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More, smaller bacteria in response to ocean's warming?

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Heterotrophic bacteria play a major role in organic matter cycling in the ocean. Although the high abundances and relatively fast growth rates of coastal surface bacterioplankton make them suitable sentinels of global change, past analyses have largely overlooked this functional group. Here, time series analysis of a decade of monthly observations in temperate Atlantic coastal waters revealed strong seasonal patterns in the abundance, size and biomass of the ubiquitous flow-cytometric groups of low (LNA) and high nucleic acid (HNA) content bacteria. Over this relatively short period, we also found that bacterioplankton cells were significantly smaller, a trend that is consistent with the hypothesized temperature-driven decrease in body size. Although decadal cell shrinking was observed for both groups, it was only LNA cells that were strongly coherent, with ecological theories linking temperature, abundance and individual size on both the seasonal and interannual scale. We explain this finding because, relative to their HNA counterparts, marine LNA bacteria are less diverse, dominated by members of the SAR11 clade. Temperature manipulation experiments in 2012 confirmed a direct effect of warming on bacterial size. Concurrent with rising temperatures in spring, significant decadal trends of increasing standing stocks (3% per year) accompanied by decreasing mean cell size (–1% per year) suggest a major shift in community structure, with a larger contribution of LNA bacteria to total biomass. The increasing prevalence of these typically oligotrophic taxa may severely impact marine food webs and carbon fluxes by an overall decrease in the efficiency of the biological pump.

1. Introduction

Climate change is significantly affecting the oceans. Either directly or indirectly [1], the effects of warmer temperatures on marine biota are multiple, but most reports either tackle poleward displacement of lower-latitude species [2] or changes in physiological properties resulting in ecosystem rearrangements [3–5]. Recently, changes in various components of marine food webs, from phytoplankton to mammals [6–8], have been documented, but few reports to date [4] have included the smallest life forms, in spite of their overwhelming importance for standing stocks and biogeochemical cycles [9]. Microbial long-term observations [10] are strongly needed to complement data previously available only for larger groups (phytoplankton and zooplankton).

Heterotrophic prokaryotes dominate the ocean's living biomass [11], mostly comprising Bacteria rather than Archaea in upper layers [12]. In flow cytometric analyses, bacterioplankton cluster into two groups of cells with different nucleic acid content [13,14]. These low (LNA) and high nucleic acid (HNA) groups match bimodal distributions of bacterial genome size [15]. Following universal size–genome relationships [16], HNA bacteria are generally bigger than their LNA counterparts [13,17]. Their ecological significance is still debated, but the emerging consensus is that they represent different lineages [18,19].

Biomass is the combination of abundance and individual size. Although relationships between abundance, body size and temperature are complex [20], two general ecological principles apply: (i) higher abundance is associated with smaller size [21], referred to hereinafter as the abundance–size rule (ASR); and (ii) higher ambient temperature results in smaller individuals according to the temperature–size rule (TSR) [22], little studied in unicellular organisms [23]. While the underlying mechanism for the ASR is clear (constant biomass at a given level of resources implies that if there are more organisms these should be smaller), the causes for the TSR are more elusive, with several alternative hypotheses (e.g. [24]). We recently assessed the validity of these two rules to explain changes in phytoplankton size-structure using data collected across the North Atlantic [25]. Similar to that study, in which changes in the overall phytoplankton size community composition rather than intraspecific changes were addressed, we provide here one of the first attempts to detect shifts in the composition and size of planktonic heterotrophic bacteria. Although claims have been made that body size will universally decrease as a consequence of climate change [26,27], studies targeting the smallest life forms are lacking. Here, we explored the seasonal and interannual patterns of LNA and HNA bacteria, focusing on their temperature responses through the ASR and TSR, in a 10-year oceanographic dataset from the southern Bay of Biscay continental shelf, in order to shed light on future directions of change of microbial plankton.

2. Material and methods

(a) Environmental variables

The study site on the Southern Bay of Biscay continental shelf (43.67° N, 5.58° W, bottom depth 110 m) off Xixón, Spain, has been monitored monthly since 2001 as part of a time series programme. Physico-chemical and environmental characterization is detailed in [17]. The depth of the upper mixed layer was determined as that where more than or equal to 0.05 kg m⁻³ density increase over 5 m was first observed. A stratification index was calculated as the per-metre difference in temperature between the surface and 75 m.

(b) Bacterioplankton abundance, cell size and biomass

Bacterioplankton samples were taken at 10–25 m intervals from April 2002 to March 2012, and analysed with a FACSCalibur (BD) flow cytometer as described in detail in [17]. Briefly, LNA and HNA cells were distinguished after nucleic acid staining in green fluorescence versus right-angle light scatter (RALS) cytograms (electronic supplementary material, figure S1a). Bacterial abundance (cells ml⁻¹) was estimated after daily calibrating the flow rate. When present, the high natural fluorescence of *Prochlorococcus* due to photosynthetic pigments prevented overlap with the HNA cluster in red versus green fluorescence

cytograms (electronic supplementary material, figure S1b). Cell size (μm³) was obtained with an empirical calibration between cell diameter and mean RALS, because of its higher sensitivity [28] compared with forward angle light scatter, assuming spherical shape [17]. This assumption may have introduced biases especially in rods or curved rods such as most SAR11 cells, abundant in our samples (see below). Cell size was converted into biomass using [29]: pg C cell⁻¹ = 0.12 × cell size^{0.72}. LNA and HNA bacterial biomass (μg C l⁻¹) was fundamentally driven by changes in abundance.

(c) Quantification of SAR11 phylotype in environmental samples

The contribution of the SAR11 clade to total abundance was assessed by catalysed reporter deposition fluorescence *in situ* hybridization (CARDFISH). For CARDFISH analysis, 4.5 ml samples were collected monthly in 2012, fixed with 3.7% formaldehyde for 3 h, filtered onto 0.2 μm pore-size polycarbonate filters and frozen until analysis. Hybridization was performed as described in [30] using the probe SAR11-441R targeting the SAR11 cluster [31]. Counterstaining of CARDFISH preparations was done with 4,6-diamidino-2-phenylindole (DAPI) at 50 μg ml⁻¹. Cells were counted with a Leica DM 5500 B epifluorescence microscope and pictures were taken with a Leica DFC 360FX monochromatic camera. The abundance and size of SAR11-positive cells were calculated using ACMETool2 image analysis software [32] and the algorithm by Massana *et al.* [33], which yielded length and width of each cell subsequently used for estimating biovolume.

(d) Time series analysis

Time series analysis was conducted with bacterial and environmental properties averaged for the upper mixed layer, with minima usually observed in summer (15 ± 1 m s.e.) and maxima in winter (58 ± 8 m). We used an additive decomposition time series model [34] to detect seasonal and long-term components. The variability of the time series (X) was expressed as

$$X = \bar{x} + bt + a + \sum A_i \cos \left[\left(\frac{2\pi t}{T_i} \right) + \theta_i \right] + \varphi_{it} + \varepsilon_t,$$

where the time series was represented by the climatological mean (\bar{x}), the linear trend with the slope (b) and intercept (a) of the linear regression with time (t), the periodic components amplitude (A_i), period (T_i) and phase (θ_i), the autocorrelation coefficients between consecutive values (φ_{it}) and the unexplained residuals (ε_t). We used the Fisher G-test for assessing the significance of periodic components. Correlation analysis between variables was subsequently made with the pre-whitened residuals (i.e. once we had filtered out the seasonal and long-term components and adjusted for autocorrelation). Decadal trends were also computed with annual averages.

(e) Temperature–size experiments

In 2012, we performed 12 monthly incubations with surface samples from the same site aimed at determining the bacterial response to temperature, both in the presence of the whole microbial community and with bacteria only, after pre-filtering the sample through 0.8 μm pore-size filters. Triplicate bottles (2 l) were incubated at three temperatures (*in situ*, -3°C and +3°C), and samples for estimating LNA and HNA abundance and cellular properties were taken twice per day for 5–10 days. Although seasonally very similar, we will include here only the results of the filtered incubations to exclude trophic interactions with other components. Healthy bacterial populations were invariably found, which started to grow shortly after confinement. HNA cells grew always faster than their LNA counterparts. The slopes of

model I linear regressions of mean cell size versus experimental temperature were used as estimates of the monthly responses of LNA and HNA bacterial sizes to warming ($\mu\text{m}^3\text{ }^\circ\text{C}^{-1}$).

3. Results

Overall, bacterial abundance ranged from 1.2 to 31×10^5 cells ml^{-1} , with HNA cells being slightly more abundant (mean $54 \pm 1\%$ s.e.). The vertical decrease in abundance was more marked from 40 m downwards (electronic supplementary material, figure S2a). Mean bacterial cell size ranged from 0.032 to $0.115 \mu\text{m}^3$ (electronic supplementary material, figure S2b), with HNA cells significantly higher than LNA (0.056 and $0.050 \mu\text{m}^3$, respectively, paired t -test, $p < 0.001$, $n = 114$). Mean cell sizes reached a minimum at 40 m and then increased slightly down to the seafloor. Larger sizes were significantly correlated with higher nucleic acid content (electronic supplementary material, figure S3).

(a) Seasonal patterns

Temperature displayed a marked seasonality (figure 1a), with the 12-month periodic component explaining 80% of total variance (table 1). HNA bacteria had no persistent cyclical components, whereas LNA bacteria showed a statistically significant annual cycle explaining 25% of total variance (figure 1b and table 1). The spring and autumn peaks in total bacterial abundance (approx. 10^6 cells ml^{-1} ; figure 1b) and biomass (approx. $16 \mu\text{g C l}^{-1}$; figure 1d) were characterized by a different relative contribution of both groups, resulting in a strong seasonality of the percentage contribution of LNA cells to bacterial biomass (figure 1e). Earlier work had already shown a very strong seasonal signal of the percentage of HNA cell abundance, with approximately 40% minima in summer and maxima more than 80% around April [17], which is definitely confirmed with this larger dataset. However, little was known about the temporal variability in HNA and LNA bacterial cell size, and no prior attempt had been made at relating their seasonal and long-term patterns to temperature. Bacterial cell size had significant seasonal components for LNA and total bacteria (figure 1c and table 1), with maxima and minima lagged by roughly one month because of the marked summer peak in HNA cell size. For LNA bacteria, seasonal patterns of abundance and size were roughly opposite, and pooled LNA cell abundance and size were negatively correlated ($r = -0.33$, $p < 0.001$, $n = 114$). LNA cell size was also negatively correlated with temperature ($r = -0.19$, $p = 0.044$, $n = 114$). Other variables concurrently measured and potentially relevant for bacteria include total chlorophyll (size-fractionated also since 2003), inorganic nutrient concentrations and stratification index. Briefly, marked stratification from June to October was accompanied by strong nutrient limitation, resulting in low chlorophyll and picophytoplankton dominance. Chlorophyll usually peaked around March–May, with greater contributions of the larger size-fractions. The variance decomposition of these ancillary variables is shown in electronic supplementary material, table S1.

(b) Comparison between SAR11 and flow-cytometric groups

Both the abundance and individual size of SAR11 bacteria covaried positively with flow-cytometric LNA cell values but not with those of HNA cells (electronic supplementary material,

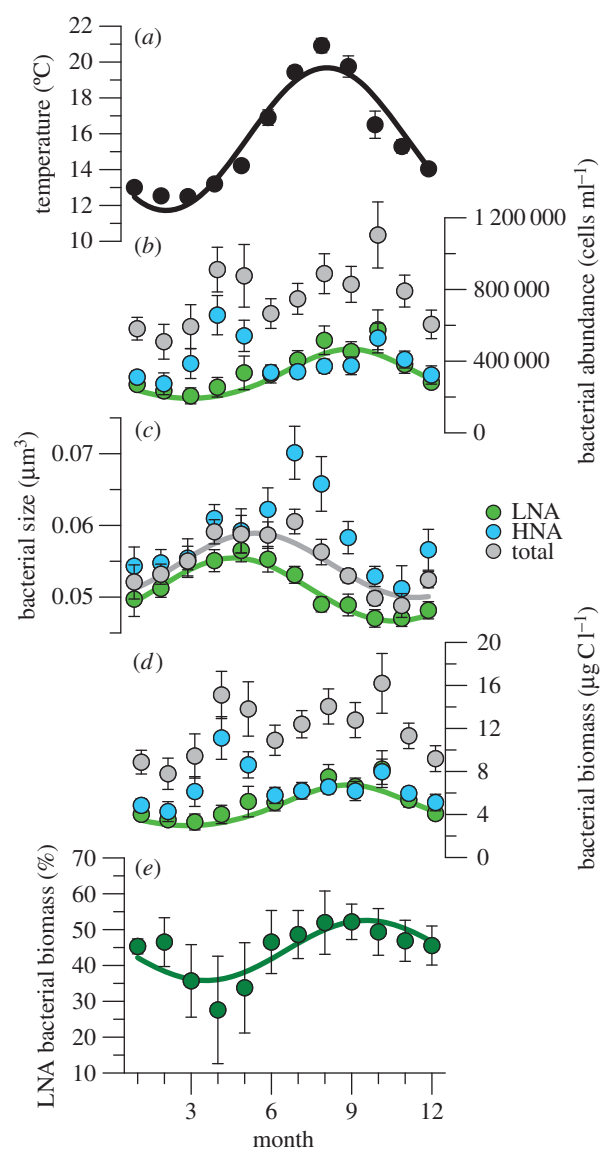


Figure 1. Seasonal variations of temperature and bacterioplankton. Monthly mean \pm s.e. values of (a) temperature, and (b–d) total, LNA and HNA (b) bacterial abundance, (c) bacterial size, (d) bacterial biomass and (e) per cent contribution of LNA bacteria to total biomass in the upper mixed layer of the study site for the April 2002–March 2012 period. Fitted curves represent statistically significant seasonality detailed in table 1.

figure S4a–d), strongly suggesting that LNA cells were mainly composed of SAR11 bacteria. If all SAR11 cells were in the LNA cluster, as current knowledge supports (see Discussion), their mean annual contribution to LNA bacterial abundance would be $74\% \pm 16\%$. With regard to size, although strongly correlated (electronic supplementary material, figure S4c), the mean 28% lower size of LNA compared with that of SAR11 cells should not be attributed, in our opinion, to the presence of smaller bacteria, but to the different methods used (flow cytometry versus microscopic image analysis).

(c) Long-term trends

In addition to the seasonal patterns, we also identified conspicuous decadal trends (table 1). Figure 2 shows them as linear regressions of annual mean values (electronic supplementary material, table S2). In two cases (total bacterial abundance and LNA bacterial biomass), annual mean values increased significantly with time although the variance decomposition of the whole time series failed to find linear trends. The increase in annual mean temperatures was not significant, partly

Table 1. Variance decomposition of the upper mixed layer bacterial times series (April 2002–March 2012) at the study site for total, LNA and HNA cells, and the percentage contribution of LNA cells to total biomass (%LNA biomass). Abundance (cells ml⁻¹), size (μm³) and biomass (μg C l⁻¹) variables were log₁₀ transformed. Only significant ($p < 0.05$) variance components are shown, indicating the fraction of total variance accounted for (%var). Slope (b), intercept (a), period in months (T), amplitude (A), phase in radians (θ), time when maximum value occurs in months (T_{\max}), autocorrelation coefficient (Φ). Lag in months.

variable	mean	linear trend			periodic components					autocorrelation			total %var
		b	a	%var	T	A	θ	T_{\max}	%var	lag	Φ	%var	
temperature	15.7	—	—	—	12	3.89	4.26	8.13	80.43	1	0.34	2.28	82.71
total abundance	5.83	—	—	—	—	—	—	—	—	1	0.20	4.32	4.32
LNA abundance	5.47	0.016	-32.50	2.63	12	0.19	4.69	8.95	25.50	—	—	—	28.13
HNA abundance	5.55	—	—	—	—	—	—	—	—	—	—	—	—
total size	-1.27	-0.005	10.53	8.73	12	0.04	2.92	5.58	26.31	—	—	—	35.04
LNA size	-1.29	-0.004	7.25	4.19	12	0.04	2.44	4.67	31.84	—	—	—	36.05
HNA size	-1.24	-0.007	13.17	7.07	—	—	—	—	—	—	—	—	7.07
total biomass	1.02	—	—	—	—	—	—	—	—	1	0.21	4.77	4.77
LNA biomass	0.65	—	—	—	12	0.18	4.56	8.72	23.01	—	—	—	23.01
HNA biomass	0.76	—	—	—	—	—	—	—	—	—	—	—	—
%LNA biomass	44.2	0.59	-1179	2.58	12	8.40	5.00	9.54	31.32	—	—	—	33.90

because of the large weight of outliers in low-frequency monitoring. Examining monthly temperature changes (data not shown), the only consistent period of warming extended from April through July, with October and November presenting cooling trends. Once in September and twice in October, coastal upwelling caused notable cooling. Consequently, we considered also mean temperatures for the April–July period, coincident with the steepest yearly increase (figure 1a). A significant decadal warming of approximately 1.5°C became apparent for April–July mean temperatures (figure 2a), while mean values for the rest of the year showed a non-significant decrease. The large differences between the annual and spring–early summer temperatures in 2007 and 2011 (figure 2a) were largely due to the above-mentioned strong upwelling events, decreasing almost 4°C the expected monthly values. The abundance of LNA cells increased significantly during the decade (figure 2b), adding approximately 3% to the variance explained by seasonality (table 1). Although HNA cells did not show persistent cyclical components, the negative decadal trend in HNA cell size explained a larger percentage of variance than declining LNA cell size, also reflected in steeper slopes (table 1; electronic supplementary material, table S2). Total bacterioplankton cell size decreased consistently from 2002 to 2012 (equivalent to approx. 1% shrinking per year) as a combined consequence of the sustained decrease in the size of both groups (figure 2c) and the positive trend in the abundance of the smaller LNA bacteria, which also increased significantly in biomass when annual values were considered (figure 2d; electronic supplementary material, table S2). The mean contribution of LNA cells to total bacterial biomass significantly increased from 40 to 47% between 2002 and 2012 (figure 2e and table 1; electronic supplementary material, table S2).

(d) Residual analysis

The correlations between pre-whitened residuals of bacterial properties and selected environmental variables are shown in table 2. Temperature residuals were significantly correlated with both LNA and HNA cell size residuals, but

negatively in the former and positively in the latter. Although the relationship between temperature and the contribution of LNA bacteria to total biomass was not significant, a significant correlation was found with LNA cells contribution to total numbers ($r = 0.21$, $p = 0.021$, $n = 120$). The residuals of the contribution of LNA bacteria to total biomass were also positively correlated with those of stratification index and nitrate concentrations, and negatively with total chlorophyll. The latter correlation became more negative with the absolute and relative concentrations, of chlorophyll in the microplankton size class. Total and size-fractionated chlorophyll residuals were also variably associated with the residuals of LNA and HNA cells, summarized by a negative effect of total chlorophyll on HNA cell size (and positive on HNA abundance) and a positive effect on LNA cell size, more marked with large than small phytoplankton.

(e) Temperature–size experiments

The 6°C gradient in experimental temperature generally resulted in smaller bacterial cell sizes, both for LNA (eight out of 12 experiments) and HNA cells (nine experiments), as indicated by negative slopes in the cell size versus temperature linear regressions. The respective monthly cell size versus temperature changes were significantly correlated for both groups (figure 3a), with an average 2.2-fold higher decrease in HNA than in LNA cells. Both groups showed highly coherent cell size decreases with temperature from March to September (data not shown), particularly marked for spring–early summer ($n = 4$), with mean values of -0.00041 and $-0.00091 \mu\text{m}^3\text{°C}^{-1}$ in LNA and HNA bacteria, respectively. Figure 3b shows that mean values of bacterial cell size and temperature for the April–July period of the time series were also significantly and negatively correlated.

4. Discussion

The decadal time series analysis of the LNA and HNA bacterial groups indicates that seasonal changes in the abundance

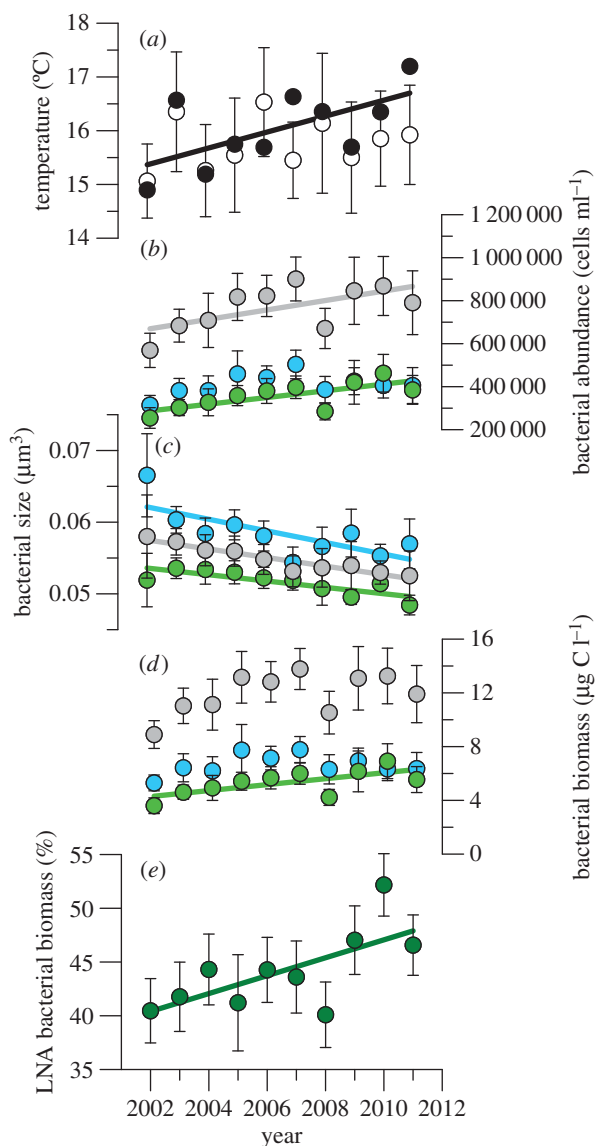


Figure 2. Long-term trends of temperature and bacterioplankton. Annual (April–March) mean \pm s.e. values of (a) temperature, and (b–d) total, LNA and HNA (b) bacterial abundance, (c) bacterial size, (d) bacterial biomass and (e) percentage contribution of LNA bacteria to total biomass in the upper mixed layer of the study site. Filled symbols in (a) represent average temperatures for the April–July period. Error bars for this period are not shown but were on average 48% higher than the annual mean s.e. values. Fitted continuous lines represent significant trends detailed in electronic supplementary material, table S1.

and cell size of the LNA group were coherent with both the TSR and the ASR (i.e. more abundant and smaller LNA bacteria were found during the warmest months). Although the ASR is based on energetic equilibrium, which probably does not hold in a seasonally varying environment, the analysis of the entire LNA bacteria dataset is consistent with the general ASR prediction that the higher the abundance, the smaller the size of an organism [21]. Based on the good agreement between the size of LNA and SAR11 bacteria, the latter derived from the more precise image analysis of DAPI-stained samples (electronic supplementary material, figure S4c), our assumption of spherical form in flow cytometry data probably had a minor effect on biovolume estimates, at least regarding the relative differences between groups and over time that were the object of this study.

We explain the coherent behaviour of the LNA cells by their lower diversity (electronic supplementary material, figure S5), probably dominated by a single alphaproteobacterial clade, SAR11 (electronic supplementary material, figure S4), as previously found [18,31,35]. Genome sizes of cultured SAR11 strains would place them neatly in the LNA rather than the HNA cluster [36,37]. In a large cross-Atlantic survey, LNA populations were invariably dominated by SAR11, while HNA cells were phylogenetically diverse, including members of Bacteroidetes, Gammaproteobacteria and other Alphaproteobacteria lineages [18]. Numerous ecotypes with different metabolic functions and temperature dependences are grouped under the SAR11 lineage [38,39], which could help explain the slightly higher presence of SAR11 in winter in the Pacific [40], contrary to common observations of summer abundance maxima [38,41] (this study). At our site, two SAR11 ecotypes were found among the top 10 most abundant bacterial OTUs (plus eight more in the top 100) in a 3.5-year survey [42]. Their biogeochemical roles and the existence of differing long-term trends exceed the scope of this study, but are part of ongoing efforts. On the contrary, the higher diversity (electronic supplementary material, figure S5) and marked species succession within HNA cells [18,19] may be related to the absence of seasonality and of significant correlation between abundance and size. Lack of bacterial time series through sufficiently extended periods precludes assessing the generality of our observations. Yet there is evidence of seasonally recurring patterns in phylogenetic community composition in temperate systems [43,44], including our own site [42]. These cyclical changes can also be discerned in the much coarser flow cytometric classification (figure 1).

While 10 years were too short to find sustained warming of the upper mixed layer, in recent decades, the oceans have warmed up at an unprecedented pace (e.g. [45]). Extended analysis (electronic supplementary material, figure S6) and longer records just 50 km eastwards show that the southern Bay of Biscay is no exception, with an approximate $0.05^{\circ}\text{C yr}^{-1}$ increase [46] comparable with other latitudes. This increase has not been homogeneous and our results indicate that significant spring–early summer warming (figure 4a) was accompanied by non-significant cooling for the rest of the year. Seasonally uneven warming is widespread, with reports for our study area consistently agreeing on the preponderance of higher summer temperatures over winter values, including the whole northwestern European continental shelf [47], the Iberian Peninsula [48] and the above-mentioned nearby site [49]. Based on monthly snapshots rather than continuous measurements, decadal trends were more compelling for bacterial characteristics than for temperature, probably due to the fact that bacterioplankton integrate environmental forcing over periods longer than days. Nevertheless, the inverse relationship found between spring–early summer mean temperatures and bacterial sizes (figure 3b) might be linked to the period of the year in which larger decreases in LNA and HNA cell volume with experimental temperature were observed (figure 3a).

We are not aware of previous studies concurrently assessing temporal trends of marine bacterioplankton abundance and cell size. The few reported time series of heterotrophic bacteria are mostly restricted to coastal temperate waters, for which consistent interannual decreases [50] and increases in abundance [51] were associated with parallel changes in phytoplankton standing stocks. Recent increases in heterotrophic bacteria and picophytoplankton in polar environments were

Table 2. Pearson correlation coefficients between the pre-whitened residuals of heterotrophic bacterioplankton and selected environmental variables. SI, stratification index; NO_3 , nitrate concentration; Chl, total chlorophyll; pChl, chlorophyll smaller than $2\ \mu\text{m}$; nChl, chlorophyll $2\text{--}20\ \mu\text{m}$; mChl, chlorophyll larger than $20\ \mu\text{m}$; %pChl, %nChl and %mChl, percentage contribution to Chl of pChl, nChl and mChl, respectively. $n = 114$.

	total abund	LNA abund	HNA abund	total size	LNA size	HNA size	total biomass	LNA biomass	HNA biomass	%LNA biomass
temperature	—	—	—	—	-0.19^*	0.26^{**}	—	—	—	—
SI	—	—	-0.23^{**}	0.19^*	—	0.34^{***}	—	—	—	0.23^*
NO_3	-0.23^*	—	-0.31^{**}	—	—	—	-0.22^*	—	-0.30^{**}	0.31^{**}
Chl	—	—	0.18^*	—	0.19^*	-0.25^{**}	—	—	—	-0.19^*
pChl	—	—	—	—	—	—	—	—	—	—
nChl	0.22^*	—	—	—	—	—	0.23^*	—	0.18^*	—
mChl	—	—	0.25^{**}	—	0.27^{**}	—	—	—	0.23^*	-0.27^{**}
%pChl	—	—	—	-0.19^*	-0.23^*	—	-0.19^*	—	—	—
%nChl	—	—	—	—	—	—	—	—	—	0.22^*
%mChl	—	—	0.19^*	—	0.20^*	—	—	—	—	-0.27^{**}

*** $p < 0.001$; ** $p < 0.01$, * $p < 0.05$.

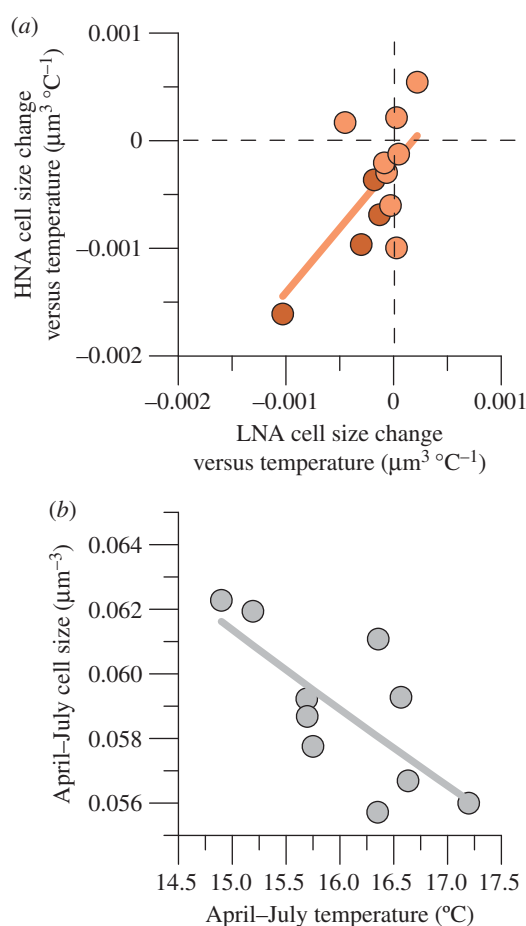


Figure 3. (a) Comparison between cell size change versus temperature (ccc-temp) for LNA and HNA bacteria in experimental incubations with surface samples taken in 2012. Darker symbols represent results from April through July. Fitted line: $\text{HNA}_{\text{ccc-temp}} = -0.0002 + 1.19 \text{ LNA}_{\text{ccc-temp}}$, $r^2 = 0.41$, $p = 0.024$, $n = 12$. Dashed lines represent no change (0 slope) of the linear regression, more details given in the text. (b) Relationship between mean values of bacterial cell size and temperature for the period extending from April through July of the 10 years of available data. Fitted line: $\text{April–July cell size} = 0.098 - 0.024 \text{ April–July temperature}$, $r^2 = 0.52$, $p = 0.017$, $n = 10$.

explained by enhanced sea ice melting and decreased nutrient supply [52], which can still lead to higher primary production, largely channelled through dissolved organic compounds usable by bacteria [53]. Availability of resources for bacterial uptake could be an alternative hypothesis to temperature explaining the observed trends. Using phytoplankton as a proxy for resource availability, high values should result in high bacterial abundances [54]. This could explain the spring peaks in HNA bacterial abundance (figure 1b), positively correlated with chlorophyll ($r = 0.18$, $p = 0.048$, $n = 114$; see also table 2). However, no correlation was found with LNA cells, and neither total chlorophyll nor any size-fraction showed decadal trends (electronic supplementary material, table S1). Recent work concurs that LNA cells are spatio-temporally independent of phytoplankton, at least at short scales [38,55], although a study in the Pacific has found positive correlations of SAR11 with DOC and primary production [40]. LNA/SAR11 cells could also have benefitted from environmental conditions related to high temperature, such as low inorganic nutrients content and high light, foreseen to expand in the future ocean [1,56]. Although environmental variables other than temperature also showed seasonal and long-term trends (electronic supplementary material, table S1), the only ones that changed intra- and interannually coherently with bacterial properties were inorganic nutrient concentrations. However, while the seasonal patterns of NO_3 and PO_4 and LNA cell abundance were roughly opposite (cf. table 1; electronic supplementary material, table S1), both nutrients increased rather than decreased with time, but this increment did not translate into larger phytoplankton biomass.

Because of the covariance and linear trends of most variables, correlation analysis of residual variation (table 2) is more suitable to search for mechanistic explanations. The preferential association of HNA cells with phytoplankton (large rather than small) was confirmed by residual analysis. Although nitrate concentration residuals showed correlations of opposite sign with HNA cell abundance and %LNA biomass, it is difficult to see nitrate availability as the direct driver when phosphate residuals were not correlated (data not shown) and without concomitant changes in

phytoplankton. The negative correlation between temperature and LNA cell size residuals confirms the compliance with the TSR of LNA bacteria, which also prevailed in abundance in warmer waters. Their positive association also with the water-column thermal gradient (table 2) suggests that indirect effects of global warming such as enhanced stratification [46,47,57] might exacerbate the future prevalence of LNA cells. Altogether, the match between seasonal and decadal trends and residual analysis identify temperature as the candidate for the dominance of increasingly smaller LNA bacteria in these waters. Finer-scale variations within the group (e.g. changes in the relative contribution of SAR11 ecotypes with differing temperature ranges [38,39]) may be behind the observed patterns.

Smaller phytoplankton and zooplankton in response to rising temperatures have been demonstrated both experimentally and through sustained observations [4,25,58,59], but as far as we know this is the first study documenting a systematic long-term decline in bacterioplankton size, thus supporting the view of decreasing body size as a universal ecological response to warming [26,27] alongside changes in phenology [60] and distributional ranges [2]. The decadal trends of declining total, LNA and HNA bacterial sizes (figure 2c) are highly relevant, particularly considering that only LNA cells consistently followed the ASR and TSR on the two temporal scales (seasonal and interannual) compared here. Seasonality of HNA cells, bimodal for abundance and unimodal for size (figure 1b,c), did not preclude a decadal compliance with the TSR. While the seasonal patterns of LNA and HNA bacterial size differed markedly, probably linked in the latter case to the recurrent appearance of large species in spring and summer [42], both groups decreased consistently in size as temperature rose. In the 10 years covered by this study, surface warming was better detected in the period extending from April through July than in the annual average (figure 2a).

Bearing in mind that a gradual ocean warming of 0.5°C per decade cannot be directly comparable with a quasi-instantaneous 3°C increase in an incubation, the hypothesis that temperature is the ultimate cause for the marked decrease in bacterial mean size (approx. 1% yr⁻¹) is further strengthened by the experimental results with natural samples in which temperature was the only factor manipulated (figure 3a). Interestingly, the size of HNA cells tended to decrease more markedly than LNA cells both interannually (figure 2c; electronic supplementary material, table S2) and experimentally (figure 3a). Applying the 2012 mean spring–early summer decrease of LNA and HNA cell size per °C (figure 3a, darker symbols) to the 2002–2011 April–July warming (figure 2a) would result in an overall size decrease of 3%. This value is smaller than the observed 10% (figure 2c), possibly related to the hugely different temporal scales compared (days versus years). Negative associations between temperature and bacterial size have been documented in large spatial

surveys [61] as well as experimentally through reductions in genome size [62], usually strongly correlated with cell volume [16] (electronic supplementary material, figure S3).

Atlantic cross-ocean analysis [25] hints at a rapid increase in the foreseen contribution of cyanobacteria to planktonic communities [63]. Residual variation analysis (table 2) suggests that the future prevalence of small phytoplankton may also enhance LNA bacterial contribution to total stocks. The implications of a future dominance of small autotrophic and heterotrophic bacteria for oceanic food webs and biogeochemistry are multiple. The size structure of planktonic communities affects ecosystem functions such as predator–prey interactions, efficiency of energy transfer across trophic levels and ultimately carbon export [5,64]. Gradual replacement of bacteria by typically oligotrophic taxa such as the SAR11 clade and others included within the LNA cluster [14,18,35] will be likely to alter carbon fluxes and trophic relationships as these bacteria grow slowly [65] and, contrary to HNA cells, appear not to respond simultaneously to changes in phytoplankton primary production [55]. We document here a rapid change in bacterioplankton community composition that probably follows fundamental ecological and metabolic rules. This study adds to evidence from recent and geological times [4,59] to strongly support that, regardless of the trophic level and biological organization, a shift towards lower sizes can be a widespread response of marine organisms to global change. Additional studies are needed to test these associations between warming, bacterial abundance and cell size, but only longer oceanographic time series will confirm the generality of shrinking marine bacteria. The consequences for ecosystem functioning are far-reaching, but can be summarized as a lower transfer efficiency of primary production towards higher trophic levels together with a decrease in the importance of the biological pump for efficient carbon sequestration in the ocean's interior.

Data accessibility. All data are deposited in Dryad (manuscript Dryad ID: <http://dx.doi.org/10.5061/dryad.kh7nt>).

Authors' contributions. L.D.-P., A.C.-D. and X.A.G.M. collected the samples and performed flow-cytometric analysis. X.A.G.M. and E.N. performed time series analysis. N.A.-G. and L.A.-S. performed CARDFISH and image analyses. H.W.D. and N.G. were involved in the study design. X.A.G.M., E.N., A.L.-U. and L.A.-S. designed the study. X.A.G.M. wrote the paper. All authors discussed the results and commented on the manuscript.

Competing interests. We declare we have no competing interests.

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References

1. Sarmiento JL *et al.* 2004 Response of ocean ecosystems to climate warming. *Glob. Biogeochem. Cycles* **18**, GB3003. (doi:10.1029/2003GB002134)
2. Chen IC, Hill JK, Ohlemüller R, Roy DB, Thomas CD. 2011 Rapid range shifts of species associated with high levels of climate warming. *Science* **333**, 1024–1026. (doi:10.1126/science.1206432)
3. López-Urrutia Á, San Martín E, Harris RP, Irigoien X. 2006 Scaling the metabolic balance of the oceans. *Proc. Natl Acad. Sci. USA* **103**, 8739–8744. (doi:10.1073/pnas.0601137103)

4. Daufresne M, Lengfellner K, Sommer U. 2009 Global warming benefits the small in aquatic ecosystems. *Proc. Natl Acad. Sci. USA* **106**, 12 788–12 793. (doi:10.1073/pnas.0902080106)
5. Wohlers J, Engel A, Zöllner E, Breithaupt P, Jürgens K, Hoppe HG, Sommer U, Riebesell U. 2009 Changes in biogenic carbon flow in response to sea surface warming. *Proc. Natl Acad. Sci. USA* **106**, 7067–7072. (doi:10.1073/pnas.0812743106)
6. Behrenfeld MJ *et al.* 2006 Climate-driven trends in contemporary ocean productivity. *Nature* **444**, 752–755. (doi:10.1038/nature05317)
7. Richardson AJ, Schoeman DS. 2004 Climate impact on plankton ecosystems in the Northeast Atlantic. *Science* **305**, 1609–1612. (doi:10.1126/science.1100958)
8. Montes-Hugo M, Doney SC, Ducklow HW, Fraser W, Martinson D, Stammerjohn SE, Schofield O. 2009 Recent changes in phytoplankton communities associated with rapid regional climate change along the western Antarctic Peninsula. *Science* **323**, 1470–1473. (doi:10.1126/science.1164533)
9. Azam F. 1998 Microbial control of oceanic carbon flux: the plot thickens. *Science* **280**, 694–696. (doi:10.1126/science.280.5364.694)
10. Ducklow HW, Doney SC, Steinberg DK. 2009 Contributions of long-term research and time-series observations to marine ecology and biogeochemistry. *Annu. Rev. Mar. Sci.* **1**, 279–302. (doi:10.1146/annurev.marine.010908.163801)
11. Whitman WB, Coleman DC, Wiebe WJ. 1998 Prokaryotes: the unseen majority. *Proc. Natl Acad. Sci. USA* **95**, 6578–6583. (doi:10.1073/pnas.95.12.6578)
12. Karner MB, DeLong EF, Karl DM. 2001 Archaeal dominance in the mesopelagic zone of the Pacific Ocean. *Nature* **409**, 507–510. (doi:10.1038/35054051)
13. Bouvier T, del Giorgio PA, Gasol JM. 2007 A comparative study of the cytometric characteristics of high and low nucleic-acid bacterioplankton cells from different aquatic ecosystems. *Environ. Microbiol.* **9**, 2050–2066. (doi:10.1111/j.1462-2920.2007.01321.x)
14. Gasol JM, Zweifel UL, Peters F, Fuhrman JA, Hagström Å. 1999 Significance of size and nucleic acid content heterogeneity as measured by flow cytometry in natural planktonic bacteria. *Appl. Environ. Microbiol.* **65**, 4475–4483.
15. Koonin EV, Wolf YI. 2008 Genomics of bacteria and archaea: the emerging dynamic view of the prokaryotic world. *Nucleic Acids Res.* **36**, 6688–6719. (doi:10.1093/Nar/Gkn668)
16. Hessen DO, Daufresne M, Leinaas HP. 2013 Temperature–size relations from the cellular-genomic perspective. *Biol. Rev.* **88**, 476–489. (doi:10.1111/Brv.12006)
17. Calvo-Díaz A, Morán XAG. 2006 Seasonal dynamics of picoplankton in shelf waters of the southern Bay of Biscay. *Aquat. Microb. Ecol.* **42**, 159–174. (doi:10.3354/ame042159)
18. Schattnerhofer M, Wulf J, Kostadinov I, Glockner FO, Zubkov MV, Fuchs BM. 2011 Phylogenetic characterisation of picoplanktonic populations with high and low nucleic acid content in the North Atlantic Ocean. *Syst. Appl. Microbiol.* **34**, 470–475. (doi:10.1016/j.syapm.2011.01.008)
19. Vila-Costa M, Gasol JM, Sharma S, Moran MA. 2012 Community analysis of high- and low-nucleic acid-containing bacteria in NW Mediterranean coastal waters using 16S rDNA pyrosequencing. *Environ. Microbiol.* **14**, 1390–1402. (doi:10.1111/j.1462-2920.2012.02720.x)
20. Kingsolver JG, Huey RB. 2008 Size, temperature, and fitness: three rules. *Evol. Ecol. Res.* **10**, 251–268.
21. White EP, Ernest SKM, Kerkhoff AJ, Enquist BJ. 2007 Relationships between body size and abundance in ecology. *Trends Ecol. Evol.* **22**, 323–330. (doi:10.1016/j.tree.2007.03.007)
22. Atkinson D. 1994 Temperature and organism size – a biological law for ectotherms. *Adv. Ecol. Res.* **25**, 1–58. (doi:10.1016/S0065-2504(08)60212-3)
23. Forster J, Hirst AG, Esteban GF. 2013 Achieving temperature-size changes in a unicellular organism. *ISME J.* **7**, 28–36. (doi:10.1038/ismej.2012.76)
24. Walters RJ, Hassall M. 2006 The temperature–size rule in ectotherms: may a general explanation exist after all? *Am. Nat.* **167**, 510–523. (doi:10.1086/501029)
25. Morán XAG, López-Urrutia A, Calvo-Díaz A, Li WKW. 2010 Increasing importance of small phytoplankton in a warmer ocean. *Glob. Change Biol.* **16**, 1137–1144. (doi:10.1111/j.1365-2486.2009.01960.x)
26. Sheridan JA, Bickford D. 2011 Shrinking body size as an ecological response to climate change. *Nat. Clim. Change* **1**, 401–406. (doi:10.1038/Nclimate1259)
27. Gardner JL, Peters A, Kearney MR, Joseph L, Heinsohn R. 2011 Declining body size: a third universal response to warming? *Trends Ecol. Evol.* **26**, 285–291. (doi:10.1016/j.tree.2011.03.005)
28. Felip M, Andreaa S, Sommaruga R, Straskrbova V, Catalan J. 2007 Suitability of flow cytometry for estimating bacterial biovolume in natural plankton samples: comparison with microscopy data. *Appl. Environ. Microbiol.* **73**, 4508–4514. (doi:10.1128/AEM.00733-07)
29. Norland S. 1993 The relationship between biomass and volume of bacteria. In *Handbook of methods in aquatic microbial ecology* (eds PF Kemp, BF Sherr, EB Sherr, JJ Cole), pp. 303–307. Boca Raton, FL: Lewis Publishers.
30. Pernthaler A, Pernthaler J, Amann R. 2004 Sensitive multicolor fluorescence in situ hybridization for the identification of environmental microorganisms. In *Molecular microbial ecology manual*, vol. 3, 2nd edn (eds GA Kowalchuk, FJ de Bruijn, IM Head, AD Akkermans, JD van Elsas), pp. 711–726. Dordrecht, The Netherlands: Kluwer Academic Publishers.
31. Morris RM, Rappe MS, Connon SA, Vergin KL, Siebold WA, Carlson CA, Giovannoni SJ. 2002 SAR11 clade dominates ocean surface bacterioplankton communities. *Nature* **420**, 806–810. (doi:10.1038/nature01240)
32. Zeder M, Ellrott A, Amann R. 2011 Automated sample area definition for high-throughput microscopy. *Cytometry Part A* **79A**, 306–310. (doi:10.1002/Cyto.A.21034)
33. Massana R, Gasol JM, Bjørnsen PK, Blackburn N, Hagström Å, Hietanen S, Hygum BH, Kuparinen J, Pedrós-Alió C. 1997 Measurement of bacterial size via image analysis of epifluorescence preparations: description of an inexpensive system and solutions to some of the most common problems. *Sci. Mar.* **61**, 397–407.
34. Bode A, Anadón R, Morán XAG, Nogueira E, Teira E, Varela M. 2011 Decadal variability in chlorophyll and primary production off NW Spain. *Clim. Res.* **48**, 293–305. (doi:10.3354/Cr00935)
35. Mary I, Heywood JL, Fuchs BM, Amann R, Tarran GA, Burkill PH, Zubkov MV. 2006 SAR11 dominance among metabolically active low nucleic acid bacterioplankton in surface waters along an Atlantic meridional transect. *Aquat. Microb. Ecol.* **45**, 107–113. (doi:10.3354/ame045107)
36. Giovannoni SJ *et al.* 2005 Genome streamlining in a cosmopolitan oceanic bacterium. *Science* **309**, 1242–1245. (doi:10.1126/science.1114057)
37. Grote J, Thrash JC, Huggett MJ, Landry ZC, Carini P, Giovannoni SJ, Rappe MS. 2012 Streamlining and core genome conservation among highly divergent members of the SAR11 Clade. *mBio* **3**, e00252-12. (doi:10.1128/mBio.00252-12)
38. Carlson CA, Morris R, Parsons R, Treusch AH, Giovannoni SJ, Vergin K. 2009 Seasonal dynamics of SAR11 populations in the euphotic and mesopelagic zones of the northwestern Sargasso Sea. *ISME J.* **3**, 283–295. (doi:10.1038/ismej.2008.117)
39. Brown MV *et al.* 2012 Global biogeography of SAR11 marine bacteria. *Mol. Syst. Biol.* **8**, 595. (doi:10.1038/msb.2012.28)
40. Eiler A, Hayakawa DH, Church MJ, Karl DM, Rappé MS. 2009 Dynamics of the SAR11 bacterioplankton lineage in relation to environmental conditions in the oligotrophic North Pacific subtropical gyre. *Environ. Microbiol.* **11**, 2291–2300. (doi:10.1111/j.1462-2920.2009.01954.x)
41. Morris RM, Vergin KL, Cho JC, Rappé MS, Carlson CA, Giovannoni SJ. 2005 Temporal and spatial response of bacterioplankton lineages to annual convective overturn at the Bermuda Atlantic time-series study site. *Limnol. Oceanogr.* **50**, 1687–1696. (doi:10.4319/lo.2005.50.5.1687)
42. Alonso-Sáez L, Díaz-Pérez L, Morán XAG. In press. The hidden seasonality of the rare biosphere in coastal marine bacterioplankton. *Environ. Microbiol.* (doi:10.1111/1462-2920.12801)
43. Fuhrman JA, Hewson I, Schwalbach MS, Steele JA, Brown MV, Naeem S. 2006 Annually reoccurring bacterial communities are predictable from ocean conditions. *Proc. Natl Acad. Sci. USA* **103**, 13 104–13 109. (doi:10.1073/pnas.0602399103)
44. Gilbert JA *et al.* 2012 Defining seasonal marine microbial community dynamics. *ISME J.* **6**, 298–308. (doi:10.1038/ismej.2011.107)
45. Hansen J, Sato M, Ruedy R, Lo K, Lea DW, Medina-Elizade M. 2006 Global temperature change. *Proc. Natl Acad. Sci. USA* **103**, 14 288–14 293. (doi:10.1073/pnas.0606291103)

46. Llope M, Anadon R, Viesca L, Quevedo M, Gonzalez-Quiros R, Stenseth NC. 2006 Hydrography of the southern Bay of Biscay shelf-break region: integrating the multiscale physical variability over the period 1993–2003. *J. Geophys. Res.* **111**, C09021. (doi:10.1029/2005jc002963)
47. Holt J *et al.* 2012 Multi-decadal variability and trends in the temperature of the northwest European continental shelf: a model-data synthesis. *Prog. Oceanogr.* **106**, 96–117. (doi:10.1016/j.pocean.2012.08.001)
48. El Kenawy A, Lopez-Moreno JI, Vicente-Serrano SM. 2012 Trend and variability of surface air temperature in northeastern Spain (1920–2006): linkage to atmospheric circulation. *Atmos. Res.* **106**, 159–180. (doi:10.1016/j.atmosres.2011.12.006)
49. González-Gil R, González Taboada E, Höfer J, Anadón R. 2015 Winter mixing and coastal upwelling drive long-term changes in zooplankton in the Bay of Biscay (1993–2010). *J. Plankton Res.* **37**, 337–351. (doi:10.1093/plankt/fbv001)
50. Sarmento H, Montoya JM, Vazquez-Dominguez E, Vaque D, Gasol JM. 2010 Warming effects on marine microbial food web processes: how far can we go when it comes to predictions? *Phil. Trans. R. Soc. B* **365**, 2137–2149. (doi:10.1098/rstb.2010.0045)
51. Verity PG, Borkman DG. 2010 A decade of change in the Skidaway River Estuary. III. Plankton. *Estuaries Coasts* **33**, 513–540. (doi:10.1007/s12237-009-9208-2)
52. Li WKW, McLaughlin FA, Lovejoy C, Carmack EC. 2009 Smallest algae thrive as the Arctic Ocean freshens. *Science* **326**, 539. (doi:10.1126/science.1179798)
53. Kirchman DL, Morán XAG, Ducklow H. 2009 Microbial growth in the polar oceans: role of temperature and potential impact of climate change. *Nat. Rev. Microbiol.* **7**, 451–459. (doi:10.1038/nrmicro2115)
54. Gasol JM, Duarte CM. 2000 Comparative analyses in aquatic microbial ecology: how far do they go? *FEMS Microbiol. Ecol.* **31**, 99–106. (doi:10.1111/j.1574-6941.2000.tb00675.x)
55. Morán XAG, Ducklow HW, Erickson M. 2011 Single-cell physiological structure and growth rates of heterotrophic bacteria in a temperate estuary (Waquoit Bay, Massachusetts). *Limnol. Oceanogr.* **2011**, 37–48. (doi:10.4319/lo.2011.56.1.0037)
56. Doney SC. 2006 Oceanography: plankton in a warmer world. *Nature* **444**, 695–696. (doi:10.1038/444695a)
57. Capotondi A, Alexander MA, Bond NA, Curchitser EN, Scott JD. 2012 Enhanced upper ocean stratification with climate change in the CMIP3 models. *J. Geophys. Res.* **117**, C04031. (doi:10.1029/2011jc007409)
58. Finkel ZV, Katz ME, Wright JD, Schofield OME, Falkowski PG. 2005 Climatically driven macroevolutionary patterns in the size of marine diatoms over the cenozoic. *Proc. Natl Acad. Sci. USA* **102**, 8927–8932. (doi:10.1073/pnas.0409907102)
59. Falkowski PG, Oliver MJ. 2007 Mix and match: how climate selects phytoplankton. *Nat. Rev. Microbiol.* **5**, 813–819. (doi:10.1038/nrmicro1751)
60. Parmesan C, Yohe G. 2003 A globally coherent fingerprint of climate change impacts across natural systems. *Nature* **421**, 37–42. (doi:10.1038/Nature01286)
61. Straza TRA, Cottrell MT, Ducklow HW, Kirchman DL. 2009 Geographic and phylogenetic variation in bacterial biovolume as revealed by protein and nucleic acid staining. *Appl. Environ. Microbiol.* **75**, 4028–4034. (doi:10.1128/Aem.00183-09)
62. Sabath N, Ferrada E, Barve A, Wagner A. 2013 Growth temperature and genome size in bacteria are negatively correlated, suggesting genomic streamlining during thermal adaptation. *Genome Biol. Evol.* **5**, 966–977. (doi:10.1093/Gbe/Evt050)
63. Flombaum P *et al.* 2013 Present and future global distributions of the marine Cyanobacteria *Prochlorococcus* and *Synechococcus*. *Proc. Natl Acad. Sci. USA* **110**, 9824–9829. (doi:10.1073/pnas.1307701110)
64. Dickman EM, Newell JM, Gonzalez MJ, Vanni MJ. 2008 Light, nutrients, and food-chain length constrain planktonic energy transfer efficiency across multiple trophic levels. *Proc. Natl Acad. Sci. USA* **105**, 18 408–18 412. (doi:10.1073/pnas.0805566105)
65. Teira E, Martinez-Garcia S, Lonborg C, Alvarez-Salgado XA. 2009 Growth rates of different phylogenetic bacterioplankton groups in a coastal upwelling system. *Environ. Microbiol. Rep.* **1**, 545–554. (doi:10.1111/j.1758-2229.2009.00079.x)