

# Carbonate production of an emergent reef platform, Warraber Island, Torres Strait, Australia

DEIRDRE E. HART<sup>1</sup> (✉)

PAUL S. KENCH<sup>2</sup>

<sup>1</sup>*Department of Geography, University of Canterbury, Private Bag 4800 Christchurch 8020, New Zealand*

deirdre.hart@canterbury.ac.nz  
Phone +64 3 379 2987 ext 7916  
Fax +64 3 379 2907

<sup>2</sup>*School of Geography and Environmental Science, University of Auckland, Private Bag 92019, Auckland, New Zealand*

## Abstract

Complex relationships exist between tropical reef ecology, carbonate (CaCO<sub>3</sub>) production and carbonate sinks. This paper investigates census-based techniques for determining the distribution and carbonate production of reef organisms on an emergent platform in central Torres Strait, Australia, and compares the contemporary budget with geological findings to infer shifts in reef productivity over the late Holocene. Results indicate that contemporary carbonate production varies by several orders of magnitude between and within the different reef-flat sub-environments depending on cover type and extent. Average estimated reef flat production was  $1.66 \pm 1.78 \text{ kg m}^{-2} \text{ yr}^{-1}$  (mean  $\pm$  SD) although only 23% of the area was covered by carbonate producers. Collectively, these organisms produce  $17,399 \pm 18,618 \text{ t CaCO}_3 \text{ yr}^{-1}$ , with production dominated by coral (73%) and subordinate contributions by encrusting coralline algae (18%) articulated coralline algae, molluscs, foraminifera and *Halimeda* (<4%). Comparisons between these organisms production across the different reef flat zones, surface sediment composition and accumulation rates calculated from cores indicate that it is necessary to understand the spatial distribution, density and production of each major organism when considering the types and amounts of carbonate available for storage in the various reef carbonate sinks. These findings raise questions as to the reliability of using modal production rates in global models independent of ecosystem investigation, in particular, indicating that current models may overestimate reef productivity in emergent settings.

**Key words** *calcification, carbonate production, reef flat, Torres Strait, coral, coralline algae, Halimeda, foraminifera, molluscs*

## Introduction

Carbonate production by coral reefs located throughout the tropical and sub-tropical oceans is an important component of the global carbon cycle (Vecsei 2004). Recent carbonate studies have focussed on estimating global values of reef production to support climate modelling (Milliman 1993; Kleypas et al. 1999; Vecsei 2004). Such global estimates are dependent on

up-scaling from a small number of individual coral reef studies that represent limited coverage of the world's reefs.

At the reef platform scale carbonate production estimates are also of critical importance in understanding the geological and geomorphic development of coral reefs and islands. Production by primary frame builders (corals and encrusting coralline algae) is an important component in reef development (Hubbard et al. 1990). Furthermore, carbonate production by primary frame builders and secondary benthic organisms, along with mechanical and biological erosion, control the generation of detrital sediment on reef platforms, sediment which is subsequently: reincorporated into reef framework (Hubbard et al. 1990); stored on reef surfaces; transported off-reef (Hughes 1999); or transferred to infill lagoons (Macintyre et al. 1987; Kench 1998; Purdy and Gischler 2005) or build islands (Maragos et al. 1973; Hopley 1982; Woodroffe et al. 1999; Yamano et al. 2000, 2002). To date few studies have attempted to quantify how fine spatial variations in organism density and production influence the character of both reef framework and sediment reservoirs. Notable exceptions include Stearn et al. (1977) and Scoffin et al. (1980) in Barbados; Sadd (1984) and Hubbard et al. (1990) in the Caribbean; and Harney and Fletcher (2003) in Hawaii. Collectively these papers highlight the limited geographical and physiographic coverage of such studies. All were conducted in non-emergent, fringing reef environments potentially influenced by high-island silicate or hydrological inputs so that their results have limited applicability in interpreting the emergent, carbonate environments of Great Barrier Reef (GBR) platforms and Pacific atolls.

Vecsei (2004) identifies four principal approaches used to quantify carbonate production on reefs: i) hydrochemical techniques based on water chemistry changes (Davies and Kinsey 1977; Smith and Harrison 1977; Smith and Kinsey 1976, 1978; Smith 1981, 1983; Kinsey 1985); ii) the census-based approach, which uses data on reef organism cover and extension/production rates (Chave et al. 1972; Stearn et al. 1977; Sadd 1984; Hubbard 1985; Vecsei 2001; Yamano et al. 2000; Harney and Fletcher 2003); iii) geological estimates from net accumulations of carbonate on individual reefs (Ryan et al. 2001); and iv) modelling techniques focussed on net reef accumulation (Kleypas 1997). All these approaches yield aggregate estimates of production at the total reef scale but only the first two, hydrochemical and census-based methods, are applicable at sub-reef scales or in evaluating organism-level production differences.

Productivity rates calculated from hydrochemical (alkalinity-reduction) measurements alone include both the carbonate precipitation and early dissolution occurring in shallow reef waters (Kinsey 1985). These measurements commonly represent carbonate production by entire reef communities and not the relative contributions of different producer types. The widespread adoption of hydrochemical methods over the last three decades has led to significant advances in understanding the productivity of different reef-habitat assemblages (Kinsey 1983; Milliman 1993; Vecsei 2004). These advances have, however, been at the expense of detailed knowledge of the relative contributions of different organism types to gross reef production.

In contrast, census methods afford the opportunity to determine the relative contributions of different carbonate producers to total reef productivity as well as opportunities for detailed

spatial comparisons between carbonate contributions and sediment composition, and between patterns of 'framework' versus 'direct-sediment' production (Harney and Fletcher 2003) at sub-reef scales. This study documents detailed patterns of carbonate production on an emergent reef-flat as determined using census-based techniques.

Smith and Kinsey (1976) criticised census techniques for their potential for error from the accumulation of contributions by individual biological components. However, there have been sufficient, significant advances in the extent and accuracy of published carbonate producer growth rates to warrant reconsideration of the accuracy of census techniques, and a review of the rates that underlie them. Further, the robustness of census-based results may be easily tested via error estimates and comparisons drawn with hydrochemically-determined modal production rates.

This paper presents results from the application of census techniques to construct a high spatial and organism-type resolution budget of carbonate production (types, quantities and distribution) on the emergent reef flat of Warraber Reef, Torres Strait, Australia (10°12'S, 142°49'E). Estimates of production are evaluated via error analyses and comparisons with published estimates from other reef environments. Results highlight the improved spatial resolution that the census approach provides in understanding how several types of reef organisms contribute differently to sediment and framework sinks in reef platform environments. The implications of such differences are explored with regard to the geological development of reef platforms, global carbon budgets, and the generation of detrital sediments. While production on the platform slope is likely to rival that produced on the reef top, this paper focuses on the latter environment since the Warraber reef flat is a relatively closed system in terms of carbonate and sediment generation.

## **Materials and methods**

### **Field setting**

Torres Strait consists of a shallow (15-25 m deep) shelf with scattered islands, reefs and shoals, situated between north-eastern Australia and southern Papua New Guinea (Fig. 1). Reefs grow throughout the Strait fringing high islands and as large platforms and coral shoals (Woodroffe et al. 2000).

Warraber (Sue) Reef comprises a small cay and large platform system, with a total area of 11 km<sup>2</sup>, situated in the central Strait. A planar-type reef (Hopley 1982), Warraber is flanked by two parallel, slightly-deeper (1-2 m) platforms, Burrar (Bet) and Guijar (Poll) Reefs, which together are referred to as the Three Sisters (Fig. 1b).

Warraber Island comprises a 750 m by 1,500 m wide, oval-shaped, low-elevation (2-8 m above mean sea level, MSL) cay fringed by sandy beaches and situated towards the north-western corner of the reef platform (Fig. 1a). The island and surrounding reef flat are Holocene in origin, having formed over a shallow Pleistocene platform (Woodroffe et al. 2000). The present reef flat comprises two distinct areas separated by differences in gross elevation but both fringed by an elevated, youthful coral-algal rim (Fig. 1c): a large, elevated, central platform in the east, with extensive sand flats covering fossil microatolls and

branching coral; and smaller, lower, western reef flat, characterised by inner muddy sandflats and outer coral patches interspersed with sandy channels plus a boat channel, constructed in 1991, dividing the area in two. Woodroffe et al. (2000) interpret the western reef flat as more youthful than the central, emergent fossil reef flat.

## Climate and oceanographic regime

Strong tidal currents (up to  $4 \text{ ms}^{-1}$ ) scour the bed of Torres Strait affecting the form of reef development and the area is subject to wind-generated surges developed locally, in the Indian Ocean, and in the Coral Sea (Amin 1978). Torres Strait lies north of the main cyclone belt of the GBR, with the central Strait having experienced seven Category 1-2 cyclones since 1910 (Puotinen 2004).

Warraber Reef experiences a semi-diurnal, meso-tidal regime with a maximum range of 4 m above the lowest astronomical tide (ALAT). The entire reef flat is submerged at high tide, at which time offshore wave energy propagates across the platform. Conversely, at lower stages of the tide, the elevated central area and reef rim are largely exposed while the outer reef flat experiences ponding (Brander et al. 2004). Significant wave heights outside the platform range from 0-1 m during the wet season, when north-westerly winds prevail, to 0-2 m during the dry season, when south-easterly winds prevail (Young and Holland 1996).

## Methodology

Aerial photographs and initial field investigations were used to construct preliminary physiographic maps of the reef flat including a network of seven, 0.6-3.5 km long transects spaced around the island and radiating out to the reef edge. Transects were surveyed using a staff and level and, within sets of three  $1 \text{ m}^2$  quadrats at 37 sites along the transects, observations were made of sediment depth and type, and living organism type and planimetric cover or abundance (Appendix 1). Rugosity was gauged as the ratio between the length of chain required to cover the cross-sectional profile of a quadrat and the 1 m aerial width of the quadrat profile (with 1-4 indicating flat to very rugose surfaces respectively). Data from the transects were supplemented with additional quadrat surveys in each broad reef zone between them.

Coral species were identified using Veron (1986, 2000) and Wood (1983) and growth forms recorded. Molluscs were identified using Short and Potter (1987), Cernohorsky (1978), Hilton (1978, 1979) and Wilson and Gillett (1971) while foraminifera were identified using Jones (1994).

Analysis of variance (Single Factor ANOVA and Monte Carlo Randomisation Tests) was performed on the quadrat cover data for each organism type to compare inter- versus intra-site variation (Zar 1999). Cover types were mapped along RT1-RT7 and observations from intervening areas were used to classify the remaining reef flat. A map of ecological zones was constructed and the area occupied by each zone was calculated.

In this paper, the term 'calcification' refers to the potential carbonate production rates ( $\text{g m}^{-2} \text{ yr}^{-1}$ ) of individual organisms, while 'production' or 'carbonate production' refers to the estimated rates ( $\text{g m}^{-2} \text{ yr}^{-1}$ ) or amounts ( $\text{t yr}^{-1}$ ) of calcium carbonate ( $\text{CaCO}_3$ ) produced. A

review of published figures was conducted to determine the calcification rates, skeletal densities and aerial adjustment factors to apply to carbonate-producing organisms found on the reef flat in order to calculate per-quadrat production rates (Appendix 1).

Calculated potential production rates for organisms found in each quadrat (average cover >1%) were then summed and results from replicate quadrats were averaged to give gross carbonate production estimates per site ( $\text{g m}^{-2} \text{yr}^{-1}$ ). The quadrat data and calculated zonal areas were summed to produce gross carbonate production rates and standard deviations ( $\sigma$ ) ( $\text{g m}^{-2} \text{yr}^{-1}$ ). Maximum, minimum and best-estimate carbonate production figures were compared in order to test sensitivity to variation in species' production rates. This variation was then compared to that which resulted from the patchy nature of cover in survey quadrats (spatial variation).

## Results

### Reef flat morphology

Transects RT1 to RT6 show marked differences in reef flat width, elevation and topographic complexity around Warraber Island (Fig. 2). Reef flat width from island shore to reef rim ranges from 2,700 m to 600 m from east to west, with a 0.7-1 m difference in elevation between the elevated eastern (RT1-RT3) and lower western (RT4-RT6) transects. The broad eastern reef can be divided into three morphological components: an elevated inner-reef platform (0-1,000 m from shore, predominantly above MSL) that reflects higher reef growth during the mid-Holocene; a central basin (1,000-2,200 m from shore, predominantly below MSL); and a higher-elevation windward reef rim (2,200-2,700 m from shore, situated AMSL). In contrast, the narrower, deeper, western reef flats do not exhibit distinct morphological differences but rather possess more-varied local-scale topography, reflecting the presence of muddy sandflats versus large live microatolls, interspersed by sandy hollows and dense, branching-coral thickets.

### Spatial variability in living cover

The live cover data from quadrats along RT1-RT5 were analysed for spatial variability (Table 1): analysis of variance tests (Single Factor ANOVA and Monte Carlo Randomisation) indicate that for all cover types, except coral-massive and coral-fol/encr/mu, data from the three replicate quadrats at each site were more similar than any random combination of quadrats ( $P < 0.05$ ) (Table 1a-b). The non-significance of results for coral-massive ( $P > 0.1$ ) and coral-fol/encr/mu ( $P > 0.4$ ) reflects the highly-variable local distribution patterns of these cover types. For example, massive corals commonly covered 80-100% of quadrats they occupied, but 0% of adjacent replicate quadrats, which frequently contained 100% sandy substrate. Alone neither type of quadrat accurately represents local ecological cover but in combination they indicate the types and densities of organisms present. It was, thus, deemed suitable to group cover data from each set of three replicate quadrats into averaged 'per site' values.

Inter-transect variation was not greater than intra-transect variation (Table 1c) with P-values significant ( $< 0.05$ ) only for the category 'coral-ramose-other', indicating that variation

between sites across the reef flat as a whole was no greater than the variation found along each transect for all other cover types. Accordingly, it was deemed inappropriate to further group the cover data by transect. This finding indicates a pattern of ecological zones running across, rather than along, the reef transects. For the cover type 'coral-ramose-other' the higher inter-transect variation is consistent with the division of the reef flat into a series of elevated eastern and lower western zones, the latter of which is based primarily on variation in the cover of *Montipora digitata*. West of the boat channel *M. digitata* is sparse, occurring in outer reef flat zones as small to medium sized branching colonies while east of the boat channel *M. digitata* forms a number of tall, wide, dense, monospecific bands mid-way across the reef flat which are replaced by a moderately-tall, dense, mixed cover of *M. digitata* and *Acropora* species and massive coral colonies.

Based on the 'per site' census data and observations from intervening areas, ten ecological reef flat zones were determined according to percentage living cover and substrate type, each zone being characterised by a distinct combination of biological assemblages and substrate types (Fig. 3). All of the eastern zones were  $>1 \text{ km}^2$ , with those in the west and north  $<0.5 \text{ km}^2$  each, while the total area covered by the zones (i.e. excluding the boat channel and island) was  $10.46 \text{ km}^2$ . The island, boat channel and whole reef platform cover  $0.81 \text{ km}^2$ ,  $0.06 \text{ km}^2$  and  $11.33 \text{ km}^2$  respectively (Fig. 3).

## Reef flat carbonate productivity

Table 2 summarises the organism-level calcification rates used to calculate carbonate production on the Warraber reef flat. The results of these calculations are summarised for each zone in Table 3a, which shows the 'best estimate' carbonate production rates ( $\text{g m}^{-2} \text{ yr}^{-1}$ ) of the different assemblages of organisms with standard deviations indicating levels of inter-quadrat variation in each zone.

Productivity estimates vary between the ten zones by two orders of magnitude, from  $65 \text{ g m}^{-2} \text{ yr}^{-1}$  in Zone 5 to  $3,999 \text{ g m}^{-2} \text{ yr}^{-1}$  in Zone 6. Such large differences are expected and are due to spatial variability in live cover and the composition of carbonate-producing assemblages found in each zone (e.g., mollusc versus coral dominated). The most productive areas ( $1,764$ – $3,999 \text{ g m}^{-2} \text{ yr}^{-1}$ ) are located on the central to outer reef flat zones characterised by massive and mixed-branching/massive coral cover (Zones 3, 6, 9), while moderate amounts of carbonate ( $566$ – $1,081 \text{ g m}^{-2} \text{ yr}^{-1}$ ) are produced in the dense-branching and reef-rim zones (4, 7a-7b). The least productive areas of the reef flat ( $65$ – $161 \text{ g m}^{-2} \text{ yr}^{-1}$ ) are those dominated by sandy or muddy substrate located close to the island and on the elevated eastern reef flat (Zones 1-2, 5, Table 3a).

In addition to between-zone differences in estimated carbonate production rates, a large degree of within-zone variation was found as demonstrated by the standard deviations associated with each rate (Table 3a). This variation is explained by the patchy nature of the ecosystems. In Zone 3, for example, quadrats with 100% cover of highly-productive massive and branching corals occurred adjacent to quadrats containing only bare sand. Analysis of variance performed on the per quadrat carbonate-production data (One-way ANOVA and Monte Carlo Randomisation Tests) confirm that, despite the high degree of intra-zonal variation, the zonal grouping of quadrats is highly significant ( $P < 0.001$ , degrees of freedom between groups = 9 and within groups = 101,  $n = 111$ ).

## Total reef flat carbonate production

The average rate of estimated carbonate production for the reef flat as a whole was  $1,663 \pm 1,780 \text{ g m}^{-2} \text{ yr}^{-1}$  (mean  $\pm$  SD) (Table 3a) with the total area covered by carbonate-producing organisms  $2.41 \text{ km}^2$  or 23% of the total reef flat. These organisms produce an estimated  $17,399 \pm 18,618 \text{ t CaCO}_3 \text{ yr}^{-1}$  (mean  $\pm$  SD) (Table 3b). Total production varies by three orders of magnitude between zones, from  $30 \text{ t yr}^{-1}$  in Zone 5 to  $12,575 \text{ t yr}^{-1}$  in Zone 3, as a function of the area of each zone, as well as of the calcification and cover rates of organisms present.

Table 3b presents estimates of the amount of carbonate contributed annually by the different producer types, highlighting spatial variability in the importance of carbonate producing organisms on the reef surface. Production was dominated by molluscs in Zones 1-2, by coral in Zones 3, 6, 8 and 9 and by encrusting coralline algae in Zones 4 and 7b. Despite the importance of each of these organisms within individual zones, some are quantitatively of little importance to total production. The dominance of coral in Zone 3, for example, represents far more carbonate ( $11,120 \text{ t yr}^{-1}$ ) than the dominance of molluscs in Zones 1-2 ( $100\text{--}193 \text{ t yr}^{-1}$ ). Total production on the Warraber reef flat is dominated by Zone 3 (70% of total or  $12,575 \text{ t yr}^{-1}$ ) while the majority (87%) of carbonate produced on the reef flat is contributed by only three producers: massive corals (59%), encrusting coralline algae (18%), and branching *Acropora* (10%). Other producers contribute  $\leq 4\%$  each to total carbonate production (Table 3b).

Comparisons between the percentage of total carbonate produced by each type of organism and cover<sup>b</sup> (the cover an organism relative to the total area occupied by carbonate producing organisms,  $2.41 \text{ km}^2$ ) reveals a markedly non-linear relationship (Table 3) due to the differential growth and production rates in Table 2. Most notably production by massive, branching-*Acropora* and fol/encr/mu corals is large relative to their areal cover, whilst production by encrusting coralline algae, other branching corals, *Halimeda* and articulated coralline algae is small relative to their cover. Further, the dominant carbonate producer at each site was rarely the dominant cover type. This is due to the predominance of the non-carbonate producing brown algae as well as to the large amount of carbonate produced by corals per unit area compared to encrusting coralline algae.

## Sensitivity of production results to growth rates

The sensitivity of carbonate production results to variations in organism growth rates was modelled for the main carbonate producers using mean (best-estimate), minimum and maximum calcification rates (Table 2) which, as indicated in Appendix 1, are conservative and excessive production values respectively. By comparison with the mean scenario, the proportion of carbonate produced by individual organism types changes by  $<4\%$  and  $<2\%$  under the minimum and maximum scenarios respectively (Table 4) with estimated total carbonate produced under these scenarios  $7,726$  and  $26,347 \text{ t yr}^{-1}$  respectively, corresponding to average production rates of  $738$  to  $2,518 \text{ g m}^{-2} \text{ yr}^{-1}$ .

Such comparisons do not indicate the effect of each organism experiencing different growth conditions, some finding them average, others optimal or suboptimal. Under these circumstances greater variation in the proportion of carbonate produced by each organism

could be expected, with total production somewhere between 7,726 and 26,347 t yr<sup>-1</sup>. Using the maximum coral scenario but minimum scenario for other organisms, the proportion of carbonate produced by coral would increase 14%, while decreasing 8% for coralline algae and 0-4% for other organisms. Total production under this scenario is 12,763 t yr<sup>-1</sup>.

Potential variation in carbonate production with differing growth conditions may be contextualised relative to actual variation resulting from the patchy nature of reef ecosystems (adjacent quadrat cover variation) as indicated by standard deviations in Table 3. Spatial variation in the best-estimate carbonate production on Warraber (i.e.  $\pm$  SD) is of the order of  $\pm 18,618$  t yr<sup>-1</sup> ( $\pm 107\%$ ), or  $\pm 1,780$  g m<sup>-2</sup> yr<sup>-1</sup>. Thus the potential variation in carbonate production resulting from the patchy nature of reef ecosystems is greater than that which might result from variable growth conditions.

## Discussion

The census approach yields estimates of carbonate production by organism-type at a number of spatial scales, including the entire reef platform, eco-morphological zones on a reef platform, and within eco-morphological zones. Of interest are: how these results compare with carbonate rates calculated for reef platforms elsewhere; how the census approach compares with commonly-used alkalinity reduction and geological techniques; whether the census approach provides improved accuracy and resolution of carbonate productivity on reefs and; implications for interpreting reef-flat carbonate productivity, geological development of reef platforms and sediment budgets.

### Warraber carbonate productivity in global context

Table 5 contains examples of published gross carbonate-production estimates alongside those calculated for Warraber (for a comprehensive review of carbonate-production rates up to 1985 see Kinsey 1985). The estimates of Smith and Kinsey (1976) and Kinsey (1981) are given as examples of rates calculated using alkalinity-reduction techniques which, as noted, include both carbonate precipitation and early dissolution. In recognition of methodological differences, comparisons focus on the order of magnitude of estimates. The average estimated production rate for the Warraber reef flat, 1.66 kg m<sup>-2</sup> yr<sup>-1</sup>, is lower than the majority of estimates from other reef environments, which range between 0.8-30.5 kg m<sup>-2</sup> yr<sup>-1</sup>. The Warraber rate is, however, of the same order of magnitude as those from other reef-flat studies (c.4 kg m<sup>-2</sup> yr<sup>-1</sup>, Table 5).

A number of factors could contribute to the low production value for Warraber and reef flats in general. First, Warraber is an emergent reef flat where productivity is constrained across broad tracts of elevated reef. Second, it is important to note that Table 5 lists studies undertaken in different reef environments and where production was dominated by different organisms. On the Warraber reef flat, the total estimated carbonate production was dominated by coral (74%), with subordinate proportions produced by encrusting coralline algae (18%) and other organisms in minor proportions (Table 3b).

In reef flat environments, low production rates have also been reported by Yamano et al. (2000) on a coral, *Halimeda* and foraminifera dominated reef surface and by Eakin (1996) with production dominated by coralline algae (56%) and coral (44%). A few studies of other

environments have found similar low productivity values: for example, Stearn et al (1977) on Bellairs fore-reef slope where coral was responsible for 71%, and coralline algae for 29%, of production, and Hubbard et al. (1990) on a shelf-edge reef where coral comprised 93%, and coralline algae 7%, of production.

## **Comparison of census-based and alkalinity-reduction techniques**

From an extensive review of alkalinity-reduction studies, Kinsey (1983; 1985) proposed that a series of absolute carbonate production rates were applicable to reefs in the latitudinal range 23°S to 23°N: 4 kg m<sup>-2</sup> yr<sup>-1</sup> on high-energy Pacific coral/algal reef-flat and rim environments; 0.5 kg m<sup>-2</sup> yr<sup>-1</sup> in sheltered sandy back-reef environments; and 2 kg m<sup>-2</sup> yr<sup>-1</sup> in shallow coral environments.

For comparison with the census approach used here the Kinsey modes were applied to Warraber, with Zones 1, 2 and 5 classified as sandy back-reef; Zones 4, 7a and 7b as high-activity rim; and Zones 8 and 9 as shallow-coral environments. Due to the patchy nature of coral and sand cover in Zones 3 and 6 these were characterised as intermediate between Kinsey's sandy reef-flat and shallow-coral environments. Using these modes, total annual carbonate production for the entire reef flat is 16,540 t yr<sup>-1</sup>, with an average production rate of 1.58 kg m<sup>-2</sup> yr<sup>-1</sup>. At the level of the entire reef flat these results compare well with the census-based estimates, providing a first approximation of reef flat carbonate productivity.

Important differences exist, however, at the individual reef-zone scale. Fig. 4a-b compares the census results with those calculated using Kinsey's modes across each zone: major differences occur in zones characterised as reef rim (4, 7a and 7b) or sandy reef flat with patchy mixed coral cover (3 and 6). Zones 3 and 4, for example, contribute 24% and 52% of total carbonate produced when calculated using Kinsey's modes versus census-based estimates of 72% and 13%. These comparisons highlight the need for more-than-superficial classification of reef ecosystems when using alkalinity-reduction based modes to estimate carbonate productivity and raise questions of accuracy regarding their application in global models of reef productivity independent of ecosystem investigation.

Fig. 4a also differentiates the carbonate produced across the reef flat by 'framework' versus 'direct sediment' producers (after Harney and Fletcher 2003), a distinction not possible with alkalinity-reduction results. Direct sediment production is shown to comprise a minor proportion of total carbonate production on the reef flat (8%).

In addition to the above inter-zone insights, the census-based approach provides improved resolution of spatial variations in productivity within individual reef-flat zones. As outlined in Appendix 1, with the exception of molluscs, the contemporary growth rates of carbonate-producing organisms are well documented from a number of reefs and reef environments and show that growth rates vary according to environmental conditions and the age and health of organisms.

However, Scoffin and Garrett (1974) and Vecsei (2001, 2004) show that growth rates are sufficiently similar within species for slight variations in the cover of one species relative to another to drastically alter the constituent composition of carbonate sinks. Both the rates expressed in Table 2, and the comparisons made between productivity variations due to the

patchy nature of reef ecosystems and potential variation in growth conditions support this assertion. The potential for variation in carbonate sinks due to variation in species cover rates is ultimately expressed in the estimates summarised in Table 3, which show that productivity varies considerably between the different zones or sub-environments of the reef flat (0.065-3.99 kg m<sup>-2</sup> yr<sup>-1</sup>) depending on the cover of different producers.

For example, these figures are particularly sensitive to the presence of massive coral, which is highly-productive and largely limited to the deeper, central to outer reef flat (Zones 3, 6, 9). Within these zones estimated production rates (1.76-3.99 kg m<sup>-2</sup> yr<sup>-1</sup>) approach average rates published for other reef flats (Table 5). In contrast, estimated production rates for the moderately-productive, monospecific branching-coral Zone 8 and raised coralgall-rim Zones 4, 7a and 7b (0.57-1.08 kg m<sup>-2</sup> yr<sup>-1</sup>) are comparable to those recorded for sand areas (0.4-1.2 kg m<sup>-2</sup> yr<sup>-1</sup>) in Kinsey (1985). Estimates from the least-productive, sandy, inner reef-flat zones on Warraber (0.065-0.15 kg m<sup>-2</sup> yr<sup>-1</sup>) are amongst the lowest-recorded carbonate production rates.

## Geological context of the census-based findings

The spatial variations (intra-reef-flat) in carbonate production highlighted by the census-based estimates provide a basis for evaluating long-term changes in reef production from the mid-late Holocene at the sub-reef scale. Gross vertical framework accretion on Warraber may be estimated using average carbonate production rates for each type of framework builder (corals, encrusting coralline algae) present in a zone divided by their individual densities (listed in Appendix 1). This gives a gross vertical framework accretion rate for the reef flat as a whole of 1.15 mm yr<sup>-1</sup>, with rates varying from 2.25 to 6.34 mm yr<sup>-1</sup> on the outer reef flat, from 0.35 to 0.56 mm yr<sup>-1</sup> on the reef rim, and at 0.0 mm yr<sup>-1</sup> on the inner reef flat (Fig. 4c). Accumulation rates vary across the ten ecological zones, independent of zone size, depending on the types of organisms that dominate production and the densities of their skeletons. Alternatively, if gross accumulation is calculated according to Harney and Fletcher's (2003) average framework density (1.48 g cm<sup>-3</sup>), the reef flat average is 1.03 mm yr<sup>-1</sup>, a similar figure to the 1.15 mm yr<sup>-1</sup> estimate derived using detailed organism densities.

Harney and Fletcher (2003) and Hubbard et al. (1990) report average framework erosion rates (biological plus mechanical) of 27% and 21% respectively. Using a 25% erosion value for Warraber, the mean net vertical framework accumulation on the reef flat is 0.86 mm yr<sup>-1</sup>, the same order of magnitude as Harney and Fletcher's (2003) 0.60 mm yr<sup>-1</sup> rate for Kailua Bay and that calculated from Hubbard et al.'s (1990) results for Cane Bay, 0.61 mm yr<sup>-1</sup>, but less than Smith's (1983) 3 mm yr<sup>-1</sup> rate for Holocene margin reefs, Stearn et al.'s (1977) 11 mm yr<sup>-1</sup> rate for a rapidly-growing reef, and Buddemeier and Smith's (1988) 10 mm yr<sup>-1</sup> sustained-maximum consensus rate.

Using core samples from the inner reef flat and island, Woodroffe et al. (2000) indicate that the Holocene reef started to grow over Pleistocene foundations 6 m below the present reef surface around 6,700 yr ago, reaching its present elevation 5300 yr ago, when sea level was 0.8-1.0 m higher than today, thereafter ceasing vertical accumulation. 6 m of vertical framework accumulation over the 1,400 yr between 6,700 and 5,300 yr ago corresponds to a net vertical accumulation rate on the inner reef flat of 4.29 mm yr<sup>-1</sup>, including both framework and sediment material. Assuming the ratio of framework to sediment within the

reef is around 50:50, as roughly indicated by core composition, and consistent with Buddemeier and Smith (1988), the inner reef flat accumulated framework at an average net rate of  $2.14 \text{ m yr}^{-1}$  from 6,700 to 5,300 yr ago. Both the  $2.14 \text{ mm yr}^{-1}$  'framework' and  $4.29 \text{ mm yr}^{-1}$  'total' mid-Holocene accumulation rates for the inner-reef flat are well within the range of contemporary gross framework accumulation rates calculated for the outer reef flat but above those of the inner zones (Fig. 4c). The finding that the emergent inner-reef flat is not presently accumulating is consistent with Woodroffe et al.'s (2000) results.

The contrast between the contemporary lack of inner reef-flat accumulation and the core-derived mid-Holocene rates may be explained by intra-platform and regional changes in carbonate production conditions as constrained by sea level and pace of reef development. At the reef scale, Woodroffe et al. (2000) describe how the now-central zones 'caught up' with sea level about 5,300 yr ago, followed by stepwise extensions south up to 4,500 yr ago, and subsequent infilling of central areas with ongoing extensions northward. This morphological development would have induced intra-platform changes in physical and growing conditions with increasing distance to the rim for inner areas - at the same time as the regional 0.8-1 m fall in sea level led to the emergence of the central reef flat, a characteristic which is common on the fringing reefs of the inner GBR. Together, these changes would have caused the now-central zones to experience a succession from reef-rim, to lagoon, to outer reef-flat and, finally, to emergent inner-reef-flat environment. Results illustrated in Fig. 4 indicate that such a succession would have been accompanied by large lateral shifts in reef flat ecology and shifts in carbonate production and framework accumulation.

Determining the high-level of variation in framework accumulation rates that exists across the contemporary Warraber reef flat (Fig. 4c) was only made possible using census-based techniques. Results show that the largely-intertidal Warraber reef flat produces approximately two orders of magnitude less carbonate than typical back reef (subtidal) settings. Largely-intertidal reef surfaces are common in the Indo-Pacific, particularly where reefs accreted in keep-up or catch-up growth mode in the mid-Holocene, subsequently being emerged through relative sea-level fall in the late Holocene. Given the spatial extent of the emergent reef platform on Warraber ( $3.76 \text{ km}^2$  or 33 % of reef platform surface), results suggest that global estimates of carbonate productivity should be revised in light of the low productivity of these surfaces and the likelihood that such surfaces cover a substantial proportion of Indo-Pacific reefs.

Extending the census-based geological model into the future, it is possible that climate-change induced sea-level and storminess changes could lead to a partial reversal of Holocene changes in growth conditions across Warraber platform. The extent of this reversal will likely be determined by reef community response to the latter two physical factors, to temperature changes (Buddemeier and Smith 1988) and to the sediment accumulations now occupying the mid-Holocene growth surface.

Buddemeier and Smith (1988) establish an apparent global match between reef growth rates and sea level rise, questioning whether this match is coincidental or functional. The variation in growth rates found between the emergent-inner and deeper-outer reef flat zones on Warraber indicates that the match is, in large part, functional in this setting. This is consistent with Kinsey's (1981) suggestion that the currently-shallow and slow-growing Holocene reef

flats of the Pacific possess the same potential to increase growth rates in response to accelerated sea level rise as the currently fast-growing Holocene reefs of the Caribbean.

## Sediment Implications

The results of this study have significant implications for interpreting carbonate sinks and their relationship to detrital sediment budgets within reef systems. Fig. 5 contrasts the proportion of total carbonate production contributed by the different reef-flat producers with their occurrence as constituents in the surficial sediments of each zone and, ultimately, in the island beach (Hart 2003). The overall dominance of coral (74%) and small contribution of molluscs (4%) to reef-flat production contrast markedly with their representation in beach sediments (coral 8%, molluscs 55%). *Halimeda* is also over-represented in the beach (7%) relative to its production of carbonate (1%). Coralline algae (19% CaCO<sub>3</sub>, 16% sediment) and foraminifera (2% CaCO<sub>3</sub>, 5% sediment) are similar in terms of their carbonate contributions and beach sediment representation. Organism differences between carbonate contributions and surficial reef-flat sediment composition are slightly less than, but of a similar order of magnitude to, those for the island beach (Fig. 5). These differences may be explained by variations in framework accommodation space (Fig. 2), in the spatial distribution of 'direct sediment' and 'framework' production across the reef zones (Fig. 4a), in processes of sediment-particle production (Chave 1964) and taphonomic processes (Scoffin 1992), in particular transportability (Folk and Robles 1964; Maiklem 1968), and by spatial separation between carbonate production and sediment-deposition zones (Yamano et al. 2000; Purdy and Gischler 2005).

For example, gastropod tests immediately contribute to the detrital sediment reservoir upon organism mortality and they have higher turnover rates than coral, helping explain their high abundance in reef flat sediments. Furthermore, gastropod tests are of a suitable size for beach nourishment, are predominantly produced in areas close to the island (Zones 1, 2, 5) where there is little accommodation space and their skeletal architecture make them highly susceptible to transport (Maiklem 1968; Kench and McLean 1996). Together these factors could explain mollusc dominance of reef-flat and island-beach deposits (Fig. 5) despite their small contribution to total reef flat carbonate production (Table 3).

By contrast, most carbonate production and the vast majority of coral production on Warraber occurs on the outer reef flat (Fig. 4a), which has up to 1 m of accommodation space (Fig. 2) and, thus, potential for carbonate to be retained as framework. And, although not well quantified, coral has high durability properties related to its architecture (Chave 1964; Folk and Robles 1964; Scoffin 1987), implying relatively-slow conversion rates to sediment, contributing to the dilution of coral in surficial sediments by organisms with higher turnover and sediment-conversion rates (Scoffin 1992).

Furthermore, when coral is eventually broken down it may not be into particle sizes suitable for island-beach nourishment. Coral bioeroders observed on the Warraber reef flat comprised grazing gastropods (Zones 1, 4, 7a-7b), boring bivalves (Zone 3, 6) and *Echinometra* urchins (Zones 6 and 9). With the exception of bivalves, which can break off large skeletal blocks, these organisms tend to reduce coral to very fine sediment bypassing the sand sizes that comprise the island beach.

Mechanical erosion of branching coral is more likely to produce sediment suitable for island-beach maintenance on Warraber. West of the island delicate branches of *Montipora digitata* and *Seriatopora hystrix* were observed to be broken off and swept islandward from Zones 8-9 during storm-wave conditions. The amount of carbonate produced on Warraber by branching corals (12%) which may break into sand-sizes particles is, however, small versus that produced by microatolls (59%, Table 3) which likely erode into finer particle sizes and which must traverse up to 2 km of reef surface to contribute to the island deposits.

These initial comparisons indicate that the total amount of carbonate produced on the reef flat is a poor indicator of both the amount and type of carbonate available to be turned into sediment and contribute to sedimentary deposits on reef platforms (e.g., islands, sand aprons and reef-flat sand reservoirs). Clues as to the potential production of beach-nourishing sediment are provided by teasing out the distributions and types of carbonate produced in the different zones of the reef flat. It is recommended that the next step in understanding the relationship between the rates and types of carbonate produced, and the ultimate nature of sink deposits, is to make detailed comparisons between the types, amounts and distribution of carbonate production, and the types and amounts of material found in each reef sink.

Over the longer-term the shifts in reef top ecology discussed and subsequent changes in dominant producers (and rates of production) have major implications for the sediment reservoir and development of geomorphic deposits on reef surfaces. For example, reef islands are unconsolidated accumulations of reef sediment. The accumulation of such islands and their ongoing maintenance is directly dependent on the generation of reef sediments and their transport to island shorelines. However, shifts in reef top ecology and carbonate production as identified at Warraber indicate that sediment type and abundance has likely changed over the past 5,000 years. Such shifts may be critical in 'turning on' and 'turning off' reef island formation and in understanding future changes in reef island stability. As shown by and Yamano et al. (2000) ecological shifts in the late Holocene as a consequence of sea level fall leading to reef flat emergence allowed increased production of foraminifera on the Green Island reef surface, possibly triggering the late Holocene development of this foraminifera-rich island. In contrast, coral is the dominant constituent comprising many reef islands in the Indo-Pacific (Stoddart and Steers 1977). Of relevance is whether or not islands in settings with emergent reef surfaces, such as Warraber, which currently produce only small volumes of coral, are still able to supply sediment to islands in sufficient quantities to maintain island shorelines. In conclusion, the census-based approach examined in this paper has been shown to allow carbonate production values to be established at sub-reef flat scales, thereby providing critical information for evaluating changes in production and organism type available to contribute to the sediment reservoir at locations proximal to reef islands.

## **Acknowledgements**

Research was supported by an Australian Research Council Grant to CD Woodroffe, PJ Cowell, and RF McLean and University of New South Wales ADFA Postgraduate Research Scholarship to DEH. We thank RF McLean for conceptual advice and field assistance, B Billy, B Samosorn, RW Brander, A Coutts-Smith and GA Stewart for field assistance, W Anderson for statistical advice, T Billy, C Tamu and the people of Warraber Island and

Beverly and Bill Stephens for their help and hospitality, A Vecsei for supplying raw data from Vecsei (2001), and P Bealing and M Brosnan for assistance with Figures 1 and 3.

## References

- Amin M (1978) A statistical analysis of storm surges in Torres Strait. *Aust J Mar Freshw Res* 29:479-496
- Brander RW, Kench PS, Hart DE (2004) Spatial and temporal variations in wave characteristics across a reef platform, Warraber Island, Torres Strait, Australia. *Mar Geol* 207:169-184
- Buddemeier RW and Smith SV (1988) Coral reef growth in an era of rapidly rising sea level: predictions and suggestions for long-term research. *Coral Reefs* 7:51-56
- Cernohorsky WO (1978) *Tropical Pacific Marine Shells*. Pacific Publications (Aust.) Pty Ltd, Sydney
- Chave K (1964) Skeletal Durability and Preservation. In: Imbrie J, Newell N (eds) *Approaches to Palaeoecology*. John Wiley and Sons Inc, Sydney, pp 377-387
- Chave KE, Smith SV, Roy KJ (1972) Carbonate production by coral reefs. *Mar Geol* 12:123-140
- Davies PJ, Kinsey DW (1977) Holocene reef growth – One Tree Island, Great Barrier Reef. *Mar Geol* 24:1-11
- Eakin CM (1996) Where have all the carbonates gone? A model comparison of calcium carbonate budgets before and after the 1982-1983 El Niño at Uva Island in the eastern Pacific. *Coral Reefs* 15:109-119
- Folk R, Robles P (1964) Carbonate sands of Isla Perez, Alacran Reef Complex, Yucatan. *J Geol* 72:255-292
- Harney JN (2000) Carbonate Sedimentology of a Windward Shoreface: Kailua Bay, Oahu, Hawaiian Islands. PhD thesis, University of Hawaii, p274
- Harney JN, Fletcher CH (2003) A budget of carbonate framework and sediment production, Kailua Bay, Oahu, Hawaii. *J Sediment Res* 73:856-868
- Hart DE (2003) Eco-sedimentological environments of an inter-tidal reef platform, Warraber Island, Torres Strait. Ph.D. thesis, University of New South Wales, p220
- Hilton AG (1978) *Guide to Australian Shells*. Robert Brown and Associates Pty Ltd, Port Moresby
- Hilton AG (1979) *Guide to Shells of Papua New Guinea*. Robert Brown and Associates Pty Ltd, Port Moresby
- Hopley D (1982) *The Geomorphology of the Great Barrier Reef - Quaternary Development of Coral Reefs*. Wiley-Interscience Publication, John Wiley and Sons Ltd, New York
- Hubbard DK (1985) What do we mean by reef growth? *Proc 5th Int Coral Reef Symp* 6:433-438
- Hubbard DK, Miller AI, Scaturro D (1990) Production and cycling of calcium carbonate in shelf-edge reef systems (St Croix, U.S. Virgin Islands): Applications to the nature of reef systems in the fossil record. *J Sediment Petrol* 60:335-360
- Hughes TP (1999) Off-reef transport of coral fragments at Lizard Island, Australia. *Mar Geol* 157:1-6
- Jones RW (1994) *Challenger Foraminifera*. Oxford University Press
- Kench PS (1998) Physical controls on development of lagoon sand deposits and lagoon infilling in an Indian Ocean atoll. *J Coastal Res* 14:1014-1024

- Kench PS, McLean RF (1996) Hydraulic characteristics of heterogeneous bioclastic deposits: new possibilities for interpreting environmental processes. *Sedimentology* 43:531-540
- Kinsey DW (1981) The Pacific/Atlantic Reef growth Controversy. *Proc 4th Int Coral Reef Symp* 1:493-498
- Kinsey DW (1983) Standards of performance in coral reef primary production and carbon turnover. In: Barnes DJ (ed) *Perspectives on Coral Reefs*. Brian Clouster Publisher, ACT, Australia, pp 209-218
- Kinsey DW (1985) Metabolism, calcification and carbon production I. System level studies. *Proc 5th Int Coral Reef Symp* 6:505-526
- Kleypas JA (1997) Modeled estimates of global reef habitat and carbonate production since the last glacial maximum. *Paleoceanography* 12:533-54
- Kleypas JA, Buddemeier RW, Archer D, Gattuso JP, Langdon C, Opdyke BN (1999) Geochemical consequences of increased atmospheric carbon dioxide on coral reefs. *Science* 284:118-120
- Macintyre IG, Graus RR, Reinthal PN, Litter MM, Litter DS (1987) The Barrier Reef sediment apron: Tobacco Reef, Belize. *Coral Reefs* 6:1-12
- Maiklem WR (1968) Some hydraulic properties of bioclastic carbonate grains. *Sedimentology* 10:101-109
- Maragos JE, Baines GBK, Beveridge PJ (1973) Tropical Cyclone Bebe creates a new landform on Funafuti Atoll. *Science* 181:1161-1164
- Milliman JD (1993) Production and accumulation of calcium carbonate in the ocean: budget of a nonsteady state. *Global Biogeochem Cycles* 7:927-957
- Odum HT, Odum EP (1955) Trophic structure and productivity of windward coral reef community on Eniwetok Atoll. *Ecol Monogr* 25:291-320
- Puotinen ML (2004) Tropical cyclone impacts on reef communities: Modelling the disturbance regime in the Great Barrier Reef region, 1969-2003. PhD thesis, James Cook University, Townsville, p229
- Purdy EG, Gischler E (2005) The transient nature of the empty bucket model of reef sedimentation. *Sediment Geol* 175:35-47
- Ryan DA, Opdyke BN, Jell JS (2001) Holocene sediments of Wistari Reef: towards a global quantification of coral reef related neritic sedimentation in the Holocene. *Palaeogeogr Palaeoclimatol Palaeoecol* 175:173-184
- Sadd JL (1984) Sediment transport and CaCO<sub>3</sub> budget on a fringing reef, Cane Bay, St Croix, U.S. Virgin Islands. *Bull Mar Sci* 35:221-238
- Scoffin TP (1987) *Introduction to Carbonate Sediments and Rocks*. Blackwell, Glasgow
- Scoffin TP (1992) Taphonomy of coral reefs: a review. *Coral Reefs* 11:57-77
- Scoffin TP, Garrett P (1974) Processes in the formation and preservation of internal structure in Bermuda patch reefs. *Proc 2nd Int Coral Reef Symp* 2:429-448
- Scoffin TP, Stearn CW, Boucher D, Frydl P, Hawkins CM, Hunter IG, MacGeachy JK (1980) Calcium carbonate budget of a fringing reef on the west coast of Barbados. *Bull Mar Sci* 30:475-508
- Short JW, Potter DG (1987) *Shells of Queensland and the Great Barrier Reef*. Golden Press Pty. Limited, Drummoyne, NSW
- Smith SV (1981) The Houtman Abrolhos Islands: carbon metabolism of coral reefs at high latitudes. *Limnol Oceanogr* 26:612-621
- Smith SV (1983) Coral reef calcification. In: Barnes DJ (ed) *Perspectives on Coral Reefs*. Brian Clouster Publisher, ACT, Australia, pp 240-247

- Smith SV, Harrison JT (1977) Calcium carbonate production of the Mare Incognitum, the upper windward reef slope, at Enewetak Atoll. *Science* 197:556-559
- Smith SV, Kinsey DW (1976) Calcium carbonate production, coral reef growth and sea level change. *Science* 194:937-939
- Smith SV, Kinsey DW (1978) Calcification and organic carbon metabolism as indicated by carbon dioxide. In: Stoddart DR, Johannes RE (eds) *Coral Reefs: Research Methods*. UNESCO, Monographs on Oceanographic Methodology. Paris 5:469-484
- Stearn CW, Scoffin TP, Martindale W (1977) Calcium carbonate budget of a fringing reef on the west coast of Barbados. *Bull Mar Sci* 27:479-510
- Stoddart DR, Steers JA (1977) The nature and origin of coral reef islands. In: Jones OA, Endean R (ed) *Biology and geology of coral reefs*, vol 4, Geol 2. Academic Press, New York, pp 59-105
- Vecsei A (2001) Fore-reef carbonate production: development of a regional census-based method and first estimates. *Palaeogeogr Palaeoclimatol Palaeoecol* 175:185-200
- Vecsei A (2004) A new estimate of global reefal carbonate production including the fore-reefs. *Global Planet Change* 43:1-18
- Veron JEN (1986) *Corals of Australia and the Indo-Pacific*. Angus and Robertson Publishers, North Ryde, NSW, Australia
- Veron JEN (2000) *Corals of the World* (3 vol.). Australian Institute of Marine Science, Townsville, Australia
- Wilson BR, Gillett K (1971) *Australian Shells*. AH and AW Reed, Sydney
- Wood EM (1983) *Reef Corals of the World*. T.F.H. Publications, Neptune City
- Woodroffe C, McLean RF, Smithers SG, Lawson EM (1999) Atoll reef-island formation and response to sea-level change: West Island, Cocos (Keeling) Islands. *Mar Geol* 160:85-104
- Woodroffe CD, Kennedy DM, Hopley D, Rasmussen CE, Smithers SG (2000) Holocene reef growth in Torres Strait. *Mar Geol* 170:331-346
- Yamano H, Miyajima T, Koike I (2000) Importance of foraminifera for the formation and maintenance of a coral sand cay: Green Island, the Great Barrier Reef, Australia. *Coral Reefs* 19:51-58
- Yamano H, Kayanne H, Matsuda F, Tsuji Y (2002) Lagoonal facies, ages, and sedimentation in three atolls in the Pacific. *Mar Geol* 185:233-247
- Young IR, Holland GJ (1996) *Atlas of the Oceans: Wind and Wave Climate*. Elsevier Science Ltd, Oxford
- Zar JH (1999) *Biostatistical Analysis* 4<sup>th</sup> Edn. Prentice Hall, New Jersey

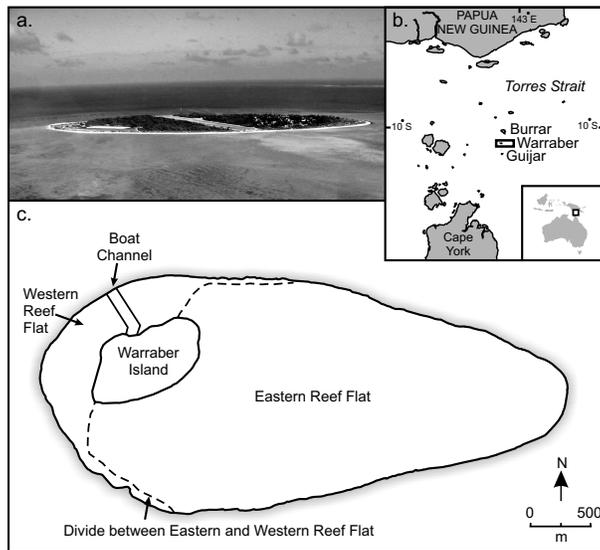


Fig. 1 Image and maps showing (a) Warraber Island, (b) its location in The Three Sisters reef group, Central Torres Strait, Australia, and (c) the main features of the Warraber Reef platform

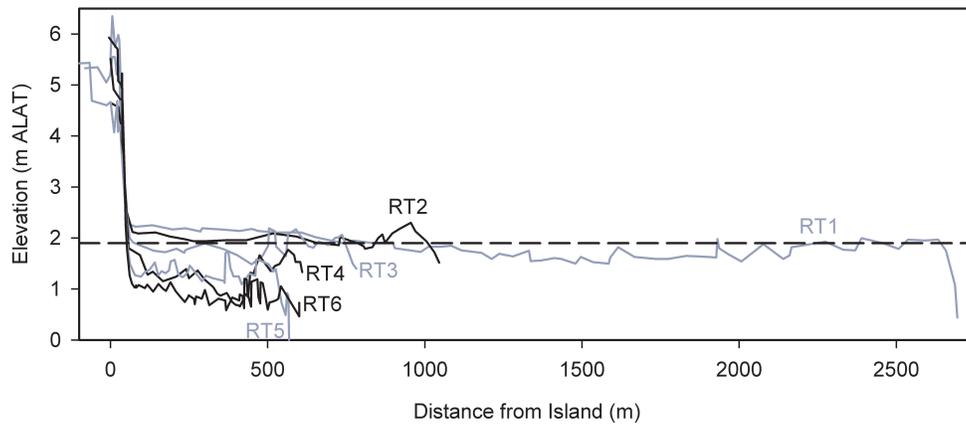


Fig. 2 Topographic surveys of the reef flat transects in relation to mean sea level (i.e. the dotted line at 1.9 m above the lowest astronomical tide, ALAT)

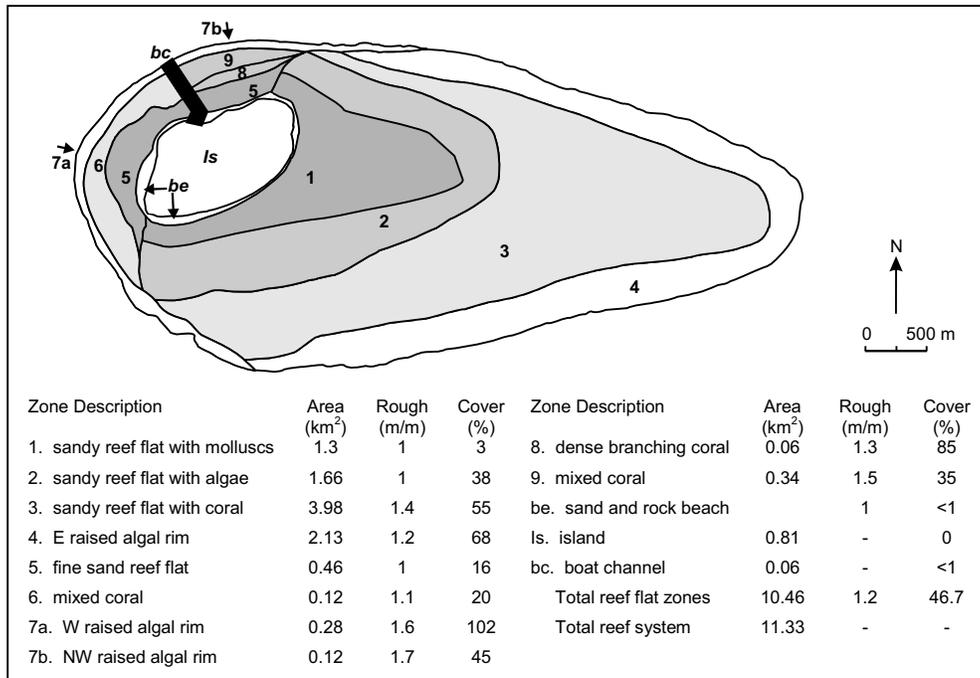


Fig. 3 Map and description of the ten identified ecological zones of the Warraber reef flat, including area, roughness and cover characteristics

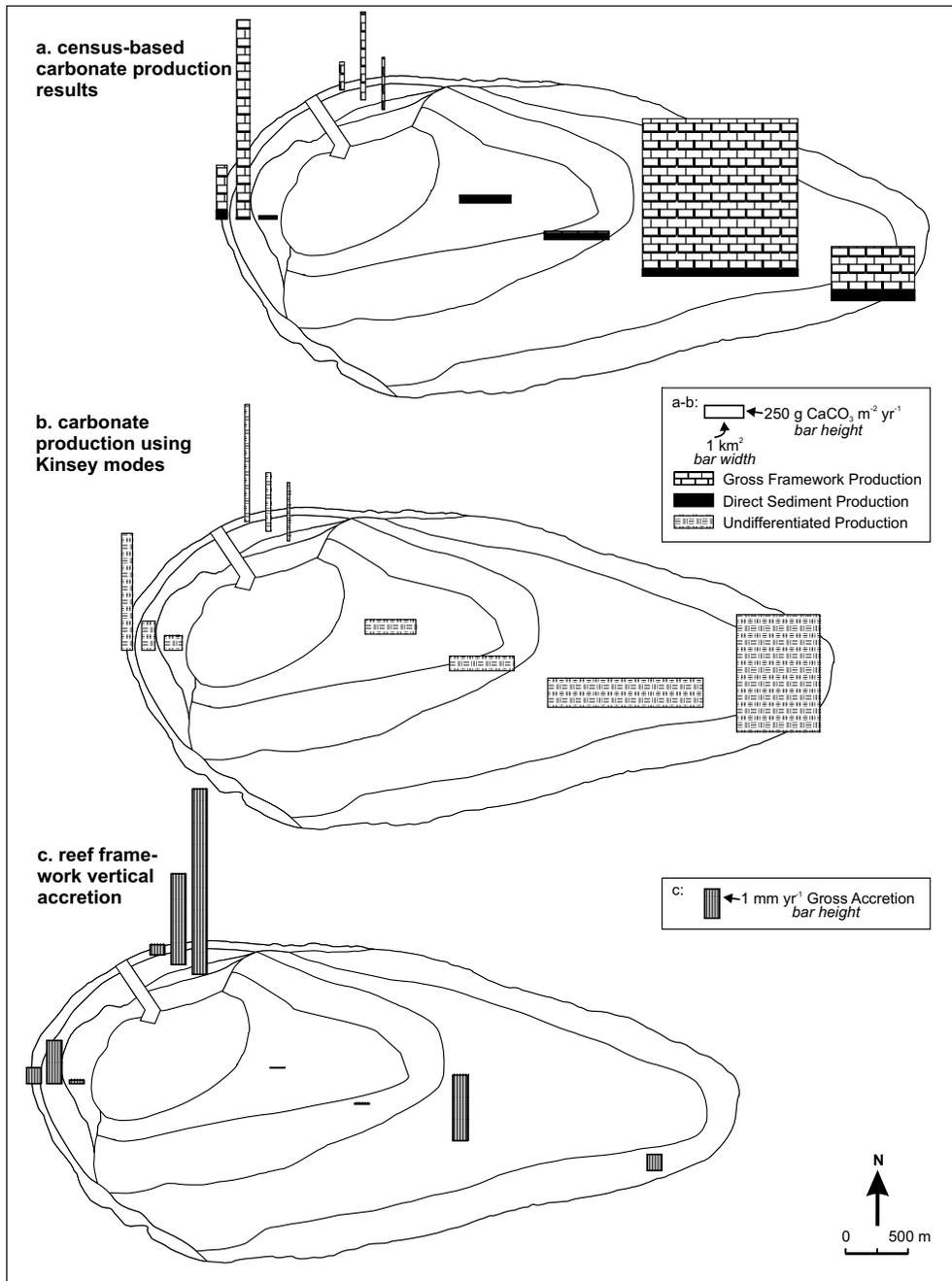


Fig. 4 Carbonate production rates (bar height) and amounts (bar area) in the ten ecological zones of the Warraber reef flat estimated using census-based techniques (a) versus modes derived from Kinsey's (1983, 1985) alkalinity-reduction review (b), and gross vertical framework accumulation rates calculated for the ten ecological zones based on framework building organism cover, production and density

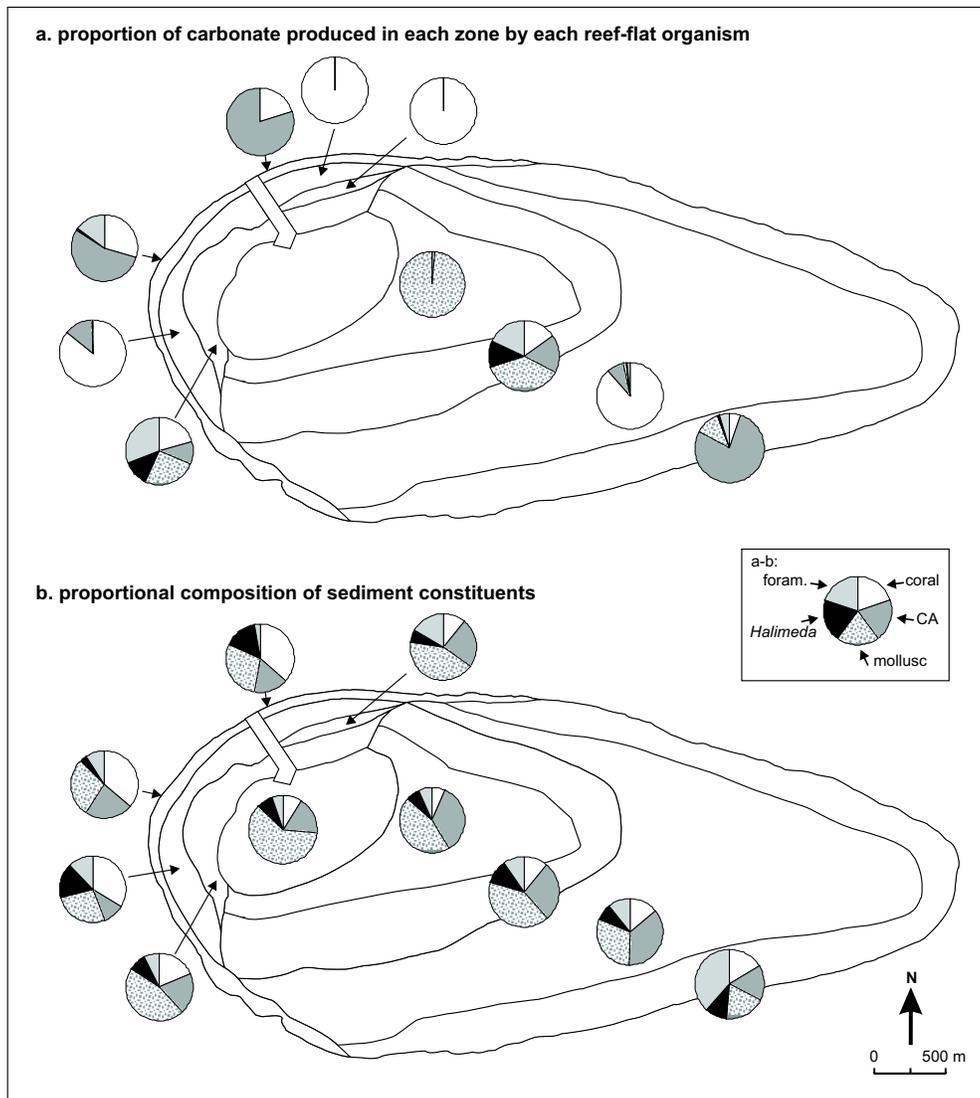


Fig. 5 Proportion of carbonate production contributed by the five main producers (a) versus the constituent composition of sediments (b) within each ecological zone of the reef flat. Note foram. abbreviates foraminifera

Table 1 Results from analysis of variance tests for (a) cover by quadrat versus by site, (b) cover by site within transect and (c) cover by site versus by transect

Cover type	a) Single Factor ANOVA*		a) Monte Carlo Randomisation	b) Single Factor ANOVA*		c) Single Factor ANOVA**		c) Monte Carlo Randomisation
	Site between transect		P-value	Site within transect		Site versus transect		P-value
	F	P-value		F	P-value	F	P-value	
Coral-ramose-other	18.079	<0.001	0	15.162	<0.001	2.732	0.046	0.041
Coral-ramose- <i>Acropora</i>	2.870	<0.001	0.003	2.578	<0.001	2.022	0.115	0.102
Coral-massive	1.358	0.133	0.089	1.428	0.106	0.559	0.694	0.733
Coral- <i>fol/encr/mu</i>	1.007	0.476	0.188	0.948	0.555	1.564	0.208	0.162
<i>Halimeda</i>	1.885	0.011	0.022	1.834	0.017	1.247	0.311	0.298
CA-encr	12.452	<0.001	0	12.566	<0.001	0.919	0.465	0.515
CA-artic	3.875	<0.001	<0.001	3.936	<0.001	0.861	0.498	0.493
Mollusc	9.957	<0.001	0	9.138	<0.001	1.807	0.152	0.138
Foraminifera	1.714	0.026	0.039	1.590	0.052	1.701	0.174	0.166
Brown algae	6.125	<0.001	0	5.906	<0.001	1.333	0.279	0.274
Sponge	2.696	<0.001	0.002	2.774	<0.001	0.745	0.569	0.588
Sea grass	1.891	0.011	0.019	2.019	0.007	0.43	0.786	0.902

\* degrees of freedom within groups = 74, degrees of freedom between groups = 36, n = 111, confidence level = 0.95

\*\* degrees of freedom within groups = 32, degrees of freedom between groups = 4, n = 37, confidence level = 0.95

Table 2 Estimated calcification rates used to calculate carbonate production on the Warraber reef flat as derived from a review of published rates (see Appendix 1 for details)

Organism	Best-estimate calcification rate (g m <sup>-2</sup> yr <sup>-1</sup> )	Minimum to maximum calcification rates (g m <sup>-2</sup> yr <sup>-1</sup> )	Adjustment factor
Coralline algae	1,872	1,500-2,500	multiplied by the square of quadrat rugosity for crustose species
Coral massive	16,160	7,680-24,640	
Coral- <i>fol/encr/mu</i>	17,000	3,000-31,000	
Coral <i>ramose-Acropora</i>	19,242	10,818-27,666	multiplied by an effective cover factor of 0.25
Coral <i>ramose-other</i>	1,394	767-2,021	
<i>Halimeda</i>	1,066	400-1,667	multiplied by a branch extension factor of 0.4
foraminifera	120	30-230	
molluscs	100	10-200	
			multiplied by a factor of between 0-3 depending on organism density
			multiplied by a factor of between 0-3 depending on organism density

Table 3 Gross carbonate production rates (a) and amounts (b) by producer type across the ten ecological zones of the Warraber reef flat calculated using the best-estimate calcification rates. Cover<sup>b</sup> is the percentage of area occupied by carbonate producers, which comprises 23% or 2.41 km<sup>2</sup> of the reef flat. SD is standard deviation

Zone	1	2	3	4	5	6	8	9	7a	7b	Reef Flat	Reef Flat
<b>a. Carbonate Production Rate</b>												
	<b>(g m<sup>-2</sup> yr<sup>-1</sup>)</b>										<b>Cover<sup>b</sup></b>	<b>(%)</b>
Coral ramose-other	0	0	70	0	0	21	1,046	376	0	0	38	12
Coral ramose-Acrop	0	0	321	0	0	994	0	577	321	0	170	4
Coral massive	0	24	2,343	18	13	1,966	0	754	0	0	974	26
Coral fol/encr/mu	0	0	57	38	0	453	0	57	0	113	46	1
CA encrusting	1	22	234	821	7	541	0	0	586	453	299	43
CA articulated	1	6	32	21	0	3	0	0	0	0	18	4
molluscs	148	61	39	122	17	17	0	0	0	0	69	3
Halimeda	0	19	27	8	8	4	0	0	14	0	16	6
Foraminifera	0	29	33	53	20	0	0	0	160	0	33	1
Framework	1	46	3,026	876	21	3,976	1,046	1,764	906	566	1,527	86
Direct Sediment	149	115	131	205	45	23	0	0	174	0	136	14
Average Production	149	161	3,157	1,081	65	3,999	1,046	1,764	1,081	566	1,663	100
SD	115	187	3,931	819	113	1,343	304	887	352	98	1,780	18
<b>b. Gross Carbonate Production</b>												
	<b>(t yr<sup>-1</sup>)</b>										<b>(%)</b>	
Coral ramose-other	0	0	281	0	0	7	66	47	0	0	401	2
Coral ramose-Acrop	0	0	1,278	0	0	341	0	72	90	0	1,780	10
Coral massive	0	41	9,335	38	6	674	0	94	0	0	10,187	59
Coral fol/encr/mu	0	0	226	81	0	155	0	7	0	13	482	3
CA encrusting	1	36	934	1,751	3	185	0	0	164	54	3,128	18
CA articulated	1	10	128	44	0	1	0	0	0	0	185	1
molluscs	193	100	155	261	8	6	0	0	0	0	722	4
Halimeda total	0	32	106	18	4	1	0	0	4	0	165	1
Foraminifera	0	48	133	114	9	0	0	0	45	0	349	2
Framework	1	76	12,053	1,870	9	1,363	66	219	253	67	15,978	92
Direct Sediment	194	191	522	437	20	8	0	0	49	0	1,421	8
Total Production	195	267	12,575	2,306	30	1,371	66	219	302	67	17,399	100
SD	150	309	15,660	1,748	51	460	19	110	98	12	18,618	107
Total Production (%)	1	2	72	13	0	8	0	1	2	0	100	-

Table 4 Estimated amounts and proportions of carbonate produced by the different types of organism on Warraber determined using the best estimate (mean), minimum and maximum calcification rates

	CaCO <sub>3</sub> production (t yr <sup>-1</sup> )			CaCO <sub>3</sub> production (%)		
	Mean	Maximum	Minimum	Mean	Maximum	Minimum
CA encrusting	3,128	4,177	1,282	18	16	17
CA articulated	185	248	76	1	1	1
Coral ramose-other	401	581	221	2	2	3
Coral ramose-Acrop	1,780	2,559	1,001	10	10	13
Coral massive	10,187	15,533	4,842	59	59	63
Coral fol/encr/mu	482	879	85	3	3	1
Halimeda total	165	258	62	1	1	1
molluscs	722	1,444	72	4	5	1
Foraminifera	349	668	87	2	3	1
Total	17,399	26,347	7,726	100	100	100

Table 5 Comparison of carbonate-production estimates for the Warraber reef flat with published rates from several reef environments

Location	Reef Environment	Method	Carbonate Production (kg CaCO <sub>3</sub> m <sup>-2</sup> yr <sup>-1</sup> )	Study area	Total production (t yr <sup>-1</sup> )	Source
Warraber Island, Torres Strait	0-4 m deep inter-tidal reef flat	census-based	1.66	10,462,700 m <sup>2</sup>	17,399 ± 18,618	present study
Green Island, Great Barrier Reef	reef flat and slope	census-based	1.6-3.9*	410,000 m <sup>2</sup>	656-1,606	Yamano et al. 2000
Mode of Several Pacific Reefs	1-3 m deep, seaward reef flat	alkalinity-reduction	4	-	-	Smith and Kinsey 1976
Mode of Several Pacific Reefs	protected 5-6 m deep lagoon/bank	alkalinity-reduction	0.8	-	-	Smith and Kinsey 1976
Japtan Inter-Island Reef, Eniwetok	inter-island reef flat	census-based	30.5	455 m long transect	-	Odum and Odum 1955
Kailua Bay, Hawaii	fringing reef with large sand bodies and a diverse benthic community	census-based	3.18	10,000,000 m <sup>2</sup>	74,810 ± 7,440	Harney 2000
Kaneohe Bay, Hawaii	shallow (<3m) reef flat	alkalinity-reduction	4.7 (3.5-8)	200 m long transect	-	Kinsey 1981
Discovery Bay, Jamaica	shallow (<3m) reef flat	alkalinity-reduction	4.4 (1.2-10)	200 m long transect	-	Kinsey 1981
Bellairs Reef, Barbados	5 m deep fringing reef	census-based	15	10,800 m <sup>2</sup>	163	Stearn <i>et al.</i> 1977
Cane Bay, St. Croix Virgin Islands	fringing reef 2-60 m deep incl. hard ground, reef flat and slope	census-based	1.9 (0.85-5.0)	30,000 m <sup>2</sup>	57.5	Sadd 1984
Cane Bay, St. Croix Virgin Islands	fringing reef shelf, 0-40 m deep incl. reef flat, slope and shelf	census-based	1.21 (0-5.78)	412,200 m <sup>2</sup>	499*	Hubbard et al. 1990
hypothetical	reef flat	census-based	3	-	-	Chave et al. 1972
hypothetical	lagoon	census-based	5	-	-	Chave et al. 1972
hypothetical	algal ridge	census-based	9	-	-	Chave et al. 1972
hypothetical	upper slope	census-based	60	-	-	Chave et al. 1972
hypothetical	lower slope	census-based	8	-	-	Chave et al. 1972

\* calculated from production and areal figures given in the source

## Appendix 1. Overview of organism calcification rates and factors used to estimate carbonate production on the Warraber reef flat

The published growth and carbonate production rates that underpin the census-based method are reviewed below for the five main producers present on Warraber, corals, coralline algae, molluscs, foraminifera and *Halimeda*, in order to determine appropriate calcification rates and factors.

### Coralline algae

Published estimates of the coralline algal calcification rates vary by an order of magnitude, although most rates lie between 1,500-2,500 g m<sup>-2</sup> yr<sup>-1</sup> (Table A1). Variation arises due to differences in (a) the growth rates of individual species, (b) physical habitat suitability, and (c) predation intensity (Adey and Vassar 1975; Stearn et al. 1977). In general, coralline algal calcification is highest in shallow-water habitats and in areas with minimal grazing, particularly by parrot fish (Eakin 1996; Adey and Vassar 1975).

Table A1 Published estimates of coralline algae extension and gross calcification for various reef environments

<sup>a</sup> calculated using Stearn et al.'s (1977) density of 1.56 g cm<sup>-3</sup>

Reef Environment	Extension rate (mm yr <sup>-1</sup> )	Calcification (g CaCO <sub>3</sub> m <sup>-2</sup> yr <sup>-1</sup> )	Source
algal ridge and reef, Virgin Islands	1-5.2	1,560-8,112 <sup>a</sup>	Adey & Vassar 1975
fore-reef, Barbados	0.1-1.5	167-2,378	Stearn et al. 1977
upper fore-reef and reef crest, Japan	1.0-1.2	1,560-1,872 <sup>a</sup>	Matsuda 1989
mostly reef flat, Panama	1.9	1,872	Eakin 1992

The flat-surface calcification rate 1,872 g m<sup>-2</sup> yr<sup>-1</sup> was used in this study to calculate coralline algal production rates. This value was derived by multiplying Matsuda's (1989) moderate, Pacific, reef-flat extension rate (1.2 mm yr<sup>-1</sup>) by Stearn et al.'s (1977) measure of the bulk skeletal density of coralline algae (1.56 g cm<sup>-3</sup>) and is similar to that found in other Pacific shallow reef environments (Table A1). Reflecting the majority of published estimates, the calcification rates 1,500 g m<sup>-2</sup> yr<sup>-1</sup> and 2,500 g m<sup>-2</sup> yr<sup>-1</sup> were used as minimum and maximum estimates of calcification. Calcification of crustose coralline algae in a given quadrat (g m<sup>-2</sup> yr<sup>-1</sup>) was calculated by multiplying this flat-surface calcification rate (g m<sup>-2</sup> yr<sup>-1</sup>) by the percentage cover and the square of quadrat rugosity, as suggested by Eakin (1996).

### Coral

The numerous extension rates of Pacific corals reported by Vecsei (2001), and other authors, form the primary source of data on coral calcification employed in this study (Table A2). These data indicate that Pacific coral growth rates vary by two orders of magnitude depending on the species, growth form and environment studied. Published shallow-water (<10 m) coral extension rates were grouped into four categories: i) massive; ii) foliaceous, encrusting and mushroom (fol/encr/mu); iii) ramose-*Acropora*; and iv) ramose-other (after Vecsei 2001). Best-estimate (mean), minimum (mean - 1σ) and maximum (mean + 1σ) extension rates were calculated for each category (Table A2). Ramose-*Acropora*

rates were further multiplied by a factor of 0.4 (Graus et al. 1977; Bottjer 1980) to account for the difference between extension perpendicular to the cover surface (ie. linear), and that along branches.

The observed cover data from Warraber were multiplied by an estimate of the effective cover for each growth form (1 for massive, foliaceous, encrusting and mushroom colonies and 0.25 for ramose-*Acropora* and ramose-other) (after Vecsei 2001). Carbonate production rates were then calculated by multiplying the adjusted cover data by the published densities and calculated extension rates of each coral category (Table A2) and results were summed to give the estimated production of corals in each quadrat ( $\text{g m}^{-2} \text{yr}^{-1}$ ).

### ***Halimeda***

Table A3 presents published growth rates for the green alga *Halimeda* and indicates that rates vary depending upon the measurement technique, species and plant density or biomass per unit area. Alkalinity-reduction methods can lead to over-estimation of calcification, while those reliant on segment staining and tagging produce minimum estimates since any new segments lost are not accounted for (Multer 1988; Payri 1988). Discrete, rhipsalian, sand-dwelling *Halimeda* species produce up to an order of magnitude less calcium carbonate per square meter than sprawling, lithophytic species (Hillis-Colinvaux 1974; Drew 1983; Multer 1988; Payri 1988).

The majority (>95%) of *Halimeda* plants found on Warraber were of the rhipsalian, sand-dwelling, slow-growing variety while the maximum observed plant density found in quadrats was 40 plants or 15% cover. An estimated calcification rate of  $160 \text{ g m}^{-2} \text{yr}^{-1}$  per 15% cover (or  $1,066 \text{ g m}^{-2} \text{yr}^{-1}$  for 100% cover) was used to calculate the production rate of *Halimeda* in survey quadrats. Minimum and maximum estimates of 60 and  $250 \text{ g m}^{-2} \text{yr}^{-1}$  of calcification per 15% cover ( $400$  and  $1,667 \text{ g m}^{-2} \text{yr}^{-1}$  for 100% cover) were used to test the sensitivity of the total carbonate production figures to *Halimeda* production rates, taking into account potential variation in local growth rates and in the exact ratio of rhipsalian to lithophytic species present in the different reef flat sub-environments.

Table A2 Coral extension rates, skeletal density and calcification rates used in calculating carbonate production on Warraber. Most values are from sites in the Pacific Ocean with a few values from sites in the Indian Ocean, Red Sea and, for foliaceous corals, from the Atlantic. Standard deviations ( $\sigma$ ) are given after  $\pm$  and the number of observations is indicated by n

<sup>a</sup> cited in Huston (1985, 22)

<sup>b</sup> multiplied by an effective cover factor of 0.25

<sup>c</sup> multiplied by a branch extension factor of 0.4

Category	Extension (mm yr <sup>-1</sup> )	Range	Sources	Density (g CaCO <sub>3</sub> cm <sup>-3</sup> )	Source	Calcification (g CaCO <sub>3</sub> m <sup>-2</sup> yr <sup>-1</sup> )
massive	10.1 $\pm$ 5.3	1.4-32 (n=110)	Mayor 1924 <sup>a</sup> ; Knutson et al. 1972; Buddemeier et al. 1975; Highsmith 1979; Grigg 1982; Wellington 1982; Wellington and Glynn 1983; Charuchinda and Chansang 1985; Hughes 1987; Guzman and Cortes 1989; Scoffin et al. 1992; Klein et al. 1993; Glynn et al. 1996; Stimson 1996; Vecsei 2001	1.6	Hughes 1987	16,160 $\pm$ 8,480
foliaceous, mushroom and encrusting	8.5 $\pm$ 7	0.8-23 (n=20)	Mayor 1924 <sup>a</sup> ; Edmondson 1929 <sup>a</sup> ; Wellington 1982; Hughes and Jackson 1985; Huston 1985; Stimson 1996	2.0	Hughes 1987	17,000 $\pm$ 14,000
ramose – <i>Acropora</i>	106.9 $\pm$ 46.8	4-185 (n=18)	Mayor 1924 <sup>a</sup> ; Crossland 1981; Oliver et al. 1983; Charuchinda and Hylleberg 1984; Yap and Gomez 1984; Marsh 1993; Stimson 1996; Harriott 1998; Vecsei 2001	1.8	Schuhmacher and Plewka 1981	19,242 $\pm$ 8,424 <sup>bc</sup>
ramose – other	33.8 $\pm$ 15.2	6-72 (n=21)	Mayor 1924 <sup>a</sup> ; Glynn 1976; Neudecker 1977; Glynn and Stewart 1973; Wellington 1982; Vecsei 2001	0.165	Eakin 1992, 1996	1,394 $\pm$ 627 <sup>b</sup>

Table A3 Published mean growth and calcification rates of various types of *Halimeda* species at different densities in different types of environment

Species	Type	Environment	Method	Biomass or Density	Calcification (g CaCO <sub>3</sub> m <sup>-2</sup> yr <sup>-1</sup> )	Source
<i>H. incrassata</i>	rhipsalian	shallow lagoon, Bermuda	segment staining	6.7 g m <sup>-2</sup>	50	Wefer 1980
<i>H. incrassata</i> and <i>H. monile</i>	rhipsalian	barrier reef, Antigua	segment staining	36 plants m <sup>-2</sup>	114	Multer 1988
<i>H. incrassata</i> and <i>H. monile</i>	rhipsalian	fringing lagoon, Antigua	segment staining	30 plants m <sup>-2</sup>	62	Multer 1988
<i>H. incrassata</i> and <i>H. monile</i>	rhipsalian	open lagoon, Antigua	segment staining	26 plants m <sup>-2</sup>	61	Multer 1988
<i>H. opuntia</i>	lithophytic	lagoon, Florida	segment staining	1 plant m <sup>-2</sup> (c.1,200 g/m <sup>2</sup> )	1,088	Hudson 1985

<i>H. opuntia</i> and <i>H. copiosa</i>	lithophytic	lagoon, Davies Reef, Australia	segment tagging	167 g dry weight/m <sup>2</sup>	2,234	Drew 1983
<i>H. incrassata</i> f. <i>ocata</i> , <i>H. opuntia</i> and <i>H. discoidea</i>	mostly lithophytic	lagoon, Tahiti	alkalinity reduction	111 g dry weight/m <sup>2</sup>	1,400	Payri 1988
<i>H. copiosa</i> and <i>H. opuntia</i> v. <i>hederacea</i> (+15 other minor species)	mostly lithophytic	inter-reefal seabed, Australia	census based	503 g dry weight/m <sup>2</sup>	2,519	Drew and Abel 1985
several, incl. <i>H. incrassata</i> , <i>H. copiosa</i> and <i>H. goreauii</i>	mixed	bank margin, Bahamas	census based	1,000 plants/m <sup>2</sup>	2,400	Freile et al. 1995

Table A4 Published calcification rates of foraminifera in various reef environments

Species	Environment	Calcification (g CaCO <sub>3</sub> m <sup>-2</sup> yr <sup>-1</sup> )	Source
numerous	reef - global	230 (30-1,000)	Langer et al. 1997
numerous	lagoon - global	30.4 (1.2-120)	Langer et al. 1997
<i>Amphistegina lessonii</i> , <i>Baculogypsina sphaerulata</i> and <i>Calcarina hispida</i>	reef flat - Green Is, GBR	210-480	Yamano et al. 2000
numerous	continental shelf - north GBR	40 (Holocene rate)	Tudhope and Scoffin 1988
<i>Amphistegina</i> , <i>Baculogypsina</i> , <i>Heterostegina</i> and <i>Calcarina</i> species	reef flat - Caroline Islands	187-2,762	Hallock 1981
<i>Amphistegina</i> , <i>Baculogypsina</i> , <i>Heterostegina</i> and <i>Calcarina</i> species	reef slope - Caroline Islands	57-568	Hallock 1981
<i>Amphistegina</i> and <i>Heterostegina</i> species	reef flat and slope - Hawaii	38-145	Hallock 1981
<i>Amphistegina</i> species	rock pool - Hawaii	305-512	Muller 1974
predominantly <i>Amphistegina</i> (plus <i>Heterostegina</i> , <i>Marginopora</i> and 90 other species)	nearshore - Hawaii	263	Muller 1976
<i>Baculogypsina sphaerulata</i>	shallow tide pool - Japan	500-700	Sakai and Nishihira 1981
<i>Archaias angulatus</i>	shallow lagoon - Florida	60-100	Hallock et al. 1986

## Foraminifera

In the majority of reef carbonate budget studies foraminiferal contributions are excluded (Stearn et al. 1977; Eakin 1996), determined from the volume of tests in sediments (Hubbard et al. 1990), or gauged from global estimates (Vecsei 2001). Published estimates of reef foraminifera calcification rates vary considerably depending on the species and environment studied (Table A4). On Warraber, live tests of the larger, more productive foraminifera (*Marginopora vertebralis*, *M. vertebralis* var. *plicata*, *Amphisorus hemprichii*) were observed in small numbers in quadrats where they were present while abundant tests of the smaller, less-productive species (*Amphistegina lessonii*, *Baculogypsina sphaerulata*, *Calcarina spengleri*) were found where these species were present. Overall, a mean productivity rate of  $120 \text{ g CaCO}_3 \text{ m}^{-2} \text{ yr}^{-1}$  was used to estimate the contribution of foraminifera to the carbonate produced in each quadrat where they were recorded, a figure which lies mid-way between Langer et al.'s (1997) low- and high-productivity environment estimates. Langer et al.'s (1997) mean rates for these two types of environment, 30 and  $230 \text{ g m}^{-2} \text{ yr}^{-1}$ , were used as minimum and maximum estimates. For each quadrat the best-estimate  $120 \text{ g m}^{-2} \text{ yr}^{-1}$  figure (as well as the 30 and  $230 \text{ g m}^{-2} \text{ yr}^{-1}$  minimum and maximum figures) was multiplied by a factor of between 0 and 3 depending on the density of live foraminifera counted during the ecological census.

## Molluscs

Although their shells often comprise a significant proportion of sediments, most reef carbonate budgets omit mollusc production (eg. Stearn et al. 1977; Sadd 1984; Eakin 1996; Vecsei 2001) and, as a result, there is a paucity of literature on their calcification. When mollusc production is included, it is often estimated using the abundance of shells in sediment deposits (eg. Hubbard et al. 1990; Yamano et al. 2000). However, the age of such sediments is highly variable (Roy 1991; Harney et al. 2000) and often spans the period of recent high sea levels, which exceeds 2000 years in Torres Strait (Woodroffe et al. 2000). Further, the shells of dead molluscs in any one area (eg. a quadrat) are a poor indicator of immediate live assemblages due to the influence of taphonomic processes, such as post-mortem transport and deposition out of production areas (Cummins et al. 1986a-b; Zuschin et al. 2000).

Table A5 Published estimates of shell production rates measured from living mollusc assemblages  
<sup>a</sup> estimated annual shell production rates calculated from the soft tissue to total weigh ratios and soft-tissue and total production figures given in the original

Type and/or Species	Calcification ( $\text{g CaCO}_3 \text{ m}^{-2} \text{ yr}^{-1}$ )	Source
micromolluscs - on hard substrates	9-180	Harney & Fletcher 2003
micromolluscs - in <i>Halimeda</i> beds	210-540	Harney & Fletcher 2003
micromolluscs - average	70	Harney & Fletcher 2003
macromollusc - <i>T. maxima</i>	4.8	Richard 1981 <sup>a</sup>
macromollusc - <i>C. fragum</i>	0.22	Richard 1981 <sup>a</sup>

In the present study, a rate  $100 \text{ g CaCO}_3 \text{ m}^{-2} \text{ yr}^{-1}$  is used to indicate mollusc production. This rate lies between the few published rates for micro- and macromolluscs (Table A5). Comparisons between field observations of the volume of molluscs found in individual quadrats, and the weight of an equivalent

volume of empty shells (which require no correction for organic content, Harney et al. 2000), indicates that this estimate is realistic. Minimum and maximum estimates of 10-200 g CaCO<sub>3</sub> m<sup>-2</sup> yr<sup>-1</sup> are used to test the sensitivity of total carbonate production figures to variation in mollusc production rates. For each quadrat the best-estimate 100 g m<sup>-2</sup> yr<sup>-1</sup> figure (as well as the 10 and 200 g m<sup>-2</sup> yr<sup>-1</sup> minimum and maximum figures) was multiplied by a factor of between 0 and 3 depending on the density of live molluscs counted during the ecological census.

## References

- Adey WH, Vassar JM (1975) Colonization, succession and growth rates of tropical crustose coralline algae (Rhodophyta, Cryptonemiales). *Phycologia* 14:55-69
- Bottjer DJ (1980) Branching morphology of the reef coral *Acropora cervicornis* in different hydraulic regimes. *J Paleontol* 54:1102-1107
- Buddemeier RW, Smith SV, Kinzie RA (1975) Holocene windward reef-flat history, Enewetak Atoll. *Geol Soc Am Bull* 86:1581-1584
- Charuchinda M, Chansang H (1985) Skeleton extension and banding formation of *Porites lutea* of fringing reefs along the south and west coasts of Phuket Island (Thailand). *Proc 5th Int Coral Reef Symp* 6:83-87
- Charuchinda M, Hylleberg J (1984) Skeletal extension of *Acropora formosa* at a fringing reef in the Andaman Sea. *Coral Reefs* 3:215-219
- Crossland CJ (1981) Seasonal growth of *Acropora* cf. *formosa* and *Pocillopora damicornis* on a high latitude reef (Houtman Abrolhos, Western Australia). *Proc 4th Int Coral Reef Symp* 1:663-667
- Cummins H, Powell EN, Stanton RJ Jr., Staff G (1986a) The rate of taphonomic loss in modern benthic habitats: how much of the potentially preservable community is preserved? *Palaeogeogr Palaeoclimatol Palaeoecol* 52:291-230
- Cummins H, Powell EN, Stanton RJ Jr., Staff G (1986b) The size frequency distribution in palaeoecology: effects of the taphonomic process during formation of molluscan death assemblages in Texas Bays. *Palaeontology* (Oxford) 29:495-518
- Drew EA (1983) *Halimeda* biomass, growth rates and sediment generation on reefs in the Great Barrier Reef Province. *Coral Reefs* 2:101-110
- Drew EA, Abel KM (1985) Biology, sedimentology and geography of the vast inter-reefal *Halimeda* meadows within the Great Barrier Reef Province. *Proc 5th Int Coral Reef Symp* 5:15-20
- Eakin CM (1992) Post-El Niño Panamanian reefs: less accretion, more erosion and damselfish protection. *Proc 7th Int Coral Reef Symp* 387-396
- Eakin CM (1996) Where have all the carbonates gone? A model comparison of calcium carbonate budgets before and after the 1982-1983 El Niño at Uva Island in the eastern Pacific. *Coral Reefs* 15:109-119
- Freile D, Milliman JD, Hillis L (1995) Leeward bank margin *Halimeda* meadows and draperies and their sedimentary importance on the western Great Bahama Bank slope. *Coral Reefs* 14:27-33
- Glynn PW (1976) Some physical and biological determinants of coral community structure in the eastern Pacific. *Ecol Monogr* 46:431-456
- Glynn PW, Stewart RH (1973) Distribution of coral reefs in the Pearl Islands (Gulf of Panama) in relation to thermal conditions. *Limnol Oceanogr* 18:367-379

- Glynn PW, Veron JEN, Wellington GM (1996) Clipperton Atoll (eastern Pacific): oceanography, geomorphology, reef-building coral ecology and biogeography. *Coral Reefs* 15:71-99
- Graus RG, Chamberlain JA, Boker AM (1977) Structural modification of corals in relation to waves and currents. In: Frost SH, Weiss MP, Saunders JB (eds) *Reefs and Related Carbonates – Ecology and Sedimentology*, American Society of Petroleum Geologists, *Studies in Geology* 4:135-153
- Grigg RW (1982) Darwin point: a threshold for atoll formation? *Coral Reefs* 1:29-34
- Guzman HM, Cortes J (1989) Growth rates of eight species of scleractinian corals in the eastern Pacific (Costa Rica). *Bull Mar Sci* 44:1186-1194
- Hallock P (1981) Production of carbonate sediments by selected large benthic foraminifera on two Pacific coral reefs. *J Sediment Petrol* 51:467-474
- Hallock P, Cottey TL, Forward LB, Halas J (1986) Population biology and sediment production of *Archaias angulatus* (Foraminiferida) in Largo Sound, Florida. *J Foraminifer Res* 16:1-8
- Harney JN, Fletcher CH (2003) A budget of carbonate framework and sediment production, Kailua Bay, Oahu, Hawaii. *J Sediment Res* 73:856-868
- Harney JN, Grossman EE, Richmond BM, Fletcher CH (2000) Age and composition of carbonate shoreface sediments, Kailua Bay, Oahu, Hawaii. *Coral Reefs* 19:141-154
- Harriott VJ (1998) Growth of the staghorn coral *Acropora formosa* at Houtman Abrolhos, Western Australia. *Mar Biol* 132:319-325
- Highsmith R (1979) Coral growth rates and environmental control of density banding. *J Exp Mar Biol Ecol* 37:105-125
- Hillis-Colinvaux L (1974) Productivity of the coral reef alga *Halimeda* (order Siphonales). *Proc 2nd Int Coral Reef Symp* 1:35-42
- Hubbard DK, Miller AI, Scaturro D (1990) Production and cycling of calcium carbonate in shelf-edge reef systems (St Croix, U.S. Virgin Islands): Applications to the nature of reef systems in the fossil record. *J Sediment Petrol* 60:335-360
- Hudson JH (1985) Growth rate and carbonate production in *Halimeda opuntia*: Marquesas Keys, Florida. In: Toomey DF, Nitecki MH (eds) *Paleoalgology: contemporary research and applications*. Springer-Verlag, Berlin, pp 257-263
- Hughes T P (1987) Skeletal density and growth form of corals. *Mar Ecol Prog Ser* 35:259-266
- Hughes TP, Jackson JBC (1985) Population dynamics and life histories of foliaceous corals. *Ecol Monogr* 55:141-166
- Huston M (1985) Variation in coral growth rates with depth at Discovery Bay, Jamaica. *Coral Reefs* 4:19-25
- Klein R, Patzold J, Wefer G, Loya Y (1993) Depth-related timing of density band formation in *Porites* spp. Corals from the Red Sea inferred from X-ray chronology and stable isotope composition. *Mar Ecol Prog Ser* 97:99-104
- Knutson D, Buddemeier R, Smith S (1972) Coral chronometers: seasonal growth bands in coral reefs. *Science* 177:270-272
- Langer MR, Silk MT, Lipps JH (1997) Global ocean carbonate and carbon dioxide production: The role of reef foraminifera. *J Foraminifer Res* 27:271-277
- Marsh LM (1993) The occurrence and growth of *Acropora* in extra-tropical waters off Perth, Western Australia. *Proc 7th Int Coral Reef Symp* 1233-1238

- Matsuda S (1989) Succession and growth rates of encrusting crustose coralline algae (Rhodophyta, Cryptonemiales) in the upper fore-reef environment off Ishigaki Island, Ryukyu Islands. *Coral Reefs* 7:185-195
- Muller PH (1974) Sediment production and population biology of the benthic foraminifer *Amphistegina madagascariensis*. *Limnol Oceanogr* 19:802-809
- Muller PH (1976) Sediment production by shallow-water, benthic foraminifera at selected sites on Oahu, Hawaii. *Maritime Sediments Special Publication* 1:263-265
- Multer HG (1988) Growth rate, ultrastructure and sediment contribution of *Halimeda incrassata* and *Halimeda monile*, Nonsuch and Falmouth Bays, Antigua, W.I. *Coral Reefs* 6:179-186
- Neudecker S (1977) Transplant experiments to test the effect of fish grazing on coral distribution. *Proc 3rd Int Coral Reef Symp* 1:317-323
- Oliver JK, Chalker BE, Dunlap WC (1983) Bathymetric adaptations of reef-building corals at Davies Reef, Great Barrier Reef, Australia. I. Long-term growth responses of *Acropora formosa* (Dana 1846). *J Exp Mar Biol Ecol* 73:11-35
- Payri CE (1988) *Halimeda* contribution to organic and inorganic production in a Tahitian reef system. *Coral Reefs* 6:251-262
- Richard G (1981) A first evaluation of the findings on the growth and production of lagoon and reef molluscs in French Polynesia. *Proc 4th Int Coral Reef Symp* 2:637-641
- Roy PS (1991) Shell hash dating and mixing models for palimpsest marine sediments. *Radiocarbon* 33:283-289
- Sadd JL (1984) Sediment transport and CaCO<sub>3</sub> budget on a fringing reef, Cane Bay, St Croix, U.S. Virgin Islands. *Bull Mar Sci* 35(2):221-238
- Sakai K, Nishihira M (1981) Population study of the benthic foraminifer *Baculogypsina sphaerulata* on the Okinawan Reef flat and preliminary estimation of its annual production. *Proc 4th Int Coral Reef Symp* 2:763-766
- Schuhmacher H, Plewka M (1981) The adaptive significance of mechanical properties versus morphological adjustments in skeletons of *Acropora palmata* and *Acropora cervicornis* (Cnidaria, Scleractinia). *Proc 4th Int Coral Reef Symp* 2:121-128
- Scoffin TP, Tudhope AW, Brown BE, Chandang H, Cheeney RF (1992) Patterns and possible environmental controls of skeletogenesis of *Porites lutea*, South Thailand. *Coral Reefs* 11:1-11
- Stearn CW, Scoffin TP, Martindale W (1977) Calcium carbonate budget of a fringing reef on the west coast of Barbados. *Bull Mar Sci* 27:479-510
- Stimson J (1996) Wave-like outward growth of some table- and plate-forming corals and a hypothetical mechanism. *Bull Mar Sci* 58:301-313
- Tudhope AW, Scoffin TP (1988) The relative importance of benthic foraminifera in the production of carbonate sediment on the Central Queensland Shelf. *Proc 6th Int Coral Reef Symp* 2:583-588
- Vecsei A (2001) Fore-reef carbonate production: development of a regional census-based method and first estimates. *Palaeogeogr Palaeoclimatol Palaeoecol* 175:185-200
- Wefer G (1980) Carbonate production by algae *Halimeda*, *Penicillus* and *Padina*. *Nature* 285:323-324
- Wellington GM (1982) Depth zonation of corals in the Gulf of Panama: control and facilitation by reef fishes. *Ecol Monogr* 52:223-241

- Wellington GM, Glynn PW (1983) Environmental influences on skeletal banding in eastern Pacific (Panama) corals. *Coral Reefs* 1:215-222
- Woodroffe CD, Kennedy DM, Hopley D, Rasmussen CE, Smithers SG (2000) Holocene reef growth in Torres Strait. *Mar Geol* 170:331-346
- Yamano H, Miyajima T, Koike I (2000) Importance of foraminifera for the formation and maintenance of a coral sand cay: Green Island, the Great Barrier Reef, Australia. *Coral Reefs* 19:51-58
- Yap HT, Gomez ED (1984) Growth of *Acropora pulchra*. II. Responses of natural and transplanted colonies to temperature and day length. *Mar Biol* 81:209-215
- Zuschin M, Hohenegger J, Steininger FF (2000) A comparison of living and dead molluscs on coral reef associated hard substrata in the northern Red Sea – implications for the fossil record. *Palaeogeogr Palaeoclimatol Palaeoecol* 159:167-190