

New Zealand Aquatic
Environment and Biodiversity
Report No. 82
2011
ISSN 1176-9440

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

I.W.G. Smith

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

I.W.G. Smith

Department of Anthropology and Archaeology
University of Otago
PO Box 56
Dunedin 9054

**Published by Ministry of Fisheries
Wellington
2011**

ISSN 1176-9440

©
**Ministry of Fisheries
2011**

Smith, I.W.G. (2011).
Estimating the magnitude of pre-European Maori marine harvest in two New Zealand study areas
New Zealand Aquatic Environment and Biodiversity Report No. 82.

This series continues the
Marine Biodiversity Biosecurity Report series
which ceased with No. 7 in February 2005.

EXECUTIVE SUMMARY

Smith, I.W.G. (2011). Estimating the magnitude of pre-European Maori marine harvest in two New Zealand study areas.

New Zealand Aquatic Environment and Biodiversity Report No. 82.

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 AD, *ca.* 1550 AD, and *ca.* 1750 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 well-supported 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 *ca.* 1400 AD in Greater Hauraki the best estimate of harvested biomass of snapper (*Pagrus auratus*) is 72.1 ± 21.6 tonnes (minimum 12.1 ± 3.6 t; maximum 107.5 ± 32.3 t), which rose to 938.8 ± 281.7 t (min. 354.8 ± 106.4 t; max. 1354.7 ± 406.4 t) at *ca.* 1550 AD, and then 997.2 ± 299.2 (min. 560.1 ± 168.0 t; max. 1393.9 ± 418.2 t) at *ca.* 1750 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 ± 85.5 t (min. 47.9 ± 14.4 t; max. 425.1 ± 127.5 t) at *ca.* 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 ± 71.1 t (min. 55.2 ± 16.6 t; max. 383.5 ± 115.1 t) to 103.3 ± 31.0 t (min. 24.0 ± 7.2 ; max. 173.8 ± 52.1 t) then 77.0 ± 23.1 t (min. 57.4 ± 17.2 ; max. 99.7 ± 29.9 t) in *ca.* 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 Otago-Catlins, 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), *ca.* 1750 AD (before European whaling and sealing), *ca.* 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 18th century AD (*ca.* 1400, 1550, 1750 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 well-explored 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 18th and 19th 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 10 000 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 δR set at -7 ± 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. AD 1400, ca. AD 1550 and ca. 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 1σ and 2σ 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 under- or 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 %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 Diomedidae 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 %MNI for species in this family were reduced by half and all other taxon frequencies increased proportionately.

The adjusted %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 (%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 under-representation 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 %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 %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 %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 *ca.* 100,000 for the total New Zealand population in 1769, with *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 100 000 respectively, giving estimates for Greater Hauraki of 10 800 and 13 500. Anderson (1998: 196–7) has estimated that in the early 19th 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 *ca.* 1750 AD for Otago-Catlins of about 1800, with ± 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 20 000 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 *ca.* 1550 AD the population would have reached about 90% of its *ca.* 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 72 000, best estimate of 80 000, and a maximum of 90 000. 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, 10 200 and 11 700.

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 *ca.* 1550 AD was no greater than at *ca.* 1400 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 *ca.* 1400 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 $\delta^{13}\text{C}$, $\delta^{15}\text{N}$ and $\delta^{34}\text{S}$ isotopes from the small number of archaeologically-derived 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 AD, 1550AD, 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 *ca.* 1400 AD 22 of these taxa were included in a total fish harvest best estimated at 128.5 ± 44.3 tonnes (Table 5). The taxonomic range narrowed to 16 at *ca.* 1550 AD in a total harvest of 2328.2 ± 698.5 t (Table 6). By *ca.* 1750 AD, the total fish take of 2607.1 ± 782.1 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 ± 21.6 t to 938.8 ± 281.6 t and then 997.2 ± 299.2 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 ± 5.8 t (15%), wrasses (Labridae) 13.5 ± 4.0 t (10.5%), and leatherjacket (*Meuschenia scaber*) 11.3 ± 3.4 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 ± 153.4 t (22%); horse mackerel (*Trachurus novaezelandiae*), 323.1 ± 96.9 t (13.9%); kahawai, 227.9 ± 68.4 t (9.8%); and barracouta (*Thyrsites atun*) 169.3 ± 50.8 t (7.3%). The harvest of wrasses (23.1 ± 6.9 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 ± 287.8 t, 36.8%) was almost equal to that of snapper, and would just outstrip it if combined with the 39.9 ± 12 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 ± 95.8 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 ± 23.5 t at ca. 1400 AD (Table 8). The taxonomic range narrowed to 14 at ca. 1550 AD in a total harvest of 797.5 ± 239.3 t (Table 9). By ca. 1750 AD, the total fish take of 725.1 ± 217.5 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 ± 13.0 t (55.2%) at ca. 1400 AD, 361.4 ± 108.4 t (45.3%) at ca. 1550 AD, and 363.3 ± 109.0 t (50.1%) at ca. 1750 AD. Three further species made important contributions to harvested biomass at ca. 1400 AD; ling (*Genypterus blacodes*) 7.8 ± 2.3 t (9.9%), hapuku (*Polyprion oxygenios*) 7.7 ± 2.3 t (9.8%), and red cod (*Pseudophycis bachus*) 6.8 ± 2.0 t (8.7%). The same three species were the major components in addition to barracouta at ca. 1550 AD, although in differing order; hapuku 141.0 ± 42.3 t (17.7%), red cod 128.3 ± 38.5 t (16.1%), and ling 68.2 ± 20.5 t (8.5%). This order was maintained at ca. 1750 AD; hapuku 230.9 ± 69.3 t (31.8%), red cod 68.4 ± 20.5 t (9.4%), and ling 56.8 ± 17 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 ± 2.0 t at ca. 1400 AD (Table 11), 66 in the estimate of 1736.3 ± 520.9 t at ca. 1550 AD (Table 12), and 38 in the estimate of 2294.7 ± 688.4 t at ca. 1750 AD (Table 13).

Three species dominate the shellfish biomass harvested at ca. 1400 AD, making up two thirds of the total; paua (*Haliotis iris*) 2.2 ± 0.7 t (34%), cats eye (*Lunella smaragdus*) 1.2 ± 0.4 t (18.2%), and green-lipped mussel (*Perna canaliculus*) 0.9 ± 0.3 t (14.3%). At ca. 1550 AD two species made up more than 84% of the total biomass harvested: cockle (*Austrovenus stutchburyi*) 1103.2 ± 331.0 t (63.5%), and pipi (*Paphies australis*) 358.2 ± 107.5 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 ca. 1750 AD as they did at ca. 1550; cockle providing 1358.0 ± 407.4 t (59.2%), and pipi 594.5 ± 178.4 t (25.9%). Tuatua (*Paphies subtriangulata*) with 250.5 ± 75.2 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 ± 2.7 t (Table 14), with just 12 taxa in the estimated 43.0 ± 12.9 t harvested at ca. 1550 AD (Table 15), but 29 were included in the estimated harvest of 51.4 ± 15.4 t at ca. 1750 AD (Table 16). The reduced taxonomic range at ca. 1550 is almost certainly due to the restricted sample of study sites available for that time period; ten study assemblages were available for ca. 1400 AD and nine for ca. 1750 AD, but only two for ca. 1550 AD.

At ca. 1400 AD four species made up 82.4% of the total harvested biomass; cockle (*Austrovenus stutchburyi*) 2.4 ± 0.7 t (26.9%), paua (*Haliotis iris*) 2.3 ± 0.7 t (24.8%), blue mussel (*Mytilus galloprovincialis*) 1.5 ± 0.5 t (16.6%), and pipi (*Paphies australis*) 1.3 ± 0.3 t (14.1%). Two of these four species also dominate harvested shellfish biomass at ca. 1550 AD; blue mussels providing $16.1 \pm$

4.8 t (37.4%), and paua 14.2 ± 4.3 t (33%). Mudsnaills (*Amphibola crenata*) added another 4.2 ± 1.3 t (9.8%), cockles 2.3 ± 0.2 t (5.4%), and pipi 1.7 ± 0.5 t (4%) to the total harvested biomass in *ca.* 1550 AD. At *ca.* 1750 AD the rank order of the two leading shellfish species reverses; paua yielding 35.1 ± 10.5 t (68.4%), and blue mussels 8.2 ± 2.5 t (15.9%). None of the other previously important species contributed more than 2.5% of total harvested biomass at *ca.* 1750 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 ± 5.8 t in *ca.* 1400 AD (Table 17). Only seven taxa were among the harvest of $28.3 \pm$ t at *ca.* 1550 AD (Table 18), while at *ca.* 1750 AD there were ten in the 56.2 ± 16.9 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 ± 1.4 t (24.4%), spotted shag (*Stictocarbo punctatus*) 4.0 ± 1.2 t (20.7%), albatrosses (Diomedeidae) 2.5 ± 0.8 t (13.2%), and pied shag (*Phalacrocorax varius*) 2.3 ± 0.7 t (11.9%). Three of these taxa together yielded 90.2% of the harvested biomass in *ca.* 1550 AD; albatrosses 17.6 ± 5.3 t (62.2%), little penguin 5.1 ± 1.5 t (18.5%), and spotted shag 2.8 ± 0.8 t (10%). At *ca.* 1750 AD the same three taxa provided 71.6% of the harvested biomass; albatrosses 17.6 ± 5.3 t (31.3%), spotted shag 14.1 ± 4.2 t (25%), and little penguin 8.6 ± 2.6 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.* 1400 25 marine and coastal bird taxa formed part of a total estimated harvest of 32.1 ± 9.6 t (Table 20). Only sixteen were among the harvest of 46.9 ± 14.1 t in *ca.* 1550 AD (Table 21), while 18 contributed to the 80.3 ± 24.1 t harvested in *ca.* 1750 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 ± 2.6 t (27.2%), Fiordland crested penguin (*Eudyptes pachyrhynchus*) 8.5 ± 2.5 t (26.4%), white-capped albatross (*Thalassarche cauta*) 4.7 ± 1.4 t (14.5%), and little penguin (*Eudyptula minor*) 2.5 ± 0.8 t (7.8%). Five taxa, including the four above, made up 74% of the biomass harvested at *ca.* 1550 AD; spotted shag (*Stictocarbo punctatus*) 9.9 ± 3.0 t (21%), white-capped albatross 6.9 ± 2.1 t (14.6%), *Megadyptes* sp. penguin 6.8 ± 2.1 t (14.6%), Fiordland crested penguin 5.8 ± 1.8 t (12.5%), and little penguin 5.3 ± 1.6 t (11.3%). At *ca.* 1750 AD the leading four taxa contributed 67.9% of the total biomass harvested; white capped albatross 20.6 ± 6.2 t (25.6%), Stewart Island shag (*Leucocarbo chalconotus*) 15.1 ± 4.5 t (18.8%), spotted shag 9.9 ± 3.0 t (12.3%), and *Megadyptes*, by this time probably yellow-eyed penguin, 9.0 ± 2.7 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 ± 124.3 t in *ca.* 1400AD (Table 23). Fur seals made up the largest part of this, 284.9 ± 85.5 t (68.7%), followed by sea lions, 63.4 ± 19.0 t (15.3%), and pilot whales, 57.5 ± 17.3 t (13.9%). By *ca.* 1550 AD the total harvest had fallen to 1.79 ± 0.54 t, derived entirely from dolphins (Table 24). Dolphins were also the sole contributor to the marine mammals harvest of 2.13 ± 0.64 t in *ca.* 1750 AD (Table 25).

3.4.1 Otago-Catlins

The archaeological record shows that at least eight marine mammal taxa were harvested in Otago-Catlins, 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 Hector's 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 *ca.* 1400 AD five taxa were included in the total harvest of 371.3 ± 111.4 t (Table 26). This was dominated by two species; fur seals providing 237.0 ± 71.1 t (63.8%), and sea lions 126.2 ± 37.8 t (34%). Four of the same taxa, plus another, made up the considerably smaller harvest of 144.5 ± 43.4 t at *ca.* 1550 AD (Table 27). Fur seals, at 103.3 ± 31 t (71.5%), and sea lions, at 47.2 ± 11.2 t (25.8%), were again dominant at *ca.* 1550 AD. At *ca.* 1750 AD the total harvest had fallen further to 121.6 ± 36.5 t derived from just three taxa (Table 28). The main contributors to the total harvested biomass at *ca.* 1750 AD were fur seals, at 77.0 ± 23.1 t (63.4%), and pilot whales at 44.0 ± 13.2 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 ± 1612 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 *ca.* 1750 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 *ca.* 1750 AD is 6.1 times greater than that at *ca.* 1400 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 *ca.* 1400 and *ca.* 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 *ca.* 1750 AD ranges between 20% and 72% of those exploited at *ca.* 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 be 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 *ca.* 1550 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.

7. ACKNOWLEDGMENTS

This research was funded by the Ministry of Fisheries through the *Taking Stock* Project ZBD200505. I am grateful to Alison MacDiarmid for the opportunity to participate in this project, Foss Leach for inspiration and support, and Tiffany James-Lee for assistance with data collection. Richard Walter and Richard Ford provided useful comments on the manuscript.

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. 1650-1850*. 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.; Snodgrass, 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 Auckland. 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, Berkeley. 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. 76–108.
- 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>.
- Ulrich, 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 age-determinations - 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: 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
<i>Greater Hauraki</i>								
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
<i>Otago-Catlins</i>								
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
<i>Greater Hauraki</i>									
ca. 1400	500	2 000	2 500	1 800	2 150	2 172	0.53	0.66	0.78
ca. 1550	6 100	10 200	11 700	1 800	2 150	2 172	0.40	0.53	0.66
ca. 1750	10 800	12 150	13 500	1 800	2 150	2 172	0.40	0.53	0.66
<i>Otago-Catlins</i>									
ca. 1400	500	1 800	2 500	2 150	2 172	2 193	0.66	0.78	0.90
ca. 1550	500	1 800	2 600	2 150	2 172	2 193	0.66	0.78	0.90
ca. 1750	1 600	1 800	2 000	2 150	2 172	2 193	0.66	0.78	0.90

Table 4: Estimated total energy (Kcal x 10⁶) harvested from animals during focal years in each study area.

Area/year	Minimum estimate	Best estimate	Maximum estimate
<i>Greater Hauraki</i>			
ca. 1400	174.11	1 035.87	1 545.92
ca. 1550	1 603.08	4 242.36	6 121.85
ca. 1750	2 838.24	5 053.40	7 063.67
<i>Otago-Catlins</i>			
ca. 1400	258.97	1 113.06	1 801.00
ca. 1550	388.45	1 236.74	1 873.04
ca. 1750	828.70	1 113.06	1 440.80

Table 5: Estimated numbers (N) and biomass (tonnes) of fish harvested in Greater Hauraki at ca. 1400 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass	N	Biomass	N	Biomass
<i>Carcharhiniforme ?sp</i>	24	0.485	144	2.886	215	4.307
<i>Myliobatis tenuicaudatus</i>	3	0.051	15	0.304	23	0.453
<i>Elasmobranchii ?sp</i>	27	0.268	159	1.595	238	2.380
<i>Callorhynchus milii</i>	3	0.011	15	0.064	23	0.095
<i>Anguilla</i> spp.	27	0.024	159	0.144	238	0.214
<i>Pseudophycis bachus</i>	27	0.040	159	0.239	238	0.357
<i>Zeus faber</i>	52	0.073	311	0.436	465	0.651
<i>Chelidonichthys kumu</i>	92	0.064	547	0.383	816	0.571
<i>Pseudocaranx dentex</i>	6	0.010	38	0.061	57	0.091
<i>Trachurus declivis</i>	3	0.003	15	0.020	23	0.029
<i>Trachurus novaezelandiae</i>	3	0.003	15	0.015	23	0.023
<i>Arripis trutta</i>	1 795	3.231	10 679	19.221	15 937	28.686
<i>Pagrus auratus</i>	5 504	12.110	32 750	72.049	48 875	107.525
<i>Nemadactylus macropterus</i>	448	0.358	2 666	2.133	3 978	3.183
<i>Latridopsis ciliaris</i>	79	0.214	471	1.271	703	1.897
<i>Mugil cephalus</i>	3	0.003	15	0.015	23	0.023
<i>Aldrichetta forsteri</i>	544	0.109	3 235	0.647	4 829	0.966
Labridae	1 508	2.261	8 970	13.455	13 386	20.079
<i>Parapercis colias</i>	119	0.083	706	0.494	1 054	0.738
<i>Thyrssites atun</i>	111	0.255	661	1.520	986	2.268
<i>Serollela brama</i>	24	0.056	144	0.332	215	0.495
<i>Meuschenia scaber</i>	2 365	1.892	14 074	11.259	21 003	16.803
TOTAL	12 765	21.605	75 950	128.542	113 347	191.835
		7.448		44.316		66.137

Table 6: Estimated numbers (N) and biomass (tonnes) of fish harvested in Greater Hauraki at ca. 1550 AD.

	Minimum estimate			Best estimate			Maximum estimate		
	N	Biomass	± 30%	N	Biomass	± 30%	N	Biomass	± 30%
<i>Carcharhiniforme</i> ?sp	9 663	193.267	57.980	25 572	511.436	153.431	36 901	738.017	221.405
<i>Anguilla</i> spp.	5 815	5.234	1.570	15 389	13.850	4.155	22 207	19.987	5.996
<i>Chelidonichthys kumu</i>	1 355	0.949	0.285	3 587	2.511	0.753	5 176	3.623	1.087
<i>Pseudocaranx dentex</i>	12 724	20.359	6.108	33 671	53.874	16.162	48 589	77.742	23.323
<i>Trachurus declivis</i>	219	0.284	0.085	579	0.752	0.226	835	1.085	0.326
<i>Trachurus novaezelandiae</i>	122 082	122.082	36.625	323 061	323.061	96.918	466 186	466.186	139.856
<i>Arripis trutta</i>	47 836	86.104	25.831	126 586	227.855	68.357	182 667	328.801	98.640
<i>Pagrus auratus</i>	161 260	354.772	106.432	426 736	938.820	281.646	615 793	1354.745	406.423
<i>Mugil cephalus</i>	5 815	5.815	1.745	15 389	15.389	4.617	22 207	22.207	6.662
<i>Aldrichetta forsteri</i>	9 663	1.933	0.580	25 572	5.114	1.534	36 901	7.380	2.214
Labridae	5 815	8.723	2.617	15 389	23.084	6.925	22 207	33.311	9.993
<i>Genyagnus monopterygius</i>	44	0.022	0.007	116	0.058	0.017	167	0.083	0.025
<i>Parapercis colias</i>	44	0.031	0.009	116	0.081	0.024	167	0.117	0.035
<i>Thyrsites atun</i>	27 809	63.962	19.189	73 591	169.260	50.778	106 194	244.247	73.274
<i>Scomber australasicus</i>	11 631	11.631	3.489	30 779	30.779	9.234	44 415	44.415	13.324
<i>Rhombosolea</i> ?sp	15 479	4.644	1.393	40 961	12.288	3.687	59 108	17.732	5.320
TOTAL	437 256	879.811	263.943	1 159 561	2 328.213	698.464	1 669 721	3 359.679	1 007.904

Table 7: Estimated numbers (N) and biomass (tonnes) of fish harvested in Greater Hauraki at ca. 1750 AD.

	Minimum estimate			Best estimate			Maximum estimate		
	N	Biomass	± 30%	N	Biomass	± 30%	N	Biomass	± 30%
<i>Squalus blainvillei</i>	8 966	22.415	6.725	15 964	39.910	11.973	22 314	55.786	16.736
Carcharhiniforme ?sp	26 941	538.829	161.649	47 968	959.369	287.811	67 051	1 341.011	402.303
<i>Myliobatis tenuicaudatus</i>	8 966	179.322	53.797	15 964	319.278	95.783	22 314	446.289	133.887
<i>Anguilla</i> spp.	20 950	18.855	5.656	37 300	33.570	10.071	52 139	46.925	14.077
<i>Chelidonichthys kumu</i>	8 966	6.276	1.883	15 964	11.175	3.352	22 314	15.620	4.686
<i>Pseudocaranx dentex</i>	20 950	33.519	10.056	37 300	59.680	17.904	52 139	83.422	25.026
<i>Trachurus declivis</i>	8 966	11.656	3.497	15 964	20.753	6.226	22 314	29.009	8.703
Carangidae ?sp	8 966	9.863	2.959	15 964	17.560	5.268	22 314	24.546	7.364
<i>Arripis truttia</i>	17 975	32.356	9.707	32 005	57.608	17.282	44 736	80.525	24.158
<i>Pagrus auratus</i>	254 586	560.089	168.027	453 283	997.222	299.166	633 601	1 393.922	418.177
<i>Mugil cephalus</i>	8 966	1.793	0.538	15 964	3.193	0.958	22 314	4.463	1.339
Labridae	8 966	13.449	4.035	15 964	23.946	7.184	22 314	33.472	10.041
<i>Odax pullus</i>	8 966	8.070	2.421	15 964	14.368	4.310	22 314	20.083	6.025
<i>Thyrsites atun</i>	8 966	20.622	6.187	15 964	36.717	11.015	22 314	51.323	15.397
<i>Meuschenia scaber</i>	8 966	7.173	2.152	15 964	12.771	3.831	22 314	17.852	5.355
TOTAL	431 063	1 464.288	439.286	767 495	2 607.119	782.136	1 072 809	3 644.247	1 093.274

Table 8: Estimated numbers (N) and biomass (tonnes) of fish harvested in Otago-Catlins at ca. 1400 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass	N	Biomass	N	Biomass
<i>Elasmobranchii</i> ?sp	1	0.008	3	0.034	6	0.055
<i>Anguilla</i> spp.	1	0.016	3	0.068	6	0.110
<i>Pseudophycis bachus</i>	1 058	1.587	4 546	6.819	7 356	11.033
<i>Lotella rhacinus</i>	1	0.001	3	0.003	6	0.006
<i>Genypterus blacodes</i>	362	1.810	1 556	7.779	2 517	12.587
<i>Helicolenus barathris</i>	3	0.001	14	0.005	22	0.009
<i>Scorpaena cardinalis</i>	32	0.032	139	0.139	225	0.225
<i>Chelidonichthys kumu</i>	0	0.000	2	0.001	3	0.002
<i>Polyprion oxygenios</i>	89	1.789	385	7.691	622	12.444
<i>Trachurus declivis</i>	1	0.001	2	0.002	3	0.004
<i>Trachurus novaezelandiae</i>	16	0.016	69	0.069	112	0.112
<i>Pagrus auratus</i>	238	0.525	1 025	2.255	1 658	3.649
<i>Nemadactylus macropterus</i>	34	0.027	146	0.117	236	0.189
<i>Latridopsis ciliaris</i>	28	0.074	118	0.320	192	0.517
<i>Latris lineata</i>	191	0.572	820	2.459	1 326	3.979
<i>Aldrichetta forsteri</i>	1	0.000	3	0.001	6	0.001
Labridae	212	0.318	911	1.366	1 474	2.211
<i>Parapercis colias</i>	750	0.525	3 222	2.256	5 214	3.650
Nototheniidae	524	0.838	2 252	3.603	3 644	5.830
<i>Thyrssites atun</i>	4 373	10.058	18 796	43.232	30 414	69.952
<i>Rexea solandri</i>	2	0.007	10	0.031	17	0.050
<i>Hyperoglyphe antarctica</i>	1	0.003	3	0.014	6	0.022
TOTAL	7 917	18.209	34 029	78.264	55 061	126.636
		5.463	23.479	37.991		

Table 9: Estimated numbers (N) and biomass (tonnes) of fish harvested in Otago-Catlins at ca. 1550 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass ± 30%	N	Biomass ± 30%	N	Biomass ± 30%
<i>Elasmobranchii</i> ?sp	238	2.375	1 021	10.209	1 718	17.180
<i>Pseudophycis bachus</i>	19 906	29.859	85 559	128.338	143 977	215.965
<i>Gemypterus blacodes</i>	3 172	15.861	13 634	68.170	22 943	114.716
<i>Helicolenus barathris</i>	31	0.012	132	0.053	222	0.089
<i>Scorpaena papillosus</i>	31	0.031	132	0.132	222	0.222
<i>Scorpaena cardinalis</i>	460	0.460	1 976	1.976	3 325	3.325
<i>Neophrynichthys latus</i>	31	0.058	132	0.250	222	0.421
<i>Polyprion oxygenios</i>	1 640	32.794	7 048	140.951	11 860	237.190
<i>Nemadactylus macropterus</i>	46	0.037	198	0.158	333	0.266
<i>Latris lineata</i>	590	1.770	2 536	7.607	4 267	12.802
Labridae	4 682	7.022	20 122	30.183	33 861	50.791
<i>Parapercis colias</i>	3 984	2.789	17 125	11.987	28 817	20.172
Nototheniidae	5 256	8.410	22 592	36.147	38 017	60.827
<i>Thyrsites atun</i>	36 556	84.079	157 121	361.379	264 401	608.121
TOTAL	76 622	185.557	329 325	797.541	554 183	1 342.086
		55.667		239.262		402.626

Table 10: Estimated numbers (N) and biomass (tonnes) of fish harvested in Otago-Catlins at ca. 1750 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass	N	Biomass	N	Biomass
<i>Elasmobranchii</i> ?sp	170	1.702	229	2.286	296	2.960
<i>Callorhynchus milii</i>	170	0.715	229	0.960	296	1.243
<i>Pseudophycis bachus</i>	3 3928	50.891	45 570	68.355	58 988	88.481
<i>Genypterus blacodes</i>	8461	42.303	11 364	56.819	14 710	73.550
<i>Polyprion oxygenios</i>	8597	171.936	11 547	230.935	14 947	298.934
<i>Latridopsis ciliaris</i>	51	0.138	69	0.185	89	0.240
Labridae	1021	1.532	1 372	2.058	1 776	2.664
<i>Paraperca colias</i>	221	0.155	297	0.208	385	0.269
<i>Thyrstites atun</i>	11 7614	270.513	157 974	363.339	204 488	470.323
TOTAL	17 0233	539.885	228 649	725.146	295 974	938.663
		161.965		217.544		281.599
		± 30%		± 30%		± 30%

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

	Minimum estimate		Best estimate		Maximum estimate	
	$N \times 10^3$	Biomass	$N \times 10^3$	Biomass	$N \times 10^3$	Biomass
<i>Mytilus galloprovincialis</i>	1.77	0.0053	10.52	0.0316	15.71	0.0471
<i>Perna canaliculus</i>	39.35	0.1574	234.14	0.9366	349.43	1.3977
oyster ?sp.	20.22	0.0404	120.27	0.2405	179.50	0.3590
<i>Paphies australis</i>	6.81	0.0068	40.50	0.0405	60.44	0.0604
<i>Paphies subtriangulata</i>	22.68	0.0454	134.94	0.2699	201.39	0.4028
<i>Austrovenus stutchburyi</i>	6.44	0.0129	38.32	0.0766	57.19	0.1144
<i>Dosinia</i> ?sp.	2.71	0.0054	16.11	0.0322	24.05	0.0481
<i>Cellana denticulata</i>	30.02	0.0600	178.59	0.3572	266.52	0.5330
<i>Cellana radicans</i>	17.55	0.0351	104.40	0.2088	155.81	0.3116
<i>Haliotis iris</i>	2.50	0.3748	14.86	2.2297	22.18	3.3275
<i>Diloma aethiops</i>	2.90	0.0058	17.24	0.0345	25.72	0.0514
<i>Diloma</i> ?sp	1.21	0.0181	7.17	0.1076	10.71	0.1606
<i>Lunella smaragdus</i>	50.20	0.2008	298.68	1.1947	445.75	1.7830
<i>Nerita atramentosa</i>	6.59	0.0066	39.20	0.0392	58.50	0.0585
<i>Maoricolpus roseus</i>	1.84	0.0018	10.94	0.0109	16.32	0.0163
<i>Maoricrypta costata</i>	3.79	0.0076	22.52	0.0450	33.61	0.0672
<i>Maoricrypta monoxyla</i>	1.44	0.0014	8.58	0.0086	12.81	0.0128
<i>Cominella maculosa</i>	2.07	0.0041	12.31	0.0246	18.37	0.0367
<i>Haustrum haustorium</i>	3.46	0.0519	20.57	0.3085	30.70	0.4605
<i>Dicathais orbita</i>	2.54	0.0381	15.13	0.2270	22.58	0.3387
<i>Amalda australis</i>	1.43	0.0014	8.52	0.0085	12.71	0.0127
<i>Amphibola crenata</i>	9.54	0.0095	56.77	0.0568	84.72	0.0847
all others	9.46	0.0107	56.31	0.0639	84.04	0.0954
TOTAL	246.50	1.1015	1 466.61	6.5535	2 188.75	9.7803
		0.3304	1.9660	2.9341		

Table 13: Estimated numbers (N x 10³) and biomass (tonnes) of shellfish harvested in Greater Hauraki at ca. 1750 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N x 10 ³	Biomass	N x 10 ³	Biomass	N x 10 ³	Biomass
<i>Perna canaliculus</i>	723.6	2.895	1 288.4	5.154	1 800.9	7.204
<i>Purpurocardia</i> ?sp.	11 131.7	11.132	19 819.6	19.820	27 704.0	27.704
<i>Paphies australis</i>	333 923.2	333.923	594 539.9	594.540	831 051.4	831.051
<i>Paphies subtriangulata</i>	70 348.8	140.698	125 253.8	250.508	175 080.5	350.161
<i>Austrovenus stutchburyi</i>	381 349.4	762.699	678 980.7	1 357.961	949 083.4	1 898.167
<i>Macra discors</i>	4 783.9	14.352	8 517.5	25.553	11 905.8	35.718
<i>Dosinia anus</i>	92.0	0.184	163.8	0.328	229.0	0.458
<i>Dosinia</i> ?sp.	261.8	0.524	466.1	0.932	651.5	1.303
<i>Myadora striata</i>	261.8	0.262	466.1	0.466	651.5	0.652
<i>Haliotis iris</i>	8.2	1.231	14.6	2.192	20.4	3.064
<i>Zethalia zelandica</i>	2 879.7	2.880	5 127.2	5.127	7 166.8	7.167
<i>Cookia sulcata</i>	8.2	0.123	14.6	0.219	20.4	0.306
<i>Lunella smaragdus</i>	386.5	1.546	688.2	2.753	962.0	3.848
<i>Zeacumantus lutulentus</i>	1 914.7	1.915	3 409.0	3.409	4 765.2	4.765
<i>Struthiolaria papulosa</i>	772.6	1.545	1 375.6	2.751	1 922.8	3.846
<i>Aethocola glans</i>	115.0	0.230	204.7	0.409	286.1	0.572
<i>Cominella adpersa</i>	679.0	1.358	1 209.0	2.418	1 690.0	3.380
<i>Cominella glandiformis</i>	460.0	0.460	819.0	0.819	1 144.8	1.145
<i>Cominella virgata</i>	574.8	0.575	1 023.4	1.023	1 430.5	1.431
<i>Xymene ambiguous</i>	1 580.2	1.580	2 813.4	2.813	3 932.6	3.933
<i>Dicathais orbita</i>	8.2	0.123	14.6	0.219	20.4	0.306
<i>Amphibola crenata</i>	834.4	0.834	1 485.7	1.486	2 076.7	2.077
all others	7 363.0	7.726	13 109.5	13.755	18 324.5	19.227
TOTAL	820 708.4	1 288.793	1 461 245.6	2 294.655	2 042 537.9	3 207.483
		386.638		688.397		962.245

Table 14: Estimated numbers (N x 10³) and biomass (tonnes) of shellfish harvested in Otago-Catlins at ca. 1400 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N x 10 ³	Biomass	N x 10 ³	Biomass	N x 10 ³	Biomass
<i>Mytilus galloprovincialis</i>	116.9	0.351	502.3	1.507	812.8	2.438
<i>Perna canaliculus</i>	11.5	0.046	49.3	0.197	79.8	0.319
<i>Aulacomya maoriana</i>	0.4	0.001	1.7	0.003	2.8	0.006
<i>Macra discors</i>	1.0	0.003	4.4	0.013	7.1	0.021
<i>Paphies australis</i>	298.0	0.298	1 280.7	1.281	2 072.2	2.072
<i>Austrovenus stutchburyi</i>	284.8	0.570	1 224.2	2.448	1 980.8	3.962
<i>Prothaca crassicostrata</i>	1.2	0.001	5.2	0.005	8.3	0.008
<i>Dosinia</i> ?sp.	2.0	0.004	8.8	0.018	14.2	0.028
<i>Cellana ornata</i>	0.4	0.001	1.7	0.003	2.7	0.005
<i>Cellana strigilis</i>	7.6	0.015	32.5	0.065	52.5	0.105
<i>Patelloida corticata</i>	1.7	0.002	7.2	0.007	11.7	0.012
<i>Haliotis iris</i>	3.5	0.525	15.0	2.255	24.3	3.648
<i>Haliotis australis</i>	0.7	0.069	3.0	0.296	4.8	0.479
<i>Diloma aethiops</i>	24.3	0.049	104.5	0.209	169.1	0.338
<i>Diloma</i> ?sp	1.0	0.015	4.4	0.066	7.1	0.107
<i>Cookia sulcata</i>	4.2	0.062	17.8	0.268	28.9	0.433
<i>Lunella smaragdus</i>	4.1	0.017	17.8	0.071	28.8	0.115
<i>Maoricrypta</i> ?sp	2.6	0.003	11.4	0.011	18.4	0.018
<i>Argobuccinum pustulosum tumidum</i>	1.0	0.002	4.4	0.009	7.1	0.014
<i>Paratrophon patens</i>	0.4	0.000	1.7	0.002	2.7	0.003
<i>Amphibola crenata</i>	78.6	0.079	337.7	0.338	546.5	0.546
<i>Siphonaria</i> ?sp	1.0	0.001	4.2	0.004	6.9	0.007
all others	4.0	0.004	17.3	0.017	28.0	0.028
TOTAL	850.9	2.116	3 657.1	9.094	5 917.5	14.714
				± 30%		± 30%

Table 15: Estimated numbers (N x 10³) and biomass (tonnes) of shellfish harvested in Otago-Catlins at ca. 1550 AD

	Minimum estimate		Best estimate		Maximum estimate	
	N x 10 ³	Biomass	N x 10 ³	Biomass	N x 10 ³	Biomass
<i>Mytilus galloprovincialis</i>	1 246.0	3.738	5 355.5	16.066	9 012.1	27.036
<i>Perna canaliculus</i>	25.1	0.100	107.9	0.432	181.6	0.727
<i>Aulacomya maoriana</i>	3.8	0.008	16.2	0.032	27.2	0.054
<i>Ostrea chilensis</i>	2.6	0.005	11.1	0.022	18.7	0.037
<i>Purpurocardia</i> ?sp	36.4	0.036	156.4	0.156	263.2	0.263
<i>Paphies australis</i>	402.1	0.402	1 728.1	1.728	2 908.0	2.908
<i>Paphies subtriangulata</i>	14.8	0.030	63.6	0.127	107.0	0.214
<i>Austrovenus stutchburyi</i>	269.5	0.539	1 158.5	2.317	1 949.6	3.899
<i>Cellana strigilis</i>	13.0	0.026	55.7	0.111	93.8	0.188
<i>Notoacmea elongata</i>	3.6	0.004	15.3	0.015	25.8	0.026
<i>Patelloida corticata</i>	11.5	0.011	49.3	0.049	83.0	0.083
<i>Haliotis iris</i>	22.0	3.298	94.5	14.177	159.0	23.857
<i>Haliotis australis</i>	1.0	0.099	4.2	0.425	7.1	0.715
<i>Cantharidus</i> ?sp	14.9	0.015	64.0	0.064	107.6	0.108
<i>Diloma aethiops</i>	43.6	0.087	187.2	0.374	315.1	0.630
<i>Diloma subrostrata</i>	3.5	0.004	15.2	0.015	25.6	0.026
<i>Diloma</i> ?sp	12.7	0.190	54.5	0.818	91.7	1.376
<i>Cookia sulcata</i>	13.9	0.208	59.7	0.896	100.5	1.507
<i>Lunella smaragdus</i>	35.3	0.141	151.9	0.608	255.6	1.023
<i>Maoricrypta</i> ?sp	3.2	0.003	13.6	0.014	22.8	0.023
<i>Amphibola crenata</i>	977.3	0.977	4 200.5	4.201	7 068.5	7.069
<i>Siphonaria</i> ?sp	4.0	0.004	17.0	0.017	28.6	0.029
all others	27.3	0.080	117.5	0.344	197.8	0.579
TOTAL	3 186.9	10.007	13 697.6	43.009	23 050.1	72.375
		3.002	12.903	0.008	21.713	0.009
		0.024	0.103	0.005	0.174	0.007
		0.001	0.005	0.002	0.007	0.007
		0.001	0.005	0.001	0.001	0.001
		0.057	0.245	0.015	0.015	0.015
		0.063	0.269	0.015	0.015	0.015
		0.042	0.182	0.015	0.015	0.015
		0.001	0.004	0.001	0.001	0.001
		0.293	1.260	0.005	0.005	0.005
		0.001	0.005	0.001	0.001	0.001
		0.024	0.103	0.005	0.005	0.005
		3.002	12.903	0.008	0.008	0.008
		0.001	0.005	0.001	0.001	0.001
		0.004	0.017	0.001	0.001	0.001
		0.080	0.344	0.005	0.005	0.005
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0.001	0.001	0.001
		10.007	43.009	0.008	0.008	0.008
		3.002	12.903	0.008	0.008	0.008
		0.007	0.026	0		

Table 17: Estimated numbers (N) and biomass (tonnes) of marine and coastal birds harvested in Greater Hauraki at ca. 1400 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass	N	Biomass	N	Biomass
<i>Eudiptula minor</i>	714	0.785	4 248	4.673	6 340	6.974
Diomedidae ?sp	94	0.425	562	2.530	839	3.775
<i>Macronectes halli</i>	30	0.134	177	0.798	264	1.190
<i>Pterodroma macroptera</i>	115	0.057	682	0.341	1 018	0.509
<i>Pachyptila vitata</i>	51	0.010	306	0.061	457	0.091
<i>Puffinus griseus</i>	177	0.142	1 054	0.844	1 574	1.259
<i>Puffinus gavia</i>	327	0.098	1 944	0.583	2 901	0.870
<i>Puffinus assimilis</i>	144	0.029	854	0.171	1 274	0.255
<i>Pelecanoides urinatrix</i>	181	0.024	1 076	0.140	1 606	0.209
<i>Morus serrator</i>	8	0.017	45	0.103	67	0.154
<i>Phalacrocorax melanoleucos</i>	48	0.034	288	0.202	430	0.301
<i>Phalacrocorax carbo</i>	102	0.225	609	1.339	908	1.998
<i>Phalacrocorax varius</i>	191	0.383	1 139	2.277	1 699	3.398
<i>Stictocorbo punctatus</i>	555	0.667	3 305	3.966	4 932	5.918
<i>Calidras canutus rogersi</i>	4	0.000	23	0.002	35	0.003
<i>Charadrius obscurus</i>	4	0.001	23	0.003	35	0.005
<i>Anarhynchus frontalis</i>	4	0.000	23	0.001	35	0.002
<i>Larus dominicanus</i>	202	0.171	1 199	1.019	1 790	1.521
<i>Larus novaehollandiae</i>	4	0.001	23	0.006	35	0.009
<i>Hydroprogne caspia</i>	25	0.017	149	0.104	222	0.155
<i>Chlidonias albobstriata</i>	25	0.002	149	0.012	222	0.018
<i>Sterna striata</i>	4	0.001	23	0.004	35	0.006
TOTAL	3 009	3.223	17 902	19.178	26 716	28.620
		0.967		5.753		8.586

Table 18: Estimated numbers (N) and biomass (tonnes) of marine and coastal birds harvested in Greater Hauraki at ca. 1550 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass	N	Biomass	N	Biomass
<i>Eudiptula minor</i>	1 753	1.928	4 639	5.103	6 694	7.364
Diomedidae ?sp	1 480	6.659	3 916	17.622	5 651	25.430
<i>Pterodroma macroptera</i>	823	0.411	2 177	1.088	3 141	1.571
<i>Phalacrocorax melanoleucos</i>	370	0.259	979	0.686	1 413	0.989
<i>Stictocarbo punctatus</i>	892	1.070	2 360	2.832	3 405	4.086
<i>Larus dominicanus</i>	370	0.315	979	0.833	1 413	1.201
<i>Sterna striata</i>	370	0.059	979	0.157	1 413	0.226
TOTAL	6 057	10.701	16 030	28.320	23 132	40.867
				± 30%		± 30%
				8.496		12.260

Table 19: Estimated numbers (N) and biomass (tonnes) of marine and coastal birds harvested in Greater Hauraki at ca. 1750 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass	N	Biomass	N	Biomass
<i>Eudiptula minor</i>	4 392	4.831	7 819	8.601	10 929	12.022
Diomedidae ?sp	2 196	9.881	3 909	17.593	5 465	24.591
<i>Pterodroma macroptera</i>	4 392	2.196	7 819	3.909	10 929	5.465
<i>Puffinus gavia</i>	6 587	1.976	11 728	3.519	16 394	4.918
<i>Pelecanoides urinatrix</i>	2 196	0.285	3 909	0.508	5 465	0.710
Procellariidae ?sp	2 196	0.751	3 909	1.337	5 465	1.869
<i>Phalacrocorax melanoleucos</i>	2 196	1.537	3 909	2.737	5 465	3.825
<i>Stictocarbo punctatus</i>	6 587	7.905	11 728	14.074	16 394	19.673
<i>Larus dominicanus</i>	2 196	1.866	3 909	3.323	5 465	4.645
<i>Sterna striata</i>	2 196	0.351	3 909	0.626	5 465	0.874
TOTAL	35 132	31.579	62 552	56.226	87 435	78.593
				± 30%		± 30%
				16.868		23.578

Table 22: Estimated numbers (N) and biomass (tonnes) of marine and coastal birds harvested in Otago-Catlins at ca. 1750 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass	N	Biomass	N	Biomass
<i>Eudyptes pachyrrhynchus</i>	318	1.272	427	1.709	553	2.212
<i>Megadyptes</i> ?sp	1 272	6.680	1 709	8.972	2 212	11.614
<i>Diomedea exulans</i>	590	4.723	793	6.344	1 026	8.212
<i>Diomedea epomorpha</i>	405	3.239	544	4.350	704	5.631
<i>Thalassarche chrysostoma</i>	648	2.917	871	3.918	1 127	5.072
<i>Thalassarche bulleri</i>	70	0.014	94	0.019	121	0.024
<i>Thalassarche cauta</i>	3 403	15.311	4 570	20.565	5 916	26.621
<i>Pterodroma cookii</i>	824	0.107	1 107	0.144	1 433	0.186
<i>Pachyptila vitata</i>	1 031	0.206	1 385	0.277	1 793	0.359
<i>Puffinus gavia</i>	318	0.095	427	0.128	553	0.166
<i>Pelagodroma marina</i>	181	0.008	243	0.011	314	0.014
<i>Pelecanoides urinatrix</i>	496	0.065	667	0.087	863	0.112
<i>Phalacrocorax carbo</i>	916	2.015	1 230	2.706	1 592	3.503
<i>Phalacrocorax varius</i>	359	0.718	482	0.965	624	1.249
<i>Leucocarbo chalconotus</i>	4 506	11.265	6 052	15.131	7 835	19.586
<i>Sitotocarbo punctatus</i>	6 118	7.342	8 218	9.861	10 637	12.765
<i>Larus dominicanus</i>	1 627	3.578	2 185	4.806	2 828	6.222
<i>Larus novaehollandiae</i>	1 014	0.264	1 363	0.354	1 764	0.459
TOTAL	24 097	59.819	32 366	80.346	41 896	104.004
		17.946		24.104		31.201

Table 23: Estimated numbers (N) and biomass (tonnes) of marine mammals harvested in Greater Hauraki at ca. 1400 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass	N	Biomass	N	Biomass
<i>Arctocephalus forsteri</i>						
- pup	74	0.639	442	3.799	659	5.670
- juvenile	233	5.837	1 389	34.729	2 073	51.830
- sub adult male	202	20.166	1 200	119.979	1 791	179.055
- adult female	42	2.125	253	12.642	377	18.866
- adult male	127	19.111	758	113.705	1 131	169.692
- total	679	47.877	4 042	284.853	6 032	425.112
<i>Phocartus hookeri</i>	95	10.663	564	63.443	842	94.682
<i>Mirounga leonina</i>	1	0.790	5	4.700	7	7.013
<i>Globicephala ?sp</i>	7	9.668	42	57.522	63	85.845
Dolphin ?sp	8	0.667	47	3.971	70	5.926
Total	790	69.666	4 700	414.489	7 013	618.579

Table 24: Estimated numbers (N) and biomass (tonnes) of marine mammals harvested in Greater Hauraki at ca. 1550 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass	N	Biomass	N	Biomass
Dolphin ?sp	8	0.667	21	1.791	31	2.584
Total		0.200		0.537		0.775

Table 25: Estimated numbers (N) and biomass (tonnes) of marine mammals harvested in Greater Hauraki at ca. 1750 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass	N	Biomass	N	Biomass
Dolphin ?sp	14	1.198	25	2.133	35	2.981
Total		0.359		0.640		0.894

Table 26: Estimated numbers (N) and biomass (tonnes) of marine mammals harvested in Otago-Catlins at ca. 1400 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass ± 30%	N	Biomass ± 30%	N	Biomass ± 30%
<i>Arctocephalus forsteri</i>						
- pup	64	0.549	274	2.355	443	3.810
- juvenile	492	12.303	2 112	52.795	3 417	85.426
- sub adult male	191	19.140	821	82.135	1 329	132.899
- adult female	82	4.104	352	17.612	570	28.497
- adult male	128	19.140	548	82.135	886	132.899
- total	957	55.235	4 107	237.030	6 645	383.529
<i>Phocarcus hookeri</i>						
- pup	260	29.262	1 121	126.163	1 815	204.140
- juvenile	1	1.227	5	5.265	9	8.519
- adult female	1	0.153	5	0.658	9	1.065
- adult male	6	0.518	26	2.224	43	3.599
- total	1 226	86.396	5 265	371.341	8 519	600.852
Total		25.919		111.402		180.256

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

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass ± 30%	N	Biomass ± 30%	N	Biomass ± 30%
<i>Arctocephalus forsteri</i>						
- pup	22	0.192	96	0.826	162	1.390
- juvenile	215	5.374	924	23.098	1 555	38.869
- sub adult male	84	8.416	362	36.171	609	60.867
- adult female	30	1.509	130	6.484	218	10.911
- adult male	57	8.538	245	36.696	412	61.751
- total	409	24.028	1 756	103.274	2 955	173.788
<i>Phocartus hookeri</i>	77	8.664	331	37.240	557	62.666
<i>Hydrurga leptonyx</i>	0.5	0.061	2	0.263	4	0.442
<i>Globicephala</i> ?sp	0.5	0.665	2	2.858	4	4.810
Dolphin ?sp	2	0.207	11	0.888	18	1.494
Total	489	33.625	2 102	144.523	3 537	243.201

Table 28: Estimated numbers (N) and biomass (tonnes) of marine mammals harvested in Otago-Catlins at ca. 1750 AD.

	Minimum estimate		Best estimate		Maximum estimate	
	N	Biomass ± 30%	N	Biomass ± 30%	N	Biomass ± 30%
<i>Arctocephalus forsteri</i>						
- juvenile	555	13.880	745	18.613	964	24.093
- sub adult male	218	21.802	292	29.235	378	37.843
- adult male	145	21.756	194	29.173	252	37.763
- total	918	57.437	1 231	77.021	1 594	99.700
<i>Globicephala</i> ?sp	24	32.772	32	44.018	42	56.979
Dolphin ?sp	5	0.390	6	0.523	8	0.677
Total	947	90.600	1 270	121.562	1 644	157.356

Table 29: Taxa harvested at *ca.* 1750 AD as a proportion of taxa harvested at *ca.* 1400 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

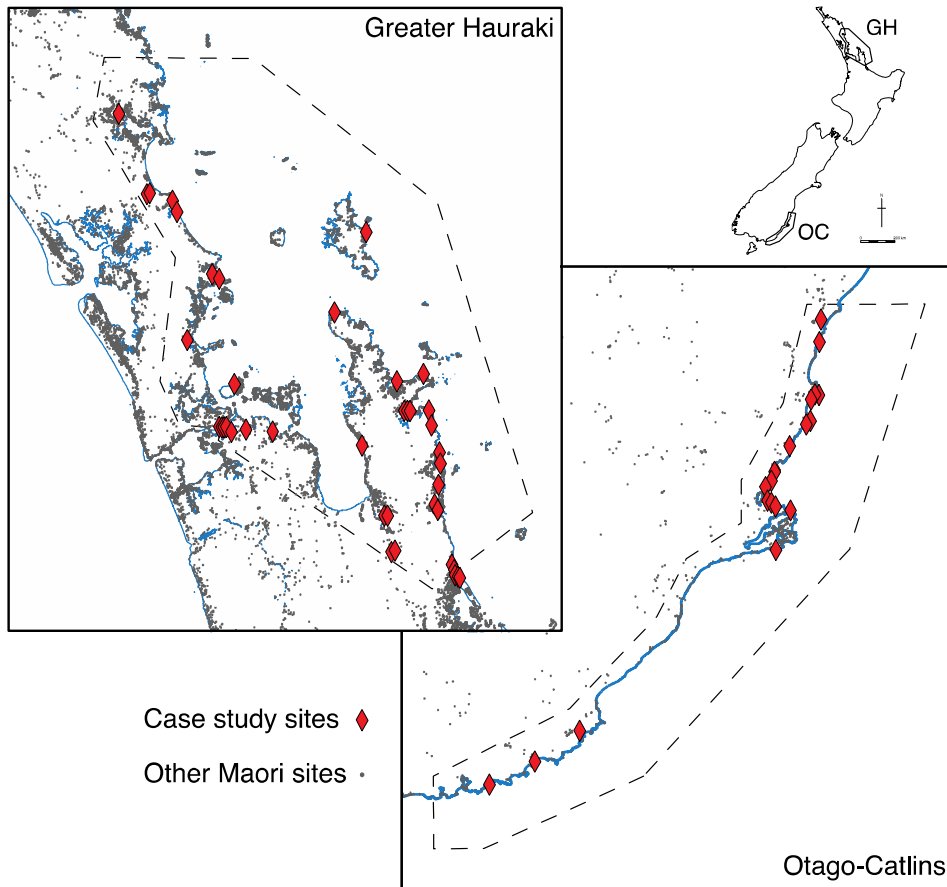


Figure 1: Greater Hauraki and Otago-Catlins, showing location of study sites and other pre-European Maori sites.

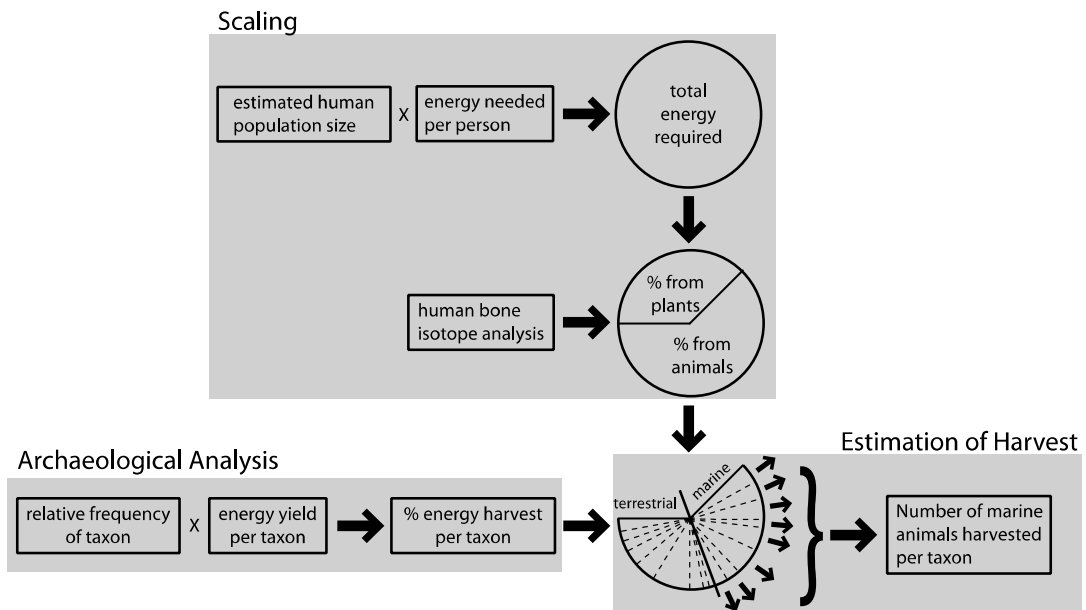


Figure 2: Simplified model of analytical procedure.

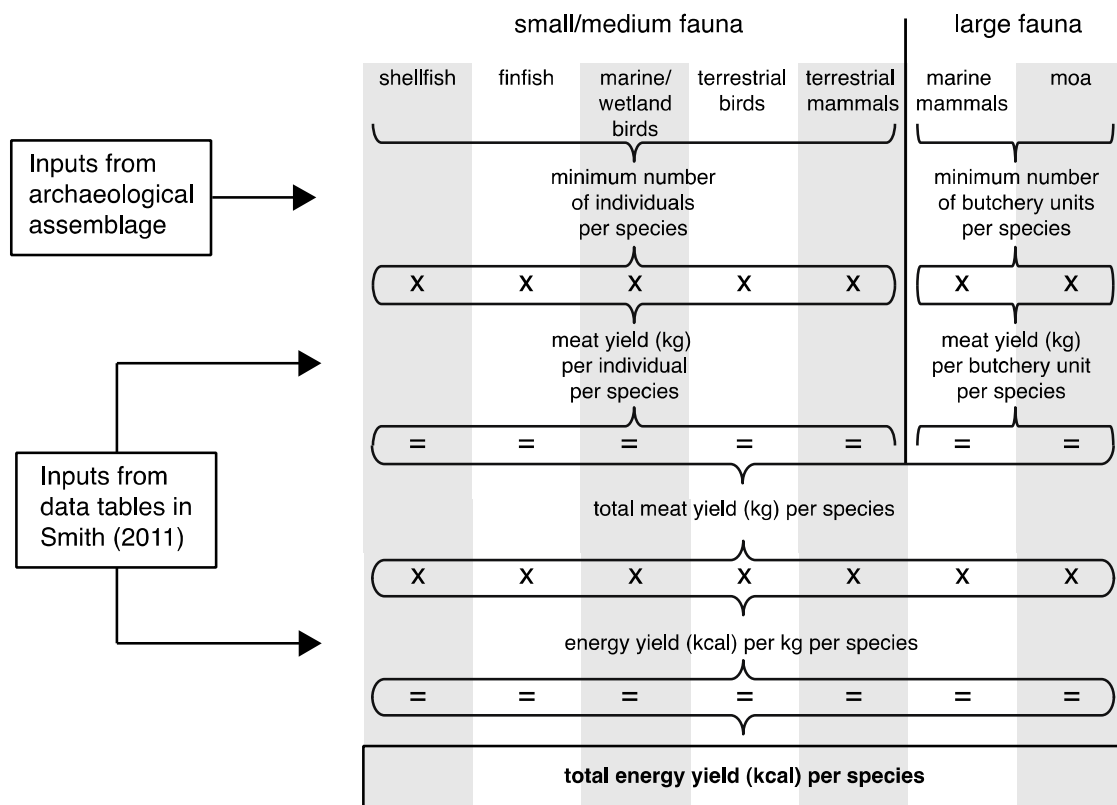


Figure 3: Procedures used in determining energy yield per species in study assemblages.

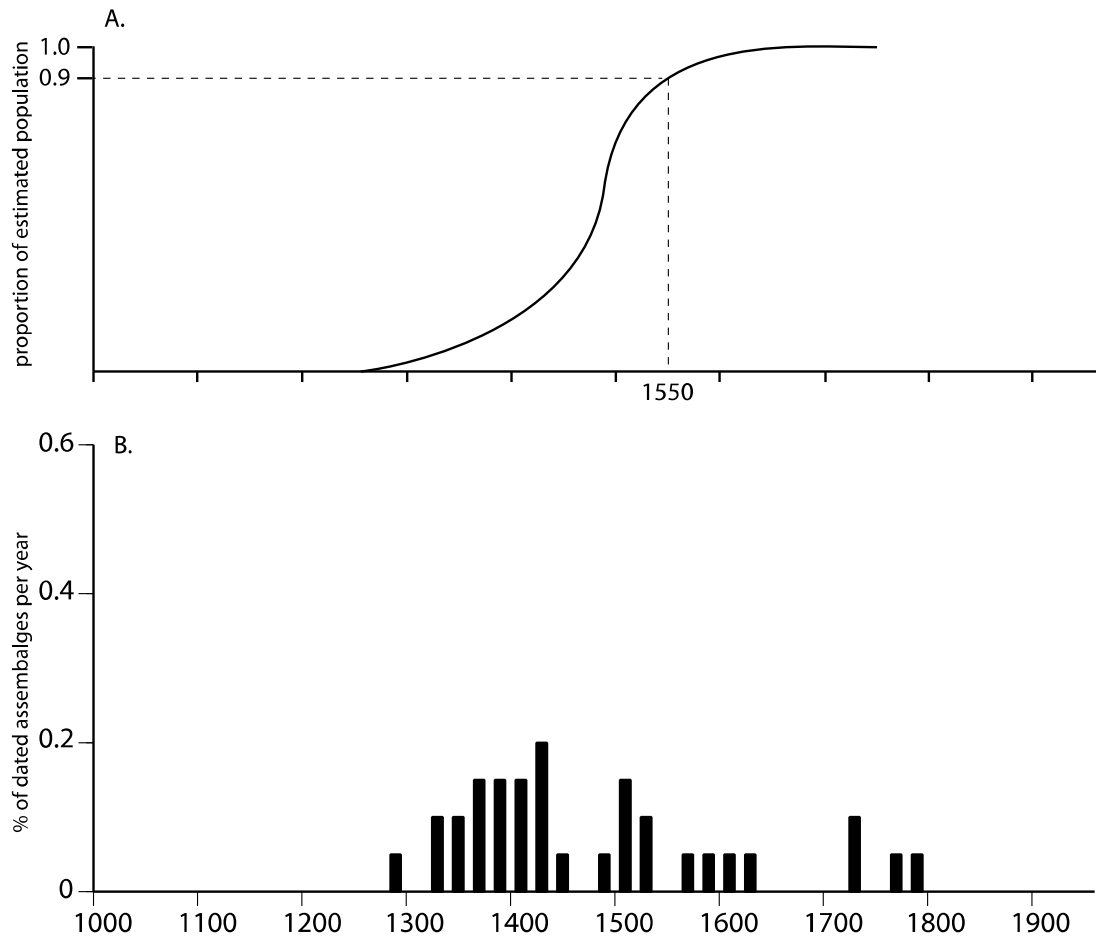


Figure 4: (A) Estimated growth rate for total New Zealand population based on cumulative frequency of radiocarbon dates (after McFadgen *et al.* 1994: Figure 4) showing proportion of population at *ca.* 1550 AD. (B) Frequency of radiocarbon dated assemblages in Otago-Catlins study dataset.

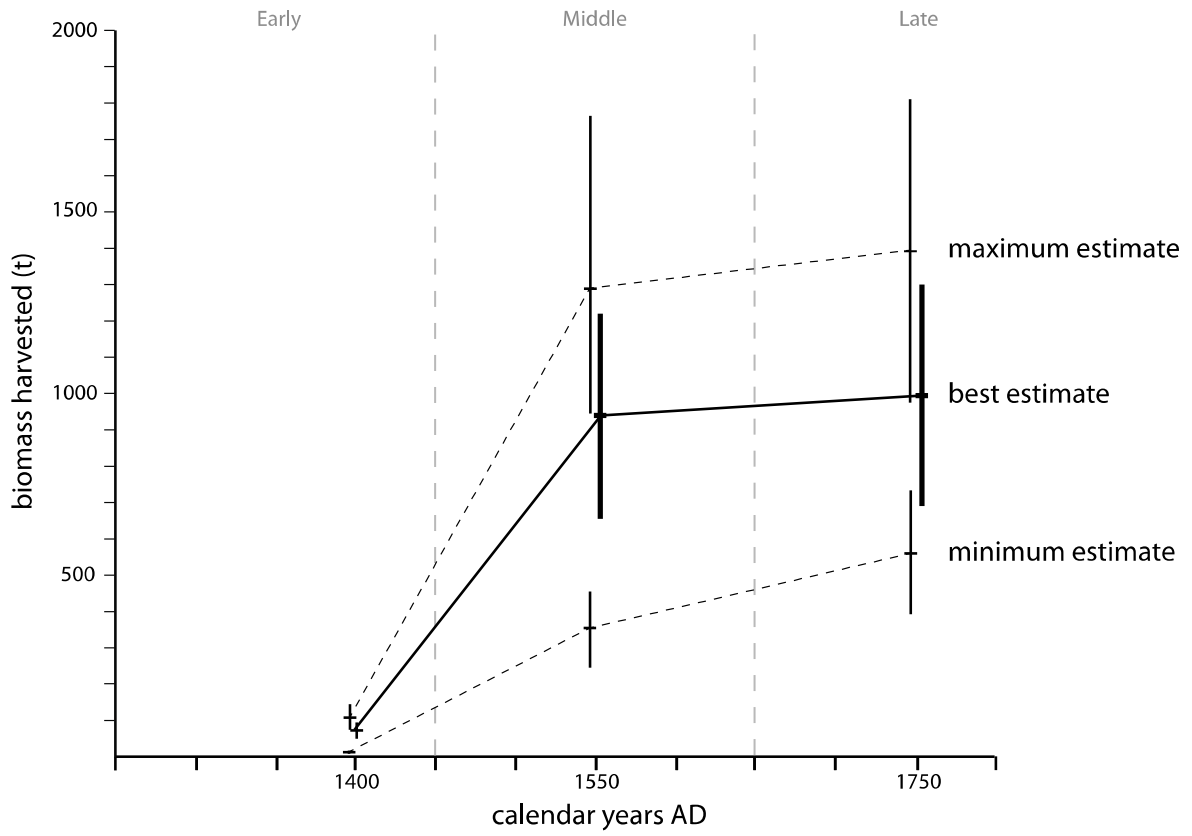


Figure 5: Minimum, best and maximum estimates for biomass of snapper harvested in Greater Hauraki.

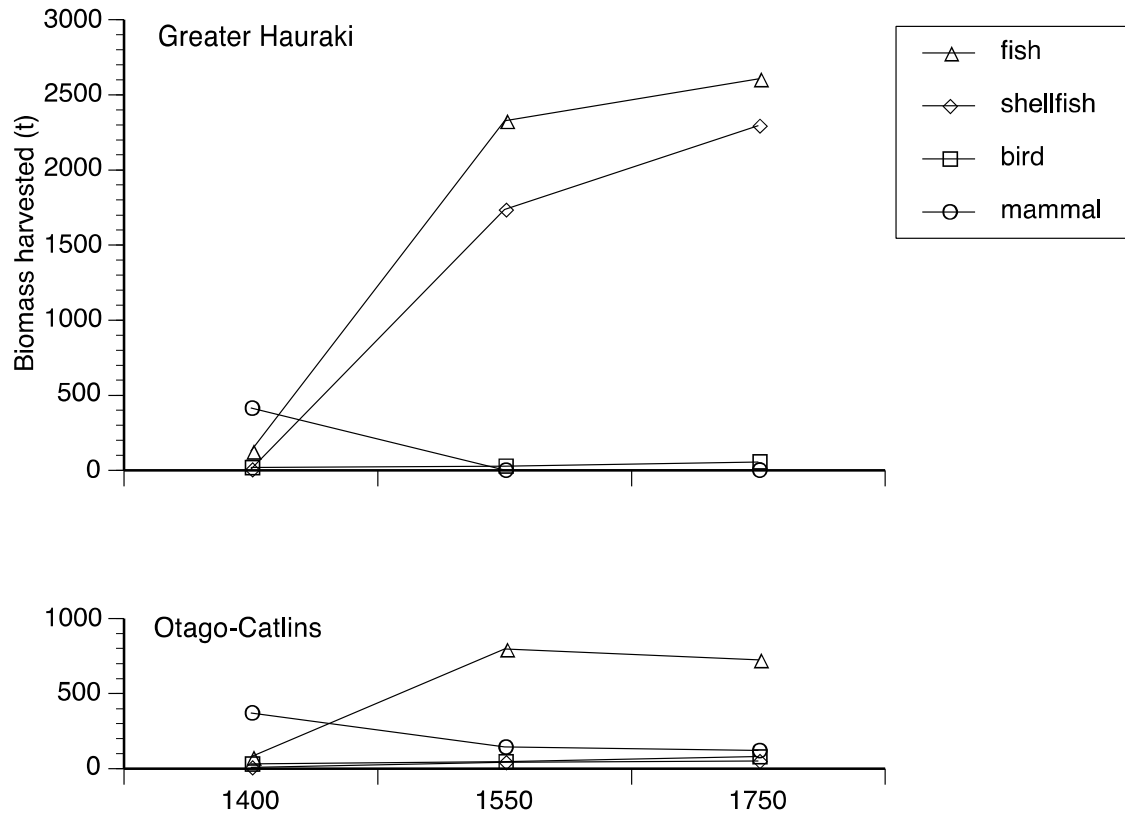


Figure 6: Best estimates for biomass harvested from four classes of marine fauna in each study area

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).

Appendix 1.1 Fish – Greater Hauraki

Taxon	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Squatulus blainvillei</i>						
Carcharhiniforme ?sp	0.19	0.2231	2.21	9.6544	2.08	0.8475
<i>Myliobatis tenuicaudatus</i>	0.02	0.0315			6.25	15.2029
Elasmobranchii ?sp	0.21	0.1443			2.08	6.7799
<i>Callorhynchus milii</i>	0.02	0.0066				
<i>Anguilla</i> spp	0.21	0.0104	1.33	0.2461	4.86	0.5008
<i>Pseudophycis bachus</i>	0.21	0.0131				
<i>Zeus faber</i>	0.41	0.0300				
<i>Chelidonichthys kumu</i>	0.72	0.0262	0.31	0.0420	2.08	0.1568
<i>Pseudocaranx dentex</i>	0.05	0.0049	2.91	1.0570	4.86	0.9829
<i>Trachurus declivis</i>	0.02	0.0017	0.05	0.0159	2.08	0.3688
<i>Trachurus novaezelandiae</i>	0.02	0.0013	27.92	6.8394		
Carangidae ?sp					2.08	0.2882
<i>Aripis trutta</i>	14.06	2.3744	10.94	6.8730	4.17	1.4587
<i>Pagrus auratus</i>	43.12	5.7160	36.88	18.1870	59.06	16.2171
<i>Nemadactylus macropterus</i>	3.51	0.2502				

Appendix 1.1 continued

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Latridopsis ciliaris</i>	0.62	0.1114				
<i>Mugil cephalus</i>	0.02	0.0018	1.33	0.4446		
<i>Aldrichetta forsteri</i>	4.26	0.0503	2.21	0.0971	2.08	0.0509
Labridae	11.81	0.9092	1.33	0.3809	2.08	0.3317
<i>Odax pullus</i>					2.08	0.1990
<i>Genyagnus monopterygius</i>			0.01	0.0009		
<i>Parapercis colias</i>			0.01	0.0012		
<i>Thyrsites atun</i>	0.93	0.0301	6.36	3.2091	2.08	0.5844
<i>Scomber australasicus</i>	0.87	0.1180	2.66	0.9502		
<i>Serollela brama</i>	0.19	0.0395				
<i>Meuschenia scaber</i>	18.53	0.6459			2.08	0.1502
<i>Rhombosolea ?sp</i>			3.54	0.1932		
Total	100	10.74	100	48.19	100	44.12

Appendix 1.2 Fish – Otago-Catlins

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Elasmobranchii</i> ?sp	0.010	0.0029	0.31	0.8597	0.10	0.1925
<i>Callorhynchus milii</i>					0.10	0.0926
<i>Anguilla</i> spp	0.010	0.0066				
<i>Pseudophycis bachus</i>	13.359	0.3474	25.98	6.5376	19.93	3.4820
<i>Lotella rhacinus</i>	0.010	0.0002				
<i>Genypterus blacodes</i>	4.572	0.4320	4.14	3.7856	4.97	3.1553
<i>Helicolenus barathris</i>	0.040	0.0002	0.04	0.0023		
<i>Scorpaena papillosus</i>			0.04	0.0058		
<i>Scorpaena cardinalis</i>	0.408	0.0061	0.60	0.0864		
<i>Neophrynichthys latus</i>			0.04	0.0170		
<i>Chelidonichthys kumu</i>	0.005	0.0001				
<i>Polyprion oxygenios</i>	1.130	0.6379	2.14	11.6921	5.05	19.1564
<i>Trachurus declivis</i>	0.005	0.0002				
<i>Trachurus novaezelandiae</i>	0.204	0.0056				
<i>Pagrus auratus</i>	3.012	0.1665				
<i>Nemadactylus macropterus</i>	0.428	0.0127	0.06	0.0173		
<i>Latridopsis ciliaris</i>	0.348	0.0261			0.03	0.0151
<i>Latris lineata</i>	2.409	0.2006	0.77	0.6205		
<i>Aldrichetta forsteri</i>	0.010	0.0000				
Labridae	2.677	0.0859	6.11	1.8982	0.60	0.1294
Nototheniidae	6.618	0.2039	6.86	2.0459		
<i>Parapercis colias</i>	9.469	0.1277	5.20	0.6785	0.13	0.0118

Appendix 1.2 continued

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Thyrsites atun</i>	55.236	3.1239	47.71	26.1132	69.09	26.2549
<i>Rexia solandri</i>	0.030	0.0025				
<i>Hyperoglyphe antarctica</i>	0.010	0.0010				
Total	100.000	5.39	100.000	54.36	100.000	52.49

barracouta
gemfish
bluenose warehou

Appendix 1.3 Shellfish – Greater Hauraki

Taxon	ca 1400		ca 1550		ca 1750	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Linucula hartvigiana</i>	0.008	0.000010				
<i>Mytilus galloprovincialis</i>	0.718	0.002722	0.001	0.000642		
<i>Perna canaliculus</i>	15.965	0.080051	0.029	0.043792	0.088	0.153952
<i>Linnoperna pulex</i>	0.004	0.000006				
<i>Barbatia novaezealandiae</i>	0.002	0.000002				
<i>Glycermis modesta</i>	0.001	0.000001				
<i>Tugetona laticostata</i>	0.015	0.000019				
<i>Ostrea chilensis</i>	0.080	0.000101	0.028	0.026970	0.001	0.001138
<i>Saccostrea cucullata glomerata</i>	8.201	0.020743	0.086	0.032395		
oyster ?sp.			0.030	0.022491		

nut shell
blue mussel
green-lipped mussel
little black mussel
ark shell
small dog cockle
large dog cockle
mud oyster/bluff oyster
rock oyster

Appendix 1.3 continued

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Anomia trigonopsis</i>	0.043	0.000054				
<i>Pecten novaezelandiae</i>	0.055	0.000073	0.085	0.033485	0.011	0.005141
<i>Pecten</i> ?sp.	0.022	0.000028	0.093	0.036623		
<i>Talochlamys zelandiae</i>	0.002	0.000002				
<i>Divalucina cumingi</i>	0.062	0.000079				
<i>Cardita aoteana</i>			0.069	0.026117		
<i>Purpurocardia purpurata</i>	0.025	0.000032				
<i>Purpurocardia</i> ?sp.			0.106	0.040190	1.356	0.597315
<i>Mytila stowei</i>	0.002	0.000002				
<i>Cyclomactra ovata</i>			0.006	0.004210		
<i>Mactra discors</i>			0.003	0.003548	0.583	0.770092
<i>Mactra</i> ?sp.			0.007	0.004937		
<i>Paphies australis</i>	2.761	0.001796	34.385	6.696830	40.687	9.217530
<i>Paphies subtriangulata</i>	9.201	0.032120	5.146	5.377563	8.572	10.419895
<i>Paphies ventricosa</i>			0.018	0.027131		
<i>Diplodonta striatula</i>	0.001	0.000001				
<i>Felaniella zelandica</i>	0.004	0.000006				
<i>Austrovenus stutchburyi</i>	2.613	0.003566	52.954	21.632550	46.474	22.084135
<i>Protothaca crassicosata</i>	0.406	0.000513			0.001	0.000440
<i>Tawera spissa</i>	0.096	0.000121				
<i>Tawera</i> ?sp.	0.031	0.000039				
<i>Dosinia anus</i>	0.261	0.000660	0.028	0.010587		
<i>Dosinia subrosea</i>			0.016	0.011878	0.011	0.009873
<i>Dosinia</i> ?sp.			0.014	0.010593		
<i>Irus reflexus</i>	1.099	0.002779	0.001	0.000513	0.032	0.028095
<i>Myadora striata</i>	0.025	0.000031			0.032	0.014047

Appendix 1.3 continued

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Cellana denticulata</i>	12.177	0.030800				
<i>Cellana ornata</i>	0.028	0.000035				
<i>Cellana radians</i>	7.119	0.018006	0.124	0.094021		
<i>Asteracmea suteri</i>	0.001	0.000001				
<i>Patelloida corticata</i>	0.315	0.000398				
<i>Haliotis iris</i>	1.014	0.238822	0.000	0.031834	0.001	0.082054
<i>Haliotis australis</i>	0.013	0.002075				
<i>Haliotis</i> ?sp.	0.140	0.000178				
<i>Scutus breviculus</i>	0.013	0.000251	0.004	0.021550		
<i>Tugali elegans</i>	0.043	0.000054				
<i>Coelotrochus viridius</i>	0.016	0.000020				
<i>Cantharidus</i> ?sp.	0.003	0.000003				
<i>Diloma aethiops</i>	1.175	0.002972	1.478	1.119211	0.001	0.000881
<i>Diloma bicanaliculata</i>	0.001	0.000001				
<i>Diloma subrostrata</i>	0.317	0.000400				
<i>Diloma</i> ?sp	0.489	0.009279				
<i>Antisolarium egenum</i>	0.194	0.000245				
<i>Zethalia zelandica</i>			0.010	0.003876	0.351	0.154521
Trochidae ?sp.			0.003	0.000969		
<i>Calliostoma pellucidum</i>	0.002	0.000002				
<i>Calliostoma punctulatum</i>			0.004	0.001437		
<i>Cookia sulcata</i>	0.140	0.002664			0.001	0.006606
<i>Lunella smaragdus</i>	20.366	0.103023	2.090	3.165479	0.047	0.082961
<i>Herpetopoma bella</i>	0.002	0.000002				
<i>Nerita atramentosa</i>	2.673	0.003380				
<i>Zeacumantus lutulentus</i>	0.006	0.000008	0.031	0.011752	0.233	0.102740
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 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Zeacumantus subcarinatus</i>						
<i>Zeacumantus</i> ?sp.			0.005	0.002078		
<i>Maoricolpus roseus</i>	0.746	0.000943	0.063	0.023673		
<i>Maoricrypta</i> ?sp.			0.040	0.015299		
<i>Rissoina</i> ?sp.	0.003	0.000003	0.081	0.030801		
<i>Struthiolaria papulosa</i>			0.274	0.207	0.094	0.082912
<i>Maoricrypta costata</i>	1.536	0.003884				
<i>Maoricrypta monoxyla</i>	0.585	0.000740				
<i>Sigapatella novaezelandiae</i>	0.108	0.000272				
<i>Sigapatella tenuis</i>	0.131	0.000166				
<i>Lamellaria ophione</i>	0.003	0.000003				
<i>Argobuccinum pustulosum</i>			0.051	0.038758		
<i>Cabestana spengleri</i>			0.005	0.005814		
<i>Janthina janthina</i>			0.280	0.105870		
<i>Aethocola glans</i>	0.258	0.000653	0.004	0.002924	0.014	0.012337
<i>Buccinulum pallidum powelli</i>	0.001	0.000002				
<i>Buccinulum vittatum vittatum</i>	0.002	0.000002				
<i>Buccinulum</i> ?sp.			0.029	0.010833		
<i>Cominella adpersa</i>	0.008	0.000020	0.160	0.121263	0.083	0.072874
<i>Cominella glandiformis</i>	0.086	0.000108	0.013	0.004937	0.056	0.024682
<i>Cominella maculosa</i>	0.839	0.002122	0.023	0.017365		
<i>Cominella quoyana quoyana</i>	0.008	0.000010				
<i>Cominella virgata</i>	0.018	0.000022	0.042	0.015803	0.070	0.030844
<i>Cominella</i> ?sp.	0.124	0.000157	0.236	0.089501		
<i>Penion sulcatus</i>	0.001	0.000003	0.008	0.008580		
<i>Zemitrella</i> ?sp.	0.001	0.000001				

Appendix 1.3 continued

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Taron</i> ?sp.	0.005	0.000007	0.002	0.000676	0.013	0.005750
<i>Haustrum haustorium</i>	1.402	0.026605	0.018	0.100685	0.001	0.006606
<i>Haustrum scobina</i>	0.226	0.000286				
<i>Murexsul mariae</i>	0.002	0.000002				
<i>Murexsul octogonus</i>	0.002	0.000004				
<i>Xymene ambiguus</i>	0.064	0.000081	0.000	0.000169	0.193	0.084790
<i>Xymene plebeius</i>	0.055	0.000070				
<i>Xymene traversi</i>	0.070	0.000088				
<i>Dicathais orbita</i>	1.032	0.019570	0.018	0.100710	0.001	0.006606
<i>Austromitra rubiginosa</i>	0.043	0.000054				
<i>Alcithoe arabica</i>			0.041	0.046954	0.001	0.001321
<i>Amalda australis</i>	0.581	0.000735				
<i>Phenatoma rosea</i>	0.003	0.000003				
<i>Turbonilla zealandica</i>	0.065	0.000082				
Acteonidae ?sp	0.129	0.000163				
<i>Amphibola crenata</i>	3.871	0.004895	1.568	0.593590	0.102	0.044774
<i>Amurochiton glaucus</i>	0.022	0.000027				
<i>Antalis nana</i>	0.028	0.000036				
gastropod ?sp.			0.193	0.073	0.897	0.395088
Total	100	0.62	100	40.11	100	44.50

Appendix 1.4 Shellfish – Otago-Catlins

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Mytilus galloprovincialis</i>						
<i>Perna canaliculus</i>						
<i>Aulacomya maoriana</i>						
<i>Limnoperna pulex</i>						
<i>Modiolus areolatus</i>						
Mytilidae sp.						
<i>Barbatia novaezealandiae</i>						
<i>Glycermis modesta</i>						
<i>Ostrea chilensis</i>						
<i>Purpurocardia</i> ?sp.						
<i>Lasaea hinemoana</i>						
<i>Macra discors</i>						
<i>Zenatia acinaces</i>						
<i>Paphies australis</i>						
<i>Paphies subtriangulata</i>						
<i>Paphies ventricosa</i>						
<i>Macomona liliana</i>						
<i>Serratina charlottae</i>						
<i>Austrovenus stutchburyi</i>						
<i>Protothaca crassicosata</i>						
<i>Tawera spissa</i>						
<i>Tawera</i> ?sp.						
<i>Dosinia</i> ?sp.						
blue mussel	13.736	0.107401	39.098	1.154553	46.206	0.586416
green-lipped mussel	1.348	0.013925	0.788	0.030769	1.570	0.026332
ribbed mussel	0.047	0.000244	0.118	0.002324	0.297	0.002509
little black mussel			0.006	0.000062		
bearded mussel			0.003	0.000093		
ark shell	0.000	0.000000			2.250	0.009519
small dog cockle	0.001	0.000004	0.003	0.000028		
mud oyster	0.000	0.000001	0.081	0.002050	0.124	0.001357
false cockle ?sp			1.142	0.011240		
large trough shell	0.120	0.000940	0.004	0.000038		
scimitar maetra			0.003	0.000084		
pipi	35.018	0.046950	0.000	0.000004		
tuatua			12.616	0.063885	0.786	0.001711
toheroa			0.464	0.012617	0.767	0.008951
large wedge shell	0.007	0.000036				
wedge shell	0.003	0.000014				
cockle	33.473	0.094134	8.458	0.089839	2.481	0.011325
ribbed venus	0.141	0.000367				
morning star						
	0.240	0.001253				

Appendix 1.4 continued

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Ruditapes largillierti</i>	0.025	0.000064	0.028	0.000278		
<i>Myadora striata</i>			0.000	0.000004		
<i>Cellana ornata</i>	0.046	0.000121	0.031	0.000305		
<i>Cellana radians</i>	0.035	0.000182	0.023	0.000457	9.710	0.082152
<i>Cellana strigilis</i>	0.888	0.004627	0.407	0.008012	0.017	0.000140
<i>Cellana</i> ?sp.	0.020	0.000052	0.007	0.000064	10.629	0.044966
<i>Notoacmea elongata</i>	0.012	0.000030	0.112	0.001105		
<i>Notoacmea pileopsis</i>			0.019	0.000190		
<i>Notoacmea scopulina</i>	0.012	0.000030	0.004	0.000038		
<i>Patelloida corticata</i>	0.197	0.000515	0.360	0.003543		
<i>Radiacmea inconspicua</i>	0.004	0.000011	0.009	0.000084		
<i>Haliotis iris</i>	0.411	0.199726	0.690	1.266069	3.965	3.125533
<i>Haliotis virginea</i>	0.012	0.001880	0.004	0.002366		
<i>Haliotis australis</i>	0.081	0.026326	0.031	0.037352		
<i>Haliotis</i> ?sp.	0.032	0.000083	0.014	0.000140		
<i>Emarginula</i> ?sp			0.000	0.000002		
<i>Scutus breviculus</i>	0.012	0.000454	0.010	0.001441	3.283	0.208336
<i>Coelotrochus viridius</i>	0.012	0.000030	0.011	0.000107	0.008	0.000035
<i>Cantharidius sanguineus</i>			0.035	0.000343		
<i>Cantharidius</i> ?sp.			0.432	0.004257	0.008	0.000035
<i>Diloma aethiops</i>	2.858	0.014896	1.367	0.026917	0.017	0.000140
<i>Diloma arida</i>			0.031	0.000305		
<i>Diloma bicanaliculata</i>			0.004	0.000038		
<i>Diloma nigerrima</i>	0.012	0.000030	0.063	0.000622		
<i>Diloma subrostrata</i>	0.001	0.000004	0.111	0.001096		
<i>Diloma zelandica</i>	0.025	0.000064	0.025	0.000243		

Appendix 1.4 continued

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Amphibola crenata</i>	9.235	0.024070	30.666	0.301855	11.632	0.049210
<i>Benhamina obliquata</i>	0.001	0.000004	0.004	0.000080	0.008	0.000070
<i>Siphonaria australis</i>			0.000	0.000002		
<i>Siphonaria propria</i>	0.002	0.000006				
<i>Siphonaria</i> ?sp	0.116	0.000303	0.124	0.001219		
<i>Trimusculus conicus</i>			0.004	0.000040		
<i>Antalis</i> ?sp	0.001	0.000004			0.008	0.000035
gastropod ?sp.	0.032	0.000084				
<i>Patellacea</i> ?sp			0.047	0.000466	0.008	0.000035
Polyplacophora ?sp.	0.035	0.000091	0.178	0.001752	0.008	0.000035
Total	100	0.57	100	3.20	100	4.28

Appendix 1.5 Marine and Coastal Birds – Greater Hauraki

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Euodyptula minor</i>	23.73	1.014	28.94	0.270	12.50	0.382
Diomedeidae ?sp	3.14	0.549	24.43	0.933	6.25	0.782
<i>Macronectes halli</i>	0.99	0.173				
<i>Pterodroma macrotpera</i>	3.81	0.074	13.58	0.058	12.50	0.174
<i>Pachyptila vitata</i>	1.71	0.013				

Appendix 1.5 continued

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Puffinus griseus</i>	5.89	0.183				
<i>Puffinus gavia</i>	10.86	0.127			18.75	0.156
<i>Puffinus assimilis</i>	4.77	0.037				
<i>Pelecanoides urinatrix</i>	6.01	0.030			6.25	0.023
Procellariidae ?sp					6.25	0.059
<i>Morus serrator</i>	0.25	0.022				
<i>Phalacrocorax melanoleucos</i>	1.61	0.044	6.11	0.036	6.25	0.122
<i>Phalacrocorax carbo</i>	3.40	0.290				
<i>Phalacrocorax varius</i>	6.36	0.494				
<i>Stictocarbo punctatus</i>	18.46	0.860	14.72	0.150	18.75	0.626
<i>Calidras canutus</i>	0.13	0.000				
<i>Charadrius obscurus</i>	0.13	0.001				
<i>Anarhynchus frontalis</i>	0.13	0.000				
<i>Larus dominicanus</i>	6.70	0.221	6.11	0.044	6.25	0.148
<i>Larus novaehollandiae</i>	0.13	0.001				
<i>Hydroprogne caspia</i>	0.83	0.023				
<i>Childonias albobristata</i>	0.83	0.003				
<i>Sterna striata</i>	0.13	0.001	6.11	0.008	6.25	0.028
Total	100	4.16	100	1.50	100	2.50

Appendix 1.6

Marine and Coastal Birds – Otago-Catlins

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Eudyptes pachyrhynchus</i>	13.568	1.713	4.56	1.179	1.32	0.345
<i>Megadyptes</i> ?sp	10.664	1.767	4.07	1.381	5.28	1.811
<i>Eudyptula minor</i>	14.654	0.509	15.01	1.067		
<i>Diomedea exulans</i>	0.512	0.129			2.45	1.281
<i>Diomedea epomorpha</i>					1.68	0.878
<i>Thalassarche chrysostoma</i>					2.69	0.791
<i>Thalassarche bulleri</i>	0.443	0.003			0.29	0.004
<i>Thalassarche cauta</i>	6.644	0.943	4.76	1.384	14.12	4.152
<i>Pterodroma inexpecta</i>	1.587	0.016				
<i>Pterodroma cookii</i>	1.518	0.010			3.42	0.029
<i>Pachyptila vittata</i>	1.168	0.007			4.28	0.056
<i>Pachyptila turtur</i>	5.120	0.020	3.20	0.026		
<i>Puffinus griseus</i>	8.641	0.218	3.35	0.173		
<i>Puffinus tenuirostris</i>	1.152	0.022				
<i>Puffinus gavia</i>	1.888	0.018	4.69	0.091	1.32	0.026
<i>Pelagodroma marina</i>	0.606	0.001	2.55	0.115		
<i>Pelecanoides urinatrix</i>	6.189	0.025	2.35	0.007	0.75	0.002
<i>Pelecanoides georgicus</i>	2.688	0.010	13.04	0.110	2.06	0.017
<i>Phalacrocorax melanoleucos</i>	0.780	0.017	2.87	0.130		
<i>Phalacrocorax carbo</i>	3.641	0.230	3.14	0.406	1.49	0.195
<i>Phalacrocorax varius</i>	0.794	0.055	3.49	0.496	3.80	0.546
<i>Leucocorax chalconotus</i>	4.674	0.369	4.76	0.769	18.70	3.055

Appendix 1.6 continued

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Stictocarbo punctatus</i>	9.304	0.352	25.67	1.990	25.39	1.991
<i>Larus dominicanus</i>	1.632	0.044	2.49	0.137	6.75	0.970
<i>Larus novaehollandiae</i>					4.21	0.072
<i>Larus bulleri</i>	0.577	0.005				
<i>Childonias albostrigata</i>	0.309	0.001				
<i>Sterna striata</i>	1.247	0.006				
Total	100.00	6.49	100.00	9.46	100.00	16.22

Appendix 1.7

Marine Mammals – Greater Hauraki

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Arctocephalus forsteri</i>	9.40	0.5385				
<i>Arctocephalus forsteri</i>	29.56	5.0526				
<i>Arctocephalus forsteri</i>	25.53	17.9041				
<i>Arctocephalus forsteri</i>	5.38	1.9185				
<i>Arctocephalus forsteri</i>	16.13	18.1182				
<i>Arctocephalus forsteri</i>	86.00					
<i>Phocarcus hookeri</i>	12.00	9.4682				
<i>Mirounga leonina</i>	0.10	0.6887				
<i>Globicephala</i> ?sp	0.90	0.6571				
Dolphin ?sp	1.00	0.4542	100.00	0.05	100.00	0.05
Total	100	54.80	100	0.05	100	0.05

Appendix 1.8

Marine Mammals – Otago-Catlins

	ca 1400 AD		ca 1550 AD		ca 1750 AD	
	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution	Taxon frequency	Proportional energy contribution
<i>Arctocephalus forsteri</i>	5.20	0.3111	4.57	0.1090		
<i>Arctocephalus forsteri</i>	40.11	7.1595	43.96	3.1273	60.10	2.5242
<i>Arctocephalus forsteri</i>	15.60	11.4247	17.21	5.0233	23.60	4.0668
<i>Arctocephalus forsteri</i>	6.69	2.4912	6.17	0.9157		
<i>Arctocephalus forsteri</i>	10.40	12.1993	11.64	5.4418	15.70	4.3333
<i>Arctocephalus forsteri</i>	78.00		83.55		99.40	
<i>Phocarcus hookeri</i>	21.20	17.4680	15.75	5.1722		
<i>Mirounga leonina</i>	0.10	0.7192				
<i>Hydrurga leptonyx</i>	0.10	0.0899	0.10	0.0358		
<i>Globicephala</i> ?sp	0.10	0.0762	0.10	0.0304	0.10	0.4680
Dolphin ?sp	0.50	0.2371	0.50	0.0945	0.50	0.0558
Total	100.00	52.10	100.00	19.95	100.00	10.98

