# Estimating the magnitude of pre-European Maori marine harvest in two New Zealand study areas 

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## EXECUTIVE SUMMARY

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This report forms part of the Taking Stock project which is seeking to understand the long-term effects of climate variations and human impacts on the structure and functioning of New Zealand's marine shelf ecosystems. It uses archaeological data to estimate the numbers and biomass of marine animals used as food by people in two New Zealand study areas during the period before European settlement. The study areas are Greater Hauraki, on the northeast coast of the North Island, and Otago-Catlins, on the southeast coast of the South Island. For each area estimates are made for three points in time; ca. $1400 \mathrm{AD}, c a .1550 \mathrm{AD}$, and $c a .1750 \mathrm{AD}$.

The method used to generate the estimates is novel, and involves three steps. First, archaeological data were analysed to determine the range of marine animal species harvested by Maori at each focal date in each study area, and the proportional contribution of each species to the total energy harvest from animals. Second, the scale of the total energy harvest from animals was approximated by estimating the size of human population, the energy required to sustain them, and the proportion of that derived from animal foods. Finally, the number of each marine animal species required to contribute the proportions derived in the first step to the total energy harvest from animals estimated in the second step was calculated using an estimate of the calorific value of the meat derived from an individual of the species concerned. Three estimates were made for each species; a minimum estimate which used the lowest probable values for each variable in the second step of analysis, a maximum estimate using the highest probable values, and a best estimate based on the most realistic or wellsupported values. Cumulative calculation errors were applied to each set of estimates.

Estimates are presented for 101 taxa from the Greater Hauraki study area ( 46 shellfish, 28 fish, 22 birds and 5 mammals) and 96 from the Otago-Catlins study area ( 36 shellfish, 25 fish, 28 birds and 7 mammals). In a small number of cases estimates are available for only one or two of the focal dates, either because of changes in the availability of the species concerned or the pattern of harvesting by Maori.

For the majority of species the minimum, best and maximum estimates all indicate increased harvesting over time. For example, at $c a .1400$ AD in Greater Hauraki the best estimate of harvested biomass of snapper (Pagrus auratus) is $72.1 \pm 21.6$ tonnes (minimum 12.1 $\pm 3.6 \mathrm{t}$; maximum $107.5 \pm$ 32.3 t ), which rose to $938.8 \pm 281.7 \mathrm{t}(\mathrm{min} .354 .8 \pm 106.4 \mathrm{t}$; max. $1354.7 \pm 406.4 \mathrm{t}$ ) at $c a .1550 \mathrm{AD}$, and then $997.2 \pm 299.2(\min .560 .1 \pm 168.0 \mathrm{t}$; max. $1393.9 \pm 418.2 \mathrm{t})$ at $c a .1750 \mathrm{AD}$. In contrast, marine mammals and some marine and coastal bird exhibit declining trends under all three sets of estimates. For example, the fur seal (Arctocephalus forsteri) which yielded a best estimate of $284.9 \pm$ $85.5 \mathrm{t}(\mathrm{min} .47 .9 \pm 14.4 \mathrm{t}$; max. $425.1 \pm 127.5 \mathrm{t})$ at $c a .1400$ AD in Greater Hauraki was no longer harvested at the two later focal dates. In Otago-Catlins harvests of the same species fall successively from best estimates of $237.0 \pm 71.1 \mathrm{t}(\mathrm{min} .55 .2 \pm 16.6 \mathrm{t} ; \max .383 .5 \pm 115.1 \mathrm{t})$ to $103.3 \pm 31.0 \mathrm{t}(\mathrm{min}$. $24.0 \pm 7.2 ; \max .173 .8 \pm 52.1 \mathrm{t})$ then $77.0 \pm 23.1 \mathrm{t}(\min .57 .4 \pm 17.2 ; \max .99 .7 \pm 29.9 \mathrm{t})$ in $c a .1400$, 1550 and 1750 AD , respectively.

In the Greater Hauraki region growth of the human population was one of two main reasons for the increased harvests of most marine animals. In addition, the demand placed on most fish, shellfish and some bird species increased over time because seals, moas and some of the marine and coastal birds that made important contributions to earlier Maori diets ceased to be available. This process of replacement of one food source by another was the single most important driver of change in OtagoCatlins, where the human population remained more or less stable throughout the study period.

As well as contributing to understanding of long-term changes in the structure and functioning of New Zealand's marine shelf ecosystems, the estimates provide baseline data against which modern customary and commercial harvests of marine animals can be measured.

## 1. INTRODUCTION

### 1.1 Overview

New Zealand was the last major land mass to be settled by humans, their arrival dated to around 1280 AD (Wilmshurst et al. 2011). Consequently, New Zealand has a short and reasonably complete archaeological, historical and contemporary record of human exploitation of marine resources compared to most other places where the earliest evidence of human impacts on marine ecosystems is difficult to discern because of climate fluctuations and changes in sea level (MacDiarmid 2011). The collaborative multi-disciplinary Taking Stock project, funded by the Ministry of Fisheries, has the overall objective of determining the effects of climate variation and human impact on the structure and functioning of New Zealand shelf ecosystems over the timescale of human occupation. To achieve this it set out to build a mass balance model of current coastal and marine shelf ecosystems in each of two study areas, and then to estimate how each of these operated at five earlier intervals: ca. 1950 AD (before modern industrial fishing), $c a .1750 \mathrm{AD}$ (before European whaling and sealing), $c a$. 1550 AD (about the middle of the Maori period of occupation), ca. 1400 AD (soon after Maori arrival in New Zealand) and ca. 1000 AD (before human settlement). For each of the earlier intervals reconstruction depends in part on estimation of the taxonomic composition and biomass of removals from the marine environment through human activities. This report draws upon archaeological data for the exploitation of marine resources by Māori in the Hauraki Gulf and along the Otago-Catlins coast to estimate what they harvested from the sea at three intervals between earliest settlement and European arrival at the end of the $18^{\text {th }}$ century $\mathrm{AD}(c a .1400,1550,1750 \mathrm{AD})$.

### 1.2 Previous Research

There is a long history of using archaeological data to infer changes in past ecosystems. Numerous instances of animal extinctions and distributional changes have been detected, dated and examined through archaeological research (e.g. Grayson 2001, Rick \& Erlandson 2008). In New Zealand, as elsewhere in the world, most of these examples concern terrestrial fauna, and in many cases predation by people and their commensals or human-induced habitat modification have been implicated (Anderson 1989, Anderson 1997, Holdaway 1999, McGlone 1989).

For the marine environment, evidence is much more equivocal. New Zealand seals provide one wellexplored case of pre-industrial human impacts (Smith 1989, Smith 2005), and internationally there are others for terrestrial-breeding marine mammals (Bryden et al. 1999, Burton et al. 2001). Equally, there are cases of apparently stable, long-term exploitative relationships (Etnier 2007), and for marine-breeding animals there is little undisputed evidence of dramatic human impact before the emergence of commercial whaling in the $18^{\text {th }}$ and $19^{\text {th }}$ centuries (Reeves \& Smith 2006) and more recent industrial-scale fisheries (Myers \& Worm 2003, Pauly et al. 1998). Indeed, the New Zealand data for pre-European shell and fin fisheries show that it is difficult to separate potential effects of human predation and climate without intensive and closely targeted archaeological research (Leach 2006).

Direct estimation of animal population biomass from archaeological data is not generally possible. The archaeological window into past ecological systems is blurred by transformative processes that influence the creation of the archaeological record. These include harvesting and carcass processing, which are largely determined by cultural patterns, along with natural taphonomic processes of decay, and variations in the accuracy with which different items are amenable to archaeozoological analysis (Reitz
\& Wing 2008). Furthermore archaeologists are primarily concerned with determining long-term patterns and regularities in human behaviour, rather than reconstructing past ecosystems, and their data acquisition and analytical methods are designed accordingly. In order to be useful in palaeoecological reconstruction, archaeozoological data must be interpreted with due regard to the cultural and natural formation processes that shaped them, and the archaeological filters through which they have passed.

In the case of pre-European New Zealand, one of the key cultural factors that must be accommodated is the mobile nature of human settlement, whereby members of a community are hypothesised (Anderson \& Smith 1996, Walter et al. 2006) to have made regular intra-annual shifts of residence to facilitate the exploitation of dispersed, seasonally available resources, and communities made occasional territorial shifts over time. Thus no single site can be considered to provide a complete picture of the pattern of marine resource exploitation by a community, and some sites may represent multiple phases of exploitation with differing return intervals. In these circumstances it is essential to aggregate data at a regional level from a judiciously selected range of sites.

With these cautions in mind, some inferences about palaeoecology are possible. The presence of physical remains of an animal species in a regional set of archaeological sites can generally be used to infer that this species occurred in the catchment area of those sites at the time of their occupation, and thus provides a basis for reconstructing the distribution of that species in the past. Similarly, where age or sex can be determined from physical remains, the presence of animals of specific age or sex classes allows some inferences to be made about the age composition and breeding status of exploited populations. Potential confounding factors include long-distance transportation of preserved food remains, industrial usage of bones, teeth or shells from distant sources or older archaeological deposits, and disturbance of archaeological deposits introducing taxa from earlier or later time periods. Where recovery and analytical procedures are adequate, problems of this kind can usually be identified and ameliorated. In contrast, the absence of a species, age or sex class in the archaeological record is not so clear cut. Cultural factors such as dietary preference and harvesting technology, or analytical factors such as sample size, may have intervened. These must be accounted for before archaeological absences can be used to infer lacunae in past animal distributions.

Caution is required in making inferences about the abundance of various species in the past from archaeological data. The relative abundance of taxa in archaeozoological assemblages is primarily a record of the frequency with which they were harvested, modified over time by taphonomic decay. Nonetheless it is reasonable to infer that species which are regularly represented in high frequencies in a regional sample of sites were relatively commonly available. Furthermore, where there is a significant decline in the relative abundance of a species over time, without any evidence for changes in harvesting technology, a decline in their availability can be inferred. Relative abundances of archaeofauna can also be used to derive quantitative assessments of the relative importance of various animals in the diet of the people exploiting them by converting them into the weights of meat that they represent, from which can be derived estimates of the nutritional and energy yields derived from each exploited species (Smith 2004, Smith 2011).

Two broad approaches were taken to generate information useful to the Taking Stock project. First, an overview of marine resource utilisation was constructed for each study area based upon the presence/absence of marine taxa in archaeozoological assemblages and, where suitable data was available, their relative abundance. The second approach involved estimating the magnitude of marine biomass removals through human exploitation in each study area. This was undertaken using estimates of the size of human populations, their energy requirements and the relative contributions of marine foods to their diet.

### 1.3 Objectives

The purpose of this report is to assess and collate existing archaeological data on human removals from the marine environment in two New Zealand study areas in order to address the archaeological aspects of Objective 2 of the Taking Stock Project ZBD200505, which was "To assess and collate existing archaeological, historical and contemporary data (including catch records and stock assessments) on relevant components of the marine ecosystem to provide a detailed description of change in the shelf marine ecosystem in two areas of contrasting human occupation over last 1000 years". The specific objective for the archaeological component of this project was to provide detailed estimates of the magnitude of human removals from the marine shelf ecosystems of two study areas at three points in times between first human arrival in New Zealand and European settlement.

## 2. METHODS

### 2.1 Study Areas and Selection of Study sites

Two study areas were utilized for the Taking Stock project, Greater Hauraki on the east coast of the North Island and Otago-Catlins on the east coast of the South Island, in each case extending from mean high water out to a depth of 250 m (Figure 1). To derive archaeological information relevant to human exploitation of these areas data was drawn from archaeological sites on the adjacent coasts. More than 10000 sites presumed to derive from the pre-European period have been recorded adjacent to the Greater Hauraki coast and some 800 in Otago-Catlins (CINZAS 2008), the difference reflecting the marked concentration of pre-European Maori population in the northern third of the country.

Only a small proportion of recorded sites in each area have been investigated by archaeological excavation, and information from these was assessed through a review of published literature, theses and dissertations in archaeology from University of Auckland and University of Otago, and excavation reports lodged in the NZ Historic Places Trust's Archaeological Report Digital Library. A sample for each study area was selected for detailed analysis on the basis of two criteria: the availability of data on faunal remains suitable for the methodology described below; and the availability of reliable chronological information enabling the sites, or specific assemblages from them to be placed securely in time.

The requirements for data on faunal remains were (a) that taxonomic identifications had been reported for all animal remains in the excavated sample, and (b) the number of animals assigned to each taxon was reported. In this context identifications include determinations to species level along with assignments to genus, family or higher-level designations as necessitated by the nature of archaeozoological material. Identification data were accepted as reported, except where assemblages or components of them had been re-examined (e.g. Leach \& Boocock 1993, Millener 1981, Smith 1985, Worthy 1998) any revisions of identification were incorporated. Where necessary identifications were updated to accommodate revisions of nomenclature, based on the following sources: for shellfish, Spencer et al. (2009); finfish, Froese \& Pauly (2010); birds, Checklist Committee (OSNZ) (2010); and mammals, King (1995) and Baker et al. (2010). Quantification of identified taxa was in terms of the minimum number of individuals (MNI): the smallest number of individual animals necessary to account for all of the remains of a taxon in an archaeological assemblage (Reitz \& Wing 2008). While NISP (number of identified specimens) is sometimes preferred for inter-assemblage comparisons (Lyman 2008), this measure was not reported for the majority of assemblages under consideration here. Initially assemblages were selected for analysis only if MNI were reported for all classes of fauna represented at the site. However this did not provide sufficiently large datasets for some time periods in each study area, and additional
assemblages for which some taxa were noted as present but not quantified were also included. All the taxonomic identifications, MNI values and presence data utilised in this study, along with the data sources that they were drawn from, are reported in detail by Smith \& James-Lee (2010).

With regard to chronology, the primary requirement was for radiocarbon determinations that closely dated formation of the occupation deposit from which the faunal sample under analysis derived. All dates reported in publications were treated according to protocols that are set out in detail elsewhere (Smith 2010). In brief these involved checking data accuracy; culling dates that did not meet sample suitability criteria (Anderson 1991, Petchey 1999, Schmidt 2000a); recalibration using the SH04 calibration curve (McCormac et al. 2004) for terrestrial samples, and for marine samples the Marine 04 calibration curve (Hughen et al. 2004) with delta $R$ set at $-7 \pm 45$, as recommended by the Waikato Radiocarbon Dating Laboratory (Petchey pers. com.); and, where appropriate, combining multiple determinations from the same context into pooled mean ages (Ward \& Wilson 1978). The 126 admissible dates and 29 pooled mean ages employed in this study are reported in Smith \& James-Lee (2010).

Together the selection criteria admitted a total of 107 assemblages from 67 sites for analysis. For the Greater Hauraki area a total of 75 assemblages from 48 sites were included and 32 assemblages from 19 sites in the Otago-Catlins area (Figure 1). In order to provide data relevant to Taking Stock's ca. $\mathrm{AD} 1400, c a$. AD 1550 and $c a$. AD 1750 time slices, and bearing in mind the uncertainties inherent in radiocarbon dating, assemblages were grouped according to three broad period designations: Early (ca. AD 1250-1450), Middle (AD 1450-1650), and Late (AD 1650-1800). Assemblages were allocated to time periods using Smith's (2010) protocol which uses both $1 \sigma$ and $2 \sigma$ calibrated age ranges to distinguish those that can be assigned with confidence to a discrete period from those that overlap the period boundaries. On this basis almost two thirds of the assemblages were assigned to one of the target periods, and the remainder to one of the two overlap zones (Table 1). Although the latter do not represent discrete time spans, they usefully group assemblages that cluster in age around the arbitrary period boundaries, and for assessment of general trends in faunal assemblage composition during the initial steps of the analysis described below (2.3.1, 2.3.2) they are used as if they were discrete periods. However estimation of the magnitude of Maori marine harvest was undertaken only for the three formal periods.

### 2.2 Analytical Method: Overview

Estimates of the magnitude of marine biomass removals by people in the prehistoric past are necessarily speculative, as they depend upon variables that cannot be known with precision. However it is possible to deduce reasonable bounds for these, and thereby constrain the limits of speculation. The approach used here builds upon that employed by Leach (2006: 277-279) to calculate the scale of the Maori snapper fishery in Northland. It has been adapted to incorporate the full range of marine taxa harvested by Maori in each study region. There were three major steps involved in this analysis (Figure 2).

1. The archaeological assemblage data discussed above were scrutinized to establish the range of animal species harvested by Maori in each study area during each study period, the relative frequencies with which this occurred, and proportional contributions of each species to the total energy harvest from animals.
2. The scale of the total energy harvest from animals during each period in each study area was approximated by estimating the size of the human population, the energy required to sustain them, and the proportion of this derived from animal foods.
3. The number of each marine animal species required to contribute the proportions derived in (1) to the total energy harvest from animals estimated in (2) was calculated using an estimate of the calorific value of meat derived from an individual of the species concerned.

Each of these steps is discussed further below.

### 2.3 Analysis of archaeological data

Although the final outputs of this project concern the harvesting of marine animals, preliminary analysis of archaeological data incorporated both marine and terrestrial fauna, as it is necessary to reconstruct the contributions of all sources of food in pre-European Maori diet in order to apply the model described above. Archaeological data from each time period in each study area were analysed in five steps.

### 2.3.1 Determining the range of species harvested.

The species of animals harvested was established by summarising the full suite of assemblages in each area/period data set. This enabled incorporation of taxa represented only in assemblages for which quantified data were incomplete. The proportion of assemblages in which each species was represented during each study period was calculated to provide the broadest level analysis of changes in harvest patterns over time. Analysis of these data disclosed taxa that are likely to be either underor over-represented in quantified archaeofaunal samples, and also highlighted variations in species representation between sites of different functional types, both factors relevant to defining overall regional patterns. Further details and results of this analysis are presented elsewhere (Smith in press).

### 2.3.2 Determining the frequency with which species were harvested.

The frequency with which species were harvested was investigated through analysis of the data sets quantified via MNI. This is a derived measure, and presents some difficulties as variations in calculation methodology can influence data outputs, especially when, as here, information is drawn from a range of different researchers (Reitz \& Wing 2008). However it was the only measure available for a wide range of assemblages, and the only one from which estimates of harvested biomass across all classes of fauna could be generated. The uncertainties that this introduces into calculations are considered further below (Section 2.3.5).

MNI for each assemblage were first subdivided into seven faunal classes (fish, shellfish, marine and coastal birds, marine mammals, moas, smaller terrestrial birds, and terrestrial mammals) and \%MNI within each class calculated. This procedure provides a much finer-grained assessment of within-class frequency than is possible when assemblages are treated as a whole, due to wide variations in faunal class abundance between sites of different functional types, such as specialised shell fishing camps or generalised occupation sites (Smith in press).

Mean \%MNI across all assemblages in each period/area data set were then calculated. For two data sets (Greater Hauraki, Middle/Late and Late period marine/coastal birds) comprising a small number of assemblages with low species MNI, \%sum MNI was preferred as a better estimate of relative abundance. Where the taxa list for a period/area set included imprecisely identified taxa (e.g. Phalacrocorax melanoleucos?; Trachurus ?sp.; 'elephant seal or leopard seal'; etc.) in addition to positively identified species of the same sort(s), frequencies for the former were redistributed to the latter. Where the latter included more than one positively identified species, their relative abundance was factored into the redistribution so as to maintain proportionality of positively identified taxa.

For several data sets adjustments were made to incorporate taxa known to be present from analysis in 2.3.1 above, but not present in the quantified samples. This was required for all classes of fauna for the Middle period in Otago-Catlins for which there were only two study assemblages, neither having quantified data. In these cases the average of mean $\% \mathrm{MNI}$ in each of the adjacent overlap periods (Early/Middle, Middle/Late) was calculated for the species noted as present in the Middle period assemblages, and the resulting values were adjusted so that each faunal class summed to $100 \%$. For the Middle period in Greater Hauraki none of the assemblages, quantified or otherwise, yielded marine or coastal birds, but as argued elsewhere (Smith in press) this is more likely to reflect sampling variation than lack of harvesting. For the present analysis, the range of species represented in the subsequent overlap period (Middle/Late) was utilised, with their frequency derived from the average of mean $\%$ MNI in the Early/Middle and Middle/Late overlap periods, adjusted to total $100 \%$.

Finally, adjustments were made to the Late period frequencies for marine/coastal bird in both study areas where Diomedeidae appeared to be over-represented through bones from beach wrecks being collected as raw material for artefact manufacture (Smith in press). Comparison with earlier periods suggests remains of this family are about twice as common as expected, and on this basis $\% \mathrm{MNI}$ for species in this family were reduced by half and all other taxon frequencies increased proportionately.

The adjusted $\% \mathrm{MNI}$ values for each taxon are referred to as taxon frequencies when incorporated in subsequent steps of the analysis. Values for the Early, Middle and Late periods in each study area are listed in Appendix 1.

### 2.3.3 Determining the energy yield per species in study assemblages

The importance of each species as a source of energy in the human diet was evaluated for each assemblage using procedures set out in detail elsewhere (Smith 2011). In brief this involved converting the frequency of each species in an assemblage first to the weight of meat that they represented and then to the calorific value of energy that this would produce (Figure 3).

For smaller-sized classes of fauna it was presumed that all usable meat on each animal would have been available for consumption, and MNI were used as the starting point for calculation. Mean adult body weights, and a conservative estimate of the proportion likely to have been usable meat in the prehistoric New Zealand setting were used to generate meat yields, and data from proximate composition analysis for the species concerned or the nearest comparable taxon were used to derive calorific yields. Details of the values adopted for each species, the rationale for their selection, and the sources that they are drawn from are given elsewhere (Smith 2011).

For larger-sized animals, archaeological evidence frequently discloses only partial carcass representation, hence assuming that complete animals were present risks over-estimating their dietary importance. Except where skeletal element representation in an assemblage indicated that a complete or near complete individual was present, species frequency for the larger animal classes was measured in terms of the minimum number of butchery units (MNBU): the smallest number of butchery units necessary to account for all of the remains of a taxon in an archaeological assemblage. Age and sex related size differences are also relevant to accurate estimation of dietary importance in large animals, and where adequate data were available for archaeological remains (in the present context, only for some seal species) these were incorporated into calculations (Smith 2011).

These procedures were applied to the subset of study assemblages with quantified data for all classes of fauna. In each case the proportion of total energy from animals (\%Kcal) derived from each of the seven major classes of fauna was calculated, and incorporated into the following step.

### 2.3.4 Determining the relative energy yields of each faunal class per area/period

The relative contributions to the energy harvest from each faunal class during each formal period (i.e. Early, Middle, Late) in each study area was estimated from mean energy contributions ( $\% \mathrm{Kcal}$ ) of each major faunal class in either the total area/period data set, or a subset thereof. A series of adjustments were then made to each set of calculated values to take account of probable underrepresentation of some classes of fauna. Both the rationale for the selection of most appropriate data suites and justification for subsequent adjustments are discussed in detail in a manuscript currently in preparation (Smith n.d.). Key elements of these are summarised below.

For the Early period in both study areas all the sample sites appear to have been villages at which a wide range of subsistence activities were represented, indicating that the best estimate of the relative importance of each faunal classes could be derived from its mean $\% \mathrm{Kcal}$ in the total data set for each area. Sites from the Middle period in Greater Hauraki included a small number of generalised village sites along with a greater number of specialised shellfish gathering and/or fishing camps. Most of the latter represented comparatively small volumes of food energy, and if incorporated directly into an area/period average would over-estimate the importance of shellfish and fish. The best estimate for this period was drawn from the mean $\% \mathrm{Kcal}$ of the five largest assemblages, which included two villages and three short-term camps. For the Middle period in Otago-Catlins, which had no quantified data sets, mean $\% \mathrm{Kcal}$ for each of the two adjacent overlap periods were averaged to provide a best estimate. Quantified assemblages from the Late period in both study areas were relatively few in number, of greatly unequal sizes and did not well represent the expected range of site types. The addition of Middle/Late overlap period assemblages largely resolved these problems, hence the best estimate for the Late period in each study area was derived from mean \%Kcal of all Middle/Late and Late assemblages.

Adjustments were made to the calculated values for each area/period so that they would better represent the overall makeup of human dietary patterns. These are summarised below.

1. Due to the coastal location of nearly all study sites, it is likely that terrestrial animals exploited by the Maori communities of each study area are under-represented. It is difficult to estimate the magnitude of this as there are as yet no comprehensive analyses of energy yields per taxa at inland sites. A semi-quantitative analysis of relative frequencies of the main faunal classes in Early period sites in the southern South Island (Anderson 1982: Table 5) shows that when inland sites are added to those on the coast, the weighted abundance of terrestrial taxa increases by about $7 \%$. In view of the generally lower energy yields from terrestrial animals compared to those from the sea, a lower value was used here, and $5 \%$ was added to the total calculated value for terrestrial animals, distributed proportionately to each component (moas, terrestrial birds, terrestrial mammals), and $5 \%$ deducted from the total for marine animals, subtracted proportionately from each component (fish, shellfish, marine and coastal birds, marine mammals).
2. While marine mammals were totally absent from the study assemblages from the Middle and Late periods in Greater Hauraki, analysis in 2.3.1 showed that small numbers of dolphins had almost certainly been exploited. To accommodate this $0.05 \%$ of total Kcal was transferred from fish to marine mammals in those two data sets.
3. Both marine/coastal and terrestrial small birds appear to be under-represented in the archaeological record, especially after the Early period, with the most likely explanation relating to the manner in which bird carcasses were processed for preservation and transport (Smith in press). There is as yet no quantified measure of the magnitude of under-representation, and the values used here are estimates. The calculated values for marine/coastal and terrestrial small birds were increased by $1 \%$ in Early period assemblages and $1.5 \%$ in Middle and Late period
assemblages, with equivalent amounts subtracted from the marine and terrestrial components with the highest values. This adjustment was not applied to the Otago-Catlins Late period marine/coastal birds, which were already well represented in the calculated value.
4. Dogs appeared to be under-represented in the Early period data sets from both study areas, with relative frequencies lower than those reported for other sites of this age (Allo Bay-Petersen 1979; Clark 1995). On current data it is difficult to quantify the impact of this on relative energy yields and an arbitrary value of $1 \%$ is used here, with this proportion of total Kcal transferred from the moa component to terrestrial mammals.
5. Sharks and rays are almost certainly under-represented in the archaeological record due to poor survival of their predominantly cartilaginous skeletons, while ethnohistorical evidence indicates that they were likely to have been a significant component of Late period Greater Hauraki Maori fisheries (Smith in press). If at least one taxon from this class had been represented in each Late period assemblage, the energy yield for fish would have been at least $4 \%$ higher than the calculated values. On this basis, $4 \%$ of total Kcal was transferred from shellfish to fish.
6. Shellfish appear to have been under-represented in the Late period Otago-Catlins data set, with higher relative frequencies at sites not able to be included in the present study (Davies 1980). As with dogs, it is difficult to quantify the extent of under-representation, and an estimate was employed, with $2.5 \%$ of total Kcal was transferred from fish to shellfish.

Table 2 lists both the initial calculated values and final adjusted values for the Early, Middle and Late Periods in each study area. The final adjusted values are referred to as faunal component energy proportions when incorporated into subsequent steps of the analysis.

### 2.3.5 Calculating the contributions of each species to total energy harvested from animals

The contribution of each species to the total energy harvest from animal foods during each of the three formal periods in each study area was calculated as follows (also see Figure 2).
a. Taxon frequencies determined in 2.3 .2 were multiplied by energy yield per animal using the procedures described in 2.3.3 to derive an energy output per species. Note that for fur seals, these calculations were undertaken by age-sex class, and that for Pilot whales, energy yield per animal assumed that only $10 \%$ of available meat weight was consumed, following protocols established elsewhere (Smith 1985, 2004).
b. The energy output for each species calculated in (a) was divided by the sum of energy outputs for all species represented within the major faunal component to which it belonged to determine the proportion which that species contributed to total energy derived from that faunal component.
c. The proportional contribution of each species calculated in (b) was multiplied by the relevant faunal component energy proportion determined in 2.3.4 to derive a final proportional energy contribution per species. The values calculated for each species are listed in Appendix 1.

Cumulative calculation errors involved in deriving the proportional energy contribution per species were determined from estimates of the size of error associated with the three key input variables in the calculation. The taxon frequencies were based upon MNI data, which have inherent uncertainty (section 2.3.2) and calculated as means across variable sized sample sets of assemblages suggesting an error of $\pm 5 \%$ is in order. Energy yields per individual animal were derived from a mean body
weight, an estimate of the proportion typically eaten, and a value for the energy yield per kg of flesh. Each of these is likely to have varied due to a range of factors including the age, size and condition of the animal concerned along with day-to-day variations in the butchery and consumption practices of the communities harvesting the animals. Again, $\pm 5 \%$ is allowed for each component, giving a cumulative error of $\pm 15 \%$ for the estimate of energy yield per animal. The faunal component energy proportions were the most difficult to derive from the archaeological data because they involved judgement about composition of an appropriate regional data set, and incorporated several adjustments to calculated values to correct for perceived biases in the archaeological record. For these an error of $\pm 10 \%$ is proposed. With the final proportional energy contribution per species derived from the product of these three variables, an error of $\pm 30 \%$ should be applied to the calculated values.

### 2.4 Estimating the scale of the total energy harvest from animals

The scale of the total energy harvest from animals during the focal year of each period in each study area was calculated from estimates of (a) the size of human population, (b) average energy needs per person per day, and (c) the proportion of that energy derived from animal foods. There are considerable uncertainties about these variables, described below, and different values for these produce widely different totals for the estimate of total energy from animals. For these reasons an attempt has been made to give a good indication of upper and lower limits as well as what is considered a 'best estimate'. Minimum estimates of total energy from animals use the lowest likely values for (a), (b) and (c); maximum estimates use the highest acceptable values; while best estimates are based on what are considered here to be the most well supported or realistic values. These values are summarised in Table 3 and the rationale for their selection given below in sections 2.4.1 to 2.4.3.

### 2.4.1 Estimating the size of human population

The size of the human population is most reliably determined for the Late period (ca. 1750 AD ) based upon detailed analysis by Pool (1991) who gave a best estimate of $c a .100,000$ for the total New Zealand population in 1769 , with $\mathrm{ca} .90 \%$ of that in the North Island. It is estimated here that the Greater Hauraki study area population was about $13.5 \%$ of the North Island total, based on Urlich's (1969) estimate that $5 \%$ were in Thames-Coromandel and $7 \%$ in Auckland, to which has been added one tenth of the $15 \%$ she estimated for the whole of Northland. This yielded the best estimate value of 12 150. Minimum and maximum values were calculated using total North Island populations of 80 000 and 100000 respectively, giving estimates for Greater Hauraki of 10800 and 13500 . Anderson (1998: 196-7) has estimated that in the early $19^{\text {th }}$ century the population of Ngai Tahu in the southern two thirds of the South Island was no more than 5000. Anderson's analysis of the distribution of this population, along with the distribution of Late period archaeological sites suggests a population at $c a$. 1750 AD for Otago-Catlins of about 1800, with $\pm 200$ allowed for minimum and maximum estimates.

Population estimates for the Early period (ca. 1400 AD ) are based on the likelihood that the initial colonising population would have grown rapidly from a founding population of ca. 300-500 at 1280 AD increasing exponentially at rates in the order of $2 \%$ to $3.5 \%$ per annum. This would give nationwide totals of at least 4000 , and possibly up to 20000 by 1400 AD , with the best estimate towards the upper end of this range. Based on what is known of the nationwide distribution of Early period archaeological sites, it is estimated that this population was more or less evenly distributed between the North and South Islands, and about a quarter of the North Island total was in Greater Hauraki. This gives minimum, best and maximum estimates of 500, 2000 and 2500. Likewise, it is likely that about a quarter of the South Island population were in Otago-Catlins. The same minimum and maximum estimates used for Greater Hauraki have been adopted, but a slightly lower best estimate of 1800 .

For the Middle period (ca. 1550 AD ) in Greater Hauraki population size was estimated using the trend line for population growth in prehistoric New Zealand indicated by cumulative frequency of radiocarbon dates, adjusted for calibration stochastic distortion effect (McFadgen et al. 1994: Figure 3). This suggests that at $c a .1550 \mathrm{AD}$ the population would have reached about $90 \%$ of its $c a .1750$ AD level (Figure 4). This factor was applied to the North Island population estimates for the Late period given above, giving a minimum of 72000 , best estimate of 80000 , and a maximum of 90000 . The proportion of these within Greater Hauraki is difficult to assess, but a range of values were used here; $8.5 \%$ for the minimum estimate, $12.6 \%$ for the best estimate, and $13 \%$ for the maximum; giving population estimates of 6100, 10200 and 11700 .

It is widely recognised that the population in southern New Zealand took a much lower trajectory, perhaps even declining after the Early period (Jacomb et al. 2010, McGlone et al. 1994). Examination of the median ages of radiocarbon dated study assemblages from Otago-Catlins used in this study supports this view (Figure 4), although it needs to be noted that both there are additional dated sites from both Middle and Late periods that were excluded from the present study because they lacked fully identified and/or quantified fauna. The best estimate used here (1800) assume that the population at $c a .1550 \mathrm{AD}$ was no greater than at $c a .1400 \mathrm{AD}$, while the maximum estimate (2600) is slightly higher than that of the earlier period and the minimum estimate (500) maintains the same lower limit as at $c a .1400 \mathrm{AD}$.

### 2.4.2 Estimating average energy needs per person per day

The daily energy requirements of a person depend upon a wide range of factors including gender, body weight, activity levels and the energy demands of the environmental conditions in which they live. The estimates used here are drawn from data compiled and reviewed in detail elsewhere (Leach 2006, Leach et al. 1996) which focused explicitly on determining an appropriate range for prehistoric populations in the Pacific and New Zealand. For Greater Hauraki the minimum value (1800) is that accepted by Leach et al. (1996: 24) as the lowest viable for a 70 kg woman with low activity levels. The best estimate (2150) is the mean value suggested for pre-European Maori by Leach (2006: 277). The maximum (2172) is the midpoint between that mean and the value derived by stochastic modelling of pre-European diet in the Chatham Islands (Leach et al. 2003), where the environmental conditions are likely to have placed greater energy demands on individuals than was the case in Greater Hauraki.

Values adopted for Otago-Catlins are set slightly higher to accommodate the greater energy demands of a colder climate. The minimum value (2150) is Leach's mean for pre-European Maori; the best estimate (2172) is the value adopted above for the upper limit in Greater Hauraki; and the maximum value (2193) is that derived for the Chatham Islands.

### 2.4.3 Estimating the proportion of energy derived from animal foods

Estimates of the proportion of the total energy consumed by people that derived from animal foods are based upon analysis of $\delta^{13} \mathrm{C}, \delta^{15} \mathrm{~N}$ and $\delta^{34} \mathrm{~S}$ isotopes from the small number of archaeologicallyderived samples of human bone collagen thus-far analysed from the Pacific region (Leach et al. 2000, Leach et al. 2003). These exhibit wide variation: the lowest value (0.4) is from Watom Island in the tropical Pacific with a horticultural and fishing economy; a moderately high value ( 0.66 ) derives from Wairau Bar, South Island New Zealand, where horticulture was combined with hunting of large animals; and the highest value ( 0.9 ) is from the Chatham Islands where subsistence relied upon hunting, fishing and the gathering of wild plants.

In Greater Hauraki archaeological data for the Middle and Late periods indicate an economy based on horticulture, fisheries and a very modest level of hunting (Smith in press, Smith n.d.). On this basis the Watom and Wairau Bar values are adopted for minimum and maximum values respectively, and the mid-point between these is used for the best estimate. Archaeological data for the early period indicate a much greater reliance on hunting, hence the Wairau Bar value is adopted for the best estimate, with midpoints between that and each of the higher and lower measured values used for minimum and maximum estimates.

The Otago-Catlins region was beyond the growing limits for pre-European horticultural crops, with wild plant foods contributing to a diet that archaeological data indicates relied heavily on both hunting and fishing throughout the prehistoric sequence (Smith in press, Smith n.d.). On this basis a single set of values is used for all periods with the Wairau Bar and Chatham Islands values adopted for the minimum and maximum estimates respectively, and the midpoint between these used as the best estimate.

### 2.4.4 Calculating total energy harvest from animals

Estimates for the total energy harvest (Kcal) from animals in each study area during each of the target years ( $1400 \mathrm{AD}, 1550 \mathrm{AD}, 1750$ ) were calculated as follows. The estimate of human population size was multiplied by the estimated energy needs per person per day, which was multiplied by 365 to determine the total energy needs for the year, and this was then multiplied by the estimated proportion of energy derived from animal foods. In doing this minimum input values were combined to produce minimum estimates, and likewise for best and maximum values. Calculated values, referred to as energy from animals, are shown in Table 4.

### 2.5 Estimating the number and biomass of marine animals harvested

The final step of analysis involved using the variables and values calculated or estimated in the preceding sections to estimate the number and biomass of animals harvested during the focal year in each study area. This was undertaken in three steps.
a. The proportional energy contribution per species (2.3.5) was multiplied by the energy from animals (2.4.4) to derive the total energy harvested from that species.
b. Total energy harvested from species was divided by the energy yield per individual of that species (2.3.3) to determine the number of animals harvested.
c. The number of animals harvested was multiplied by the mean body weight per individuals of that species (2.3.3) to determine the biomass harvested.

The estimated cumulative error $( \pm 30 \%)$ in calculating proportional energy per species was applied to the number and biomass estimates ( $b$ and c ), with uncertainties about the scale of the energy harvest from animals expressed in the minimum, best and maximum estimates. Results of this are presented below.

## 3. RESULTS

Analysis of the archaeological study assemblages demonstrated that at least 210 marine or coastal species had been harvested by Maori in Greater Hauraki ( 147 shellfish, 35 fish, 22 birds, 6 mammals), and at least 159 in Otago-Catlins ( 90 shellfish, 32 fish, 30 birds, 7 mammals). Estimates made of the
number and biomass harvested from each of these are presented in Tables $5-28$. Within each of these tables taxa are arranged by systematic order after the following authorities: for fish, Froese \& Pauly (2010); shellfish, Spencer et al. (2009); birds, Checklist Committee (OSNZ) (2010); and mammals, Baker et al. (2010). This enables placement of taxa identified only to genus or higher levels, and those less securely identified, alongside the positively identified species to which they are most closely related. Common names for taxa in these tables are listed in Appendix 1. For shellfish the tables list individually only the 22 species with highest harvested biomass in each area/period with the remainder grouped as 'all others'. In the following presentation of results, only the best estimate values are used in text; minimum and maximum can be found in the Tables. The latter exhibit parallel trends to those identified using the best estimate values.

### 3.1 Fish

### 3.1.1 Greater Hauraki

A total of 35 fish taxa are recorded archaeologically as harvested by Maori in Greater Hauraki. These included some genus, family or higher level groupings necessitated by the nature of archaeological data: it is virtually impossible to identify the various species of Labridae on the basis of skeletal morphology, so they are treated as a single family group; Anguilla species are also indistinguishable, so are treated as a genus group; Carangidae are often difficult to distinguish, and only sometimes identified to species level; and the poor survival of elasmobranch remains frequently precludes specific identification.

At $c a .1400$ AD 22 of these taxa were included in a total fish harvest best estimated at $128.5 \pm 44.3$ tonnes (Table 5). The taxonomic range narrowed to 16 at $c a .1550 \mathrm{AD}$ in a total harvest of $2328.2 \pm$ 698.5 t (Table 6). By $c a .1750$ AD, the total fish take of $2607.1 \pm 782.1 \mathrm{t}$ was drawn from 15 taxa (Table 7).

Snapper (Pagrus auratus) were the largest single contributor to the fish harvest at each time period, rising from $72.0 \pm 21.6 \mathrm{t}$ to $938.8 \pm 281.6 \mathrm{t}$ and then $997.2 \pm 299.2 \mathrm{t}$, although as a proportion of total harvested fish biomass this represents a falling trend from $56 \%$ to $40 \%$ then $38 \%$. At ca. 1400 AD only three other taxa other than snapper made up significant proportions of harvested biomass; kahawai (Arripis trutta) $19.2 \pm 5.8 \mathrm{t}(15 \%)$, wrasses (Labridae) $13.5 \pm 4.0 \mathrm{t}(10.5 \%)$, and leatherjacket (Meuschenia scaber) $11.3 \pm 3.4 \mathrm{t}(8.8 \%)$. At ca. 1550 AD there were four main contributors to harvested biomass in addition to snapper; sharks (Carcharhiniformes) not positively identified to species, $511.4 \pm 153.4 \mathrm{t}(22 \%)$; horse mackerel (Trachurus novaezelandiae), $323.1 \pm 96.9 \mathrm{t}(13.9 \%)$; kahawai, $227.9 \pm 68.4 \mathrm{t}(9.8 \%)$; and barracouta (Thyrsites atun) $169.3 \pm 50.8 \mathrm{t}(7.3 \%)$. The harvest of wrasses ( $23.1 \pm 6.9 \mathrm{t}$ ) was now only $6.9 \%$ of the total, while leatherjackets were not represented in any of the study sites from this era. At ca. 1750 AD the harvested biomass of sharks not positively identified to species ( $959 \pm 287.8 \mathrm{t}, 36.8 \%$ ) was almost equal to that of snapper, and would just outstrip it if combined with the $39.9 \pm 12 \mathrm{t}$ identified as from northern dogfish (Squalus blainvillei). The eagle ray (Myliobatis tenuicaudatus) was the only other significant component of the fishery at ca. 1750 AD , contributing $319.3 \pm 95.8 \mathrm{t}(12.2 \%)$.

### 1.1.2 Otago-Catlins

A total of 32 taxa were identified in study sites from the Otago-Catlins region, including the genus and higher-level groupings noted for Greater Hauraki, along with Nototheniidae, as it is difficult to distinguish various species of black cods on skeletal morphology.

In total 22 of these taxa were included in a total fish harvest best estimated at $78.3 \pm 23.5 \mathrm{t}$ at ca .1400 AD (Table 8). The taxonomic range narrowed to 14 at $c a .1550 \mathrm{AD}$ in a total harvest of $797.5 \pm 239.3$ t (Table 9). By ca. 1750 AD , the total fish take of $725.1 \pm 217.5 \mathrm{t}$ was drawn from 9 taxa (Table 10).

Barracouta (Thyrsites atun) formed the largest part of harvested fish biomass at each time period, contributing $43.2 \pm 13.0 \mathrm{t}(55.2 \%)$ at $c a .1400 \mathrm{AD}, 361.4 \pm 108.4 \mathrm{t}(45.3 \%)$ at $c a .1550 \mathrm{AD}$, and 363.3 $\pm 109.0 \mathrm{t}(50.1 \%)$ at $c a .1750 \mathrm{AD}$. Three further species made important contributions to harvested biomass at ca. 1400 AD ; ling (Genypterus blacodes) $7.8 \pm 2.3 \mathrm{t}$ ( $9.9 \%$ ), hapuku (Polyprion oxygenios) $7.7 \pm 2.3 \mathrm{t}(9.8 \%)$, and red cod (Pseudophycis bachus) $6.8 \pm 2.0 \mathrm{t}(8.7 \%)$. The same three species were the major components in addition to barracouta at $c a .1550 \mathrm{AD}$, although in differing order; hapuku $141.0 \pm 42.3 \mathrm{t}(17.7 \%)$, red cod $128.3 \pm 38.5 \mathrm{t}(16.1 \%)$, and ling $68.2 \pm 20.5 \mathrm{t}(8.5 \%)$. This order was maintained at $c a .1750 \mathrm{AD}$; hapuku $230.9 \pm 69.3 \mathrm{t}(31.8 \%)$, red $\operatorname{cod} 68.4 \pm 20.5(9.4 \%)$, and ling $56.8 \pm 17 \mathrm{t}(7.8 \%)$.

### 3.2 Shellfish

### 3.2.1 Greater Hauraki

The archaeological study sites provided records of 147 shellfish taxa harvested by Maori in the Greater Hauraki region, with 31 of these being identifications only to genus or family level leaving the possibility that the total number of species may have been slightly more than this. Of these taxa, 88 were represented in the estimated harvest of $6.6 \pm 2.0 \mathrm{t}$ at $c a .1400 \mathrm{AD}$ (Table 11), 66 in the estimate of $1736.3 \pm 520.9 \mathrm{t}$ at ca .1550 AD (Table 12), and 38 in the estimate of $2294.7 \pm 688.4 \mathrm{t}$ at ca. 1750 AD (Table 13).

Three species dominate the shellfish biomass harvested at $c a .1400 \mathrm{AD}$, making up two thirds of the total; paua (Haliotis iris) $2.2 \pm 0.7 \mathrm{t}$ (34\%), cats eye (Lunella smaragdus) $1.2 \pm 0.4 \mathrm{t}$ ( $18.2 \%$ ), and green-lipped mussel (Perna canaliculus) $0.9 \pm 0.3 \mathrm{t}(14.3 \%)$. At ca. 1550 AD two species made up more than $84 \%$ of the total biomass harvested: cockle (Austrovenus stutchburyi) $1103.2 \pm 331.0 \mathrm{t}$ ( $63.5 \%$ ), and pipi (Paphies australis) $358.2 \pm 107.5 \mathrm{t}(20.6 \%)$. Of the three formerly dominant species (paua, cats eye and green lipped mussel), only cats eyes at $5 \%$ made a notable contribution with the others providing less than $0.1 \%$ of harvested biomass. The same two species, cockle and pipi continued to dominate the shellfish harvest at $c a .1750 \mathrm{AD}$ as they did at $c a .1550$; cockle providing $1358.0 \pm 407.4 \mathrm{t}(59.2 \%)$, and pipi $594.5 \pm 178.4 \mathrm{t}(25.9 \%)$. Tuatua (Paphies subtriangulata) with $250.5 \pm 75.2 \mathrm{t}(10.9 \%)$ was the only additional species at ca. 1750 to provide more than $1 \%$ total biomass harvested.

### 3.2.2 Otago-Catlins

The shellfish identified from study sites in Otago-Catlins include 90 taxa, including 25 identifications to only to genus or family level. At ca. 1400 AD 55 taxa were represented in the total shellfish harvest estimated at $9.1 \pm 2.7 \mathrm{t}$ (Table 14), with just 12 taxa in the estimated $43.0 \pm 12.9 \mathrm{t}$ harvested at $c a$. 1550 AD (Table 15), but 29 were included in the estimated harvest of $51.4 \pm 15.4 \mathrm{t}$ at ca .1750 AD (Table 16). The reduced taxonomic range at $c a .1550$ is almost certainly due to the restricted sample of study sites available for that time period; ten study assemblages were available for $c a .1400 \mathrm{AD}$ and nine for $c a .1750 \mathrm{AD}$, but only two for $c a .1550 \mathrm{AD}$.

At ca. 1400 AD four species made up $82.4 \%$ of the total harvested biomass; cockle (Austrovenus stutchburyi) $2.4 \pm 0.7 \mathrm{t}(26.9 \%$ ), paua (Haliotis iris) $2.3 \pm 0.7 \mathrm{t}$ ( $24.8 \%$ ), blue mussel (Mytilus galloprovincialis) $1.5 \pm 0.5 \mathrm{t}(16.6 \%)$, and pipi (Paphies australis) $1.3 \pm 0.3 \mathrm{t}(14.1 \%)$. Two of these four species also dominate harvested shellfish biomass at $c a$. 1550 AD ; blue mussels providing $16.1 \pm$
$4.8 \mathrm{t}(37.4 \%)$, and paua $14.2 \pm 4.3 \mathrm{t}$ ( $33 \%$ ). Mudsnails (Amphibola crenata) added another $4.2 \pm 1.3 \mathrm{t}$ $(9.8 \%)$, cockles $2.3 \pm 0.2 \mathrm{t}(5.4 \%)$, and pipi $1.7 \pm 0.5 \mathrm{t}(4 \%)$ to the total harvested biomass in ca. 1550 AD. At $c a$. 1750 AD the rank order of the two leading shellfish species reverses; paua yielding $35.1 \pm$ $10.5 \mathrm{t}(68.4 \%)$, and blue mussels $8.2 \pm 2.5 \mathrm{t}(15.9 \%)$. None of the other previously important species contributed more than $2.5 \%$ of total harvested biomass at $c a .1750 \mathrm{AD}$.

### 3.3 Marine and Coastal Birds

### 3.3.1 Greater Hauraki

A total of 22 marine and coastal bird species were positively identified in the study sites from Greater Hauraki, along with identifications to five higher-level taxonomic groupings. Twenty two taxa were represented in the estimated harvest of $19.2 \pm 5.8 \mathrm{t}$ in $c a .1400 \mathrm{AD}$ (Table 17). Only seven taxa were among the harvest of $28.3 \pm \mathrm{t}$ at $c a .1550 \mathrm{AD}$ (Table 18), while at $c a .1750 \mathrm{AD}$ there were ten in the $56.2 \pm 16.9 \mathrm{t}$ harvested (Table 19).

At ca. 1400 AD four taxa together made up $70.1 \%$ of the biomass harvested from marine and coastal birds; little penguin (Eudyptula minor) $4.7 \pm 1.4 \mathrm{t}(24.4 \%)$, spotted shag (Stictocarbo punctatus) $4.0 \pm$ $1.2 \mathrm{t}(20.7 \%)$, albatrosses (Diomedeidae) $2.5 \pm 0.8 \mathrm{t}(13.2 \%)$, and pied shag (Phalacrocorax varius) $2.3 \pm 0.7 \mathrm{t}(11.9 \%)$. Three of these taxa together yielded $90.2 \%$ of the harvested biomass in $c a .1550$ AD ; albatrosses $17.6 \pm 5.3 \mathrm{t}(62.2 \%)$, little penguin $5.1 \pm 1.5 \mathrm{t}(185)$, and spotted shag $2.8 \pm 0.8 \mathrm{t}$ ( $10 \%$ ). At ca. 1750 AD the same three taxa provided $71.6 \%$ of the harvested biomass; albatrosses $17.6 \pm 5.3 \mathrm{t}(31.3 \%)$, spotted shag $14.1 \pm 4.2 \mathrm{t}(25 \%)$, and little penguin $8.6 \pm 2.6 \mathrm{t}(15.3 \%)$.

### 3.3.2 Otago-Catlins

Thirty taxa of marine and coastal birds species were identified in the study sites from Otago Catlins. These include penguins listed as Megadyptes sp. as virtually all were identified prior to recent separation of the smaller extinct Waitaha penguin (M. waitaha) from the larger extant yellow-eyed penguin (M. antipodes) (Bossenkool et al. 2009). The date at which the former replaced the latter in the study area is not yet clearly known. At ca. 140025 marine and coastal bird taxa formed part of a total estimated harvest of $32.1 \pm 9.6 \mathrm{t}$ (Table 20). Only sixteen were among the harvest of $46.9 \pm 14.1$ t in ca. 1550 AD (Table 21), while 18 contributed to the $80.3 \pm 24.1 \mathrm{t}$ harvested in $c a .1750 \mathrm{AD}$ (Table 22).

At ca. 1400 AD four species made up $76 \%$ of the total biomass harvested from marine and coastal birds; Megadyptes, at this time almost certainly the waitaha penguin, $8.8 \pm 2.6 \mathrm{t}$ ( $27.2 \%$ ), Fiordland crested penguin (Eudyptes pachyrhynchus) $8.5 \pm 2.5 \mathrm{t}(26.4 \%)$, white-capped albatross (Thalassarche cauta) $4.7 \pm 1.4 \mathrm{t}(14.5 \%)$, and little penguin (Eudyptula minor) $2.5 \pm 0.8 \mathrm{t}$ ( $7.8 \%$ ). Five taxa, including the four above, made up $74 \%$ of the biomass harvested at $c a .1550 \mathrm{AD}$; spotted shag (Stictocarbo punctatus) $9.9 \pm 3.0 \mathrm{t}(21 \%)$, white-capped albatross $6.9 \pm 2.1 \mathrm{t}(14.6 \%)$, Megadyptes sp . penguin $6.8 \pm 2.1 \mathrm{t}(14.6 \%)$, Fiordland crested penguin $5.8 \pm 1.8 \mathrm{t}(12.5 \%)$, and little penguin $5.3 \pm$ $1.6 \mathrm{t}(11.3 \%)$. At $c a .1750 \mathrm{AD}$ the leading four taxa contributed $67.9 \%$ of the total biomass harvested; white capped albatross $20.6 \pm 6.2 \mathrm{t}(25.6 \%)$, Stewart Island shag (Leucocarbo chalconotus) $15.1 \pm 4.5$ $\mathrm{t}(18.8 \%)$, spotted shag $9.9 \pm 3.0 \mathrm{t}(12.3 \%)$, and Megadyptes, by this time probably yellow-eyed penguin, $9.0 \pm 2.7 \mathrm{t}$ (11.2\%).

### 3.4 Marine Mammals

### 3.4.1 Greater Hauraki

At least five marine mammal taxa are recorded as part of the Maori marine harvest in Greater Hauraki. These include the fur seal (Arctocephalus forsteri), sea lion (Phocarctus hookeri) and elephant seal (Mirounga leonina), along with at least one species of pilot whale (Globicephala) and dolphins that have not been identified to species level.

All these taxa were represented in the total harvest estimated at $414.5 \pm 124.3 \mathrm{t}$ in ca . 1400AD (Table 23). Fur seals made up the largest part of this, $284.9 \pm 85.5 \mathrm{t}$ ( $68.7 \%$ ), followed by sea lions, $63.4 \pm$ $19.0 \mathrm{t}(15.3 \%)$, and pilot whales, $57.5 \pm 17.3 \mathrm{t}(13.9 \%)$. By $c a .1550 \mathrm{AD}$ the total harvest had fallen to $1.79 \pm 0.54 \mathrm{t}$, derived entirely from dolphins (Table 24). Dolphins were also the sole contributor to the marine mammals harvest of $2.13 \pm 0.64 \mathrm{t}$ in $c a .1750 \mathrm{AD}$ (Table 25).

### 3.4.1 Otago-Catlins

The archaeological record shows that at least eight marine mammal taxa were harvested in OtagoCatlins, including the fur seal (Arctocephalus forsteri), sea lion (Phocarctus hookeri), elephant seal (Mirounga leonina), leopard seal (Hydrurga leptonyx), at least one species of pilot whale (Globicephala) and dolphins that include the common dolphin (Delphinus delphis) and Hectors dolphin (Cephalorhynchus hectori). The latter are treated here as a single group as the majority of dolphin remains have not been identified to species level.

At $c a .1400 \mathrm{AD}$ five taxa were included in the total harvest of $371.3 \pm 111.4 \mathrm{t}$ (Table 26). This was dominated by two species; fur seals providing $237.0 \pm 71.1 \mathrm{t}(63.8 \%)$, and sea lions $126.2 \pm 37.8 \mathrm{t}$ (34\%). Four of the same taxa, plus another, made up the considerably smaller harvest of $144.5 \pm 43.4$ t at ca .1550 AD (Table 27). Fur seals, at $103.3 \pm 31 \mathrm{t}(71.5 \%)$, and sea lions, at $47.2 \pm 11.2 \mathrm{t}(25.8 \%)$, were again dominant at $c a .1550 \mathrm{AD}$. At $c a .1750 \mathrm{AD}$ the total harvest had fallen further to $121.6 \pm$ 36.5 t derived from just three taxa (Table 28). The main contributors to the total harvested biomass at $c a .1750 \mathrm{AD}$ were fur seals, at $77.0 \pm 23.1 \mathrm{t}(63.4 \%)$, and pilot whales at $44.0 \pm 13.2 \mathrm{t}(36.2 \%)$.

## 4. DISCUSSION

There has been only one previous attempt to calculate the numbers and biomass of marine animals harvested by Maori, and this was concerned solely with the numbers of snapper harvested at ca. 1769 in the Northland region (Leach 2006). This utilised an analytical method broadly similar to that employed here, but with some differences in input values. Leach's estimate of $1919 \pm 1612 \mathrm{t}$ of snapper harvested for the whole of Northland is almost exactly double that given by the best estimate here for the Greater Hauraki region. The latter has a coastline approximately half the length of the Northland region, suggesting that the methods produce broadly comparable results.

One of the major difference between the two approaches is the use here of minimum, best and maximum estimates to provide a narrower band of greatest likelihood within the necessarily wide error margins that must apply to calculations this sort. The differences between the minimum, best and maximum estimates of harvested biomass are illustrated for snapper in the Greater Hauraki region (Figure 5), which shows that at $c a .1750 \mathrm{AD}$ the upper end of the error limit for the maximum estimate is 4.6 times greater than the lower error limit for the minimum value. As noted in section 2.4 the extent of these differences is driven largely by the choice of values selected for human population size, daily energy requirements and proportion derived from animals. While values considered most likely to reflect the situation in each study area at each focal date were employed for calculation of the
best estimates, these still need to be considered with some caution and are best viewed as indicators of the order of magnitude of Maori harvests, rather than precise descriptors.

In both study areas the total biomass harvested for three of the four major classes of fauna increased over time, with only marine mammals exhibiting a declining trend (Figure 6). In Greater Hauraki this can be attributed partly to growth of the human population, for which the best estimate at $c a .1750$ AD is 6.1 times greater than that at $c a .1400 \mathrm{AD}$. However both the fish and shellfish harvests expanded at considerably greater rates than this (20.3 and 347.7 times respectively). The primary driver of this was the need for Maori to compensate for the loss of two major sources of food following the extinction of moa and extirpation of fur seals and sea lions from northern New Zealand between 1400 and 1550 AD (Schmidt 2000b, Smith 2005). Fish and shellfish have significantly lower energy yields per kilogram than the animals they were replacing, further increasing demand upon these fisheries. The harvest of marine and coastal birds expanded at less than half the rate of human population growth, indicating an overall diminution of their part in the human diet, which is likely to reflect reduced availability through both hunting pressure and the impact of introduced mammalian predators (Holdaway 1999, Smith in press).

Otago-Catlins provides an even clearer demonstration of the replacement process, as the best estimates of human population postulate no growth between $c a .1400$ and $c a .1750$ AD. Over the same period fish and shellfish harvests expanded by 9.9 and 5.6 times respectively, while the marine mammal harvest fell to a third of its initial level. In this case the modest growth in the harvest of marine and coastal birds indicates that they played an increasingly important part in human diet over time. This divergence from the pattern observed in Greater Hauraki may reflect the important role that growth of the human population played in depleting marine and coastal bird resources in the latter area.

There is a strong trend through time in all four major classes of fauna from both study areas towards exploitation of a narrower taxonomic range. The number of taxa within each class harvested at $c a$. 1750 AD ranges between $20 \%$ and $72 \%$ of those exploited at $c a .1400$ AD (Table 29). Detailed analysis elsewhere (Smith in press) shows that for fish and shellfish, at least, this cannot be attributed to the size of archaeological samples under analysis; there are insufficient datasets to conduct similar tests for birds and mammals. Only a small part of this trend is attributable to reduction of available taxonomic breadth. Of marine taxa represented in the study assemblages only one has become extinct, and the Waitaha penguin appears to have been rapidly replaced by the yellow-eyed penguin (Bossenkool et al. 2009). Localised depletions were somewhat more common. As already noted, fur seals and sea lions were extirpated from northern New Zealand, and at least some petrels, prions and shearwaters ceased breeding on one or both of the main islands (Holdaway 1999) restricting opportunities for anything other than opportunistic harvesting of these taxa to distant offshore islands.

Neither extinction nor serious depletion of populations has been documented for any of the fish or shellfish taxa. For these classes of fauna, explanations for the narrowing of the range of species harvested can be found in the cultural sphere. In the case of shellfish it can been attributed largely to shifts in human settlement patterns which, in Greater Hauraki focussed increasingly on land suitable for horticulture in the vicinity of estuaries, which typically support a high abundance but restricted range of edible species (Smith in press). Otago-Catlins settlements of the later period appear to be more strongly clustered than those of earlier periods, especially in the vicinity of rocky headlands, although the driving forces in this case are less clearly understood. Declining diversity of fish catches may have been influenced by these changes in settlement pattern, but there are also indications that Maori fishing practice became more specialised with the dominant species in each region, snapper in Greater Hauraki and barracouta in Otago-Catlins, forming an increasingly large proportion of the total number of fish caught. It may be that the reliability with which these species could be located and
harvested in large numbers became a critical factor when the demise of seals and moas placed increasing demand on fishing as a source of dietary energy.

Although there are broad similarities in some trends over time, the patterns of species exploitation and timing of changes in these varies considerably between the two study areas. This is not unexpected as regional differences in pre-European Maori subsistence patterns are widely recognised (Leach 2006, Smith 2004). It does indicate, however, that the harvest estimates generated here should not be applied to other areas or more generally across the country. There are also some indications of subregional variations within the datasets used here. For example the relative importance of leatherjackets at ca. 1400 AD in Greater Hauraki can be correlated with the concentration of Early period sites in the region on the east coast of the Coromandel, while the rise to prominence of mackerel fisheries at $c a .1550 \mathrm{AD}$ coincides with a greater number of sites around the shores of the Hauraki Gulf (Smith in press).

One further feature of the species-by-species estimates is worthy of further comment. In several cases there is a considerable difference between the relative abundance of taxa and their relative contributions to harvested biomass, due to differences in body size. For example sharks make up only $6.2 \%$ of the number of fish harvested in Greater Hauraki at ca. 1750 AD , but contributed $36.8 \%$ of harvested biomass, almost equal to that from snapper which made up $59 \%$ of the total number of fish. With only a small number of exceptions, previous analyses of the archaeological evidence for Maori fishing have relied solely on estimates of abundance, and for this reason they have been unable to detect the late prehistoric rise in the importance of shark fishing in northern New Zealand.

Finally, this study has relied entirely upon pre-existing data of variable and sometimes unknown quality. There are several ways in which future research could enhance the accuracy with which estimates could be made regarding marine harvests in the past. Most importantly, existing archaeological data is least adequate for the Middle and Late periods of New Zealand's prehistory, largely because so much research effort has been focussed on the study of the initial colonisation phase. Furthermore, much of what is available for later periods has derived from small-scale development-driven rescue excavations, rather than properly resourced research investigations. Targeted research on the nature and consequences of later phases of occupation has the potential to provide greater clarity regarding the breadth, magnitude and consequences of Maori exploitation of the marine environment.

## 5. MANAGEMENT IMPLICATIONS

The data presented in this report makes available for the first time a suite of numerical estimates for the levels of customary harvest of a wide range of New Zealand's marine resources covering a substantial portion of the period prior to European settlement in New Zealand. These provide baseline data against which modern customary and commercial harvests can be assessed, and will have value in future stock assessments. Once integrated with other suites of data from the Taking Stock project, they will contribute to greater understanding of the long-term structure and functioning of New Zealand's marine shelf ecosystems.

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## 8. REFERENCES

Allo Bay-Petersen, J. (1979). The role of the dog in the economy of the New Zealand Maori. In: Anderson, A.J. (ed.) Birds of a feather. Osteological and archaeological papers from the South Pacific in honour of R.J. Scarlett. BAR International Series 62. 165-181.
Anderson, A.J. (1982). A review of economic patterns during the Archaic phase in southern New Zealand. New Zealand Journal of Archaeology 4: 45-75.
Anderson, A.J. (1989). Prodigious birds: Moas and moa-hunting in prehistoric New Zealand. Cambridge University Press, Cambridge.
Anderson, A.J. (1991). The chronology of colonization in New Zealand. Antiquity 65: 767-795.
Anderson, A.J. (1997). Prehistoric Polynesian impact on the New Zealand environment: Te Whenua Hou. In: Kirch, P.V. and T.L. Hunt (eds) Historical Ecology of the Pacific Islands: Prehistoric environmental and landscape change. New Haven, Yale University Press. 271-283.
Anderson, A.J. (1998). The Welcome of Strangers; an ethnohistory of southern Maori A.D. 16501850. University of Otago Press, Dunedin.

Anderson, A.J.; Smith, I.W.G. (1996). The transient village in southern New Zealand. World Archaeology 27: 359-371.
Baker, C.; Chilvers, B.; Constantine, R.; DuFresne, S.; Mattlin, R.; van Helden, A.; Hitchmough, R. (2010). Conservation status of New Zealand marine mammals (suborders Cetacea and Pinnipedia), 2009. New Zealand Journal of Marine and Freshwater Research 44: 101-115.
Bossenkool, S.; Austin, J.J.; Worthy, T.H.; Scofield, P.; Cooper, A.; Seddon, P.J.; Waters, J.M. (2009). Relict or colonizer? Extinction and range expansion of penguins in southern New Zealand. Proceedings of the Royal Society B 276: 815-821.
Bryden, M.M.; O'Connor, S.; Jones, R. (1999). Archaeological evidence for the extinction of a breeding population of Elephant Seals in Tasmania in prehistoric times. International Journal of Osteoarchaeology 9: 430-437.
Burton, R.K.; Snodgrss, J.J.; Gifford-Gonzalez, D.; Guilderson, T.; Brown, T.; Koch, P.L. (2001). Holocene changes in the ecology of northern furseals: insights from stable isotopes and archaeofauna. Oecologia 128: 107-115.
Checklist Committee, O. (2010). Checklist of the Birds of New Zealand, Norfolk and Macquarrie Islands, and the Ross Dependency of Antarctica (4th edition). Ornithological Society of New Zealand and Te Papa Press, Wellington.
CINZAS (2008). Central Index of New Zealand Archaeological Sites. Wellington, New Zealand Archaeological Association \& Department of Conservation.
Clark, G.R. (1995). The Kuri in prehistory: a skeletal analysis of the extinct Maori dog. MA thesis, Anthropology, University of Otago.
Davies, F.J. (1980). The prehistoric environment of the Dunedin area: the approach of salvage prehistory. MA thesis, Anthropology, University of Otago.
Etnier, M. (2007). Defining and identifying sustainable harvests of resources: archaeological examples of pinniped harvests in the eastern North Pacific. Journal for Nature Conservation 15: 196-207.
Froese, R.; Pauly, D. (2010). FishBase. World Wide Web electronic publication. version $(09,2010)$. http://www.fishbase.org
Grayson, D.K. (2001). The archaeological record of human impacts on animal populations. Journal of World Prehistory 16: 1-68.
Holdaway, R.N. (1999). Introduced predators and avifaunal extinction in New Zealand. In: MacPhee, R.D.E. (ed) Extinctions in Near Time. New York, Kluwer Academic. 189-238.

Hughen, K.A.; Baillie, M.G.L.; Bard, E.; Bayliss, A.; Beck, J.W.; Bertrand, C.; Blackwell, P.G.; Buck, C.E.; Burr, G.; Cutler, K.B.; Damon, P.E.; Edwards, R.L.; Fairbanks, R.G.; Friedrich, M.; Guilderson, T.P.; Kromer, B.; McCormac, F.G.; Manning, S.; Bronk Ramsey, C.; Reimer, P.J.; Reimer, R.W.; Remmele, S.; Southon, J.R.; Stuiver, M.; Talamo, S.; Taylor, F.W.; van
der Plicht, J.; Weyhenmeyer, C.E. (2004). Marine04 marine radiocarbon calibration, 0-26 cal kyr BP. Radiocarbon 46: 959-966.
Jacomb, C.; Walter, R.K.; Jennings, C. (2010). A review of the archaeology of Foveaux Strait, New Zealand. Journal of the Polynesian Society 119: 25-59.
King, C.M. (1995). The Handbook of New Zealand Mammals. Oxford University Press, Auckland. 600 p.
Leach, B.F. (2006). Fishing in Pre-European New Zealand. New Zealand Journal of Archaeology, and Archaeofauna, Wellington. 359 p.
Leach, B.F.; Boocock, A.S. (1993). Prehistoric Fish Catches in New Zealand. British Archaeological Reports, International Series 584, Oxford. 38 p.
Leach, B.F.; Quinn, C.; Lyon, G.L. (1996). A stochastic approach to the reconstruction of prehistoric human diet in the Pacific region from bone isotope signatures. Tuhinga: Records of the Museum of New Zealand Te Papa Tongarewa 8: 1-54.
Leach, B.F.; Quinn, C.; Lyon, G.L.; Haystead, A.; Myers, D.B. (2000). Evidence of prehistoric Lapita diet at Watom Island, Papua New Guinea, using stable isotopes. New Zealand Journal of Archaeology 20: 149-159.
Leach, B.F.; Quinn, C.; Morrison, J.; Lyon, G. (2003). The use of multiple isotope signatures in reconstructing prehistoric human diet from archaeological bone from the Pacific and New Zealand. New Zealand Journal of Archaeology 23: 31-98.
Leach, B.F.; Quinn, C.; Morrison, J.; Lyon, G.L. (2003b). The use of multiple isotope signatures in reconstructing prehistoric human diet from archaeological bone from the Pacific and New Zealand. New Zealand Journal of Archaeology 23: 31--98.
Lyman, R.L. (2008). Quantitative Paleozoology. Cambridge University Press, Cambridge. 348 p.
MacDiarmid, A.B. (2011). Large, isolated late-settled islands: potential tests of human impacts on marine ecosystems. Final Research Report, Project ZBD200505, MS31 Part A, to the Ministry of Fisheries. NIWA, Wellington. 13 p .
McCormac, F.G.; Hogg, A.G.; Blackwell, P.G.; Buck, C.E.; Higham, T.F.G.; Reimer, P.J. (2004). SHCal04 Southern Hemisphere Calibration 0-1000 cal BP. Radiocarbon 46: 1087-1092.
McFadgen, B.G.; Knox, F.B.; Cole, T.R.L. (1994). Radiocarbon calibration curve variations and their implications for the interpretation of New Zealand prehistory. Radiocarbon 36: 221-236.
McGlone, M.S. (1989). The Polynesian settlement of New Zealand in relation to environmental and biotic changes. New Zealand Journal of Ecology 12: 115-129.
McGlone, M.S.; Anderson, A.J.J.; Holdaway, R.N. (1994). An Ecological Approach to the Polynesian Settlement of New Zealand. In: Sutton, D.G. (ed) The Origins of the First New Zealanders. Auckland, University of Auckland Press. 136-163.
Millener, P.R. (1981). The Quaternary Avifauna of the North Island, New Zealand. University of Auckland, University of Auckalnd. 897 p.
Myers, R.A.; Worm, B. (2003). Rapid worldwide depletion of predatory fish communities. Nature 423: 280-283.
Pauly, D.; Christensen, V.; Dalsgaard, J.; Froese, R.; Torres, F.J. (1998). Fishing down marine food webs. Science 279: 860-863.
Petchey, F.J. (1999). New Zealand bone dating revisited: A radiocarbon discard protocol for bone. New Zealand Journal of Archaeology 19: 81-124.
Pool, I. (1991). Te Iwi Maori: A New Zealand Population, Past, Present and Future. Auckland University Press, Auckland.
Reeves, R.R.; Smith, T.D. (2006). A taxonomy of world whaling: operations and eras. In: Estes, J.A., D.P. DeMaster, D.F. Doak, T.M. Williams and R.L.J. Brownell (eds) Whales, Whaling, and Ocean Ecosystems. Berkeley, University of California Press. 82-101.
Reitz, E.J.; Wing, E.S. (2008). Zooarchaeology. Cambridge University Press, Cambridge. 533 p.
Rick, T.C.; Erlandson, J.M. (2008). Human Impacts of Ancient Marine Ecosystems: A Global Perspective. University of California Press, Berkley. 336 p.

Schmidt, M.D. (2000a). Radiocarbon Dating New Zealand Prehistory Using Marine Shell. BAR International Series, Oxford.
Schmidt, M.D. (2000b). Radiocarbon dating the end of moa-hunting in New Zealand prehistory. Archaeology in New Zealand 43: 314-329.
Smith, I.W.G. (1985). Sea mammal hunting and prehistoric subsistence in New Zealand. PhD thesis, Anthropology, University of Otago. 576 p.
Smith, I.W.G. (1989). Maori impact on the marine megafauna: pre-European distributions of New Zealand sea mammals. In: Sutton, D.G. (ed) Saying So Doesn't Make It So: Papers in Honour of B. Foss Leach. Dunedin, New Zealand Archaeological Association Monograph 17. 76108.

Smith, I.W.G. (2004). Nutritional perspectives on prehistoric marine fishing in New Zealand. New Zealand Journal of Archaeology 24: 5-31.
Smith, I.W.G. (2005). Retreat and resilience: Fur seals and human settlement in New Zealand. In: Monks, G.G. (ed) The Exploitation and Cultural Importance of Marine Mammals. Oxford, Oxbow Books. 6-18.
Smith, I.W.G. (2010). Protocols for organising radiocarbon dated assemblages from New Zealand archaeological sites for comparative analysis. Journal of Pacific Archaeology 1: 184-187.
Smith, I.W.G. (2011). Meat Weights and Nutritional Yield Values for New Zealand Archaeofauna. Otago Archaeological Laboratory Report, No. 8. 23 p.
Smith, I.W.G. (in press). Pre-European Maori exploitation of marine resources in two New Zealand case study areas: species range and temporal change. Journal of the Royal Society of New Zealand
Smith, I.W.G. (n.d.). Regional and temporal variations in energy harvests from prehistoric fauna in New Zealand. Manuscript in preparation
Smith, I.W.G.; James-Lee, T. (2010). Data for an Archaeozoological Analysis of Marine Resource Use in Two New Zealand Study Areas (revised edition). Otago Archaeological Laboratory Report, No 7.85 p.
Spencer, H.G.; Willan, R.C.; Marshall, B.; Murray, T.J. (2009). Checklist of the Recent Mollusca recorded from the New Zealand Exclusive Economic Zone. http://www.molluscs.otago.ac.nz/index.html.
Urlich, D.U. (1969). The Distribution and Migrations of the North Island Maori Population about 1800-1840. MA thesis, Geography, University of Auckland,
Walter, R.K.; Smith, I.W.G.; Jacomb, C. (2006). Sedentism, subsistence and socio-political organization in prehistoric New Zealand. World Archaeology 38: 274-290.
Ward, G.K.; Wilson, S.R. (1978). Procedures for comparing and combining radiocarbon agedeterminations - critique. Archaeometry 20: 19-31.
Wilmshurst, J.M.; Hunt, T.L.; Lipo, C.P.; Anderson, A.J. (2011). High precision radiocarbon dating shows recent rapid initial human colonisation of east Polynesia. Proceedings of the National Academy of Sciences 108: 1815-1820.
Worthy, T.H. (1998). Quaternary fossil faunas of Otago, South Island, New Zealand. Journal of the Royal Society of New Zealand 28: 421-521.

Table 1: $\quad$ Number of study assemblages per area and period.

| Period | Greater Hauraki | Otago-Catlins | Total |
| :--- | ---: | ---: | ---: |
| Early | 8 | 10 | 18 |
| Early/Middle | 11 | 9 | 20 |
| Middle | 25 | 2 | 27 |
| Middle/Late | 18 | 2 | 20 |
| Late | 13 | 9 | 22 |
| Total | 75 | 32 | 107 |

Table 2: Faunal component energy proportions (\% total Kcal from animals): initial calculated values and final adjusted values.

| Area/period | Value | Shellfish | Fish | Marine/coastal bird | Marine mammal | Moa | Terrestrial bird | Terrestrial mammal |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Greater Hauraki |  |  |  |  |  |  |  |  |
| Early | initial | 0.66 | 11.50 | 3.39 | 59.77 | 21.10 | 1.46 | 2.11 |
| Early | final | 0.62 | 10.74 | 4.16 | 54.80 | 23.38 | 2.76 | 3.54 |
| Middle | initial | 43.13 | 51.72 | 0 | 0 | 0 | 0 | 5.15 |
| Middle | final | 40.11 | 48.19 | 1.50 | 0.05 | 0 | 1.50 | 8.65 |
| Late | initial | 51.95 | 43.16 | 1.05 | 0 | 0 | 0.51 | 3.33 |
| Late | final | 44.50 | 44.12 | 2.50 | 0.05 | 0 | 2.67 | 6.17 |
| Otago-Catlins |  |  |  |  |  |  |  |  |
| Early | initial | 0.61 | 5.81 | 5.92 | 57.21 | 27.05 | 0.77 | 2.63 |
| Early | final | 0.57 | 5.39 | 6.49 | 52.10 | 29.49 | 1.90 | 4.06 |
| Middle | initial | 3.38 | 59.07 | 8.42 | 21.10 | 3.15 | 1.95 | 2.93 |
| Middle | final | 3.20 | 54.36 | 9.46 | 19.95 | 4.36 | 4.66 | 4.00 |
| Late | initial | 1.89 | 57.36 | 17.73 | 12.00 | 0 | 2.25 | 8.77 |
| Late | final | 4.28 | 52.49 | 16.22 | 10.98 | 0 | 4.77 | 11.25 |

Table 3: Maximum, minimum and best estimate values for variables used in estimating total energy from animals.

| Area/year | Human population size (n) |  |  | Energy/person/day (kcal) |  |  | Proportion from animals |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Min | Best | Max | Min | Best | Max | Min | Best | Max |
| Greater Hauraki |  |  |  |  |  |  |  |  |  |
| ca. 1400 | 500 | 2000 | 2500 | 1800 | 2150 | 2172 | 0.53 | 0.66 | 0.78 |
| ca. 1550 | 6100 | 10200 | 11700 | 1800 | 2150 | 2172 | 0.40 | 0.53 | 0.66 |
| ca. 1750 | 10800 | 12150 | 13500 | 1800 | 2150 | 2172 | 0.40 | 0.53 | 0.66 |
| Otago-Catlins |  |  |  |  |  |  |  |  |  |
| ca. 1400 | 500 | 1800 | 2500 | 2150 | 2172 | 2193 | 0.66 | 0.78 | 0.90 |
| ca. 1550 | 500 | 1800 | 2600 | 2150 | 2172 | 2193 | 0.66 | 0.78 | 0.90 |
| ca. 1750 | 1600 | 1800 | 2000 | 2150 | 2172 | 2193 | 0.66 | 0.78 | 0.90 |

Table 4: $\quad \begin{aligned} & \text { Estimated total energy (Kcal } \mathrm{x} 10^{6} \text { ) harvested from animals during focal years in each } \\ & \text { study area. }\end{aligned}$

| Area/year | Minimum estimate | Best estimate | Maximum estimate |
| :--- | ---: | ---: | ---: |
| Greater Hauraki |  |  |  |
| ca. 1400 | 174.11 | 1035.87 | 1545.92 |
| ca. 1550 | 1603.08 | 4242.36 | 6121.85 |
| ca. 1750 | 2838.24 | 5053.40 | 7063.67 |
| Otago-Catlins |  |  |  |
| ca. 1400 | 258.97 | 1113.06 | 1801.00 |
| ca. 1550 | 388.45 | 1236.74 | 1873.04 |
| ca. 1750 | 828.70 | 1113.06 | 1440.80 |






Minimum estimate

Estimated numbers (N) and biomass (tonnes) of fish harvested in Greater Hauraki at ca. 1400 AD. Carcharhiniforme ?sp Elasmobranchii ?sp Callorhinchus milii Anguilla spp.
Pseudophycis bachus Chelidonichthys kumu Pseudocaranx dentex Trachurus declivis Trachurus novaezelandiae Arripis trutta Pagrus auratus Nemadactylus macropterus Latridopsis ciliaris An Aldrichetta forsteri Labridae Parapercis colias Thyrsites atun Serollela brama Meuschenia scaber TOTAL

|  |  |
| :---: | :---: |







| Maximum estimate |  |
| ---: | ---: |
| N | Biomass |
| 6 | 0.055 |
| 6 | 0.110 |
| 7356 | 11.033 |
| 6 | 0.006 |
| 2517 | 12.587 |
| 22 | 0.009 |
| 225 | 0.225 |
| 3 | 0.002 |
| 622 | 12.444 |
| 3 | 0.004 |
| 112 | 0.112 |
| 1658 | 3.649 |
| 236 | 0.189 |
| 192 | 0.517 |
| 1326 | 3.979 |
| 6 | 0.001 |
| 1474 | 2.211 |
| 5214 | 3.650 |
| 3644 | 5.830 |
| 30414 | 69.952 |
| 17 | 0.050 |
| 6 | 0.022 |

$\pm 30 \%$
0.010
0.020
2.046
0.001
2.334
0.002
0.042
0.000
2.307
0.001
0.021
0.676
0.035
0.096
0.738
0.000
0.410
0.677
1.081
12.970
0.009
0.004

$\pm 30 \%$
0.002
0.005
0.476
0.000
0.543
0.000
0.010
0.000
0.537
0.000
0.005
0.157
0.008
0.022
0.172
0.000
0.095
0.157
0.252
3.018
0.002
0.001
5.463

| Minimum estimate |  |
| ---: | :---: |
| N | Biomass |
| 1 | 0.008 |
| 1 | 0.016 |
| 1058 | 1.587 |
| 1 | 0.001 |
| 362 | 1.810 |
| 3 | 0.001 |
| 32 | 0.032 |
| 0 | 0.000 |
| 89 | 1.789 |
| 1 | 0.001 |
| 16 | 0.016 |
| 238 | 0.525 |
| 34 | 0.027 |
| 28 | 0.074 |
| 191 | 0.572 |
| 1 | 0.000 |
| 212 | 0.318 |
| 750 | 0.525 |
| 524 | 0.838 |
| 4373 | 10.058 |
| 2 | 0.007 |
| 1 | 0.003 |
| 7917 | 18.209 |


| Maximum estimate |  |
| ---: | ---: |
| N | Biomass |
| 1718 | 17.180 |
| 143977 | 215.965 |
| 22943 | 114.716 |
| 222 | 0.089 |
| 222 | 0.222 |
| 3325 | 3.325 |
| 222 | 0.421 |
| 11860 | 237.190 |
| 333 | 0.266 |
| 4267 | 12.802 |
| 33861 | 50.791 |
| 28817 | 20.172 |
| 38017 | 60.827 |
| 264401 | 608.121 |
| 554183 | 1342.086 |554183


|  |  |
| :---: | :---: |
| z |  |$\underset{\sim}{\stackrel{7}{\sim}}$0

0
U
i

+| Maximum estimate |  |
| ---: | ---: |
| N | Biomass |
| 1718 | 17.180 |
| 143977 | 215.965 |
| 22943 | 114.716 |
| 222 | 0.089 |
| 222 | 0.222 |
| 3325 | 3.325 |
| 222 | 0.421 |
| 11860 | 237.190 |
| 333 | 0.266 |
| 4267 | 12.802 |
| 33861 | 50.791 |
| 28817 | 20.172 |
| 38017 | 60.827 |
| 264401 | 608.121 |
| 554183 | 1342.086 |n$\hat{0}$

$i$
$i$

| Minimum estimate |  |  |
| :---: | :---: | :---: |
| $\mathrm{N} \quad$ Biomass |  |  |

2.375
29.859
15.861
0.012
0.031
0.460
0.058
32.794
0.037
1.770
7.022
2.789
8.410
84.079
185.557
76622



 $\xrightarrow[N]{\underset{\sim}{i}}$
Estimated numbers (N) and biomass (tonnes) of fish harvested in Otago-Catlins at ca. 1750 AD.
Table 10:

| N |  |  |
| ---: | ---: | ---: |
|  | Biomass | $\pm 30 \%$ |
| 170 | 1.702 | 0.511 |
| 170 | 0.715 | 0.214 |
| 33928 | 50.891 | 15.267 |
| 8461 | 42.303 | 12.691 |
| 8597 | 171.936 | 51.581 |
| 51 | 0.138 | 0.041 |
| 1021 | 1.532 | 0.460 |
| 221 | 0.155 | 0.046 |
| 117614 | 270.513 | 81.154 |
| 170233 | 539.885 | 161.965 |

Estimated numbers ( $\mathbf{N} \times 10^{\mathbf{3}}$ ) and biomass (tonnes) of shellfish harvested in Greater Hauraki at ca. 1400 AD.
Table 11:

| Best estimate |  | Maximum estimate |  |  |  |
| ---: | ---: | ---: | ---: | ---: | ---: |
| $\mathrm{N} \mathrm{x10}^{3}$ | Biomass | $\pm 30 \%$ | $\mathrm{~N} \mathrm{x10}^{3}$ | Biomass | $\pm 30 \%$ |
| 10.52 | 0.0316 | 0.0095 | 15.71 | 0.0471 | 0.0141 |
| 234.14 | 0.9366 | 0.2810 | 349.43 | 1.3977 | 0.4193 |
| 120.27 | 0.2405 | 0.0722 | 179.50 | 0.3590 | 0.1077 |
| 40.50 | 0.0405 | 0.0121 | 60.44 | 0.0604 | 0.0181 |
| 134.94 | 0.2699 | 0.0810 | 201.39 | 0.4028 | 0.1208 |
| 38.32 | 0.0766 | 0.0230 | 57.19 | 0.1144 | 0.0343 |
| 16.11 | 0.0322 | 0.0097 | 24.05 | 0.0481 | 0.0144 |
| 178.59 | 0.3572 | 0.1072 | 266.52 | 0.5330 | 0.1599 |
| 104.40 | 0.2088 | 0.0626 | 155.81 | 0.3116 | 0.0935 |
| 14.86 | 2.2297 | 0.6689 | 22.18 | 3.3275 | 0.9983 |
| 17.24 | 0.0345 | 0.0103 | 25.72 | 0.0514 | 0.0154 |
| 7.17 | 0.1076 | 0.0323 | 10.71 | 0.1606 | 0.0482 |
| 298.68 | 1.1947 | 0.3584 | 445.75 | 1.7830 | 0.5349 |
| 39.20 | 0.0392 | 0.0118 | 58.50 | 0.0585 | 0.0175 |
| 10.94 | 0.0109 | 0.0033 | 16.32 | 0.0163 | 0.0049 |
| 22.52 | 0.0450 | 0.0135 | 33.61 | 0.0672 | 0.0202 |
| 8.58 | 0.0086 | 0.0026 | 12.81 | 0.0128 | 0.0038 |
| 12.31 | 0.0246 | 0.0074 | 18.37 | 0.0367 | 0.0110 |
| 20.57 | 0.3085 | 0.0926 | 30.70 | 0.4605 | 0.1381 |
| 15.13 | 0.2270 | 0.0681 | 22.58 | 0.3387 | 0.1016 |
| 8.52 | 0.0085 | 0.0026 | 12.71 | 0.0127 | 0.0038 |
| 56.77 | 0.0568 | 0.0170 | 84.72 | 0.0847 | 0.0254 |
| 56.31 | 0.0639 | 0.0192 | 84.04 | 0.0954 | 0.0286 |
| 1466.61 | 6.5535 | 1.9660 | 2188.75 | 9.7803 | 2.9341 |


| Minimum estimate |  |  |
| :---: | :---: | :---: |
| $\mathrm{N} \times 10^{3} \quad$ Biomass |  |  |



| Maximum estimate |  |  |
| ---: | ---: | ---: |
| $\mathrm{N} \mathrm{x10}$ | Biomass | $\pm 30 \%$ |
| 438.5 | 1.754 | 0.526 |
| 1286.2 | 1.286 | 0.386 |
| 414.3 | 0.829 | 0.249 |
| 1276.7 | 1.277 | 0.383 |
| 1396.3 | 1.396 | 0.419 |
| 1037.0 | 1.037 | 0.311 |
| 1595.7 | 1.596 | 0.479 |
| 516872.4 | 516.872 | 155.062 |
| 77350.1 | 154.700 | 46.410 |
| 795989.3 | 1591.979 | 477.594 |
| 1866.5 | 3.733 | 1.120 |
| 22218.9 | 44.438 | 13.331 |
| 31420.9 | 125.684 | 37.705 |
| 939.9 | 0.940 | 0.282 |
| 4118.7 | 8.237 | 2.471 |
| 1223.0 | 1.223 | 0.367 |
| 769.4 | 1.539 | 0.462 |
| 4203.5 | 4.204 | 1.261 |
| 2407.3 | 4.815 | 1.444 |
| 3553.6 | 3.554 | 1.066 |
| 621.4 | 1.864 | 0.559 |
| 23568.2 | 23.568 | 7.070 |
| 2824.8 | 9.061 | 2.119 |
| 1503182.3 | 2505.585 | 751.675 |



| Best estimate |  |
| ---: | ---: |
| $\mathrm{N} \mathrm{x} 10^{3}$ | Biomass |
| 303.9 | 1.216 |
| 891.4 | 0.891 |
| 287.1 | 0.574 |
| 884.7 | 0.885 |
| 967.6 | 0.968 |
| 718.6 | 0.719 |
| 1105.8 | 1.106 |
| 358185.7 | 358.186 |
| 53602.6 | 107.205 |
| 551610.0 | 1103.220 |
| 1293.5 | 2.587 |
| 15397.4 | 30.795 |
| 21774.3 | 87.097 |
| 651.3 | 0.651 |
| 2854.2 | 5.708 |
| 847.5 | 0.847 |
| 533.2 | 1.066 |
| 2913.0 | 2.913 |
| 1668.3 | 3.337 |
| 2462.6 | 2.463 |
| 430.6 | 1.292 |
| 16332.5 | 16.332 |
| 1957.6 | 6.279 |
| 041685.3 | 1736.337 |



| Minimum estimate |  |
| ---: | ---: |
| $\mathrm{N} \mathrm{x} 10^{3}$ | Biomass |
| 114.8 | 0.459 |
| 336.8 | 0.337 |
| 108.5 | 0.217 |
| 334.3 | 0.334 |
| 365.6 | 0.366 |
| 271.5 | 0.272 |
| 417.9 | 0.418 |
| 135349.3 | 135.349 |
| 20255.1 | 40.510 |
| 208439.5 | 416.879 |
| 488.8 | 0.978 |
| 5818.3 | 11.637 |
| 8228.0 | 32.912 |
| 246.1 | 0.246 |
| 1078.5 | 2.157 |
| 320.2 | 0.320 |
| 201.5 | 0.403 |
| 1100.7 | 1.101 |
| 630.4 | 1.261 |
| 930.5 | 0.931 |
| 162.7 | 0.488 |
| 6171.6 | 6.172 |
| 739.7 | 2.373 |
| 393626.5 | 656.118 |








| $\begin{aligned} & \text { 若 } \\ & \frac{0}{n} \\ & \frac{0}{x} \\ & z \end{aligned}$ |  <br>  |
| :---: | :---: |


| $\pm$ |
| :--- |
|  |
| - |
| $n$ |
| $n$ |
| $\vdots$ |


| $\begin{aligned} & \text { ò } \\ & \text { ò } \\ & \text { + } \end{aligned}$ | N ㄱ⼦ 0000000000000000000000 |
| :---: | :---: |
|  | 人 눈 O O |
| $\begin{aligned} & 3 \\ & \frac{0}{x} \\ & z \end{aligned}$ |  |



Minimum estimate $\begin{array}{rr}\mathrm{N} \times 10^{3} & \text { Biomass } \\ 116.9 & 0.351 \\ 11.5 & 0.046 \\ 0.4 & 0.001 \\ 1.0 & 0.003 \\ 298.0 & 0.298 \\ 284.8 & 0.570 \\ 1.2 & 0.001 \\ 2.0 & 0.004 \\ 0.4 & 0.001 \\ 7.6 & 0.015 \\ 1.7 & 0.002 \\ 3.5 & 0.525 \\ 0.7 & 0.069 \\ 24.3 & 0.049 \\ 1.0 & 0.015 \\ 4.2 & 0.062 \\ 4.1 & 0.017 \\ 2.6 & 0.003 \\ 1.0 & 0.002 \\ 0.4 & 0.000 \\ 78.6 & 0.079 \\ 1.0 & 0.001 \\ 4.0 & 0.004 \\ 850.9 & 2.116\end{array}$


| Maximum estimate |  |
| ---: | ---: |
| $\mathrm{N} \mathrm{x} 10^{3}$ | Biomass |
| 9012.1 | 27.036 |
| 181.6 | 0.727 |
| 27.2 | 0.054 |
| 18.7 | 0.037 |
| 263.2 | 0.263 |
| 2908.0 | 2.908 |
| 107.0 | 0.214 |
| 1949.6 | 3.899 |
| 93.8 | 0.188 |
| 25.8 | 0.026 |
| 83.0 | 0.083 |
| 159.0 | 23.857 |
| 7.1 | 0.715 |
| 107.6 | 0.108 |
| 315.1 | 0.630 |
| 25.6 | 0.026 |
| 91.7 | 1.376 |
| 100.5 | 1.507 |
| 255.6 | 1.023 |
| 22.8 | 0.023 |
| 7068.5 | 7.069 |
| 28.6 | 0.029 |
| 197.8 | 0.579 |
| 23 | 72.375 |



| Best estimate |  |
| ---: | ---: |
| $\mathrm{N} \times 10^{3}$ | Biomass |
| 5355.5 | 16.066 |
| 107.9 | 0.432 |
| 16.2 | 0.032 |
| 11.1 | 0.022 |
| 156.4 | 0.156 |
| 1728.1 | 1.728 |
| 63.6 | 0.127 |
| 1158.5 | 2.317 |
| 55.7 | 0.111 |
| 15.3 | 0.015 |
| 49.3 | 0.049 |
| 94.5 | 14.177 |
| 4.2 | 0.425 |
| 64.0 | 0.064 |
| 187.2 | 0.374 |
| 15.2 | 0.015 |
| 54.5 | 0.818 |
| 59.7 | 0.896 |
| 151.9 | 0.608 |
| 13.6 | 0.014 |
| 4200.5 | 4.201 |
| 17.0 | 0.017 |
| 117.5 | 0.344 |
| 1399.6 | 43.009 |


Minimum estimate
$\begin{array}{rc}\mathrm{N} \mathrm{x10} & \text { Biomass } \\ 1246.0 & 3.738 \\ 25.1 & 0.100 \\ 3.8 & 0.008 \\ 2.6 & 0.005 \\ 36.4 & 0.036 \\ 402.1 & 0.402 \\ 14.8 & 0.030 \\ 269.5 & 0.539 \\ 13.0 & 0.026 \\ 3.6 & 0.004 \\ 11.5 & 0.011 \\ 22.0 & 3.298 \\ 1.0 & 0.099 \\ 14.9 & 0.015 \\ 43.6 & 0.087 \\ 3.5 & 0.004 \\ 12.7 & 0.190 \\ 13.9 & 0.208 \\ 35.3 & 0.141 \\ 3.2 & 0.003 \\ 977.3 & 0.977 \\ 4.0 & 0.004 \\ 27.3 & 0.080 \\ 3186.9 & 10.007\end{array}$
Table 15: Estimated numbers ( $\mathbf{N} \times 10^{3}$ ) and biomass (tonnes) of shellfish harvested in Otago-Catlins at ca. 1550 AD




|  |  |
| :---: | :---: |
| $\begin{aligned} & 3 \\ & \frac{0}{x} \\ & z \end{aligned}$ |  |


|  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |  |


| Minimum estimate |  |
| ---: | ---: |
| $\mathrm{N} \times 10^{3}$ | Biomass |
| 2032.6 | 6.098 |
| 69.1 | 0.276 |
| 13.0 | 0.026 |
| 99.0 | 0.099 |
| 5.5 | 0.011 |
| 34.6 | 0.035 |
| 33.7 | 0.067 |
| 5.0 | 0.020 |
| 109.1 | 0.218 |
| 427.1 | 0.854 |
| 467.6 | 0.468 |
| 174.4 | 26.164 |
| 144.4 | 2.166 |
| 2.5 | 0.003 |
| 16.7 | 0.250 |
| 244.3 | 0.977 |
| 4.7 | 0.009 |
| 511.7 | 0.512 |
| 4.0 | 0.004 |
| 4398.9 | 38.257 |


| Best estimate |  | Maximum estimate |  |  |  |
| ---: | ---: | ---: | ---: | ---: | ---: |
| N | Biomass | $\pm 30 \%$ | N | Biomass | $\pm 30 \%$ |
| 4248 | 4.673 | 1.402 | 6340 | 6.974 | 2.092 |
| 562 | 2.530 | 0.759 | 839 | 3.775 | 1.133 |
| 177 | 0.798 | 0.239 | 264 | 1.190 | 0.357 |
| 682 | 0.341 | 0.102 | 1018 | 0.509 | 0.153 |
| 306 | 0.061 | 0.018 | 457 | 0.091 | 0.027 |
| 1054 | 0.844 | 0.253 | 1574 | 1.259 | 0.378 |
| 1944 | 0.583 | 0.175 | 2901 | 0.870 | 0.261 |
| 854 | 0.171 | 0.051 | 1274 | 0.255 | 0.076 |
| 1076 | 0.140 | 0.042 | 1606 | 0.209 | 0.063 |
| 45 | 0.103 | 0.031 | 67 | 0.154 | 0.046 |
| 288 | 0.202 | 0.061 | 430 | 0.301 | 0.090 |
| 609 | 1.339 | 0.402 | 908 | 1.998 | 0.600 |
| 1139 | 2.277 | 0.683 | 1699 | 3.398 | 1.019 |
| 3305 | 3.966 | 1.190 | 4932 | 5.918 | 1.775 |
| 23 | 0.002 | 0.001 | 35 | 0.003 | 0.001 |
| 23 | 0.003 | 0.001 | 35 | 0.005 | 0.002 |
| 23 | 0.001 | 0.000 | 35 | 0.002 | 0.001 |
| 1199 | 1.019 | 0.306 | 1790 | 1.521 | 0.456 |
| 23 | 0.006 | 0.002 | 35 | 0.009 | 0.003 |
| 149 | 0.104 | 0.031 | 222 | 0.155 | 0.047 |
| 149 | 0.012 | 0.004 | 222 | 0.018 | 0.005 |
| 23 | 0.004 | 0.001 | 35 | 0.006 | 0.002 |
| 17902 | 19.178 | 5.753 | 26716 | 28.620 | 8.586 |

Minimum estimate
$\mathrm{N} \quad$ Biomass


Eudyptula minor Diomedeidae ?sp Macronectes halli Pterodroma macroptera Pachyptila vitata Puffinus griseus Puffinus gavia Puffinus assimilis

Phalacrocorax melanoleucos Phalacrocorax carbo Phalacrocorax varius Stictocarbo punctatus Calidras canutus rogersi Charadrius obscurus Anarhyncus frontalis Larus dominicanus Larus novaehollandiae Hydroprogne caspia Childonias albostriata Sterna striata

TOTAL


| Maximum estimate |  |
| ---: | ---: |
| N | Biomass |
| 6694 | 7.364 |
| 5651 | 25.430 |
| 3141 | 1.571 |
| 1413 | 0.989 |
| 3405 | 4.086 |
| 1413 | 1.201 |
| 1413 | 0.226 |
| 23132 | 40.867 |

Estimated numbers (N) and biomass (tonnes) of marine and coastal birds harvested in Greater Hauraki at ca. 1750 AD.
Minimum estimate
$\pm 30 \%$
0.578
1.998
0.123
0.078
0.321
0.094
0.018
3.210
$6057 \quad 10.701 \quad 3.210$
1753
1480

| Best estimate |  |
| ---: | ---: |
| N | Biomass |
| 4639 | 5.103 |
| 3916 | 17.622 |
| 2177 | 1.088 |
| 979 | 0.686 |
| 2360 | 2.832 |
| 979 | 0.833 |
| 979 | 0.157 |
| 16030 | 28.320 |

        \(\pm 30 \%\)
    1.531
5.287
0.327
0.206
0.849
0.250
0.047
8.49628.320
N Biomass
7531.928
$\begin{array}{rr}1480 & 6.659 \\ 823 & 0.411 \\ 370 & 0.259 \\ 892 & 1.070 \\ 370 & 0.315 \\ 370 & 0.059 \\ 6057 & 10.701\end{array}$
Eudyptula minor
Diomedeidae?sp
Pterodroma macroptera
Pterodroma macroptera
Phalacrocorax melanoleucos
Stictocarbo punctatus
Larus dominicanus
Sterna striata
TOTAL
Table 19:
Estimated numbers (N) and biomass (tonnes) of marine and coastal birds harvested in Greater Hauraki at ca. 1750 AD


| Maximum estimate |  |
| ---: | ---: |
| N | Biomass |
| 10929 | 12.022 |
| 5465 | 24.591 |
| 10929 | 5.465 |
| 16394 | 4.918 |
| 5465 | 0.710 |
| 5465 | 1.869 |
| 5465 | 3.825 |
| 16394 | 19.673 |
| 5465 | 4.645 |
| 5465 | 0.874 |
| 87435 | 78.593 |

                        40
                                \(\pm 30 \%\)
    1.449
2.964
0.659
0.593
0.086
0.225
0.461
2.371
0.560
0.105
9.474
Minimum estimate
$\mathrm{N} \quad$ Biomass

16.868

| Best estimate |  |
| ---: | ---: |
| N | Biomass |
| 7819 | 8.601 |
| 3909 | 17.593 |
| 7819 | 3.909 |
| 11728 | 3.519 |
| 3909 | 0.508 |
| 3909 | 1.337 |
| 3909 | 2.737 |
| 11728 | 14.074 |
| 3909 | 3.323 |
| 3909 | 0.626 |
| 62552 | 56.226 |

## Table 20: <br> <br> 

 <br> <br> }Estimated numbers (N) and biomass (tonnes) of marine and coastal birds harvested in Otago-Catlins at ca. 1400 AD.




Minimum estimate
$\mathrm{N} \quad$ Biomass



Estimated numbers (N) and biomass (tonnes) of marine and coastal birds harvested in Otago-Catlins at ca. 1550 AD.

| Maximum estimate |  |
| ---: | ---: |
| N | Biomass |
| 2456 | 9.825 |
| 2192 | 11.509 |
| 8085 | 8.893 |
| 2564 | 11.537 |
| 1724 | 0.215 |
| 1804 | 1.444 |
| 2526 | 0.758 |
| 1373 | 0.961 |
| 1266 | 0.057 |
| 7024 | 0.913 |
| 1546 | 1.082 |
| 1880 | 4.136 |
| 1691 | 3.383 |
| 2564 | 6.410 |
| 13826 | 16.592 |
| 1341 | 1.140 |




| Minimum estimate |  |  |
| ---: | :---: | ---: |
| N | Biomass | $\pm 30 \%$ |
| 340 | 1.358 | 0.408 |
| 303 | 1.591 | 0.477 |
| 1118 | 1.230 | 0.369 |
| 354 | 1.595 | 0.479 |
| 238 | 0.030 | 0.009 |
| 249 | 0.200 | 0.060 |
| 349 | 0.105 | 0.031 |
| 190 | 0.133 | 0.040 |
| 175 | 0.008 | 0.002 |
| 971 | 0.126 | 0.038 |
| 214 | 0.150 | 0.045 |
| 260 | 0.572 | 0.172 |
| 234 | 0.468 | 0.140 |
| 354 | 0.886 | 0.266 |
| 1912 | 2.294 | 0.688 |
| 185 | 0.158 | 0.047 |
| 7447 | 10.902 | 3.271 |


| Maximum estimate |  |
| ---: | ---: |
| N | Biomass |
| 553 | 2.212 |
| 2212 | 11.614 |
| 1026 | 8.212 |
| 704 | 5.631 |
| 1127 | 5.072 |
| 121 | 0.024 |
| 5916 | 26.621 |
| 1433 | 0.186 |
| 1793 | 0.359 |
| 553 | 0.166 |
| 314 | 0.014 |
| 863 | 0.112 |
| 1592 | 3.503 |
| 624 | 1.249 |
| 7835 | 19.586 |
| 10637 | 12.765 |
| 2828 | 6.222 |
| 1764 | 0.459 |
| $\mathbf{4 1 8 9 6}$ | 104.004 |



$\pm 30 \%$
0.382
2.004
1.417
0.972
0.875
0.004
4.593
0.032
0.062
0.029
0.002
0.019
0.604
0.215
3.380
2.203
1.074
0.079
Minimum estimate
$\mathrm{N} \quad$ Biomass
Biomass
1.272
6.680
4.723
3.239
2.917
0.014
15.311
0.107
0.206
0.095
0.008
0.065
2.015
0.718
11.265
7.342
3.578
0.264
59.819

318
1272
590
405
648
70
3403
824
1031
318
181
496
916
359
4506
6118
1627
1014
Table 23: Estimated numbers (N) and biomass (tonnes) of marine mammals harvested in Greater Hauraki at ca. 1400 AD.



Estimated numbers (N) and biomass (tonnes) of marine mammals harvested in Otago-Catlins at ca. 1400 AD.
$\pm 30 \%$

0.706
15.839
24.640
5.283
24.640
71.109
37.849
1.580
0.197
0.667
111.402

|  |  |  |
| :---: | :---: | :---: |
| \% |  |  |
| z |  | $\bar{I}^{n n}$ |

Table 27: Estimated numbers (N) and biomass (tonnes) of marine mammals harvested in Otago-Catlins at ca. 1550 AD.

| Maximum estimate |  |  |
| ---: | ---: | ---: |
| N | Biomass | $\pm 30 \%$ |
|  |  |  |
| 162 | 1.390 | 0.417 |
| 1555 | 38.869 | 11.661 |
| 609 | 60.867 | 18.260 |
| 218 | 10.911 | 3.273 |
| 412 | 61.751 | 18.525 |
| 2955 | 173.788 | 52.136 |
| 557 | 62.666 | 18.800 |
| 4 | 0.442 | 0.133 |
| 4 | 4.810 | 1.443 |
| 18 | 1.494 | 0.448 |
| 3537 | 243.201 | 72.960 |

Estimated numbers (N) and biomass (tonnes) of marine mammals harvested in Otago-Catlins at ca. $\mathbf{1 7 5 0}$ AD.

| Maximum estimate |  |
| ---: | :---: |
| N | Biomass |
|  |  |
| 964 | 24.093 |
| 378 | 37.843 |
| 252 | 37.763 |
| 1594 | 99.700 |
| 42 | 56.979 |
| 8 | 0.677 |
| 1644 | 157.356 |



| N | Biomass | $\pm 30 \%$ |
| ---: | :---: | ---: |
|  |  |  |
| 555 | 13.880 | 4.164 |
| 218 | 21.802 | 6.541 |
| 145 | 21.756 | 6.527 |
| 918 | 57.437 | 17.231 |
| 24 | 32.772 | 9.832 |
| 5 | 0.390 | 0.117 |
| 947 | 90.600 | 27.180 | $947 \quad 90.600 \quad 27.180$

[^0]
## Table 29: Taxa harvested at $c a .1750 \mathrm{AD}$ as a proportion of taxa harvested at $c a .1400 \mathrm{AD}$.

|  | Greater Hauraki | Otago-Catlins |
| :--- | ---: | ---: |
| Fish | 0.68 | 0.41 |
| Shellfish | 0.43 | 0.53 |
| Marine and coastal bird | 0.45 | 0.72 |
| Marine mammal | 0.20 | 0.60 |



Figure 1: Greater Hauraki and Otago-Catlins, showing location of study sites and other pre-European Maori sites.


Figure 2: Simplified model of analytical procedure.


Figure 3: Procedures used in determining energy yield per species in study assemblages.



Figure 4: (A) Estimated growth rate for total New Zealand population based on cumulative frequency of radiocarbon dates (after McFadgen et al. 1994: Figure 4) showing proportion of population at ca. 1550 AD. (B) Frequency of radiocarbon dated assemblages in Otago-Catlins study dataset.


Figure 5: Minimum, best and maximum estimates for biomass of snapper harvested in Greater Hauraki.


Figure 6: Best estimates for biomass harvested from four classes of marine fauna in each study area
APPENDIX 1 DATA INPUTS FOR MARINE ANIMAL SPECIES
The following tables list for each taxon at each of three focal dates in each study area (1) its frequency as a proportion of all animals in the faunal classes to which it belongs (taxon frequency, as defined in section 2.3.2), and (2) its proportional contribution to total energy derived from all animal foods (proportional energy contribution as defined in section 2.3.5).
Fish - Greater Hauraki
 frequency contribution
0.2231
0.0315
0.1443

$\stackrel{\circ}{\circ}$ | $\square$ |
| :--- |
| 0 |
| 0 | . 8

0
0
0 N 9
8
8
0
0 $\stackrel{3}{8}$

| $\ddagger 8$ |
| :--- |
| $\underset{\sim}{2}$ |
| $\underset{\sim}{i}$ |
|  | Ň

N゙
ก゙ ca 1550 AD
 Taxon
frequency
N
1.33
$\cdots \underset{\substack{3}}{\substack{0}}$
$\pm \infty$
$\circ$
0
0
ca 1750 AD

|  |  | $\begin{aligned} & \infty \\ & 0 \\ & 0 \\ & 0 \\ & 0 \end{aligned}$ |  |  |
| :---: | :---: | :---: | :---: | :---: |
|  | $\stackrel{\infty}{\infty} \stackrel{\sim}{n} \underset{\sim}{\infty}$ | $\stackrel{\infty}{+}$ | $\begin{array}{ccc} \infty \\ \stackrel{\infty}{\circ} \\ \underset{\sim}{\infty} & \infty \\ + \\ \hline \end{array}$ | $\begin{array}{ll} \infty \\ \stackrel{y}{\circ} \\ \stackrel{y}{7} & \circ \\ \hline \end{array}$ |

$\stackrel{\rightharpoonup}{N}$
$\stackrel{\rightharpoonup}{0}$
$\circ$
$i$
$i$
$\left.\begin{array}{rr}\text { ca } 1750 \mathrm{AD} \\ \text { Taxon } \\ \text { Proquertional } \\ \text { energy }\end{array}, \begin{array}{r}\text { contribution }\end{array}\right\}$

| ca 1550 AD |  |
| :---: | :---: |
| Taxon frequency | Proportional energy contribution |
| 1.33 | 0.4446 |
| 2.21 | 0.0971 |
| 1.33 | 0.3809 |
| 0.01 | 0.0009 |
| 0.01 | 0.0012 |
| 6.36 | 3.2091 |
| 2.66 | 0.9502 |
| 3.54 | 0.1932 |
| 100 | 48.19 |


| \& |  |  |  | $$ |
| :---: | :---: | :---: | :---: | :---: |
| 守 |  |  |  |  |
| \% |  | $\underset{N}{N}$ | $\widehat{N}_{0}^{\infty}$ | $\cdots \stackrel{n}{0}$ |



Fish - Otago-Catlins


Appendix 1.2

$$
\begin{aligned}
& \text { Elasmobranchii ?sp } \\
& \text { Callorhinchus milii } \\
& \text { Anguilla spp } \\
& \text { Pseudophycis bachus } \\
& \text { Lotella rhacinus } \\
& \text { Genypterus blacodes } \\
& \text { Helicolenus barathris } \\
& \text { Scorpaena papillosus } \\
& \text { Scorpaena cardinalis } \\
& \text { Neophrynichthys latus } \\
& \text { Chelidonichthys kumu } \\
& \text { Polyprion oxygenios } \\
& \text { Trachurus declivis } \\
& \text { Trachurus novaezelandiae } \\
& \text { Pagrus auratus } \\
& \text { Nemadactylus macropterus } \\
& \text { Latridopsis ciliaris } \\
& \text { Latris lineata } \\
& \text { Aldrichetta forsteri } \\
& \text { Labridae } \\
& \text { Nototheniidae } \\
& \text { Parapercis colias }
\end{aligned}
$$

| ca 1400 AD |  | ca 1550 AD |  | ca 1750 AD |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Taxon frequency | Proportional energy contribution | Taxon frequency | Proportional energy contribution | Taxon frequency | Proportional energy contribution |
| 55.236 | 3.1239 | 47.71 | 26.1132 | 69.09 | 26.2549 |
| 0.030 | 0.0025 |  |  |  |  |
| 0.010 | 0.0010 |  |  |  |  |
| 100.000 | 5.39 | 100.000 | 54.36 | 100.000 | 52.49 |
| ca 1400 |  | ca 1550 |  | ca 1750 |  |
| Taxon frequency | Proportional energy contribution | Taxon frequency | Proportional energy contribution | Taxon frequency | Proportional energy contribution |
| 0.008 | 0.000010 |  |  |  |  |
| 0.718 | 0.002722 | 0.001 | 0.000642 |  |  |
| 15.965 | 0.080051 | 0.029 | 0.043792 | 0.088 | 0.153952 |
| 0.004 | 0.000006 |  |  |  |  |
| 0.002 | 0.000002 |  |  |  |  |
| 0.001 | 0.000001 |  |  |  |  |
| 0.015 | 0.000019 |  |  |  |  |
|  |  | 0.028 | 0.026970 | 0.001 | 0.001138 |
| 0.080 | 0.000101 | 0.086 | 0.032395 |  |  |
| 8.201 | 0.020743 | 0.030 | 0.022491 |  |  |


| Thyrsites atun <br> Rexia solandri <br> Hyperoglyphe antarctica | barracouta <br> gemfish <br> bluenose warehou |
| :--- | :--- |
| Total |  |
|  |  |
|  |  |
|  |  |
|  |  |
| Appendix 1.3 |  |
|  |  |
| Taxon | nut shell |

$\left.\begin{array}{rr}\text { ca } 1750 \mathrm{AD} \\ \begin{array}{r}\text { Taxon } \\ \text { Proportional } \\ \text { energy }\end{array} \\ \text { frequency }\end{array} \begin{array}{r}\text { contribution }\end{array}\right\}$

| ca 1550 AD |  |
| ---: | ---: |
| Taxon <br> frequency | Proportional <br> energy <br> contribution |
| 0.085 | 0.033485 |
| 0.093 | 0.036623 |
|  |  |
| 0.069 | 0.026117 |
|  |  |
| 0.106 | 0.040190 |
|  |  |
| 0.006 | 0.004210 |
| 0.003 | 0.003548 |
| 0.007 | 0.004937 |
| 34.385 | 6.696830 |
| 5.146 | 5.377563 |
| 0.018 | 0.027131 |
|  |  |
| 52.954 | 21.632550 |
|  |  |
| 0.028 | 0.010587 |
| 0.016 | 0.011878 |
| 0.014 | 0.010593 |
| 0.001 | 0.000513 |



|  |  |  | : |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |

Appendix 1.3 continued

$\left.\begin{array}{cr}\text { ca 1750 AD } \\ \text { Taxon } \\ \text { frequency }\end{array} \begin{array}{r}\text { Prortional } \\ \text { energy } \\ \text { contribution }\end{array}\right\}$


| ca 1 | AD |
| :---: | :---: |
| Taxon frequency | Proportional energy contribution |
| 12.177 | 0.030800 |
| 0.028 | 0.000035 |
| 7.119 | 0.018006 |
| 0.001 | 0.000001 |
| 0.315 | 0.000398 |
| 1.014 | 0.238822 |
| 0.013 | 0.002075 |
| 0.140 | 0.000178 |
| 0.013 | 0.000251 |
| 0.043 | 0.000054 |
| 0.016 | 0.000020 |
| 0.003 | 0.000003 |
| 1.175 | 0.002972 |
| 0.001 | 0.000001 |
| 0.317 | 0.000400 |
| 0.489 | 0.009279 |
| 0.194 | 0.000245 |
| 0.002 | 0.000002 |
| 0.140 | 0.002664 |
| 20.366 | 0.103023 |
| 0.002 | 0.000002 |
| 2.673 | 0.003380 |
| 0.006 | 0.000008 |

denticulate limpet
ornate limpet
radiate limpet
encrusted limpet
paua
silver paua
shield limpet
grooved limpet
green top shell
spotted top shell
knobbed topshell
mudflat top shell
wheel shell
spotted tiger shell
cooks turban
cats eye
black nerita
horn shell
Appendix 1.3 continued


| ca 1750 AD |  |
| :---: | :---: |
| Taxon frequency | Proportional energy contribution |
| 0.094 | 0.082912 |
| 0.014 | 0.012337 |
| 0.083 | 0.072874 |
| 0.056 | 0.024682 |
| 0.070 | 0.030844 |



| ca 1400 AD |  |
| :---: | :---: |
| Taxon frequency | Proportiona energy contribution |
| 0.746 | 0.000943 |
| 0.003 | 0.000003 |
| 1.536 | 0.003884 |
| 0.585 | 0.000740 |
| 0.108 | 0.000272 |
| 0.131 | 0.000166 |
| 0.003 | 0.000003 |
| 0.258 | 0.000653 |
| 0.001 | 0.000002 |
| 0.002 | 0.000002 |
| 0.008 | 0.000020 |
| 0.086 | 0.000108 |
| 0.839 | 0.002122 |
| 0.008 | 0.000010 |
| 0.018 | 0.000022 |
| 0.124 | 0.000157 |
| 0.001 | 0.000003 |
| 0.001 | 0.000001 |

brown horn shell
turret shell
ostrich foot
ribbed slipper shell
white slipper shell
circular slipper shell
small circular slipper shell
swollen trumpet shell
spenglers trumpet shell
violet snail
knobbed whelk
lined whelk
speckled whelk
purple-mouthed whelk
spotted whelk
quoy's whelk
red mouthed whelk
siphon whelk
Appendix 1.3 continued
Zeacumantus subcarinatus Zeacumantus ?sp. Maoricolpus roseus Maoricrypta ?sp. Struthiolaria papulosa Maoricrypta costata Maoricrypta monoxyla Sigapatella novaezelandiae Sigapatella tenuis Lamellaria ophione Argobuccinum pustulosum Cabestana spengleri Janthina janthina Aethocola glans Buccinulum pallidum powelli Buccinulum vittatum vittatum Buccinulum ?sp. Cominella adspersa Cominella glandiformis Cominella maculosa
Cominella quoyana quoyana Cominella virgata Cominella ?sp. Penion sulcatus Zemitrella ?sp.
Zemitr

| ca 1750 AD |  |
| ---: | ---: |
| Taxon <br> frequency | Proportional <br> energy <br> contribution |
| 0.013 | 0.005750 |
| 0.001 | 0.006606 |
| 0.193 | 0.084790 |
|  | 0.006606 |
| 0.001 | 0.001321 |
| 0.001 |  |
| 0.102 | 0.044774 |
|  |  |
| 0.897 | 0.395088 |
| 100 | 44.50 |

$\left.\begin{array}{rr}\text { ca } 1550 \mathrm{AD} \\ \text { Taxon } \\ \text { frequency }\end{array} \quad \begin{array}{r}\text { Proportional } \\ \text { energy } \\ \text { contribution }\end{array}\right\}$

| ca 1400 AD |  |
| ---: | ---: |
| Taxon |  |
| frequency | Proportional <br> energy <br> contribution |
| 0.005 | 0.000007 |
| 1.402 | 0.026605 |
| 0.226 | 0.000286 |
| 0.002 | 0.000002 |
| 0.002 | 0.000004 |
| 0.064 | 0.000081 |
| 0.055 | 0.000070 |
| 0.070 | 0.000088 |
| 1.032 | 0.019570 |
| 0.043 | 0.000054 |
|  |  |
| 0.581 | 0.000735 |
| 0.003 | 0.000003 |
| 0.065 | 0.000082 |
| 0.129 | 0.000163 |
| 3.871 | 0.004895 |
| 0.022 | 0.000027 |
| 0.028 | 0.000036 |
|  |  |
| 100 |  |

dark rock shell
oyster borer
octagonal murex
large trophon
white rock shell
arabic volute
southern olive shell
pink tower shell mudsnail $_{\text {tusk shell }}^{\text {gastropod ?sp }} \$$
Appendix 1.3 continued

[^1]

| ca 1750 AD |  |
| ---: | ---: |
| Taxon <br> frequency | Proportional <br> energy <br> contribution |
| 46.206 | 0.586416 |
| 1.570 | 0.026332 |
| 0.297 | 0.002509 |
|  |  |
|  |  |
|  |  |
|  | 0.009519 |
|  |  |
| 0.124 | 0.001357 |
| 0.786 | 0.001711 |
| 0.767 | 0.008951 |
| 0.113 | 0.001906 |
|  |  |
| 2.481 | 0.011325 |


| ca 1 | AD |
| :---: | :---: |
| Taxon frequency | Proportional energy contribution |
| 39.098 | 1.154553 |
| 0.788 | 0.030769 |
| 0.118 | 0.002324 |
| 0.006 | 0.000062 |
| 0.003 | 0.000093 |
| 0.003 | 0.000028 |
| 0.081 | 0.002050 |
| 1.142 | 0.011240 |
| 0.004 | 0.000038 |
| 0.003 | 0.000084 |
| 0.000 | 0.000004 |
| 12.616 | 0.063885 |
| 0.464 | 0.012617 |
| 8.458 | 0.089839 |
| 0.012 | 0.000114 |
| 0.004 | 0.000038 |


| ca 1400 AD |  |
| ---: | ---: |
| Taxon <br> frequency | Proportional <br> energy <br> contribution |
| 13.736 | 0.107401 |
| 1.348 | 0.013925 |
| 0.047 | 0.000244 |
|  |  |
|  |  |
| 0.000 | 0.000000 |
| 0.001 | 0.000004 |
| 0.000 | 0.000001 |
|  |  |
| 0.120 | 0.000940 |
|  |  |
| 35.018 | 0.046950 |
|  |  |
| 0.007 | 0.000036 |
| 0.003 | 0.000014 |
| 33.473 | 0.094134 |
| 0.141 | 0.000367 |
|  |  |
| 0.240 | 0.001253 |

Shellfish - Otago-Catlins
Appendix 1.4


$\left.\left.\begin{array}{rr}\text { ca 1750 AD } \\ \text { Taxon } \\ \text { Proportional } \\ \text { energy }\end{array}\right\} \begin{array}{r}\text { contribution }\end{array}\right\}$

| ca 1550 AD |
| ---: | ---: |
| Taxon |
| Prequency | \(\left.\begin{array}{r}Prortional <br>

energy <br>

contribution\end{array}\right\}\)| 0.028 | 0.000278 |
| ---: | ---: |
| 0.000 | 0.000004 |
| 0.031 | 0.000305 |
| 0.023 | 0.000457 |
| 0.407 | 0.008012 |
| 0.007 | 0.000064 |
| 0.112 | 0.001105 |
| 0.019 | 0.000190 |
| 0.004 | 0.000038 |
| 0.360 | 0.003543 |
| 0.009 | 0.000084 |
| 0.690 | 1.266069 |
| 0.004 | 0.002366 |
| 0.031 | 0.037352 |
| 0.014 | 0.000140 |
| 0.000 | 0.000002 |
| 0.010 | 0.001441 |
| 0.011 | 0.000107 |
| 0.035 | 0.000343 |
| 0.432 | 0.004257 |
| 1.367 | 0.026917 |
| 0.031 | 0.000305 |
| 0.004 | 0.000038 |
| 0.063 | 0.000622 |
| 0.111 | 0.001096 |
| 0.025 | 0.000243 |


|  | $\begin{aligned} & \text { J } \\ & 0 \\ & \hline \\ & 0 \\ & 0 \end{aligned}$ |  |  |  | $\begin{aligned} & \circ \\ & \stackrel{\infty}{+} \\ & \stackrel{\circ}{0} \end{aligned}$ |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 8 |  |  |  |  |  |  |
|  | $\begin{aligned} & \text { N} \\ & \text { O. } \end{aligned}$ | $\begin{array}{llll} 0 & \sim & \infty & 0 \\ \vdots & N \\ 0 & 0 & \infty \\ 0 & 0 & 0 & 0 \\ 0 & 0 & 0 \end{array}$ |  | $\begin{aligned} & \approx \\ & 0 \\ & 0 \\ & 0 \end{aligned}$ | $\begin{aligned} & \infty \\ & \stackrel{\infty}{\infty} \\ & \stackrel{N}{n} \end{aligned}$ | $\begin{array}{lll} N & \approx \\ 0 & 0 \\ 0 & 0 \\ 0 \end{array}$ |


Appendix 1.4 continued
Ruditapes largillierti
Myadora striata
Cellana ornata
Cellana radians
Cellana strigilis
Cellana ?sp.
Notoacmea elongata
Notoacmea pileopsis
Notoacmea scopulina
Patelloida corticata
Radiacmea inconspicua
Haliotis iris
Haliotis virginea
Haliotis australis
Haliotis ?sp.
Emarginula ?sp
Scutus breviculus
Coelotrochus viridius
Cantharidius sanguineus
Cantharidus ?sp.
Diloma aethiops
Diloma arida
Diloma bicanaliculata
Diloma nigerrima
Diloma subrostrata
Diloma zelandica

| ca 1750 AD |  |
| ---: | ---: |
| Taxon <br> Prequency | energy <br> contribution |
| 0.058 | 0.000245 |
| 0.379 | 0.024064 |
| 5.553 | 0.093963 |



| ca 1 | AD |
| :---: | :---: |
| Taxon frequency | Proportiona energy contribution |
| 0.120 | 0.004695 |
| 0.001 | 0.000004 |
| 0.488 | 0.019082 |
| 0.486 | 0.005066 |
| 0.031 | 0.000243 |
| 0.001 | 0.000004 |
| 0.311 | 0.000810 |
| 0.001 | 0.000007 |
| 0.120 | 0.000627 |
| 0.000 | 0.000001 |
| 0.003 | 0.000007 |
| 0.003 | 0.000007 |
| 0.005 | 0.000026 |
| 0.012 | 0.000030 |
| 0.046 | 0.000121 |

Appendix 1.4 continued
wheel shell
cooks turban
cats eye
southern cats eye
turret shell
Stewart Island turret shell
circular slipper shell
swollen trumpet shell
knobbed whelk
many-lined whelk
purple-mouthed whelk
white whelk
oyster borer
large trophon
rock trophon
Diloma ?sp
Zethalia zelandica
Cookia sulcata
Lunella smaragdus
Modelia granosa
Maoricolpus roseus
Maoricrypta ?sp.
Zeaculpus symmetricus
Risellopsis varia
Littorinidae ?sp
Struthiolaria ?sp
Sigapatella novaezelandiae
Argobuccinum pustulosum
Aethocola glans
Buccinulum linea
Buccinulum vittatum littorinoides
Buccinulum ?sp.
Cominella glandiformis
Cominella ?sp.
Haustrum lacunosum
Haustrum scobina
Xymene ambiguous
Xymene huttoni
Xymene ?sp.
Paratrophon patens
Muricidae ?sp


| ca 1400 AD |  | ca 1550 AD |  | ca 1750 AD |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Taxon frequency | Proportional energy contribution | Taxon frequency | Proportional energy contribution | Taxon frequency | Proportional energy contribution |
| 5.89 | 0.183 |  |  |  |  |
| 10.86 | 0.127 |  |  | 18.75 | 0.156 |
| 4.77 | 0.037 |  |  |  |  |
| 6.01 | 0.030 |  |  | 6.25 | 0.023 |
|  |  |  |  | 6.25 | 0.059 |
| 0.25 | 0.022 |  |  |  |  |
| 1.61 | 0.044 | 6.11 | 0.036 | 6.25 | 0.122 |
| 3.40 | 0.290 |  |  |  |  |
| 6.36 | 0.494 |  |  |  |  |
| 18.46 | 0.860 | 14.72 | 0.150 | 18.75 | 0.626 |
| 0.13 | 0.000 |  |  |  |  |
| 0.13 | 0.001 |  |  |  |  |
| 0.13 | 0.000 |  |  |  |  |
| 6.70 | 0.221 | 6.11 | 0.044 | 6.25 | 0.148 |
| 0.13 | 0.001 |  |  |  |  |
| 0.83 | 0.023 |  |  |  |  |
| 0.83 | 0.003 |  |  |  |  |
| 0.13 | 0.001 | 6.11 | 0.008 | 6.25 | 0.028 |
| 100 | 4.16 | 100 | 1.50 | 100 | 2.50 |


| ca 1400 AD |  | ca 1550 AD |  | ca 1750 AD |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Taxon frequency | Proportional energy contribution | Taxon frequency | Proportional energy contribution | Taxon frequency | Proportional energy contribution |
| 13.568 | 1.713 | 4.56 | 1.179 | 1.32 | 0.345 |
| 10.664 | 1.767 | 4.07 | 1.381 | 5.28 | 1.811 |
| 14.654 | 0.509 | 15.01 | 1.067 |  |  |
| 0.512 | 0.129 |  |  | 2.45 | 1.281 |
|  |  |  |  | 1.68 | 0.878 |
|  |  |  |  | 2.69 | 0.791 |
| 0.443 | 0.003 |  |  | 0.29 | 0.004 |
| 6.644 | 0.943 | 4.76 | 1.384 | 14.12 | 4.152 |
| 1.587 | 0.016 |  |  |  |  |
| 1.518 | 0.010 |  |  | 3.42 | 0.029 |
| 1.168 | 0.007 |  |  | 4.28 | 0.056 |
| 5.120 | 0.020 | 3.20 | 0.026 |  |  |
| 8.641 | 0.218 | 3.35 | 0.173 |  |  |
| 1.152 | 0.022 |  |  |  |  |
| 1.888 | 0.018 | 4.69 | 0.091 | 1.32 | 0.026 |
|  |  | 2.55 | 0.115 |  |  |
| 0.606 | 0.001 | 2.35 | 0.007 | 0.75 | 0.002 |
| 6.189 | 0.025 | 13.04 | 0.110 | 2.06 | 0.017 |
| 2.688 | 0.010 |  |  |  |  |
| 0.780 | 0.017 | 2.87 | 0.130 |  |  |
| 3.641 | 0.230 | 3.14 | 0.406 | 1.49 | 0.195 |
| 0.794 | 0.055 | 3.49 | 0.496 | 3.80 | 0.546 |
| 4.674 | 0.369 | 4.76 | 0.769 | 18.70 | 3.055 |


| ca 1750 AD |  |
| ---: | ---: |
| Taxon <br> frequency | Proportional <br> energy <br> contribution |
| 25.39 | 1.991 |
| 6.75 | 0.970 |
| 4.21 | 0.072 |
|  |  |
|  |  |
| 100.00 | $\mathbf{1 6 . 2 2}$ |


$\stackrel{0}{9}$
8
8
8

| ca 1400 AD |  |
| ---: | ---: |
| Taxon <br> frequency | Proportional <br> energy <br> contribution |
| 9.304 | 0.352 |
| 1.632 | 0.044 |
|  |  |
| 0.577 | 0.005 |
| 0.309 | 0.001 |
| 1.247 | 0.006 |
|  |  |
| 100.00 | $\mathbf{6 . 4 9}$ |

Spotted shag
Southern black-backed gull
Red-billed gull
Black-billed gull
Black-fronted tern
White-fronted tern
Appendix 1.6 continued
Stictocarbo punctatus
Larus dominicanus
Larus novaehollandiae
Larus bulleri
Childonias albostriata
Sterna striata
Total
$\left.\begin{array}{r}\text { ca } 1750 \mathrm{AD} \\ \begin{array}{r}\text { Taxon } \\ \text { Proportional } \\ \text { energy }\end{array} \\ \text { contribution }\end{array}\right\}$
ca 1550 AD

Taxon | Proportional |
| ---: |
| energy |
| frequency |
| contribution |

100.00
100

| ca 1400 AD |
| ---: | ---: |
| Taxon |
| frequency | \(\left.\begin{array}{r}Proportional <br>

energy <br>

contribution\end{array}\right\}\)| 9.40 | 0.5385 |
| ---: | ---: |
| 29.56 | 5.0526 |
| 25.53 | 17.9041 |
| 5.38 | 1.9185 |
| 16.13 | 18.1182 |
| 86.00 |  |
| 12.00 | 9.4682 |
| 0.10 | 0.6887 |
| 0.90 | 0.6571 |
| 1.00 | 0.4542 |
|  |  |
| 100 | 54.80 |

Marine Mammals - Greater Hauraki
 fur seal - adult female fur seal - adult male fur seal - total elephant seal dolphin?sp

## Appendix 1.7

Arctocephalus forsteri
Arctocephalus forsteri
Arctocephalus forsteri Arctocephalus forsteri Arctocephalus forsteri Arctocephalus forsteri
Phocarctus hookeri
Mirounga leonina
Globicephala ?sp
Dolphin ?sp
Total

| ca 1750 AD |  |
| :---: | :---: |
| Taxon frequency | Proportional energy contribution |
| 60.10 | 2.5242 |
| 23.60 | 4.0668 |
| 15.70 | 4.3333 |
| 99.40 |  |
| 0.10 | 0.4680 |
| 0.50 | 0.0558 |
| 100.00 | 10.98 |


|  |  |  |
| :---: | :---: | :---: |
| in |  |  |
|  |  | $\frac{0}{3} \frac{0}{0} \stackrel{0}{0}$ |


\left.| ca 1400 AD |
| ---: | ---: |
| Taxon |
| Proportional |
| energy |$\right\left.\} \begin{array}{r}\text { contribution }\end{array}\right\}$|  |  |
| ---: | ---: |
| 5.20 | 0.3111 |
| 40.11 | 7.1595 |
| 15.60 | 11.4247 |
| 6.69 | 2.4912 |
| 10.40 | 12.1993 |
| 78.00 |  |
| 21.20 | 17.4680 |
| 0.10 | 0.7192 |
| 0.10 | 0.0899 |
| 0.10 | 0.0762 |
| 0.50 | 0.2371 |
|  |  |
| 100.00 | $\mathbf{5 2 . 1 0}$ |

> fur seal - pup fur seal - juvenile fur seal - sub adult male fur seal - adult female fur seal - adult male fur seal - total sea lion elephant seal leopard seal pilot whale ?sp dolphin ?sp
Arctocephalus forsteri Arctocephalus forsteri Arctocephalus forsteri Arctocephalus forsteri Arctocephalus forsteri Arctocephalus forsteri Phocarctus hookeri Mirounga leonina Hydrurga leptonyx Globicephala ?sp Dolphin ?sp
Total


[^0]:    Arctocephalus forsteri

    - juvenile
    - sub adult male - adult male

    Globicephala ?sp Total

[^1]:    Taron 9 sp. Haustrum haustorium Haustrum scobina Murexsul octogonus Xymene ambiguous Xymene plebeius Xymene traversi Dicathais orbita Austromitra rubiginosa Alcithoe arabica Amalda australis Phenatoma rosea Turbonilla zealandica Acteonidae?sp Amphibola crenata Amuarochiton glaucus Antalis nana gastropod ?sp.

    Total

