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Cyanobacterial endobionts within a major marine planktonic calcifier (Globigerina bulloides, Foraminifera) revealed by 16S rRNA metabarcoding

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Abstract. We investigated the possibility of bacterial symbiosis in Globigerina bulloides, a palaeoceanographically important, planktonic foraminifer. This marine protist is commonly used in micropalaeontological investigations of climatically sensitive subpolar and temperate water masses as well as wind-driven upwelling regions of the world’s oceans. G. bulloides is unusual because it lacks the protist algal symbionts that are often found in other spinose species. In addition, it has a large offset in its stable carbon and oxygen isotopic compositions compared to other planktonic foraminifer species, and also that predicted from seawater equilibrium. This is suggestive of novel differences in ecology and life history of G. bulloides, making it a good candidate for investigating the potential for bacterial symbiosis as a contributory factor influencing shell calcification. Such information is essential to evaluate fully the potential response of G. bulloides to ocean acidification and climate change. To investigate possible ecological interactions between G. bulloides and marine bacteria, 18S rRNA gene sequencing, fluorescence microscopy, 16S rRNA gene metabarcoding and transmission electron microscopy (TEM) were performed on individual specimens of G. bulloides (type IId) collected from two locations in the California Current. Intracellular DNA extracted from five G. bulloides specimens was subjected to 16S rRNA gene metabarcoding and, remarkably, 37–87% of all 16S rRNA gene sequences recovered were assigned to operational taxonomic units (OTUs) from the picocyanobacterium Synechococcus. This finding was supported by TEM observations of intact Synechococcus cells in both the cytoplasm and vacuoles of G. bulloides. Their concentrations were up to 4 orders of magnitude greater inside the foraminifera than those reported for the California Current water column and approximately 5% of the intracellular Synechococcus cells observed were undergoing cell division. This suggests that Synechococcus is an endobiont of G. bulloides type IId, which is the first report of a bacterial endobiont in the planktonic foraminifera. We consider the potential roles of Synechococcus and G. bulloides within the relationship and the need to determine how widespread the association is within the widely distributed G. bulloides morphospecies. The possible influence of Synechococcus respiration on G. bulloides shell geochemistry is also explored.
1 Introduction

Mutualistic associations between organisms in marine ecosystems can provide the partners involved with the capacity to adapt to environmental stresses such as energy or nutrient limitation as well as provide robustness under the challenges caused by climate change. For example, the close association between photosynthetic microalgae and planktonic foraminifera supplies valuable fixed carbon and other benefits to the host and is a common feature of oligotrophic surface waters (Decelle et al., 2015). Since Murray first proposed a symbiotic role for these intracellular phototrophs in 1897 (Murray, 1897), many cytological and ultrastructural studies using light, fluorescence, and transmission electron microscopy have confirmed the presence of intracellular photosynthetic dinoflagellates or chrysophyte algae in a wide range of planktonic foraminifera (see Gastrich, 1987; Hemleben et al., 1989). Symbiosis was demonstrated first by Bé et al. (1982) in Orbulina universa by the elimination of its dinoflagellate endobionts and their successful reinfection. Other experimental techniques focused on tracing radiolabelled C and N (Gastrich and Bartha, 1988), stable isotope analysis (Uhle et al., 1997, 1999), and microsensor studies of the chemical microenvironment (Jørgensen et al., 1985; Rink et al., 1998). However, direct microscopic observations remain an important first step in assessing potential symbiotic associations.

Since not all planktonic foraminifera harbour protist microalgae, there is obvious diversity in the requirement for this type of mutualistic relationship. Technological advances now allow the investigation of the potential diversity and physiological role of other organisms (e.g. bacteria) entering into relationships with these abundant, calcifying protists. Understanding and investigating the full range of their mutualistic relationships is of great importance in planktonic foraminifera because of their considerable importance in Earth’s biogeochemical cycles. Their calcium carbonate shells play a significant role in marine carbon cycling within the water column and ocean sediments. They contribute up to 40% of the biogenic carbonate exported from the surface ocean (Schiebel, 2002; Schiebel et al., 2007), and shell dissolution at depth provides a significant buffering of ocean carbonate chemistry and atmospheric CO₂ (Holligan and Robertson, 1996; Iglesias-Rodriguez et al., 2002; Schiebel, 2002; Feely et al., 2004; Ridgwell and Zeebe, 2005; Sarmiento and Gruber, 2006). However, calcification rates of foraminifera are now known to be affected by the continued release of anthropogenic CO₂, and are sensitive to the associated changes in surface seawater pH (Ridgwell and Zeebe, 2005; Manno et al., 2012). Assessment of the impact of climate change on planktonic foraminifera and the implications for future biogenic carbonate production is hampered currently by a lack of basic ecological information (Lombard et al., 2011; Roy et al., 2015), including information on the diversity of symbiont–foraminifera associations that might allow different species to adapt to future environmental conditions. In addition to their role in marine carbon cycling, the geochemistry of foraminiferal shells buried within the sediments provides a long-term repository of information that reveals past changes in ocean conditions. Reconstructions of past seawater temperature, pH, dissolved inorganic carbon (DIC) concentrations and other environmental parameters based on these shell geochemical proxies provide essential constraints for refining climate change projections (Kucera 2007; Katz et al., 2010; Henderson, 2002). The underlying rationale in the use of foraminiferal shell chemistry for this purpose is that it reflects conditions in bulk seawater at the time of deposition. Yet this chemical signal can be altered by protist algal symbionts within some foraminifer species, complicating the interpretation of proxy records (Spero et al., 1991; Bemis et al., 1998; 2002; Anand et al., 2003; Russell et al., 2004). Important geochemical proxies such as δ¹⁸O, δ¹³C, and δ¹¹B are influenced by the consumption or addition of CO₂ to the calcifying microenvironment through photosynthesis by algal symbionts and symbiont–host respiration (Mashiotta et al., 1997; Rink et al., 1998; Wolf-Gladrow et al., 1999; Hönsicth et al., 2003; Egginis et al., 2004). The presence of other symbiotic or endobiotic organisms like bacteria within a foraminiferal host could also potentially complicate the interpretation of the proxy record via cryptic isotope fractionation through respiration or other metabolic processes.

Although the role and importance of the protist algal symbionts within planktonic foraminifera is widely recognised and relatively well understood, the association of planktonic foraminifera with bacteria has received very little scientific attention. Apart from a few studies reporting the presence of living bacteria inside benthic foraminifera from dysoxic sediments (Bernhard et al., 2000, 2006, 2012; Tsuchiya et al., 2015), there has been little consideration of specific endosymbioses between foraminifera and prokaryotes. Indeed, there are no reports of planktonic foraminiferal relationships with bacteria other than a single report observing the external association of Globigerinella siphonifera type I with the marine nitrogen-fixing, filamentous cyanobacterium Trichodesmium (Huber et al., 1997). This oversight is surprising, since the occurrence of bacterial symbiosis within other protists is well established, as is their great potential for providing highly specialised metabolic processes to their hosts (e.g. van Hoek et al., 2000; Schweikert and Meyer, 2001; Beier et al., 2002; Ashton et al., 2003; Foikin et al., 2003; Gast, 2009; Nowack et al., 2010; Orsi et al., 2012; Gilbert et al., 2012).

G. bulloides is a spinose planktonic foraminifer lacking protist algal symbionts (Febvre-Chevalier, 1971; Spero and Lea, 1996) that is abundant in the subpolar and temperate regions and also in the lower-latitude upwelling systems (Kleijne et al., 1989; Naidu and Malmgren, 1996). In these climatically sensitive areas it dominates the downward flux of foraminiferal shells to the sea floor and, as a consequence, is of considerable importance for palaeoclimate reconstruc-
tions (Sautter and Thunell, 1991; Spero and Lea, 1996). A complication in using \textit{G. bulloides} for palaeoclimate reconstruction, however, is that despite its lack of protist algal symbionts, the shell secreted by \textit{G. bulloides} is still out of isotopic equilibrium with respect to both carbon and oxygen isotopes, deviating from predicted values by more than any other extant, surface-dwelling species (Deuser et al., 1981; Kahn and Williams, 1981; Curry and Matthews, 1981; Kroon and Darling, 1995; Spero and Lea, 1996; Bijma et al., 1999). Such deviations are difficult to explain in the absence of protist algal symbionts, although some of the disequilibrium has been potentially linked to growth and ontogeny or even to \textit{G. bulloides} respiration rates (Spero and Lea, 1996). The presence of intracellular bacteria may provide an additional or contributing explanation.

The study of \textit{G. bulloides} is further complicated by our current inability to morphologically distinguish the numerous genotypes of \textit{G. bulloides} identified within the morphospecies (Darling and Wade, 2008; Seears et al., 2012; Morard et al., 2013). The majority of the genotypes have been elevated to species level status (Andre et al., 2014) and all are potentially ecologically distinct, though they are commonly found in the same water column, where their adaptive ranges overlap (Darling and Wade, 2008; Morard et al., 2013). Where this occurs, aggregation of two ecologically distinct \textit{G. bulloides} species could introduce significant noise into palaeoclimate calibrations (Darling et al., 2000; Kucera and Darling, 2002), particularly if they exhibit genotype-specific geochemical signatures as has recently been demonstrated in the Arabian Sea (Sadakov et al., 2016).

In this study we have focussed on the cool water lineage \textit{G. bulloides} type IId. With the exception of a single specimen of type Ila appearing off Santa Barbara in January (Darling et al., 2003), type IId was the only genotype identified off the coast of California throughout the year in both upwelling and non-upwelling hydrographic regimes (Darling et al., 2003; Darling and Wade, 2008). This is also the region where the majority of experimental geochemical studies on the \textit{G. bulloides} morphospecies have been carried out, which means that current calibrations for this area should be robust (e.g. Spero and Lea, 1996; Bemis et al., 1998, 2000, 2002), but may not be globally applicable since this genotype has not been found elsewhere to date (Darling and Wade, 2008; Morard et al., 2013).

We examined the intracellular bacterial population of individual specimens of the planktonic foraminifer \textit{G. bulloides} using a multiphasic approach. We used 18S rRNA gene sequencing to identify the genotype of the host cells and fluorescence microscopy, 16S rRNA gene metabarcoding via next-generation sequencing, transmission electron microscopy (TEM), and genetic characterisation using the polymerase chain reaction (PCR) to investigate the distribution and taxonomic affiliations of the intracellular bacteria. We demonstrate that intact, viable cells of the picocyanobacterium \textit{Synechococcus} spp. accumulate in large numbers within the cytoplasm of \textit{G. bulloides} type IId. We propose that these cells are likely to be taken up directly from the surrounding water column and that \textit{Synechococcus} should be considered an endobiont of \textit{G. bulloides} type IId. We go on to consider the nature of this association and its potential metabolic and geochemical implications. We also discuss the power of the methodological approach adopted for improving ecological knowledge of planktonic foraminifera.

2 Materials and methods

2.1 Oceanographic setting

Sampling was undertaken off the Californian coast in waters influenced by two opposing currents. The California Current flows equatorward from the North Pacific Current (∼50°N to Baja California (∼15–25°N). Southerly along-shore winds drive upwelling of cold nutrient-rich waters in early spring and summer in central California, and weaker but more sustained upwelling further toward the south. The relatively warm, saline Davidson Current and California Undercurrent flow poleward over the continental shelf. During the southern California summer, the California Current moves farther offshore, and the Davidson Current predominates near shore (Checkley and Barth, 2009). For this study, samples were collected along the narrow central California shelf ∼1 km off Bodega Head, California (38.3°N, 123.0°W), and in the Southern California Bight off Santa Catalina Island (33.4°N, 118.4°W; Fig. 1). At both sites, local variation in foraminiferal abundances and species composition is well understood (Thunell and Sautter, 1992; Field, 2004, Davis et al., 2016), driven by periods of upwelling, relaxation or downwelling, and/or seasonal predominance of the Davidson Current. The Santa Catalina Island site is close to the San Pedro Ocean Time-series (SPOT; 33.55°, 118.4°W) station, where the composition of the bacterial assemblage in the water column has been monitored for over a decade (Chow et al., 2013; Cram et al., 2015).

2.2 Sample collection

The Bodega Head samples were collected during November 2014, when there is a relaxation in the upwelling before the winter storms, and also in April 2015 (Table 1). Normally, the southerly along-shore winds start to drive renewed upwelling in early spring (Garcia-Reyes and Largier, 2012) but the samples for this study were collected prior to the delayed upwelling season of 2015, an unusual phenomenon possibly due to the strong El Niño–Southern Oscillation (e.g. Jacox et al., 2015). Bodega Head samples were obtained from vertically integrated 150µm mesh-size net tows, deployed to a maximum depth of 160 m, or to 10 m above the seafloor at shallower sites. Tow material was transferred to ambient surface seawater and kept chilled during transit to shore at the Bodega Marine Laboratory, where live foraminifera were
Figure 1. Map of the Californian coast and, at higher magnification, the region within the red box (insert) showing the Bodega Head and Santa Catalina Island sampling locations, (black circles) and the SPOT sampling site (white circle). The hydrography of the region is described in Sect. 2.1, whilst the directions of flow of the two major coastal currents, the California Current and the Davidson Current, are indicated by the arrows.

wet picked. *G. bulloides* were then identified morphologically to the morphospecies level, rinsed in 0.6 µm filtered surface seawater and preserved in RNA*Later®* (Ambion™). This reagent conserves cell integrity, inhibits nucleases at ambient temperatures, and dissolves the calcite shell. The Santa Catalina Island samples were collected by scuba diving or net tows during July/August 2013 towards the end of the weaker upwelling season off southern California. Collected foraminifera were treated as at the Bodega Marine Laboratory and transferred to RNA*Later®* at the Wrigley Marine Science Center.

### 2.3 Decalcification and washing of samples

To remove the shell and shell-associated, external contaminants, each individual specimen was decalcified by exposure to RNA*Later®* (Ambion™). The cell was then washed in filter-sterilised, salt-adjusted phosphate-buffered saline (PBS) or sterile artificial seawater, transferred to a new sterile 1.5 mL tube and washed a further three times before being transferred to DOC DNA extraction buffer (Sect. 2.4; Holzman and Pawlowski, 1996) for DNA analysis, or 4 % (*w/v*) paraformaldehyde in salt-adjusted PBS for 4 h at 4°C. Fixed cells were transferred to a polylysine-coated microscope slide and dehydrated through an ethanol series of 70, 90 and 100 % ethanol. Cells were stained in 1 µg mL$^{-1}$ DAPI (dilactate, Sigma-Aldrich) in PBS for 3 min and then rinsed with sterile deionised water. The stained preparations were mounted in AF1 mountant solution (Citifluor) and bacteria and eukaryotic nuclei visualised using a Zeiss Axio Imager Fluorescence microscope equipped with a DAPI filter set.

An unstained specimen of *G. bulloides* was also examined by fluorescence microscopy to observe the background levels of autofluorescence under the DAPI filter set to compare with the appearance of DAPI stained individuals. A TRITC filter set (excitation wavelength 540 nm, emission wavelength 580 nm) was used also on unstained individuals to investigate for the presence of autofluorescent, phycoerythrin-containing cyanobacterial cells.

### 2.4 Foramin genotyping and Sanger DNA sequencing

DNA was extracted from individual foraminifer specimens using the DOC extraction method to identify the specific genotype (Holzman and Pawlowski, 1996). PCR amplification of the foraminiferal 18S rRNA gene was performed according to Secars et al. (2012). DNA sequencing was carried out using the BigDye® Terminator v3.1 Cycle Sequencing Kit and an ABI 3730 DNA sequencer (both Applied Biosystems).

### 2.5 DAPI staining and fluorescence microscopy

Foraminifer cells were stained with 4',6-diamadino-2-phenylindole (DAPI), which forms a highly fluorescent DAPI-DNA complex that allows the visualisation of bacterial cells and eukaryotic cell nuclei under fluorescence microscopy. Individual decalcified and washed foraminifera were fixed in 4 % (*w/v*) paraformaldehyde in salt-adjusted PBS for 4 h at 4°C. Fixed cells were transferred to a polylysine-coated microscope slide and dehydrated through an ethanol series of 70, 90 and 100 % ethanol. Cells were stained in 1 µg mL$^{-1}$ DAPI (dilactate, Sigma-Aldrich) in PBS for 3 min and then rinsed with sterile deionised water. The stained preparations were mounted in AF1 mountant solution (Citifluor) and bacteria and eukaryotic nuclei visualised using a Zeiss Axio Imager Fluorescence microscope equipped with a DAPI filter set.

### 2.6 DNA extraction, amplification and 16S rRNA gene metabarcoding

DNA for 16S rRNA gene metabarcoding of the bacterial population within the foraminifera was extracted from decalcified and washed planktonic foraminiferal cells by the DOC extraction method (Holzman and Pawlowski, 1996). The DNA from the six samples – three independent *G. bulloides* isolates (BUL34, BUL36, BUL37) and a single non-spinose *Neogloboquadrina dutertrei* (DUT55) collected in July/August off Santa Catalina Island and two additional *G. bulloides* cells collected in November off Bodega Head (BUL22, BUL23; Table 1) – was amplified alongside three reagent controls containing (i) no DNA template (two replicates) and (ii) DOC buffer only. The V4 region of the
Table 1. Sampling information and details of analyses performed for each planktonic foraminifer specimen collected.

<table>
<thead>
<tr>
<th>Morphospecies</th>
<th>Sample ID</th>
<th>Sampling site</th>
<th>Sampling date</th>
<th>Co-ordinates</th>
<th>Sea surface temperature</th>
<th>Analysis</th>
</tr>
</thead>
<tbody>
<tr>
<td>G. bulloides</td>
<td>BUL21</td>
<td>Santa Catalina Island</td>
<td>Jul/Aug 2013</td>
<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>Control for fluorescence microscopy</td>
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<tr>
<td>G. bulloides</td>
<td>BUL24</td>
<td>Santa Catalina Island</td>
<td>Jul/Aug 2013</td>
<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>DAPI staining and fluorescence microscopy</td>
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<td>BUL25</td>
<td>Santa Catalina Island</td>
<td>Jul/Aug 2013</td>
<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>DAPI staining and fluorescence microscopy</td>
</tr>
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<td>BUL26</td>
<td>Santa Catalina Island</td>
<td>Jul/Aug 2013</td>
<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>DAPI staining and fluorescence microscopy</td>
</tr>
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<td>G. bulloides</td>
<td>BUL28</td>
<td>Santa Catalina Island</td>
<td>Jul/Aug 2013</td>
<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>DAPI staining and fluorescence microscopy</td>
</tr>
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<td>G. bulloides</td>
<td>BUL29</td>
<td>Santa Catalina Island</td>
<td>Jul/Aug 2013</td>
<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>DAPI staining and fluorescence microscopy</td>
</tr>
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<td>BUL30</td>
<td>Santa Catalina Island</td>
<td>Jul/Aug 2013</td>
<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>TEM</td>
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<td>G. bulloides</td>
<td>BUL32</td>
<td>Santa Catalina Island</td>
<td>July/Aug 2013</td>
<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>Metabarcoding, genotyping and Synechococcus 16S and rbcL, cloning and sequencing</td>
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<td>G. bulloides</td>
<td>BUL34</td>
<td>Santa Catalina Island</td>
<td>Jul/Aug 2013</td>
<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>Metabarcoding</td>
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<td>G. bulloides</td>
<td>BUL36</td>
<td>Santa Catalina Island</td>
<td>Jul/Aug 2013</td>
<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>Metabarcoding</td>
</tr>
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<td>G. bulloides</td>
<td>BUL37</td>
<td>Santa Catalina Island</td>
<td>Jul/Aug 2013</td>
<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>Metabarcoding</td>
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<td>BUL39</td>
<td>Santa Catalina Island</td>
<td>Jul/Aug 2013</td>
<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>TEM</td>
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<td>G. bulloides</td>
<td>BUL69</td>
<td>Santa Catalina Island</td>
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<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>TEM</td>
</tr>
<tr>
<td>N. dumerili</td>
<td>DUT55</td>
<td>Santa Catalina Island</td>
<td>Jul/Aug 2013</td>
<td>33.4°N, 118.4°W</td>
<td>18–21.5°C</td>
<td>Metabarcoding and genotyping</td>
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<td>BUL04</td>
<td>Bodega Head</td>
<td>Nov 2014</td>
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<td>14–15°C</td>
<td>Synechococcus 16S and rbcL</td>
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<td>BUL05</td>
<td>Bodega Head</td>
<td>Nov 2014</td>
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<td>14–15°C</td>
<td>Synechococcus 16S and rbcL</td>
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<td>BUL13</td>
<td>Bodega Head</td>
<td>Nov 2014</td>
<td>38.3°N, 123.0°W</td>
<td>14–15°C</td>
<td>Synechococcus 16S and rbcL</td>
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<td>BUL14</td>
<td>Bodega Head</td>
<td>Nov 2014</td>
<td>38.3°N, 123.0°W</td>
<td>14–15°C</td>
<td>Synechococcus 16S and rbcL</td>
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<td>BUL15</td>
<td>Bodega Head</td>
<td>Nov 2014</td>
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<td>14–15°C</td>
<td>Synechococcus 16S and rbcL</td>
</tr>
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<td>G. bulloides</td>
<td>BUL22</td>
<td>Bodega Head</td>
<td>Nov 2014</td>
<td>38.3°N, 123.0°W</td>
<td>14–15°C</td>
<td>Synechococcus 16S and rbcL</td>
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<tr>
<td>G. bulloides</td>
<td>BUL23</td>
<td>Bodega Head</td>
<td>Nov 2014</td>
<td>38.3°N, 123.0°W</td>
<td>14–15°C</td>
<td>Metabarcoding</td>
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<td>BUL71</td>
<td>Bodega Head</td>
<td>Apr 2015</td>
<td>38.3°N, 123.0°W</td>
<td>10.5°C</td>
<td>TEM</td>
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<td>BUL73</td>
<td>Bodega Head</td>
<td>Apr 2015</td>
<td>38.3°N, 123.0°W</td>
<td>10.5°C</td>
<td>Synechococcus 16S and rbcL</td>
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<td>Bodega Head</td>
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<td>38.3°N, 123.0°W</td>
<td>10.5°C</td>
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<td>BUL82</td>
<td>Bodega Head</td>
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<td>10.5°C</td>
<td>Synechococcus 16S and rbcL</td>
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<td>BUL83</td>
<td>Bodega Head</td>
<td>Apr 2015</td>
<td>38.3°N, 123.0°W</td>
<td>10.5°C</td>
<td>Synechococcus 16S and rbcL</td>
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<td>BUL84</td>
<td>Bodega Head</td>
<td>Apr 2015</td>
<td>38.3°N, 123.0°W</td>
<td>10.5°C</td>
<td>Synechococcus 16S and rbcL</td>
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<td>G. bulloides</td>
<td>BUL85</td>
<td>Bodega Head</td>
<td>Apr 2015</td>
<td>38.3°N, 123.0°W</td>
<td>10.5°C</td>
<td>Synechococcus 16S and rbcL</td>
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<td>G. bulloides</td>
<td>BUL86</td>
<td>Bodega Head</td>
<td>Apr 2015</td>
<td>38.3°N, 123.0°W</td>
<td>10.5°C</td>
<td>Synechococcus 16S and rbcL</td>
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<td>G. bulloides</td>
<td>BUL88</td>
<td>Bodega Head</td>
<td>Apr 2015</td>
<td>38.3°N, 123.0°W</td>
<td>10.5°C</td>
<td>Synechococcus 16S and rbcL</td>
</tr>
</tbody>
</table>

* PCR amplification of Synechococcus 16S rRNA gene and rbcL (RuBisCo large subunit).

16S RNA gene was chosen for amplification using the 515F forward primer and a barcoded 806R reverse primer series (Caporaso et al., 2012). This primer set is widely used by the Earth Microbiome Project (Gilbert et al., 2010), and therefore the amplification biases are known and well documented. For example, there is a bias against amplification of the SAR11 group of marine Alphaproteobacteria and a bias towards over-amplification of Gammaproteobacteria (Apprill et al., 2015; Walters et al., 2016; Parada et al., 2016). Each DNA sample and control was PCR-amplified with a unique barcode tag that enabled demultiplexing of the samples after being pooled for sequencing. The thermal cycling conditions are detailed by Caporaso et al. (2012). PCR reactions contained 1× Taq buffer plus additional MgCl₂ (final concentration 2.5 mM), 0.2 mM of each dNTP, 0.25 µM of each primer, 1 µL of template DNA and 1.25 U of Taq DNA polymerase (Roche Applied Science), with the volume made up to 25 µL with PCR-grade water (Sigma). All PCR reactions were set up in a PCR6 vertical laminar airflow cabinet with UV sterilisation (Labcare Systems, Bristol, UK) as described by Pagaling et al. (2014). Reaction tubes and PCR mixtures were treated for 15 min with 15 W UV light (wavelength = 254 nm) to destroy contaminating DNA, prior to addition of dNTPs, Taq polymerase and template DNA (Padua et al., 1999). The six functional and three control PCR reactions were run on a 1% agarose gel and the products were purified with the Wizard® SV Gel and PCR Clean-Up System (Promega). The purified amplicons were quantified using a Quant-iT PicoGreen ds-DNA assay kit (Life Technologies) prior to pooling at equimolar concentrations for DNA sequencing. The total number of quality-filtered sequencing reads including controls was 862,954. DNA sequencing was performed at Edinburgh Genomics using an Illumina MiSeq v3 to generate 250 base pair (bp) paired-end reads.

2.6.1 Quality filtering and contaminant removal

The Quantitative Insights in Microbial Ecology (QIIME, v1.8.0; Caporaso et al., 2010) pipeline was used to assemble paired-end reads and quality-filter the sequences. Raw reads were paired with an overlap of 200 bp and quality-filtered with a minimum Phred score of 20 for maximum accuracy (Kozich et al., 2013). Reads of less than 245 bp (i.e. short reads) were removed from the dataset with the Python script filter_short_reads.py from http://gist.github.com/walterst/7602058. Chimeras were detected using USEARCH v6.1.544 default settings (Edgar et al., 2011) and version 13_8 of Greengenes 16S rRNA gene reference database (DeSantis et al., 2006). Given the low yield of endogenous bac-
terial DNA in these small-sized samples, we anticipated that amplicon contamination from PCR amplification reagents, DNA extraction reagents, and the ultra-pure water system would contribute a significant number of DNA sequences and OTUs from contaminant genera to the sample set (Salter et al., 2014; Laurence et al., 2014). Operational taxonomic units with greater than 100 sequences in any of the three control samples were considered to be potential contaminants and were removed from the sample set. Two OTUs were removed due to contamination in the two PCR controls: a Bradyrhizobiaceae of the class Alphaproteobacteria and an Acinetobacter of the class Gammaproteobacteria. Twelve contaminating OTUs were removed due to contamination via the DOC buffer, with eight of these of the class Alphaproteobacteria. Seven of the eight were from the order Rhizobiales, five of which were classified to a higher taxonomic level and included two of additional Bradyrhizobiaceae OTUs and one each of Methyllobacterium, Mesorhizobium, Pedomicrobium. The eighth OTU from the class Alphaproteobacteria was a Sphingomonas from the order Sphingomonadales. The final four OTUs were Burkholderia bryophila of the class Betaproteobacteria; two Sediminibacterium OTUs of the phylum Bacteroidetes; and lastly an OTU identified as a Streptophyta chloroplast. A single Bradyrhizobiaceae OTU was by far the largest contaminant, with a total of 284,636 sequences from all samples and controls, and it is known to be a common contaminant of next-generation sequencing data, along with other Alphaproteobacteria (Laurence et al., 2014).

2.6.2 Operational taxonomic unit (OTU) picking and taxonomic assignment

The default QIIME pipeline was used for data analysis: OTU picking and taxonomic assignment. De novo picking (pick_de_novo_otus.py) clusters DNA sequences into OTUs with 97% similarity with no external reference and selects a representative sequence of each OTU for alignment and subsequent assignment of taxonomy. This script keeps all diversity, including unknowns, in the sample set. Closed reference picking was also performed, which removes OTUs that are not closely matched (<97%) with OTUs in the Green-genes database (pick_closed_reference_otus.py). This output is required for normalisation by copy number (NBCN) using the online Galaxy tool (http://huttenhower.sph.harvard.edu/galaxy/). This corrects the abundance of each OTU to better reflect the true organism abundance by normalising predicted 16S rRNA gene copy number for each OTU. In both OTU picking methods, OTUs with fewer than 10 sequences across all samples were removed from the sample set (filter_otus_from_oder_table.py).

2.6.3 Alpha-rarefaction and sequencing depth

In QIIME, the script alpha_rarefaction.py was used to assess whether the sequencing depth was adequate to detect foraminiferal bacterial diversity. Samples were rarefied to the lowest sequencing depth observed across all samples (10,551 in closed-reference picking in sample BUL22) and OTU richness curves were generated, using the observed-species metric, which counts the number of unique OTUs found in a sample.

2.7 TEM

TEM was used to observe and document the structural relationships between the endobiotic bacteria and foraminiferal cells. Decalcified *G. bulloides* were fixed in 3% glutaraldehyde in 0.1 M sodium cacodylate buffer, pH 7.3, for 2 h followed by three 10 min washes in 0.1 M sodium cacodylate. Specimens were then post-fixed in 1% osmium tetroxide in 0.1 M sodium cacodylate for 45 min, followed by a further three 10 min washes in 0.1 M sodium cacodylate buffer. Specimens were then dehydrated in 50, 70, 90 and 100% ethanol (X3) for 15 min each, then in two 10 min changes in propylene oxide prior to being embedded in TAAB 812 resin. Sections (1 µm thick) were cut on a Leica Ultracut ultramicrotome, stained with toluidine blue, and then viewed under a light microscope to select suitable specimen areas for investigation. Ultrathin sections (60 nm thick) were cut from selected areas, stained in uranyl acetate and lead citrate and then viewed with a JEOL JEM-1400 Plus TEM.

2.8 Genetic identification of *Synechococcus* cells identified in *G. bulloides*

*Synechococcus* cells were found in large numbers inside *G. bulloides* and were genetically characterised. A 422 bp fragment of the *Synechococcus* 16S rRNA gene was amplified from total DNA extracted via the DOC method from individual specimen BUL34 (Table 1). This provided a larger, more informative fragment for phylogenetic analysis compared with the 253 bp generated by 16S rRNA metabar-coding. Cyanobacterial specific primers were used (CY A359f 5′-GGGGAATCYTTCCGCAATGGG-3′ and CY A781R a and b 5′-GACTACWGGGGTATCTAATCCCWTT-3′; Nübel et al., 1997) and thermocycler conditions were as follows: 94°C for 2 min followed by 30 cycles at 94°C for 15 s, 55°C for 15 s and 72°C for 30 s, followed by a final extension at 72°C for 5 min. PCR reactions were performed with My-Taq REDDY mix (Bioline) and 0.25 µM of each primer and 1 µL of template DNA, with the volume made up to 25 µL with PCR-grade water (Sigma). The PCR product obtained was cloned (TOPO®-TA cloning kit, Invitrogen) and Sanger sequenced.

Clone sequences were aligned with reference *Synechococcus* 16S rRNA gene sequences retrieved from the GenBank database (NCBI) using ClustalW software within the package MEGA6 (Tamura et al., 2013). Phylogenetic trees (maximum likelihood, neighbour joining, minimum evolution, UPGMA, maximum parsimony; Sect. 3.5) were gen-

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erated using the default settings of MEGA6 with 500 bootstrap resamplings to determine the closest taxonomic affiliations (i.e. clade designation sensu Fuller et al., 2003) of the G. bulloides-associated Synechococcus. Informed by this analysis, further primers were designed that target other signature genes harboured by the Synechococcus clades identified, including that for rbcL. This phylogenetically informative gene encodes the large subunit of RuBisCO (ribulose-1,5-bisphosphate carboxylase/oxygenase), the primary CO2-fixing enzyme found in cyanobacteria. Primers (SynrbcL_For 5'-CGGCAACTTCTTCAGTCAGG-3'; SynrbcL_Rev1 5'-ATGTCGGGCTTTCTTCTC-3'; SynrbcL_Rev2 5'-CCGGCTTTCCATAAGGATGTC-3') were designed with Primer 3 (http://primer3.ut.ee/) that target a 252 bp fragment of rbcL from the most closely related Synechococcus spp. (i.e. strains CC9902, CC9311 and WH8120; see below).

Purified DNA from 14 G. bulloides specimens (Table 1) generated products of the correct size on PCR amplification with the rbcL primers. The product obtained from isolate BUL34 was selected and TA-cloned and DNA-sequenced as described above. PCR reactions were performed in a Biometra Personal Thermocycler using MyTaq REDDY mix (Bioline) and 0.25 µM of each primer and 1 µL of template DNA, with the volume made up to 25 µL with PCR-grade water (Sigma). Thermocycling conditions were 94 °C for 2 min followed by 30 cycles at 94 °C for 15 s, 56 °C for 15 s and 72 °C for 30 s, followed by a final extension at 72 °C for 5 min.

3 Results

In total, 29 individual specimens of G. bulloides collected from waters off Santa Catalina Island and Bodega Head (Fig. 1) were investigated during this study. The sampling information and analyses performed on each specimen are detailed in Table 1, and the sampling strategy and genetic characterisation are described in the methods.

3.1 Genotyping of foraminifera

Partial 18S rRNA (SSU) gene sequences amplified from BUL34 and DUT55 have been submitted to Genbank (NCBI accession numbers KX816046 and KX816047 respectively). G. bulloides specimen BUL34 is type IId and N. dutertrei specimen DUT55 is type Ic and it is the first time a N. dutertrei specimen has been amplified for this genotype. Both genotypes have been submitted to Genbank (NCBI, Bioproject accession numbers PRJNA341960; SRA accession numbers SRR4271458, SRR4271479, SRR4271493, SRR4271505).

3.2 Fluorescence microscopy and DAPI staining

Fluorescence microscopy examination of an unstained, fixed G. bulloides specimen (BUL21; Table 1; Supplement Fig. S1) under the DAPI filter set demonstrated high levels of diffuse autofluorescence across the entire cell. G. bulloides cells that were first stained with DAPI (n = 6; Table 1) also showed background autofluorescence, but in addition some more highly fluorescent regions 3–10 µm in size, that correspond to vacuoles containing condensed prey items. The many ~1 µm diameter brightly stained globular structures that occur throughout the cell are consistent with the presence of intracellular bacterial cells. The white rectangle denotes the area magnified in (b) and shows examples (white arrows) of these DNA-containing, bacteria-sized structures and their cytoplasmic location.

3.3 16S rRNA gene metabarcoding

16S rRNA gene metabarcoding was carried out on five specimens of G. bulloides and a single specimen of the non-spinoso species Neogloboquadrina dutertrei for comparison (Table 1). The raw dataset has been submitted to the Sequencing Read Archive (SRA, NCBI) (Bioproject accession number PRJNA341960; SRA accession numbers SRR4271458, SRR4271479, SRR4271493, SRR4271505, SRR4271534, SRR4271535).
Figure 3. Fluorescence micrograph of a decalcified *G. bulloides* cell examined under the TRITC filter set (excitation 540 nm, emission wavelength > 580 nm). (a) The cytoplasm from two of the chambers is shown. The autofluorescence localised to the ∼1 µm diameter globular structures found throughout the cell is consistent with the presence of phycoerythrin-containing *Synechococcus*. The white rectangle denotes the area magnified in (b) in which the autofluorescing structures can be seen as discrete entities demonstrating that the cyanobacterial cell membranes are intact (see text). The white arrowheads highlight a group of bacteria that can also be identified by DAPI staining when examined under the DAPI filter set in (e).

SRR4271506, SRR4271507, Bird et al., 2017). A total of 862,954 sequences were generated from the six samples and three controls after quality filtering and removal of short reads (< 245 bp) and chimeric sequences (Sect. 2.6.1). In closed-reference picking, after removal of pynast failures, all control sequences (288,985) contaminant OTUs (including 161,282 sequences as result of the single contaminating Bradyrhizobiaceae OTU across all six samples) and OTUs with an abundance of less than 10 sequences across all samples, a total of 214,087 sequences were clustered into OTUs and assigned taxonomy (Sect. 2.6.1 and 2.6.2). The numbers of sequences and OTUs generated in individual specimens for both closed-reference picking and de novo picking are listed in Supplement Table S1. The OTU profiles within a specimen were highly similar between de novo picking and closed-reference picking with normalisation by copy number (NBCN, Sect. 2.6.2). Therefore, we present results for closed-reference picking with NBCN and indicate when de novo picking OTUs are being represented. Rarefaction curves (Fig. S2) for OTU richness confirmed that sequencing depth was sufficient to capture the full bacterial assemblage diversity.

All replicates of *G. bulloides* contained a highly distinctive assemblage of OTUs. The OTU assemblage of an individual *N. dutertrei* (Table 1) sampled at the same time and from the same water mass is shown for direct comparison of the bacterial assemblages of two morphospecies of foraminifera (Fig. 4). Within the five BUL specimens investigated (Table 1), 37–87% of all sequences belonged to five OTUs assigned to the unicellular, cyanobacterial genus *Synechococcus* (Fig. 4). This was by far the greatest abundance of a single genus of bacteria; no other genera, or indeed family or order, of bacteria were found across all five BUL specimens at relative abundances consistently more than 2%. Therefore, the next highest relative abundance group must be described at the class level. Across four (BUL22, BUL23, BUL34, BUL36) of the five BUL specimens 15–31% of sequences belonged to the class Alphaproteobacteria and were dispersed amongst 81 OTUs. The fifth and outlying specimen (BUL37, containing 87% *Synechococcus* sequences) contained only 2.6% Alphaproteobacteria across 37 OTUs. It contained marginally more phylum Actinobacteria (1.8%) and class Betaproteobacteria (1.4%) sequences, but the sequence abundances of these classes were more similar to the other BUL specimens. The high relative abundance of *Synechococcus* within this specimen might be due to a lack of feeding on other bacteria or algae immediately prior to sampling, indicative of a high turnover rate for prey bacteria and algal cells.

There were no chloroplast-affiliated OTUs in specimens BUL22 and BUL23. However, 6.4, 27 and 3% of sequences in specimens BUL34, BUL36 and BUL37 respectively were allocated to chloroplast 16S rRNA gene OTUs from a variety of sources. These were two OTUs from a mixotrophic protist belonging to the diverse, protozoan phylum Cercozoa (Cavalier-Smith and Chao 2003); three OTUs from the phylum of Haptophyte algae, which includes coccolithophores; eight OTUs from the phylum Stramenopile, which includes both diatoms and chrysophyte algae; and, finally, 13 OTUs from the group Streptophyta (an unranked clade of plants that includes green algae).

In de novo picking, *G. bulloides* specimens BUL34, BUL36 and BUL37 contained varying percentages of sequences (25.4, 0.3 and 1.6% and respectively) in three taxonomically unassigned OTUs, whereas BUL22 and BUL23 did not contain any unassigned sequences. Of all the unassigned sequences, 94% belonged to a single OTU (e.g. 5878 sequences in sample BUL34) that was 99% identical (100% coverage of 253 bp) to an unidentified marine bacterial clone (accession number HQ673258) retrieved from the northeast subarctic Pacific Ocean (Allers et al., 2013). The nearest match (87–89% similarity over 100% coverage) to an identified phylum was to a large number of uncultured Verrucomicrobiae bacteria of the phylum Verrucomicrobia from a wide range of habitats, including marine environments. Whilst this was not a particularly close match, it is within the defined limit of > 85% DNA identity that delineates a phylum (Hugenholtz et al., 1998; Rappé and Giovannoni, 2003), albeit based only on a 253 bp fragment. The variation in abundances of this OTU found in each *G. bulloides* individual analysed (25% in BUL34; 0.3% in BUL36 and 1.6% in BUL37) might indicate this bacterium has a patchy distribution and is an opportunistic food source.
of Synechococcus
tractions with those of the water column, the concentration (Fig. 5c).
the cytoplasm and also in vacuoles of all individual
Synechococcus cyanobacterium
surrounded by thylakoid membranes, characteristic of the
ing carboxysomes (Fig. 5a) within the central cytoplasm
symbionts. However, numerous intact coccoid cells contain-
endobionts were present within the cell. No intracellular
ried out on four
G. bulloides
Transmission electron microscopy (TEM) imaging was car-
Figure 4. Relative abundance of taxonomically assigned 16S rRNA
gene sequences from bacteria and chloroplasts within the cytoplasm
of six individual foraminifer specimens: five G. bulloides (BUL22, BUL23, BUL34, BUL36 and BUL37) and one N. dutertrei
specimen (DUT55). Sequences are assigned to operational taxonomic
units (OTUs) grouped at different levels of taxonomic classification
(see key). 16S rRNA gene sequences assigned to OTUs of the genus
Synechococcus are the most abundant within G. bulloides and are at
the highest level of classification when compared to the other OTUs
assigned to individual classes or phyla.

3.4 Transmission electron microscopy

Transmission electron microscopy (TEM) imaging was carried out on four G. bulloides specimens (BUL32, BUL39, BUL69, BUL71; Table 1, Fig. 5) to observe whether any endobionts were present within the cell. No intracellular eukaryotic cells were observed, confirming a lack of algal symbionts. However, numerous intact coccolid cells containing carboxysomes (Fig. 5a) within the central cytoplasm surrounded by thylakoid membranes, characteristic of the cyanobacterium Synechococcus, were observed throughout the cytoplasm and also in vacuoles of all individual G. bulloides observed (Fig. 5b). Approximately 5% of the observed Synechococcus cells were undergoing cell division (Fig. 5c).

To compare foraminiferal cellular Synechococcus concentrations with those of the water column, the concentration of Synechococcus cells per millilitre of foraminiferal cytoplasm was calculated by assuming a conservative average host cell diameter of 200 μm (Spero and Lea, 1996; Aldridge et al., 2012), a spheroid morphology (Geslin et al., 2011) and that the cytoplasm was equivalent to 75% of the shell volume (Hannah et al., 1994). Based on averaged cell counts from the TEM images, the total number of Synechococcus cells within G. bulloides occupied less than 2% of the foraminiferal cell volume but was equivalent to 3.8 × 10⁹ Synechococcus cells mL⁻¹. This is far higher than the well established range of concentrations of Synechococcus found throughout the global ocean that range from 1 × 10²–1.5 × 10⁶ Synechococcus cells per millilitre of seawater (Partensky et al., 1999; Paerl et al., 2011). In the Southern California Bight, Synechococcus cell counts are generally fewer than 1.5 × 10⁵ cells mL⁻¹ but can reach 6 × 10⁵ cells mL⁻¹ during the blooms that are generally observed in late spring to early summer (Tai and Palenik, 2009; Tai et al., 2011). The concentration of Synechococcus in the G. bulloides cell was therefore up to 4 orders of magnitude greater than peak bloom concentrations measured in the California Bight. This suggests that Synechococcus cells accumulate within the cytoplasm of G. bulloides type IId.

3.5 Genetic characterisation of intracellular Synechococcus

Figure 5. Transmission electron microscope images of Synechococcus cells inside G. bulloides. (a) A Synechococcus cell with characteristic polyhedral carboxysomes in the central cytoplasmic region (white arrow) surrounded by thylakoid membranes. (b) Numerous Synechococcus cells within a G. bulloides cell are observed in both the cytoplasm and vacuoles (black arrow). This is a region of cytoplasm rich in fibrillar bodies (white arrow) found only in planktonic foraminifera, and whose function is unknown. (e) Synechococcus cell within a G. bulloides cell undergoing cell division as indicated by the presence of a constriction at the cell midpoint (white arrow).

Five Synechococcus OTUs were assigned in 16S rRNA gene metabarcoding with closed-reference picking. However, more than 99% of the BUL Synechococcus sequences were assigned to just one of these OTUs. The representative nucleotide sequence (253 bp) of this OTU is a 100% match to the coastal clade IV Synechococcus sp. strain
CC9902, originally isolated from the California Current. Two further OTUs were highly similar to this abundant OTU and were 99% identical to *Synechococcus* sp. strain CC9902. The remaining two OTUs both had a nucleotide match of 99% with *Synechococcus* sp. strain WH8020, a clade I strain also found typically in coastal waters. In order to confirm these clade assignments, phylogenetic analysis (Fig. S3) of a larger (422 bp) fragment of the *Synechococcus* 16S rRNA gene generated from BUL34 total DNA was performed. Ten clones (GenBank accession numbers KX815969–KX815978) clustered with the clade IV *Synechococcus* sp. strain CC9902 and two clones (GenBank accession numbers KX815979 and KX815980) clustered with clade I strains CC9311 (another California Current isolate) and WH8020, in agreement with the 16S rRNA gene metabarcoding data. The topologies of the phylogenetic trees produced were all in overall agreement with well-established analyses of *Synechococcus* 16S rRNA genes (Scanlan et al., 2009), confirming the phylogenetic resolution of the sequence data included in the present study.

In addition, a 252 bp fragment of the *Synechococcus* rbcL gene was cloned and 230 bp of this clone was DNA-sequenced (GenBank accession number KX816048) from BUL34 (Table 1). A GenBank BLAST search (NCBI) found 100% nucleotide sequence identity with the RuBisCo large subunit coding region of *Synechococcus* sp. strain CC9902 and 92% identity with *Synechococcus* sp. strain WH8020, confirming the presence of *Synechococcus* sp. strain CC9902, or a very closely related clade IV strain. The DNA of 13 further *G. bulloides* specimens (Table 1) also yielded products of ~252 bp on amplification with the *Synechococcus* rbcL primers confirming the consistency of the association between *G. bulloides* type IId and *Synechococcus* strains in the California Current year round.

4 Discussion

Our results highlight a novel endobiotic association between the usually free-living, photoautotrophic picocyanobacterium *Synechococcus* and its host, *G. bulloides* type IId, a genotype of a spinose planktonic foraminiferal morphospecies, barren of protist algal symbionts. Below, we discuss the evidence for this endobiosis, and possible roles of *Synechococcus* in *G. bulloides* host metabolism and its characteristic cytoplasm colouration. A better understanding of *G. bulloides* genotype ecology will ultimately provide ecological information for modelling foraminiferal distribution, abundance and seasonality under different climate regimes and improve the accuracy of the palaeoceanographic proxy records.

4.1 Evidence for *Synechococcus* as an abundant endobiont of *Globigerina bulloides* type IId

*G. bulloides* has consistently been reported to be barren of protist algal symbionts (Febvre-Chevalier, 1971; Gastrich, 1987; Hemleben et al., 1989; Spero and Lea, 1996). The current study supports this conclusion, since no intact algal cells were found in any of the *G. bulloides* cell sections examined using TEM. However, we do have strong evidence that *G. bulloides* type IId contains large numbers of the photoautotrophic picocyanobacterium *Synechococcus*. Intact *Synechococcus* cells accumulate within the host cytoplasm in abundances far greater than those found under bloom conditions in the California Bight or in any other foraminiferal species investigated. How this association occurs is unclear, but *G. bulloides* type IId is the only foraminiferal species currently known to associate with *Synechococcus*, an observation that implies a specific potentially mutualistic benefit, particularly since *G. bulloides* does not exploit protist algal symbionts as in other spinose species. Based on the observations discussed below, we propose that these picocyanobacteria are abundant, metabolically active endobionts living within the *G. bulloides* cell, rather than prey.

4.1.1 *Synechococcus* cells are intact and viable

DNA degradation in prey items limits the success of amplification of DNA sequences greater than ~250 bp (Pompanon et al., 2012). In this study we targeted a 253 bp fragment of the 16S rRNA gene via metabarcoding, thus providing information not only on intact, undigested bacteria but also on those bacteria phagocytosed for food. Subsequent TEM imaging has enabled us to distinguish between prey and endobiont. Indeed, TEM images have demonstrated that, of the diversity of bacteria identified in 16S rRNA gene metabarcoding, only *Synechococcus* cells were observable in the *G. bulloides* cell. The *Synechococcus* cell membranes were physically intact (Fig. 5a) and, whilst some *Synechococcus* cells were observed within vacuoles, many were distributed throughout the cytoplasm of *G. bulloides* (Fig. 5b), where digestion does not occur. As many as 5% of the intracellular *Synechococcus* population were observed to be in the process of cell division (Fig. 5c), indicative of actively growing, viable individuals (Campbell and Carpenter, 1986). Significantly, Bernhard et al. (2000) considered as few as 3% dividing cells a substantial enough proportion of the population to suggest a symbiotic role for the intracellular bacteria they observed in the benthic foraminifer *Buliminella tenuata*. Further, successful PCR amplification of a longer (422 bp) fragment of the *Synechococcus* 16S rRNA gene suggests that the *Synechococcus* DNA was more intact than might be expected if it were the DNA of a prey organism (i.e. >250 bp; Pompanon et al., 2012). This and the amplification of a second short fragment of the *rbcL* gene provides additional evidence that *Synechococcus* DNA was not
grossly degraded by nucleases. In further confirmation of the intact nature of the intracellular *Synechococcus* population, autofluorescence in the orange/red spectral region arising from the photosynthetic pigment phycoerythrin was readily detected within these DNA-containing endobionts within *G. bulloides* (Fig. 3). Phycoerythrin, a water-soluble biliprotein found routinely in marine *Synechococcus*, rapidly diffuses into the aqueous surroundings if the cell membranes are compromised (Stewart and Farmer, 1984; Wyman, 1992).

### 4.1.2 Synechococcus are endobionts in marine protists

Whilst *Synechococcus* spp. are known primarily as free-living organisms (Waterbury et al., 1979; Richardson and Jackson, 2007), an endobiotic lifestyle has also been observed in association with a number of different marine protist groups. *Synechococcus* has been identified in the benthic foraminifer *Fusenkoina rotundata*, sampled from the benthos at 600 m using both fluorescence microscopy (identified as cyanobacteria by Bernhard et al., 2000) and in TEM imaging (identified as *Synechococcus*; Buck and Bernhard, 2006). At these depths, however, the *Synechococcus* endobionts would be unable to photosynthesise, which rules out the most obvious functional metabolic role for this potential symbiont. *Synechococcus* has also been found living embedded within the extracellular matrix surrounding a marine diatom (Buck and Bentham, 1998), and within a polycystine radiolarian (Yuasa et al., 2012). This study now confirms that they are also to be found within the living cells of at least one type of planktonic foraminifer.

### 4.1.3 Synechococcus cells accumulate in the *G. bulloides* cytoplasm

Intact *Synechococcus* cells accumulate within the *G. bulloides* cytoplasm at densities (~3.8 × 10⁹ cells mL⁻¹) that are 4 orders of magnitude more concentrated than those reported in the surrounding seawater (Tai and Palenik 2009; Tai et al., 2011). Whilst DNA sequences from other bacteria were identified by 16S rRNA gene metabarcoding (Fig. 4), no bacterial cells lacking carboxysomes were observed by TEM, indicating that, unlike *Synechococcus*, other bacteria were rapidly digested once taken up. Quite how *Synechococcus* accumulate in the foraminiferal cell is yet to be established. For example, does the host or endobiont instigate the association? Are the cyanobacteria passed on via parental gametes or is the association established via direct uptake of *Synechococcus* from the water column? In the case of planktonic foraminifera harbouring protist algae, a small number of symbionts are taken up directly from the water column (horizontal transmission) rather than being inherited through vertical transmission via parental gametes (Hemleben et al., 1989; Bijma et al., 1990). Juveniles with only 2 to 3 chambers already have ~3 to 5 symbionts, and it is assumed that they are taken up from the water column exclusively since no protist symbionts (5–10 µm cell diameter) have been observed within the much smaller flagellated gametes (~2.5 µm; Hemleben et al., 1989). Although picocyanobacteria such as *Synechococcus* are much smaller in size (~1 µm diameter) than algal symbionts and could potentially be inherited via parental gametes, we favour the hypothesis that the *Synechococcus* population within *G. bulloides* is similarly taken up from the water column, despite evidence for both horizontal (Ashton et al., 2003) and vertical transmission (Schweikert and Meyer, 2001) of bacteria within protist hosts.

To investigate the potential mode of transmission of the *Synechococcus*, we compared the strain assemblages within *G. bulloides* to those of the surrounding water column. If the *Synechococcus* endobionts were horizontally transferred to *G. bulloides* via uptake from the water column, we would expect that the diversity of the internal strain assemblage would mirror that of the surrounding waters closely. Alternatively, if the endobionts were vertically transmitted, a degree of genetic drift would be expected between the internal and free-living strains of *Synechococcus* as the result of genetic isolation over time (Wernegreen, 2002; Bright and Bulgheresi, 2010). Off the coast of California, the most prevalent strains of *Synechococcus* are those belonging to clades I and IV (see Fuller et al., 2003) that display seasonal population differences throughout the annual cycle (Tai and Palenik, 2009; Tai et al., 2011). The *Synechococcus* 16S rRNA gene sequences cloned from a *G. bulloides* specimen collected in July/August (Table 1) show that the strain composition strongly reflects the seasonal cladal distribution patterns that are observed in the water column at that time of year (Tai and Palenik, 2009). Up to 100 % nucleotide identity was found for the 16S rRNA gene clones and the *rbcL* gene sequences of the internal endobionts and those of the free-living clade IV *Synechococcus* sp. strain CC9902, originally isolated from waters off the California coast. Whilst there can be a high degree of diversity among strains seemingly closely related through *rbcL* and 16S rRNA gene phylogenies, this evidence supports a strategy of horizontal rather than vertical transmission for the *G. bulloides* endobionts.

### 4.1.4 Intracellular OTU relative abundances do not reflect those of the water column

The intracellular 16S rRNA gene OTU profiles of *G. bulloides* were very different from those of the water column assemblages, indicating uptake of specific bacteria from the general microbial population. The foraminifer collection site off Santa Catalina Island in the San Pedro Channel is adjacent to the SPOT sampling location, where seasonality and trophic interactions within the microbial assemblages in the water column have been studied routinely for over a decade (Chow et al., 2013; Cram et al., 2015). In both the surface waters and deep chlorophyll maximum layer, the microbial assemblage at the SPOT sampling site is dominated by OTUs
from the ubiquitous SAR11 group (Giovannoni 1990; Morris et al., 2002) of marine Alphaproteobacteria, which represent over 30% of the assemblage. In addition, members of the Actinobacteria account for approximately 15% of OTUs, while the picocyanobacteria represent just 2–5% of the total bacterioplankton. Of the latter, Prochlorococcus dominates the assemblage although Synechococcus is also present year round (Chow et al., 2013). The remaining 50% of the microbial population comprises a series of OTUs from a variety of marine bacteria each representing less than 2% of the assemblage (Chow et al., 2013). This water column assemblage contrasts strongly with the intracellular 16S rRNA gene OTUs of *G. bulloides*, where between 37 and 87% of the total number of sequences recovered belong to Synechococcus OTUs. It should be noted that no amplification bias towards Synechococcus has been reported for this primer set and data from a number of marine locations supports this (Apprill et al., 2016). Strikingly, Prochlorococcus sequences were not identified in the three *G. bulloides* specimens collected close to the SPOT sampling location (BUL34, BUL36 and BUL37), even though Prochlorococcus represents the majority of the picocyanobacteria in the water column in this region. Further, <4.5% of OTUs in the amplified *G. bulloides* specimens were assigned to the Actinobacteria (compared to ∼15% in the water column) and no OTUs of the ubiquitous SAR11 group of Alphaproteobacteria were identified in our sample set. However, in part this is likely to be a result of bias against SAR11 clades (Apprill et al., 2015; Walters et al., 2015) in the primer set used in this study (Caporaso et al., 2012).

The composition of the internal microbial population of the *G. bulloides* cells clearly does not mirror that of the surrounding water column, highlighting the genotype-specific nature of the OTU assemblages observed within the *G. bulloides* cell. This observation is reinforced by the fact that the intracellular OTU assemblage within *G. bulloides* also differs substantially from those identified within specimens of the non-spinose species *N. dutertrei* (e.g. DUT55; Fig. 4), collected at the same time and location. *N. dutertrei* contains ∼2% bacterial OTUs, with the majority of OTUs (>97%) being assigned to Stramenopiles (a group that includes diatoms and chrysophyte algae; 53%) and Cercocoz (a diverse phylum of mixotrophic protists; 44.5%). This highlights again the morphospecies/genotype-specific nature of the *G. bulloides* intracellular OTU assemblage.

### 4.1.5 Unusual cytoplasm colouration of *G. bulloides*: a role for endobiotic Synechococcus

Living *G. bulloides* cells often exhibit a distinctive brown colouration in the specimens found off the coast of California (Spero and Lea, 1996) that is not a general feature of the other spinose species in the region. The discovery of phycocerythrin-containing *Synechococcus* within the cytoplasm of the foraminifera reported here provides a plausible explanation for this unusual property. A number of *Synechococcus* strains isolated from the California Current are brown in colour owing to the production of urobilin-rich phycoerythrins (Toledo and Palenik, 1997). Many of the clade I and IV strains with which the *G. bulloides* endobionts cluster in phylogenetic analysis are type IV chromatic adapters that exhibit elevated concentrations of this urobilin-rich phycoerythrin (Six et al., 2007) as well as the photoprotective carotenoid zeaxanthin (Bidigare et al., 1989) under blue light (i.e. under the illumination conditions typical of the oligotrophic waters off the California coast from which the samples were obtained during the present study). The presence of these pigments within the *Synechococcus* endobionts therefore probably contributes to the unusual cytoplasm colouration observed in *G. bulloides* from this location.

### 4.2 Potential metabolic roles for the *G. bulloides* endobionts

There are some obvious potential metabolic benefits to each organism in a *G. bulloides*–*Synechococcus* partnership. Firstly, the foraminifer might benefit from a supply of photosynthetically fixed carbon, as is the case with the foraminifera that harbour protist algal symbionts (Caron et al., 1995; Ulhe et al., 1997, 1999). If this were the sole benefit, however, one would question why *G. bulloides* preferentially recruits *Synechococcus* for this purpose, rather than the more conventional algal symbionts found in other species. One possible explanation is that *G. bulloides* inhabits a wide range of depths that often extend below the photic zone and it is also common in unstable upwelling waters, where potential algal symbionts may not thrive. *Synechococcus* has been found alive in aphotic waters at depths of 600 m (Bernhard et al., 2000), and has been shown to assimilate carbon mixotrophically (Paoli et al., 2008). It could therefore augment phototrophy with carbon assimilated through (photo)heterotrophy, depending on the water column depth of the host. In addition, some *Synechococcus* strains within clade I and those so far characterised in clade IV, as found in *G. bulloides*, are chromatic adapters, able to modify their pigment composition and absorption properties depending on the underwater light field (Six et al., 2007). Such adaptability might make *Synechococcus* a more compatible symbiont for the *G. bulloides* lifestyle.

Alternatively, *Synechococcus* may have additional or quite separate functional roles in association with *G. bulloides* beside endobiotic photosynthetic activity within the photic zone. For example, approximately half of the nitrogen assimilated by the host cell in the *Orbulina universa* foraminifer–symbiont system is transferred via the algal symbionts, a contribution that increases further to ∼90–100% in nitrate-depleted waters (Ulhe et al., 1999). *Synechococcus* has a very high affinity for combined nitrogen (e.g. nitrate, nitrite and ammonium) and accumulates expanded stores of this element within its light-harvesting phycobilisomes un-
der N-replete conditions (Wyman et al., 1985). Likewise, *Synechococcus* sequesters large stores of P within its cells as polyphosphate, even under low external concentrations (Martin et al., 2014). These nutrient reservoirs could be readily mobilised and exploited by the foraminiferal cell, particularly prior to gametogenesis, when planktonic foraminifera require extra elemental resources for DNA production (Hemleben et al., 1989). For *Synechococcus*, being housed within a foraminiferal cell could protect it from grazers and the multitude of cyanophages present in the water column (Suttle and Chan, 1994; Mühling et al., 2005). *Synechococcus* may also benefit presumably from a supply of host metabolic by-products or from specific nutrients as products of prey digestion.

### 4.3 Feeding preferences and life strategy of *G. bulloides* type IId

TEM in combination with 16S rRNA gene metabarcoding enables identification of both bacteria and eukaryotic chloroplasts within the foraminiferal cell. This methodology does not amplify eukaryotic, nuclear-encoded (18S) rRNA genes and, as a result, does not provide any information about the non-chloroplast-bearing zooplankton prey of *G. bulloides*. Observations of large numbers of freshly collected specimens of *G. bulloides* confirm that they feed on small zooplankton prey as well as phytoplankton (Spero and Lea, 1996). Amongst the latter, a preference for some species of diatoms and chlorophytes over dinoflagellates or chrysophytes has been reported (Lee et al., 1966). Interestingly, however, two of the five *G. bulloides* specimens in this metabarcoding study (BUL22, BUL23) did not contain any chloroplast DNA, indicating that they had not fed on phytoplankton prior to sampling. However, these specimens were sampled in November (off Bodega Head) during the period of relaxation in upwelling, from vertically integrated net tows and may have been obtained from resident populations as deep as 150 m. In contrast, the three individuals in which chloroplast 16S rRNA sequences were present (BUL34, BUL36 and BUL37) were sampled from shallow water nears in July/August (off Santa Catalina Island) towards the end of the weak summer upwelling period. These differences in OTU composition therefore could be as a result of location, depth or seasonal differences in available diet. The three *G. bulloides* with chloroplast sequences (6.4, 27 and 3 %, respectively) were clearly feeding on a range of photosynthesising eukaryotes (Sect. 3.3). OTUs indicate these to be Cercozoa (mixotrophic protists), Streptophyta (which includes green algae), Haptophyta (which includes coccolithophores) and Stramenopiles (which includes both diatoms and chrysophyte algae).

Our data suggest that *G. bulloides* may also utilise bacteria as a significant food source. *G. bulloides* contained 33.7–62.5 % of non-*Synechococcus* bacterial sequences within the cell (BUL37 was an outlier with only 10 %, Fig. 4), corresponding to a diverse assemblage of 200 OTUs. We assume that these sequences are derived from prey species because no intact bacteria lacking the carboxysomes and thylakoid membranes found in *Synechococcus* were observed in TEM images of the *G. bulloides* cytoplasm or non-digestive vacuoles. The most abundant group of sequences recovered (15–31 %, outlier BUL37 contained 2.6 %) comprise 81 OTUs belonging to the class Alphaproteobacteria (including those OTUs of the family Bradyrhizobiaceae that were not excluded as contaminants), perhaps indicating a preferential selection of specific members within this class. The remaining 17–47.5 % (outlier BUL37, 7.5 %) of sequences were made up of a diverse collection representing other major phyla of bacteria (Sect. 3.3; Fig. 4), including the Acidobacteria, Actinobacteria, Bacteroidetes, Firmicutes and Plantae-mycetes and the classes Beta- and Gammaproteobacteria of the phylum Proteobacteria.

*G. bulloides* type IId is found throughout the year in the Southern California Bight, where it is exposed to cool upwelling periods of high productivity and also to warmer periods characterised by more stratified, less productive conditions (Darling et al., 2003; Darling and Wade, 2008). It has a relatively high growth rate, possibly reproducing within 2–3 weeks (Spero and Lea, 1996; Lombard et al., 2009). In combination with our data, this suggests that *G. bulloides* type IId is a generalist predator with an opportunistic feeding strategy, utilising bacterioplankton as well as phyto- and zooplankton, the proportions of which may be seasonal and depth-dependent. Such opportunism may enable *G. bulloides* to grow and reproduce rapidly within its diverse habitat. We propose two hypotheses for the life strategy of *G. bulloides* type IId to survive the challenges presented within the broad seasonal changes in the region. The first is that it is a mixotrophic feeder (Mitra et al., 2016) and that the *Synechococcus* endobionts are photosynthesising symbionts contributing fixed carbon to the foraminiferal host. In this scenario, they would be fulfilling a functional role similar to that of algal symbionts in other spinoze species, and in particular could provide additional resources during the maximum growth phase of the shallow-dwelling juveniles. Alternatively, or concurrently, *Synechococcus* may be exploited by *G. bulloides* type IId for its nutrient assimilation and storage capacity and then digested as an extra energy, nitrogen and phosphate source for DNA replication at reproduction.

### 4.4 The importance of genotype ecology

Since *G. bulloides* occurs in great abundance in cool, high latitudes and mid- to lower-latitude upwelling systems (Kleijne et al., 1989; Naidu and Malmgren, 1996), it is one of the most commonly used planktic foraminifera for palaeoclimatic reconstruction (Sautter and Thunell, 1991; Spero and Lea 1996). In order to reconstruct past changes in oceanic conditions using the shell geochemical data, it is important to obtain a thorough understanding of the relationship
between foraminiferal ecology and the geochemistry of its shell. This relationship is based on the assumption that each foraminiferal morphospecies represents a genetically continuous species with a unique habitat preference. However, since *G. bulloides* inhabits such a wide range of different ecosystems, it is not surprising that several ecologically distinct genotypes have been recognised (Darling et al., 1999; Kucera and Darling, 2002; Darling and Wade, 2008; Seears et al., 2012; Morard et al., 2013). Indeed, recent species delineation studies support species status for several of the *G. bulloides* genotypes (André et al., 2014), including *G. bulloides* type IId. Such diversity could result in genotype-specific geochemical signatures across the morphospecies (Healy-Williams, 1985; Bijma et al., 1998; de Vargas, 2001; Kucera and Kennet, 2002; Sadekov et al., 2016). Both Kucera and Darling (2002) and Morard et al. (2013) have demonstrated that, based on ecological knowledge, integrating *G. bulloides* genotypes into assemblage-based SST reconstructions significantly improves resolution. This demonstrates the value in understanding the ecology of genotypes within a morphospecies and the necessity of establishing whether the association between *G. bulloides* type IId and *Synechococcus* is universal across this foraminiferal morphospecies complex.

### 4.5 Implications for palaeoceanography

The discovery of intracellular bacteria within a palaeoceanographically significant foraminiferal host may lead to a significant improvement in our current understanding of foraminiferal shell geochemistry. The carbon isotopic composition of planktonic foraminifera has the potential to help reconstruct changes in the chemocline of the surface ocean, providing insights into changes in ocean circulation (Spero et al., 2003). However, the interpretation of $\delta^{13}C$ data is often complicated by poor understanding of the causes of offsets between shell $\delta^{13}C$ and the $\delta^{13}C$ of the dissolved inorganic carbon from which foraminifera build their shells. In particular, the $\delta^{13}C$ of *G. bulloides* shells deviates from predicted values more than that of any other extant species (Deuser et al., 1981; Kahn and Williams, 1981; Curry and Matthews, 1981; Kroon and Darling, 1995; Spero and Lea, 1996; Bijma et al., 1999), implying consistent use of metabolic carbon during calcification by this morphospecies (Deuser et al., 1981).

Symbiont photosynthesis as well as symbiont and host respiration alters the chemical microenvironment surrounding the host shell, which in turn influences their shell geochemical signatures (Rink et al., 1998; Wolf-Gladrow et al., 1999; Eggins et al., 2004). In this *G. bulloides/Synechococcus* association, respiration of both endobiont and host would contribute $^{12}C$-depleted CO$_2$ to the calcifying microenvironment (Spero and Lea, 1996), whilst *Synechococcus* photosynthesis would counteract this by preferentially removing $^{12}CO_2$ and hence elevating $^{13}C / ^{12}C$ ratios in the remaining dissolved CO$_2$, as occurs in protist algal symbiont-bearing planktonic foraminifera (Spero et al., 1997). The large offset towards $^{13}C$-depleted values measured in *G. bulloides* suggests that *Synechococcus* respiration dominates the shell geochemical signature, and implies that photosynthesis is not the primary role of *Synechococcus* in this association (Sect. 4.2).

The presence of metabolically active *Synechococcus* in *G. bulloides* type IId therefore may account for the unusual shell $\delta^{13}C$ determined via culture-based studies conducted at Santa Catalina Island. Since *G. bulloides* type IId is abundant here, it is unlikely that variation in genotype has contributed to uncertainties in these calibrations, but applying these culture-based calibrations to other regions in which different genotypes dominate may produce erroneous results (Darling et al., 2003). It is therefore of particular importance to determine whether the *Synechococcus/G. bulloides* association exists in other *G. bulloides* genotypes in order to generate and apply genotype-specific palaeoclimate calibrations.

### 5 Conclusions

This is the first report of bacterial endobionts within a planktonic foraminiferal species. Our results show that the picocyanobacteria *Synechococcus* is found in large numbers within the protist algal symbiont-barren foraminifer, *G. bulloides*. *Synechococcus* is taken up from the water column by the host and lives and divides within the host cytoplasm at substantially higher concentrations (~4 orders of magnitude) than those found in the surrounding seawater. Its role is not yet known, but its potential for both phototrophy and (photo)heterotrophy makes *Synechococcus* an ideal symbiont for *G. bulloides* as it occupies water depths both within and below the photic zone. Additionally, the ability of *Synechococcus* to store P as polyphosphate, and N within biliproteins under nitrogen-replete conditions, would be beneficial for a foraminiferal host exhibiting fast reproductive turnover, with a high nutrient and energy demand at gametogenesis. Further experiments are required on the *G. bulloides* type IId/*Synechococcus* association to elucidate the full relationship between the two organisms. More investigations are also needed of the *G. bulloides* morphospecies globally in order to determine how widespread the association is to improve understanding and accuracy of this species as a palaeoclimate proxy.

In this study we have demonstrated that 16S rRNA gene metabarcoding of the intracellular DNA of planktonic foraminifera and TEM has the potential to provide new insights into the biological associations and seasonal feeding preferences of ecologically distinct genotypes of planktonic foraminifera. With the addition of 18S rRNA gene metabarcoding to target protist and multicellular zooplankton, next-generation DNA sequencing technologies could transform the usefulness and accuracy of planktonic foraminiferal global distribution and seasonality models by providing the
essential ecological information currently unavailable (Fraile et al., 2008; Lombard et al., 2011; Roy et al., 2015).

6 Data availability


The Supplement related to this article is available online at doi:10.5194/bg-14-901-2017-supplement.

Author contributions. Clare Bird conceived and devised the project and methods, carried out the lab work and prepared the manuscript. Kate F. Darling and Bryne T. Ngwenya advised on methodology. Ann D. Russell, Catherine V. Davis and Jennifer Fehrenbacher collected planktonic foraminifera. Andrew Free contributed to next-generation sequencing and analysis. Michael Wyman contributed to analysis of Synechococcus DNA. Clare Bird and Kate F. Darling wrote the manuscript with contributions from all co-authors.

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