2002 | Spiny dogfish | 29.212 | 19.656 | 40.598 | 8.325 | 0.263 | 69.306 | 0.285 |
| 2003 | Spiny dogfish | 8.32 | 5.264 | 12.054 | 2.879 | 0.091 | 8.289 | 0.346 |
| 1984 | Spotted ratfish | 104.44 | 54.362 | 164.563 | 46.8 | 1.48 | 2190.24 | 0.448 |
| 1987 | Spotted ratfish | 72.556 | 36.038 | 119.974 | 33.905 | 1.072 | 1149.549 | 0.467 |
| 1989 | Spotted ratfish | 61.89 | 39.93 | 85.633 | 17.445 | 0.552 | 304.328 | 0.282 |
| 1991 | Spotted ratfish | 54.074 | 34.652 | 76.366 | 16.154 | 0.511 | 260.952 | 0.299 |
| 1993 | Spotted ratfish | 58.443 | 37.003 | 84.12 | 18.762 | 0.593 | 352.013 | 0.321 |
| 1995 | Spotted ratfish | 9.008 | 6.343 | 12.078 | 2.332 | 0.074 | 5.438 | 0.259 |
| 1996 | Spotted ratfish | 27.937 | 20.619 | 36.39 | 6.205 | 0.196 | 38.502 | 0.222 |
| 1998 | Spotted ratfish | 80.383 | 50.05 | 116.384 | 26.951 | 0.852 | 726.356 | 0.335 |
| 2000 | Spotted ratfish | 61.177 | 40.378 | 83.956 | 17.123 | 0.541 | 293.197 | 0.28 |
| 2002 | Spotted ratfish | 59.436 | 38.366 | 83.005 | 18.276 | 0.578 | 334.012 | 0.307 |
| 2003 | Spotted ratfish | 118.01 | 68.952 | 175.107 | 44.149 | 1.396 | 1949.134 | 0.374 |
| 1984 | Arrowtooth flounder | 483.743 | 354.372 | 637.019 | 107.665 | 3.405 | 11591.75 | 0.223 |
| 1987 | Arrowtooth flounder | 838.718 | 641.651 | 1054.165 | 162.261 | 5.131 | 26328.63 | 0.193 |
| 1989 | Arrowtooth flounder | 992.853 | 785.595 | 1217.335 | 170.4 | 5.389 | 29036.16 | 0.172 |
| 1991 | Arrowtooth flounder | 414.506 | 322.203 | 515.091 | 74.805 | 2.366 | 5595.788 | 0.18 |
| 1993 | Arrowtooth flounder | 72.257 | 43.745 | 104.733 | 24.689 | 0.781 | 609.547 | 0.342 |
| 1995 | Arrowtooth flounder | 91.777 | 55.342 | 135.16 | 31.4 | 0.993 | 985.96 | 0.342 |
| 1996 | Arrowtooth flounder | 127.775 | 76.958 | 191.911 | 46.476 | 1.47 | 2160.019 | 0.364 |


| Year | Species | Mean | $\begin{aligned} & 10 \% \\ & \text { c.i. } \end{aligned}$ | $\begin{aligned} & \hline 90 \% \\ & \text { c.i. } \\ & \hline \end{aligned}$ | Standard deviation | Standard error | Variance | Coefficient of variation |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1998 | Arrowtooth flounder | 236.274 | 175.097 | 306.325 | 52.765 | 1.669 | 2784.145 | 0.223 |
| 2000 | Arrowtooth flounder | 246.896 | 152.499 | 356.616 | 81.57 | 2.579 | 6653.665 | 0.33 |
| 2002 | Arrowtooth flounder | 498.37 | 375.894 | 635.898 | 102.914 | 3.254 | 10591.29 | 0.207 |
| 2003 | Arrowtooth flounder | 644.824 | 506.636 | 794.967 | 115.235 | 3.644 | 13279.1 | 0.179 |
| 1984 | Rex sole | 33.939 | 22.685 | 46.114 | 9.327 | 0.295 | 86.993 | 0.275 |
| 1987 | Rex sole | 57.921 | 36.456 | 82.27 | 18.66 | 0.59 | 348.196 | 0.322 |
| 1989 | Rex sole | 60.429 | 36.258 | 88.275 | 21.693 | 0.686 | 470.586 | 0.359 |
| 1991 | Rex sole | 75.423 | 55.399 | 97.055 | 16.615 | 0.525 | 276.058 | 0.22 |
| 1993 | Rex sole | 35.103 | 23.032 | 48.87 | 10.884 | 0.344 | 118.461 | 0.31 |
| 1995 | Rex sole | 64.867 | 41.611 | 89.987 | 19.174 | 0.606 | 367.642 | 0.296 |
| 1996 | Rex sole | 83.623 | 63.186 | 105.503 | 16.42 | 0.519 | 269.616 | 0.196 |
| 1998 | Rex sole | 73.757 | 52.811 | 100.037 | 18.577 | 0.587 | 345.105 | 0.252 |
| 2000 | Rex sole | 125.118 | 94.941 | 158.97 | 25.424 | 0.804 | 646.38 | 0.203 |
| 2002 | Rex sole | 199.569 | 153.07 | 248.79 | 40.211 | 1.272 | 1616.925 | 0.201 |
| 2003 | Rex sole | 57.189 | 41.42 | 74.297 | 13.767 | 0.435 | 189.53 | 0.241 |
| 1984 | Dover sole | 66.727 | 36.984 | 104.041 | 26.647 | 0.843 | 710.063 | 0.399 |
| 1987 | Dover sole | 74.423 | 31.567 | 126.083 | 40.352 | 1.276 | 1628.284 | 0.542 |
| 1989 | Dover sole | 124.268 | 69.223 | 187.066 | 47.435 | 1.5 | 2250.079 | 0.382 |
| 1991 | Dover sole | 49.742 | 34.064 | 67.701 | 13.501 | 0.427 | 182.277 | 0.271 |
| 1993 | Dover sole | 29.173 | 18.276 | 41.526 | 9.216 | 0.291 | 84.935 | 0.316 |
| 1995 | Dover sole | 22.464 | 15.064 | 30.808 | 6.502 | 0.206 | 42.276 | 0.289 |
| 1996 | Dover sole | 32.403 | 21.967 | 44.335 | 9.006 | 0.285 | 81.108 | 0.278 |
| 1998 | Dover sole | 32.726 | 19.735 | 48.594 | 11.952 | 0.378 | 142.85 | 0.365 |
| 2000 | Dover sole | 91.807 | 61.478 | 127.057 | 26.508 | 0.838 | 702.674 | 0.289 |
| 2002 | Dover sole | 118.378 | 74.767 | 164.425 | 36.625 | 1.158 | 1341.391 | 0.309 |
| 2003 | Dover sole | 84.097 | 51.085 | 123.992 | 29.323 | 0.927 | 859.838 | 0.349 |
| 1984 | English sole | 46.473 | 31.442 | 62.948 | 12.384 | 0.392 | 153.363 | 0.266 |
| 1987 | English sole | 40.598 | 18.934 | 65.645 | 19.908 | 0.63 | 396.328 | 0.49 |
| 1989 | English sole | 56.992 | 33.651 | 82.194 | 19.451 | 0.615 | 378.341 | 0.341 |
| 1991 | English sole | 87.328 | 67.279 | 109.164 | 16.882 | 0.534 | 285.002 | 0.193 |
| 1993 | English sole | 168.672 | 115.968 | 227.082 | 45.707 | 1.445 | 2089.13 | 0.271 |


| Year | Species | Mean | $10 \%$ <br> c.i. | $90 \%$ <br> c.i. | Standard <br> deviation | Standard <br> error | Variance | Coefficient <br> of variation |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 1995 | English sole | 40.024 | 26.184 | 56.132 | 11.986 | 0.379 | 143.664 | 0.299 |
| 1996 | English sole | 41.036 | 27.311 | 55.948 | 11.244 | 0.356 | 126.428 | 0.274 |
| 1998 | English sole | 52.044 | 31.418 | 75.04 | 17.613 | 0.557 | 310.218 | 0.338 |
| 2000 | English sole | 66.102 | 43.446 | 91.275 | 19.44 | 0.615 | 377.914 | 0.294 |
| 2002 | English sole | 63.514 | 38.593 | 91.008 | 21.835 | 0.69 | 476.767 | 0.344 |
| 2003 | English sole | 66.924 | 34.521 | 106.343 | 30.845 | 0.975 | 951.414 | 0.461 |

Table 4. Mean catch rates and $90 \%$ confidence limits for dominant species in the Bonilla species assemblage in Hecate Strait based on surveys conducted from 1984-2003.

|  | Year | Species | Mean | $10 \%$ <br> c.i. | $90 \%$ <br> c.i. | Standard <br> deviation | Standard <br> error | Variance |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 1984 | Spiny dogfish | 64.35 | 51.516 | 78.749 | 10.511 | 0.332 | 110.481 | Coefficient <br> of variation |
| 1987 | Spiny dogfish | 94.235 | 66.924 | 128.347 | 24.093 | 0.762 | 580.473 | 0.256 |
| 1989 | Spiny dogfish | 38.268 | 26.294 | 51.994 | 10.707 | 0.339 | 114.64 | 0.28 |
| 1991 | Spiny dogfish | 8.356 | 5.412 | 11.722 | 2.475 | 0.078 | 6.126 | 0.296 |
| 1993 | Spiny dogfish | 207.619 | 128.361 | 302.685 | 70.967 | 2.244 | 5036.315 | 0.342 |
| 1995 | Spiny dogfish | 48.536 | 25.273 | 77.658 | 22.302 | 0.705 | 497.379 | 0.459 |
| 1996 | Spiny dogfish | 22.85 | 14.341 | 32.357 | 7.549 | 0.239 | 56.987 | 0.33 |
| 1998 | Spiny dogfish | 87.019 | 41.852 | 140.478 | 43.554 | 1.377 | 1896.951 | 0.501 |
| 2000 | Spiny dogfish | 25.299 | 19.396 | 31.579 | 4.735 | 0.15 | 22.42 | 0.187 |
| 2002 | Spiny dogfish | 12.081 | 9.099 | 15.421 | 2.46 | 0.078 | 6.052 | 0.204 |
| 2003 | Spiny dogfish | 17.707 | 10.571 | 26.147 | 6.011 | 0.19 | 36.132 | 0.339 |
| 1984 | Spotted ratfish | 11.268 | 8.293 | 14.648 | 2.605 | 0.082 | 6.786 | 0.231 |
| 1987 | Spotted ratfish | 5.489 | 3.675 | 7.446 | 1.497 | 0.047 | 2.241 | 0.273 |
| 1989 | Spotted ratfish | 34.63 | 23.526 | 46.692 | 9.369 | 0.296 | 87.778 | 0.271 |
| 1991 | Spotted ratfish | 16.204 | 10.032 | 23.246 | 5.197 | 0.164 | 27.009 | 0.321 |
| 1993 | Spotted ratfish | 7.588 | 4.167 | 11.561 | 3.145 | 0.099 | 9.891 | 0.414 |
| 1995 | Spotted ratfish | 28.453 | 12.448 | 48.666 | 15.644 | 0.495 | 244.735 | 0.55 |
| 1996 | Spotted ratfish | 15.946 | 9.754 | 22.915 | 5.186 | 0.164 | 26.895 | 0.325 |
| 1998 | Spotted ratfish | 13.125 | 5.557 | 22.293 | 8.736 | 0.276 | 76.318 | 0.666 |
| 2000 | Spotted ratfish | 87.358 | 48.41 | 132.421 | 36.114 | 1.142 | 1304.221 | 0.413 |
| 2002 | Spotted ratfish | 181.814 | 114.752 | 257.908 | 58.079 | 1.837 | 3373.17 | 0.319 |
| 2003 | Spotted ratfish | 37.751 | 25.544 | 52.55 | 11.126 | 0.352 | 123.788 | 0.295 |
| 1984 | Pacific sanddab | 25.985 | 17.675 | 36.156 | 7.353 | 0.233 | 54.067 | 0.283 |
| 1987 | Pacific sanddab | 11 | 6.036 | 17.632 | 4.635 | 0.147 | 21.483 | 0.421 |
| 1989 | Pacific sanddab | 15.693 | 7.617 | 25.591 | 7.47 | 0.236 | 55.801 | 0.476 |
| 1991 | Pacific sanddab | 16.45 | 8.442 | 26.064 | 7.308 | 0.231 | 53.407 | 0.444 |
| 1993 | Pacific sanddab | 2.077 | 1.509 | 2.667 | 0.536 | 0.017 | 0.287 | 0.258 |
| 1995 | Pacific sanddab | 30.793 | 11.857 | 56.066 | 18.631 | 0.589 | 347.114 | 0.605 |
| 1996 | Pacific sanddab | 54.079 | 27.711 | 87.775 | 24.915 | 0.788 | 620.757 | 0.461 |


| Year | Species | Mean | $\begin{aligned} & 10 \% \\ & \text { c.i. } \end{aligned}$ | $\begin{aligned} & 90 \% \\ & \text { c.i. } \end{aligned}$ | Standard deviation | Standard error | Variance | Coefficient of variation |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1998 | Pacific sanddab | 31.113 | 14.589 | 53.859 | 16.935 | 0.536 | 286.794 | 0.544 |
| 2000 | Pacific sanddab | 51.834 | 29.449 | 78.358 | 20.876 | 0.66 | 435.807 | 0.403 |
| 2002 | Pacific sanddab | 30.051 | 13.147 | 52.165 | 16.703 | 0.528 | 278.99 | 0.556 |
| 2003 | Pacific sanddab | 64.711 | 30.182 | 107.988 | 32.532 | 1.029 | 1058.331 | 0.503 |
| 1984 | Arrowtooth flounder | 6.789 | 4.682 | 9.444 | 1.891 | 0.06 | 3.576 | 0.279 |
| 1987 | Arrowtooth flounder | 4.233 | 2.629 | 6.211 | 1.55 | 0.049 | 2.403 | 0.366 |
| 1989 | Arrowtooth flounder | 31.885 | 13.221 | 57.637 | 19.888 | 0.629 | 395.533 | 0.624 |
| 1991 | Arrowtooth flounder | 13.853 | 6.386 | 22.815 | 7.067 | 0.223 | 49.942 | 0.51 |
| 1993 | Arrowtooth flounder | 19.593 | 6.378 | 38.198 | 16.146 | 0.511 | 260.693 | 0.824 |
| 1995 | Arrowtooth flounder | 69.049 | 36.594 | 110.652 | 31.051 | 0.982 | 964.165 | 0.45 |
| 1996 | Arrowtooth flounder | 23.717 | 11.357 | 39.336 | 11.777 | 0.372 | 138.698 | 0.497 |
| 1998 | Arrowtooth flounder | 12.881 | 4.549 | 24.625 | 9.516 | 0.301 | 90.554 | 0.739 |
| 2000 | Arrowtooth flounder | 9.158 | 4.967 | 14.651 | 3.93 | 0.124 | 15.445 | 0.429 |
| 2002 | Arrowtooth flounder | 34.602 | 16.909 | 56.067 | 15.923 | 0.504 | 253.542 | 0.46 |
| 2003 | Arrowtooth flounder | 37.597 | 23.058 | 54.593 | 12.75 | 0.403 | 162.562 | 0.339 |
| 1984 | Rex sole | 23.35 | 15.891 | 31.608 | 6.416 | 0.203 | 41.165 | 0.275 |
| 1987 | Rex sole | 5.054 | 3.029 | 7.539 | 1.999 | 0.063 | 3.996 | 0.396 |
| 1989 | Rex sole | 36.998 | 21.76 | 55.211 | 13.271 | 0.42 | 176.119 | 0.359 |
| 1991 | Rex sole | 12.124 | 5.136 | 21.251 | 7.184 | 0.227 | 51.61 | 0.593 |
| 1993 | Rex sole | 7.729 | 3.651 | 12.504 | 4.131 | 0.131 | 17.065 | 0.534 |
| 1995 | Rex sole | 22.279 | 14.513 | 30.762 | 6.584 | 0.208 | 43.349 | 0.296 |
| 1996 | Rex sole | 22.758 | 13.644 | 33.148 | 7.82 | 0.247 | 61.152 | 0.344 |
| 1998 | Rex sole | 8.502 | 3.816 | 14.745 | 5.125 | 0.162 | 26.266 | 0.603 |
| 2000 | Rex sole | 8.606 | 5.236 | 12.637 | 3.066 | 0.097 | 9.4 | 0.356 |
| 2002 | Rex sole | 29.056 | 16.468 | 42.912 | 11.198 | 0.354 | 125.395 | 0.385 |
| 2003 | Rex sole | 74.166 | 42.63 | 114.013 | 29.331 | 0.928 | 860.308 | 0.395 |
| 1984 | English sole | 83.469 | 61.772 | 106.853 | 18.461 | 0.584 | 340.809 | 0.221 |
| 1987 | English sole | 36.934 | 22.5 | 54.915 | 13.198 | 0.417 | 174.187 | 0.357 |
| 1989 | English sole | 327.488 | 210.316 | 465.1 | 99.65 | 3.151 | 9930.123 | 0.304 |
| 1991 | English sole | 203.158 | 153.544 | 258.348 | 42.005 | 1.328 | 1764.42 | 0.207 |
| 1993 | English sole | 16.478 | 10.344 | 24.115 | 5.612 | 0.177 | 31.495 | 0.341 |


| Year | Species | Mean | $10 \%$ <br> c.i. | $90 \%$ <br> c.i. | Standard <br> deviation | Standard <br> error | Variance | Coefficient <br> of variation |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 1995 | English sole | 79.494 | 44.777 | 119.753 | 30.139 | 0.953 | 908.359 | 0.379 |
| 1996 | English sole | 187.276 | 130.309 | 248.178 | 46.449 | 1.469 | 2157.51 | 0.248 |
| 1998 | English sole | 15.163 | 6.747 | 26.62 | 8.371 | 0.265 | 70.074 | 0.552 |
| 2000 | English sole | 65.779 | 38.354 | 99.754 | 24.652 | 0.78 | 607.721 | 0.375 |
| 2002 | English sole | 113.531 | 70.684 | 158.588 | 36.452 | 1.153 | 1328.748 | 0.321 |
| 2003 | English sole | 252.592 | 164.25 | 353.269 | 75.266 | 2.38 | 5664.971 | 0.298 |

Table 5. Species diversity, richness and evenness for assemblages in Hecate Strait.


Table 6. Species richness in Hecate Strait and the B.C. coast compared to large scale marine ecosystems in the temperate zone ${ }^{1}$.

| Marine ecosystem | Richness | Area <br> $\left(\mathrm{km}^{2}\right)$ | Latitude | Longitude | Region |
| :--- | :--- | :--- | :--- | :--- | :--- |
| Oyashio Current | 37 | 532831 | 45.2 | 151.6 | North Pacific |
| Barents sea | 59 | 1865429 | 76.0 | 37.3 | North Atlantic |
| Hecate Strait | 150 | 25000 | 53.0 | 131.0 | North Pacific |
| Iceland shelf/sea | 152 | 51820 | 66.6 | -17.6 | North Atlantic |
| West Greenland shelf | 157 | 365548 | 70.2 | -57.1 | North Atlantic |
| Baltic sea | 157 | 394265 | 58.9 | 19.7 | North Atlantic |
| Greenland sea | 162 | 1171612 | 75.5 | -11.8 | North Atlantic |
| Newfoundland Labrador shelf | 171 | 681296 | 49.5 | -53.0 | North Atlantic |
| Faroe Plateau | 174 | 150558 | 60.6 | -11.2 | North Atlantic |
| North sea | 190 | 695626 | 57.4 | 2.8 | North Atlantic |
| Black sea | 193 | 461958 | 43.5 | 34.4 | North Atlantic |
| Scotian shelf | 197 | 414534 | 47.0 | -61.7 | North Atlantic |
| Norwegian sea | 232 | 1102919 | 68.0 | 3.7 | North Atlantic |
| East Bering sea | 249 | 1186827 | 58.9 | -168.8 | North Pacific |
| West Bering sea | 311 | 2170639 | 57.5 | -175.5 | North Pacific |
| Gulf of Alaska | 317 | 1474706 | 54.1 | -139.3 | North Pacific |
| B.C. shelf/slope | 325 | 110000 | 49.0 | 125.0 | North Pacific |
| Okhotsk sea | 391 | 1557816 | 53.8 | 148.9 | North Pacific |

${ }^{1}$ Information for large scale ecosystems from Fisher et al. 2010.


Figure 1. The study area. Trawling sites are depicted in yellow and represent 1020 hauls made between 1984 and 2003. The numbered white boxes correspond to landmarks: 1) Butterworth rocks, 2) Bonilla Island and 3) Reef Island.


Figure 2. Dendrogram of species assemblages from the cluster analysis of the 1984 Hecate Strait survey catch rate data.


Figure 3. Dendrogram of species assemblages from the cluster analysis of the 1987 Hecate Strait survey catch rate data.


Figure 4. Dendrogram of species assemblages from the cluster analysis of the 1989 Hecate Strait survey catch rate data.


Figure 5. Dendrogram of species assemblages from the cluster analysis of the 1991 Hecate Strait survey catch rate data.


Figure 6. Dendrogram of species assemblages from the cluster analysis of the 1993 Hecate Strait survey catch rate data.


## Haul number

Figure 7. Dendrogram of species assemblages from the cluster analysis of the 1995 Hecate Strait survey catch rate data.


Figure 8. Dendrogram of species assemblages from the cluster analysis of the 1996 Hecate Strait survey catch rate data


Figure 9. Dendrogram of species assemblages from the cluster analysis of the 1998 Hecate Strait survey catch rate data.


Figure 10. Dendrogram of species assemblages from the cluster analysis of the 2000 Hecate Strait survey catch rate data.


Figure 11. Dendrogram of species assemblages from the cluster analysis of the 2002 Hecate Strait survey catch rate data.


Figure 12. Dendrogram of species assemblages from the cluster analysis of the 2003 Hecate Strait survey catch rate data.


Figure 13. Boundaries for fish assemblages in Hecate Strait fish: A=Reef Island , $\mathrm{B}=\mathrm{Bu} t \mathrm{terworth} \mathrm{and} \mathrm{C}=$ Bonilla. The overlap of colours indicates a gradual transition rather than a sharp transition.


Figure 14. Haul locations (1984-2003) as classified components of assemblages in the cluster analysis.






Figure 15. Species proportion by weight and year for the Reef Island assemblage. The rank 1 group is the highest weight proportion, rank 2 group is the second highest weight proportion .....


Figure 16. Species weight proportions for the Reef Island assemblage combined years (1984-2003)


Figure 17. Species proportion by weight and year for the Butterworth assemblage. The rank 1 group is the highest weight proportion, rank 2 group is the second highest weight proportion ....


Figure 18. Species weight proportions for the Butterworth assemblage combined years (1984-2003)


Figure 19. Species proportion by weight and year for the Bonilla assemblage. The rank 1 group is the highest weight proportion, rank 2 group is the second highest weight proportion .....


Figure 20. Species weight proportions for the Bonilla assemblage combined years (1984-2003)


Figure 21. Proportion of biomass by assemblage for dominant species components.


Figure 22. Mean CPUE (backtransformed) and 90\% confidence interval for dominant species in the Reef Island Assemblage, 1984-2003.


Figure 23. Boxplots of $\operatorname{In}(C P U E)$ for dominant species in the Reef Island assemblage, 1984-2003. Points in red represent the mean value.


Figure 24. Boxplots of $\ln (C P U E)$ for species in the Reef Island assemblage, 1984-2003.


Figure 25. Mean CPUE (backtransformed) and 90\% confidence interval for dominant species in the Butterworth Assemblage, 1984-2003.


Figure 26. Boxplots of $\ln (C P U E)$ for dominant species in the Butterworth assemblage, 1984-2003. Points in red represent the mean value.


Figure 27. Boxplots of $\ln$ (CPUE) for all species in the Butterworth assemblage, 1984-2003.


Figure 28. Mean CPUE (backtransformed) and 90\% confidence interval for dominant species in the Bonilla Assemblage, 1984-2003.


Figure 29. Boxplots of $\ln (C P U E)$ for dominant species in the Bonilla assemblage, 1984-2003. Points in red represent the mean value.


Figure 30. Boxplots of $\operatorname{In}(C P U E)$ for all species in the Bonilla assemblage, 19842003.

Depth (m)


Figure 31. Boxplots of depth for species in all assemblages, 1984-2003. Red dots indicate the mean value. Black dots indicate pelagic or semi-pelagic species.


Figure 32. Boxplots of depth by year for the three assemblages.


Figure 33. Boxplots of depth by assemblage.


Figure 34. Boxplots of depth by dominant species for the combined Hecate Strait assemblage surveys. The colours in the box are proportional to the relative abundance of each species in each assemblage.


Figure 35. The relationship between mean depth (top panel) and median depth (bottom panel) and the number of species encountered for surveys combined.


Figure 36. Diversity index (1984-2003) by assemblage for Hecate Strait determined from Assemblage Survey catch rate data.


Figure 37. Richness index (1984-2003) by assemblage for Hecate Strait determined from Assemblage Survey data.


Figure 38. Evenness index (1984-2003) by assemblage for Hecate Strait determined from Assemblage Survey data.

