Disentangling the mechanisms shaping the surface ocean microbiota

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Institut de Ciencies del Mar

Ramon Massana
Institut de Ciencies del Mar

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Abstract
Background The ocean microbiota modulates global biogeochemical cycles and changes in its configuration may have largescale consequences. Yet, the underlaying ecological mechanisms structuring it are unclear. Here we investigate how fundamental ecological mechanisms (selection, dispersal and ecological drift) shape the smallest members of the tropical and subtropical surface-ocean microbiota: prokaryotes and minute eukaryotes (picoeukaryotes). Furthermore, we investigate the agents exerting abiotic selection on this assemblage as well as the spatial patterns emerging from the action of ecological mechanisms. To explore the previous, we analysed the composition of surface-ocean prokaryotic and picoeukaryotic communities using DNA-sequence data (16S- and 18S-rRNA genes) collected during the circumglobal expeditions Malaspina-2010 and TARA-Oceans.

Results We found that the two main components of the tropical and subtropical surface-ocean microbiota, prokaryotes and picoeukaryotes, appear to be structured by different ecological mechanisms. Picoeukaryotic communities were predominantly structured by dispersal-limitation, while prokaryotic counterparts appeared to be shaped by the combined action of dispersal-limitation, selection and drift. Temperature-driven selection appeared as a major factor influencing species co-occurrence networks in prokaryotes but not in picoeukaryotes, indicating that association patterns may contribute to understand ocean microbiota structure and response to selection. Other measured abiotic variables seemed to have limited selective effects on community structure in the tropical and subtropical ocean. Picoeukaryotes displayed a higher spatial differentiation between communities and a higher distance decay when compared to prokaryotes, consistent with a scenario of higher dispersal limitation in the former after considering environmental heterogeneity. Lastly, random dynamics or drift seemed to have a more important role in structuring prokaryotic communities than picoeukaryotic counterparts.

Conclusions The differential action of ecological mechanisms seems to cause contrasting biogeography, in the tropical and subtropical ocean, among the smallest surface plankton, prokaryotes and picoeukaryotes. This suggests that the idiosyncrasy of the main constituents of the ocean microbiota should be considered in order to understand its current and future configuration,
which is especially relevant in a context of global change, where the reaction of surface ocean
plankton to temperature increase is still unclear.

Background
The surface ocean microbiota is a pivotal underpinning of global biogeochemical cycles [1, 2]. The
smallest ocean microbes, the picoplankton, have a key role in the global carbon cycle, being
responsible for an important fraction of the total atmospheric carbon and nitrogen fixation in the
ocean [3-5], which supports ~46% of the global primary productivity [6]. Oceanic picoplankton plays
a fundamental role in processing organic matter by recycling nutrients and carbon to support
additional production as well as by channelling organic carbon to upper trophic levels through food
webs [5, 7, 8]. The ocean picoplankton includes prokaryotes (both bacteria and archaea) and tiny
unicellular eukaryotes (hereafter picoeukaryotes), which feature fundamental differences in terms of
cellular structure, feeding habits, metabolic diversity, growth rates and behaviour [9]. Even though
marine picoeukaryotes and prokaryotes are usually investigated separately, they are intimately
connected through biogeochemical and food web networks [10-12].
The underlying ecological mechanisms determining the biogeography of prokaryotes and
picoeukaryotes in the global ocean are unclear. In particular, we do not know whether these crucial
components of the ocean microbiota are structured by the action of the same or different ecological
processes. Comprehending such processes is fundamental, as their differential action can produce
changes in the ocean microbiota composition that could impact global ecosystem function [13-15]. A
recent ecological synthesis explains the structure of communities and the emergence of
biogeography as a consequence of the action of four main processes: selection, dispersal, ecological
drift and speciation [16]. Selection involves deterministic reproductive differences among individuals
from different or the same species as a response to biotic or abiotic conditions. Selection can act in
two opposite directions, it can constrain (homogeneous selection) or promote (heterogeneous
selection) the divergence of communities [17]. Dispersal is the movement of organisms across space,
and rates can be high (homogenising dispersal), moderate, or low (dispersal limitation) [17].
Dispersal limitation occurs when species are absent from suitable habitats because potential
colonizers are too far away [18], and the significance of dispersal limitation increases as geographic scale increases [19]. Ecological drift (hereafter drift) in a local community refers to random changes in species’ relative abundances derived from stochastic birth, death, offspring production, immigration and emigration [16]. The action of drift in a metacommunity, that is, local communities that are connected via dispersal of multiple species [20], may lead to neutral dynamics [19], where random dispersal is the main mechanism of community assembly. Finally, speciation is the evolution of new species [16], and it will not be considered hereafter as it is expected to have a small impact in the turnover of communities that are connected via dispersal [21], being also difficult to measure this ecological process in the wild.

The action of the previous ecological processes is typically manifested as different taxonomic or phylogenetic patterns of community turnover, that is, -diversity. At the moment, there are several estimators of -diversity which capture different aspects of community turnover [22]. Most of these indices consider taxonomic or phylogenetic aspects of communities, but not species-association patterns, which can also manifest the action of ecological processes. For example, selection exerted by an environmental variable can drive species co-occurrences generating groups of highly associated species or modules in association networks that correspond with specific environmental conditions [23]. Different members of these modules may be more abundant in specific regions of the ocean, contributing to increase -diversity estimates between these regions when based on standard compositional or phylogenetic -diversity metrics. Yet, -diversity estimates based on association-aware metrics may point to higher similarity between these regions, as taxa belong to the same modules. Furthermore, modules may display correlations with environmental heterogeneity. Thus, association aware metrics of -diversity may allow unveiling community patterns and their relationships with environmental variables (i.e. selection), which would be missed by standard approaches [24]. So far, most studies investigating the structure of the ocean microbiota have not considered species associations in their analyses of -diversity.

The differential action of selection, dispersal and drift may generate different microbial assemblages that could feature diverse metabolisms and ecologies [14, 15]. Moderate or high selection together
with moderate dispersal rates may couple environmental heterogeneity with combinations of species, leading to a spatial pattern known as species sorting [25]. In contrast, high or low levels of dispersal may decouple environmental heterogeneity (i.e. selection) from the composition of species assemblages. High dispersal rates may maintain populations in habitats to which they are maladapted [14, 20]. Inversely, low dispersal rates may promote microbial assemblages that become more different as the geographic distance between them increases (distance decay). If environmental heterogeneity and geographic distance covary, then distance decay could reflect both selection and dispersal limitation [26]. Drift is expected to cause important random effects in local community composition in cases where selection is weak and populations are small [13, 27]. Here we investigate the mechanisms that shape the surface-ocean microbiota by using DNA-sequence data collected in two of the largest circumglobal oceanographic expeditions to date, Malaspina 2010 [28] and TARA Oceans [29]. Specifically, we ask: What is the relative importance of selection, dispersal and drift in structuring the surface ocean microbiota? Do these processes act similarly on main components of this microbiota (prokaryotes and picoeukaryotes)? What are the main agents that exert abiotic selection? Do species association networks reflect the action of selection in the upper ocean microbiota? What are the main spatial-structure patterns that emerge due to the action of selection, dispersal and drift?

Results
Quantifying the mechanisms that structure the surface ocean picoplankton

We analysed 16S and 18S rRNA-genes from prokaryotes and picoeukaryotes in 120 globally-distributed tropical and subtropical stations sampled during the Malaspina 2010 expedition [28] (Figure 1A). Note that the nature of the TARA Oceans data precluded using them in these analyses (see Methods). Operational Taxonomic Units were delineated at 99% similarity (OTUs-99%) and as unique sequence variants (OTUs-ASVs, the maximum resolution for the 18S and 16S rRNA-gene). Analyses using both, OTUs-99% and OTUs-ASVs indicated that dispersal limitation was the dominant factor structuring picoeukaryotic communities, explaining 76-67% of community turnover, while this process had a lower importance in prokaryotes (~35-25%) [Figure 1B]. Note that percentage refers to
the percentage of pairs of communities that appear to be driven by dispersal limitation. In contrast, homogenizing dispersal had a very limited role in the structuring of the tropical and subtropical upper-ocean microbiota (<3% for both picoeukaryotes and prokaryotes). Drift had a limited role in the structuring of picoeukaryotic communities as indicated by both OTUs\_99 and OTUs\_ASVs, representing ~21–6% of community turnover (Figure 1B). In contrast, drift appeared as a relevant factor structuring prokaryotic communities, explaining ~44–31% of the community turnover according to OTUs\_99 and OTUs\_ASVs (Figure 1B). The role of selection was higher in prokaryotes compared to picoeukaryotes according to both OTUs\_99 and OTUs\_ASVs, explaining ~34–27% of the turnover of prokaryotic communities, and ~17–11% of that in picoeukaryotes (Figure 1B). Heterogeneous selection had a relatively higher importance in structuring picoeukaryotes as compared to prokaryotes (~16–7% vs. ~9–4%, respectively). Instead, homogeneous selection appeared more important in structuring prokaryotic (~24–23%) than picoeukaryotic (~1–4%) communities (Figure 1B).

Our quantifications indicated different roles of ecological processes in structuring communities of marine prokaryotes and picoeukaryotes populating the tropical and subtropical surface-ocean (Figure 1B). We then aimed at confirming these results using other more traditional approaches. In these analyses, considering Malaspina data, we used OTUs\_99%, given that these likely correspond to well-defined lineages, while OTUs\_ASVs may reflect, in some cases, intraspecific variation [30]. We found moderate correlations between picoeukaryotic and prokaryotic diversity (Bray Curtis: = 0.58, gUniFrac: = 0.61, p = 0.01, Mantel tests; Figure S2, Additional file 2). Given that rare species tend to occupy less sites than more abundant ones [31], communities featuring different proportions of abundant or rare species may display different spatial turnover. We found that picoeukaryotes had proportionally more regionally rare (i.e. mean abundances across all samples <0.001%) species than prokaryotes (71% vs. 48% respectively) [Table S1, Additional file 3; Figure S3, Additional file 4]. This is consistent with the observation that picoeukaryotes had more restricted species distributions (i.e., occurring in <20% of the stations) than prokaryotes (95% vs. 88% of the species respectively) [Table
Selection acting on the microbiota

We investigated the agents exerting abiotic selection on the tropical and subtropical surface-ocean microbiota by analysing -diversity together with the environmental variables included in the Meta-119 dataset (Temperature (°C), Conductivity (S m⁻¹), Fluorescence, Salinity and Dissolved Oxygen (ml L⁻¹)). We used different indices that capture distinct facets of -diversity (Bray-Curtis, TINA_w, PINA_w, gUniFrac; see Methods). Water temperature was the most important driver of selection on prokaryotes (Figure 2), ranging between 15.7 - 29.3 C, with a mean of 24.5 C and a standard deviation of 3.2 C across the whole Meta-119 Malaspina dataset (Figure 1A). Furthermore, water temperature appeared to affect prokaryotic association networks, given that TINA_w [24] explained ~50% of community variance (ADONIS R²) [Figure 2], while other used -diversity indices that do not consider species associations explained considerably lower proportions (Figure 2). In contrast, temperature had limited effects on picoeukaryotic community turnover (Figure 2). Analyses using both the Malaspina and TARA Oceans datasets indicated stronger positive correlations between TINA_w and water-temperature differences in prokaryotes (Mantel r = 0.8-0.5, p<0.01) than in picoeukaryotes [Mantel r = 0.3, p<0.05] (Figure 3). TARA Oceans samples displayed a higher correlation with water temperature than Malaspina samples (Figure 3), which likely reflects the wider temperature range covered by TARA Oceans samples (range ~0-30 C, mean ~21 C, standard deviation ~7 C) compared to Malaspina (range ~15-30 C, mean ~24 C, standard deviation ~3 C). Overall, TINA_w results indicate that locations with similar temperatures include prokaryotic species that tend to co-occur, with this pattern disappearing as the temperature difference between stations increases. The previous pattern was either weak or non-existent in microbial eukaryotes (Figure 3). We expanded the exploration of the role of abiotic selection on microbiota structuring by analysing a larger number of environmental variables (total 17) that were available for only 57 globally distributed Malaspina stations (see details in Supplementary Methods, Additional file 6; Figure S4, Additional file 7). Results supported the importance of temperature-driven selection for prokaryotic
community structuring (Figure S5, Additional file 8) and indicated that fluorescence (a proxy for Chlorophyll a concentration) explained 31% of PINA_w-based prokaryotic community variance (ADONIS $R^2$), being non-significant for picoeukaryotes (Figure S5, Additional file 8). The remaining tested abiotic variables explained a minor fraction of community variance, suggesting that abiotic selection, at the whole ocean-microbiota level, operates via few agents, mainly temperature, although we cannot rule out that other unmeasured abiotic variables may also be exerting selection. The different correlations between temperature and -diversity as measured by TINA_w in prokaryotes and picoeukaryotes suggest that they may feature different species association networks. We found that prokaryotes sampled in both Malaspina and TARA Oceans were more associated between themselves than protists (Figure S6, Additional file 9; Table S3, Additional file 10; Table S4, Additional file 11; Table S5, Additional file 12). Furthermore, the prokaryotic networks were more modular (in terms of cliques) than the picoeukaryotic counterparts (Table S3, Additional file 10), which may reflect to certain extent, temperature-driven selection [23].

Given that selection exerted by variables that lack phylogenetic signal, typically biotic variables, could inflate estimates of dispersal limitation, we have checked whether the high dispersal limitation we estimated for picoeukaryotes could reflect zooplankton grazing. For that, we have analysed globally-distributed surface TARA Oceans stations for which we could estimate both the community composition of picoeukaryotes (here defined as the 0.8–5 µm size-fraction; 36 or 38 stations) as well as that of microzooplankton (20–180 µm size-fraction; 36 stations) or mesozooplankton (180–2,000 µm size-fraction; 38 stations) based on 18S-rRNA genes32]. Analyses considering abiotic (total 6, see Supplementary Methods, Additional file 6) and biotic (estimated zooplankton abundance) variables indicated that micro- and mesozooplankton had a minor influence on picoeukaryotic community structure (~5% of the variance explained, ADONIS $R^2$). In addition, the correlation between picoeukaryotic and zooplankton -diversity was either weak (microzooplankton, $r = 0.34$) or absent (mesozooplankton) [$p<0.01$, Mantel tests]. Thus, zooplankton grazing does not appear to influence -diversity in picoeukaryotes.

Selection acting on single species
The previous analyses investigated how selection may operate on the entire assemblage of species, without considering the different responses to selection that are expected in individual species. We therefore evaluated the potential action of selection on single species by determining their individual correlations with multiple abiotic environmental variables using the Maximum Information Coefficient (MIC). In the Malaspina dataset (Figure 1A), temperature was the variable with the highest number of associated prokaryotic species (1.7%), representing ~17% of the 16S rRNA gene-sequence abundance, while picoeukaryotic species displayed limited associations with temperature (~0.3% of the species representing ~5% of the 18S rRNA gene-sequence abundance) [Figure S7, Additional file 13]. Picoeukaryotic and prokaryotic species were also associated with oxygen, conductivity and salinity (Figure S7, Additional file 13), which co-vary with temperature. The remaining variables displayed limited associations with individual prokaryotic or picoeukaryotic species (Figure S7, Additional file 13), thus agreeing with our previous results suggesting that abiotic selection on the tropical and subtropical surface-ocean microbiota operates via few variables, with a dominant role for temperature among prokaryotes. Overall, prokaryotes featured proportionally more individual-species associations with environmental parameters than picoeukaryotes (Figure S7, Additional file 13), suggesting that environmental heterogeneity in the tropical and subtropical surface-ocean has a stronger effect on prokaryotic assemblages than on picoeukaryotic counterparts. Analyses of TARA Oceans data supported the previous results by indicating that prokaryotic species were associated predominantly with temperature and oxygen in the upper global ocean, while unicellular eukaryotes had weak associations to multiple variables (Table S6, Additional file 14).

Dispersal

Abiotic environmental conditions in adjacent stations over the trajectory of the Malaspina cruise, typically separated by 250–500 km, in the tropical and sub-tropical ocean (Figure 1A) are generally comparable [33]. Therefore, compositional differences between pairs of neighbouring communities could manifest the differential capability of distinct microbial assemblages to disperse. Following these premises, we analysed the change in picoeukaryotic and prokaryotic community composition along the trajectory of the Malaspina cruise by comparing each community to the one sampled
immediately before in a sequential manner (i.e. sequential -diversity) \([\text{Figure 4}].\) Both picoeukaryotic and prokaryotic communities displayed variable amounts of sequential -diversity \((\text{Figure 4, Panels A and B}),\) although picoeukaryotes featured, on average, a higher sequential -diversity than prokaryotes \((\text{Figure 4, Panel C}).\) This agrees with the overall mean -diversity, which was significantly higher for picoeukaryotes than for prokaryotes \((\text{Figure S8, Additional file 15}).\) Tests by subsampling the number of picoeukaryotic OTUs to the same number of prokaryotic ones \((7,025)\) indicated that different numbers of OTUs in these groups did not affect mean Bray-Curtis estimates of -diversity displayed in \textit{Figure S8, Additional file 15} \([34].\)

When geographic distance covary with environmental heterogeneity, spatial community variance may be the manifestation of both selection and/or dispersal limitation. -diversity in picoeukaryotes and prokaryotes displayed positive correlations with geographic distance (i.e. distance decay) predominantly within 1,000 km \((\text{Figure 4, Panel D}).\) Yet, correlations were weaker in prokaryotes than in picoeukaryotes, pointing to stronger dispersal limitation or selection in the latter. Variance partitioning analyses considering both environmental [Temperature (ºC), Conductivity (S m\(^{-1}\)), Fluorescence, Salinity and Dissolved Oxygen (ml L\(^{-1}\))] and geographic variables (ocean basin and subdivisions, as well as Longhurst biogeographic provinces \([35], \text{Figure S1, Additional file 1}\) indicated that in prokaryotes, geographic variables explained most of the variance (24%), while environmental variables explained 10%, and 13% was explained by both variables; 53% of the variance remained unexplained. In contrast, picoeukaryotes displayed non-significant results in the same analyses. Still, after controlling for the effects of the most important environmental variables, Longhurst provinces (but not ocean basins nor subdivisions) accounted for ~20–25% of community variance in both picoeukaryotes and prokaryotes \((\text{ADONIS } R^2) [\text{Figure 2}].\) All in all, the previous analyses seem coherent with our quantifications of ecological processes \((\text{Figure 1B}),\) in the sense that they indicate that both selection and dispersal limitation (represented by geographic variables such as distance or ocean provinces), do seem to have a role in the structuring of the surface ocean picoplankton. Selection and dispersal limitation may operate more strongly in geographic areas that constitute
ecological boundaries, leading to abrupt changes in microbiota composition. We identified 14 communities where sequential -diversity displayed abrupt changes, with 11 of them coinciding for both picoeukaryotes and prokaryotes (Figure 4, Panels A & B). The Local Contributions to Beta Diversity (LCBD) index [36] (Figure S9, Additional file16) indicated that ~22% of both picoeukaryotic and prokaryotic communities (26 stations each, totaling 36 different stations) contributed the most to the -diversity, with 16 communities coinciding for both prokaryotes and picoeukaryotes (Figure S9, Additional file16; Table S7, Additional file 17). In addition, 8 of the 36 stations featuring a significant LCBD were also identified as zones of abrupt community change in sequential -diversity analyses (Table S7, Additional file 17). These zones point to selection or dispersal operating simultaneously and strongly upon both prokaryotic and picoeukaryotic communities in the surface ocean.

Discussion

Applying an innovative ecological framework [21] allowed us to quantify the mechanisms that shape the tropical and subtropical upper-ocean microbiota. Yet, this approach has limitations (summarised by Zhou & Ning [17]) that need to be considered in the context of our results. First, our results represent the overall action of ecological processes at the whole microbiota level, and not their operation on every taxonomic group or lineage (for example, different taxonomic Classes may be structured by different processes). In addition, our results reflect the action of ecological mechanisms at the global ocean level, and we expect that other spatial scales (ocean basin for example) may lead to other results. Furthermore, our results provide a snapshot of the importance of ecological processes at the global-ocean scale, and future studies should investigate how the relative importance of these mechanisms change over time [37]. Second, the measured ecological mechanisms are associated with the evolutionary diversification that is reflected by the variation in the chosen molecular markers. OTUs_{99\%} and OTUs_{ASVs} based on the 16S and 18S rRNA genes likely reflect defined species (or gene flow units [38]) or in some cases population variation [30], and therefore, the measured ecological mechanisms in the tropical and subtropical ocean apply to those evolutionary levels. Hence, our results do not reflect the mechanisms shaping intra-population variation or those shaping taxonomic ranks above the species level. Furthermore, our results indicate
that delineating OTUs based on sequence clustering (OTUs$_{99\%}$) or sequence variants (OTUs$_{ASVs}$) can affect measurements of ecological mechanisms, although in our study, main trends were maintained. It could be hypothesized that OTUs$_{99\%}$ and OTUs$_{ASVs}$ may represent different taxonomic units in prokaryotes or picoeukaryotes, especially if one group was evolving faster than the other. Yet, both prokaryotes and picoeukaryotes show a wide range of evolutionary rates [39, 40], including lineages evolving slow or fast, therefore potential differences in unit definitions associated to different evolutionary rates will likely compensate when analysing complex assemblages of species. Third, failure to detect selection could inflate estimates of dispersal limitation. We consider that our estimates indicating substantial dispersal limitation in picoeukaryotes were not inflated, as picoeukaryotes displayed more restricted spatial distributions than prokaryotes and important biotic variables, such as potential zooplankton grazing, did not seem to affect the structure of picoeukaryotic assemblages. Furthermore, another study also suggests that dispersal limitation influences protist distributions in the global ocean [32]. Altogether, the used framework [21] can be considered as a guide that can provide important insights on the ecological mechanisms structuring the global ocean microbiota, while more data (e.g. single nucleotide variants in genes or genomes) and experiments are necessary to understand such mechanisms in further detail.

Our results indicated that the differential action of ecological processes may promote different biogeographic patterns in prokaryotic and picoeukaryotic assemblages in the upper global-ocean. This is consistent with other works using similar approaches to ours indicating that protistan and bacterial assemblages are shaped by different ecological processes [37, 41–43]. In particular, selection, which is known to have an important role in structuring prokaryotic communities [25, 26], explained a higher proportion of community turnover in surface-ocean prokaryotes (~34–27% of the turnover) than in picoeukaryotes (~17–11%). This modest role of selection in structuring the tropical and subtropical sunlit-ocean microbiota is consistent with the moderate environmental gradients characterizing this habitat. In other habitats featuring a higher selective pressure, the role of selection in structuring prokaryotes is expected to be higher [41]. The quantifications of the importance of
selection are also associated to the global scale of our survey. Thus, for example, at smaller geographic scales, where dispersal limitation is expected to have a lower impact than at global scales [18], the relative importance of selection could increase. Congruently, in surface waters of the East China Sea it was found that selection was 40% more important than dispersal limitation in structuring bacterial communities [42], while in our global study, selection and dispersal limitation had a similar importance in structuring prokaryotes. Furthermore, the previous study [42] found that selection was considerably more important than dispersal limitation in structuring communities of microbial eukaryotes. In contrast, our global assessment yields dispersal limitation to be ~5 times higher than selection in structuring picoeukaryotic communities.

We found that heterogeneous selection was more important in structuring picoeukaryotic than prokaryotic communities, while homogeneous selection was more important in structuring prokaryotic than picoeukaryotic communities. This suggest that prokaryotes and picoeukaryotes respond differently to the same environmental heterogeneity, which in the tropical and subtropical surface-ocean would be preventing community divergence in prokaryotes while promoting it in picoeukaryotes. Different adaptations in prokaryotes and picoeukaryotes [9] may determine such contrasting responses to the same environmental heterogeneity. For example, a given environmental heterogeneity could select for a few species featuring wide environmental tolerance or several species that are adapted to narrow environmental conditions.

Diverse large-scale studies have indicated that temperature is one of the main variables structuring the ocean microbiota [44–49]. Consistent with our results, Sunagawa et al. [50] reported strong correlations between prokaryotic global-ocean microbiota composition and temperature, and weak correlations with nutrients. The previous agrees with our results indicating that temperature is one of the most important agents exerting abiotic selection on the surface-ocean microbiota, although we cannot rule out the selective action of other unmeasured abiotic factors. Furthermore, temperature is one of the main abiotic variables structuring microbial assemblages in seasonal time-series, pointing also to the importance of this variable at local scales over yearly cycles [51–53]. Our analyses also unveiled an additional layer of information by indicating that temperature-driven selection affects
prokaryotic taxa co-occurrences, a pattern not observed in picoeukaryotes. Such -diversity related to species associations is typically not captured by classic compositional indices like Bray Curtis, possibly due to variations in the relative abundance of the co-occurring species [54]. In contrast to prokaryotes, less is known about the effects of temperature on the community structure of ocean picoeukaryotes, which according to our results are modest. Yet, specific picoeukaryotic lineages, such as MAST-4, do seem to be affected by temperature [55], pointing to taxonomic-group specific responses to selection. One of the possible reasons why picoeukaryotes do not show co-occurrence patterns comparable to those observed in prokaryotes is dispersal limitation, which precludes picoeukaryotic species with similar niches to share the same geographic zone. Overall, our work indicates that species association patterns are informative on the -diversity of marine prokaryotes, therefore taxa association networks should be contemplated in future analyses of the ocean microbiota.

To what extent dispersal limitation affects the distribution of ocean microbes is a matter of debate. The impact of dispersal limitation is expected to increase with increasing body size [56], therefore, larger protists are expected to be more limited by dispersal than smaller prokaryotes. Ocean protists seem to follow the previous tenet, as it has been observed that dispersal limitation appears to increase with increasing cell size [32]. Furthermore, in surface open-ocean waters, prokaryotes typically display abundances of $10^6$ cells/mL, while picoeukaryotes normally have abundances of $10^3$ cells/mL [57]. Due to random dispersal alone, the more abundant prokaryotes are expected to be distributed more thoroughly than the less abundant picoeukaryotes [31]. Thus, both cell size and abundance could partially explain our results indicating a higher dispersal limitation in picoeukaryotes than in prokaryotes. Yet, multiple studies of aquatic unicellular eukaryotes point to restricted dispersal [32, 58, 59], while other studies indicate the opposite [55, 60, 61]. This could reflect different dispersal capabilities among unicellular eukaryotes [58, 62] and the generation of dormant cysts in some species [63, 64], which may increase dispersal. Yet, cyst formation has not been reported for picoeukaryotes [9] and this may partially explain their limited dispersal. Regarding prokaryotes, previous studies indicate that dispersal limitation has a modest influence in the structure
of marine communities [50, 65, 66] which is coherent with our results. In particular, Louca et al. [67] indicate that there is virtually no dispersal limitation in surface ocean prokaryotes within specific ocean regions, suggesting that the importance of dispersal limitation may increase across large oceanic regions or basins. Nevertheless, dormancy in prokaryotes seems to be more common than in picoeukaryotes [9, 68], and this may allow the former to disperse more thoroughly by reducing their metabolisms when moving through unfavorable habitats [69].

The importance of drift in structuring microbial communities is unclear [25, 70]. Our results, considering both OTUs\textsubscript{99} and OTUs\textsubscript{ASVs} indicated that drift has a modest role in structuring picoeukaryotic communities in the tropical and subtropical surface ocean. In contrast, both OTUs\textsubscript{99} and OTUs\textsubscript{ASVs} indicated that drift has a more important role in structuring prokaryotic counterparts.

Another study also found a larger importance of drift in determining the community structure of bacteria when compared with phytoplankton populating freshwater and brackish habitats [71]. In contrast, drift was the prevalent community-structuring mechanism in unicellular eukaryotes populating lakes in a relatively small geographic area that features a strong salinity gradient, having a low importance for the structuring of prokaryotic counterparts [41]. Drift is expected to have a more important role in small populations, which is normally not the case for microbes in the global ocean. Yet, other random processes could have effects similar to drift in large microbial populations. For example, the evolution or arrival of a new bacteriophage may attack abundant bacteria in a local community, randomly reshuffling species abundances. Likely, the importance of drift in structuring microbiotas is dependent on taxa adaptations (e.g. the presence of habitat generalists or specialists), selection strength, ecological redundancy, as well as on the dimensions of the analysed habitats.

A decrease in community similarity with geographic distance (distance decay) can be the manifestation of selection and/or dispersal limitation [26]. Distance decay has been evidenced in diverse studies focusing on the surface and deep ocean microbiotas [65, 72, 73]. In our study, we have used different analyses (variance partitioning and ADONIS based on measured environmental variables and geographic features; see Methods) to interpret the measured distance decay. Variance
partitioning suggested that both geography (i.e. dispersal limitation) and environmental variation (selection) likely explain distance decay in prokaryotes, with geography having potentially a more important role, which agrees with our ADONIS analyses based on Bray Curtis and gUnifrac distances (Figure 2). Interestingly, variance partitioning was not significant in picoeukaryotes, although ADONIS analyses based on Bray Curtis and gUnifrac distances indicated that geography, and to a lesser extent temperature, would partially explain picoeukaryotic distance decay (Figure 2). Overall, provincialism, as measured by Longhurst provinces (Figure S1, Additional file 1) was the most relevant spatial feature for the community structuring of both prokaryotes and picoeukaryotes (Figure 2). Possibly, this reflects dispersal limitation, as the selective effects of main environmental variables that covary with these provinces were considered during ADONIS analyses. Longhurst provinces may also reflect, to certain extent, different water masses or currents that impose restrictions to microbial dispersal. Yet, it may be possible that dispersal limitation in picoeukaryotes is only partially reflected by provincialism, thus explaining the lack of significance in variance partitioning analyses as well as the differences between the magnitudes of dispersal limitation suggested by provincialism analyses using ADONIS (Figure 2) and those estimated by analyses of ecological processes (Figure 1B). Alternatively, dispersal limitation in picoeukaryotes may be better reflected by geographic distances between communities, as suggested by sequential Bray-Curtis analyses (Figure 4C) as well as their stronger distance decay when compared to prokaryotes (Figure 4D). Interestingly, a study investigating surface marine bacteria along ~12,000 km in the Atlantic Ocean found that provincialism explained an amount of community variance comparable to our results [65]. Furthermore, and consistent with our results, a study of the sunlit global-ocean eukaryotic microbiota indicated that basin, which may be associated to provincialism and dispersal limitation, was one of the most important variables explaining community turnover [32]. In sum, Longhurst provinces, as a spatial feature, seem to partially reflect the dispersal limitation measured for upper-ocean picoplankton.

In the surface ocean, drastic changes in microbial species composition across space may point to strong changes in abiotic selection (as expected to occur across oceanographic fronts [74, 75]), or
high immigration. We identified 14 stations featuring abrupt changes in prokaryotic or picoeukaryotic community composition as well as 36 stations with a “unique” species composition. Some of these areas correspond to nutrient-rich (selection) coastal zones (the South African Atlantic coast and the South Australia Bight) or potential upwelling (dispersal) zones, such as the Equatorial Pacific and Atlantic as well as the Costa Rica Dome. These findings were coherent with Spatial Abundance Distributions (SpAD) of bacterioplankton in the tropical and subtropical surface-ocean [33]. Altogether, the previous suggests strong selective changes or immigration from deep water layers into the surface associated to upwellings, affecting both prokaryotic and picoeukaryotic community structure. Such immigration events into the surface, when random, may partially explain the measured drift.

Conclusion
Our results indicate that selection, dispersal and drift have different roles in shaping the main components of the picoplankton (prokaryotes and picoeukaryotes) in the tropical and subtropical surface ocean. This highlights the importance of comprehending the characteristics of the different constituents of microbiotas in order to understand their structure. Our results also suggest that the surface ocean picoplankton may not show a single response to global change, and that perhaps prokaryotes will display more pronounced changes in their community structure as a response to temperature increase than picoeukaryotes, considering that temperature seems to affect more prokaryotic than picoeukaryotic assemblages. Future studies on the ocean microbiota should investigate the change in the role of selection, dispersal and drift with ocean scale (from meters to kilometers), depth, latitude and longitude as well as with time, taxonomic ranks (e.g. Class, Family, etc.) and molecular markers that evolve at different rates. Such studies will likely provide a more comprehensive understanding of the underlaying mechanisms shaping the ocean microbiota at different evolutionary levels (from lineages to populations) and will also provide insights on the environmental variables that could modify its current configuration.

Methods
Sample collection
Surface waters (3 m depth) from a total of 120 globally-distributed stations located in the tropical and
sub-tropical ocean (Figure 1A) were sampled as part of the Malaspina 2010 expedition [28]. Sampling took place between December 2010 and July 2011 and the cruise was organized in a way so that most regions were sampled during similar meteorological seasons. Samples were obtained with a 20 L Niskin bottle deployed simultaneously to a CTD profiler that measured conductivity, temperature, oxygen, fluorescence and turbidity for each sample. About 12 L of seawater were sequentially filtered through a 20 µm nylon mesh, followed by a 3 µm and 0.2 µm polycarbonate filters of 47 mm diameter (Isopore, Millipore, Burlington, MA, USA). Only the smallest size-fraction (0.2 – 3 µm, here called “picoplankton” [8]) was used in downstream analyses. Samples for inorganic nutrients (NO$_3^-$, NO$_2^-$, PO$_4^{3-}$, SiO$_2$) were collected from the Niskin bottles and measured spectrophotometrically using an Alliance Evolution II autoanalyzer (Frépillon, France) [76]. Chlorophyll measurements were obtained from Estrada et al. [77]. In specific samples nutrient concentrations were estimated using the World Ocean Database [78] due to issues with the measurements. Since not all environmental parameters were available for all stations, two contextual datasets were generated: Meta–119, including 119 stations, 5 environmental parameters and 5 spatial features (all except one station in Figure 1A) and Meta–57 (Figure S4, Additional file 7), including 57 stations and 17 environmental parameters (the 5 environmental parameters included in Meta–119 were considered here as well). See Supplementary Methods, Additional file 6.

DNA extraction, sequencing and bioinformatics
DNA was extracted using a standard phenol-chloroform protocol [79]. Both the 18S and 16S rRNA-genes were amplified from the same DNA extracts. The hypervariable V4 region of the 18S rRNA gene (~380 bp) was amplified with the primers TAReukFWD1 and TAReukREV3 [80], while the hypervariable V4-V5 (~400bp) region of the 16S rRNA gene was amplified with the primers 515F-Y-926R [81], which target both Bacteria and Archaea. Amplicon libraries were then paired-end sequenced on an Illumina (San Diego, CA, USA) MiSeq platform (2x250bp) at the Research and Testing Laboratory facility (http://www.researchandtesting.com/).

Reads were processed following and in-house protocol [82]. Operational Taxonomic Units (OTUs) were delineated at 99% similarity using UPARSE V8.1.1756 [83], producing 42,505 picoeukaryotic and
10,158 prokaryotic OTUs. Taxonomic assignment of OTUs was generated by BLASTing OTU-representative sequences against different reference databases. BLAST hits were filtered prior to taxonomy assignment using an in-house python script, considering a percentage of identity >90%, a coverage >70%, a minimum alignment length of 200 bp and an e-value > 0.00001. Metazoan, Streptophyta, nucleomorphs, Chloroplast and mitochondrial OTUs were removed from the OTUs tables. See Supplementary Methods, Additional file 6 and Table S8, Additional file 18.

Additionally, to investigate the effects of clustering on the estimation of ecological mechanisms (Fig. 1B), we determined OTUs as Amplicon Sequence Variants (ASVs) using DADA2 [84]. For the 18S, we trimmed the forward reads at 240 bp and the reverse reads at 180 bp, while for the 16S, forward reads were trimmed at 220 bp and reverse reads at 200 bp. Then, for the 18S, the maximum number of expected errors (maxEE) was set to 12 and 20 for the forward and reverse reads respectively, while for the 16S, the maxEE was set to 2 for the forward reads and to 4 for the reverse reads. OTUs were assigned taxonomy using the naïve Bayesian classifier method [85] together with the SILVA version 132 [86] database as implemented in DADA2. Eukaryotic OTUs were also BLASTed [87] against the Protist Ribosomal Reference database (PR², version 4.11.1; [88]). Streptophyta, Metazoa, nucleomorphs, chloroplasts and mitochondria were removed from OTUs tables. Tables of OTUs were rarefied to 20,000 reads per sample with the function rrarefy in Vegan. Only OTUs with abundances >100 reads were used for the calculation of ecological mechanisms (Fig. 1B).

We used publicly-available data from the TARA Oceans global expedition [29] in multiple analyses. This expedition took place between September 2009 - March 2012, and includes samples from the same hemisphere during different meteorological seasons. Due to the nature of the TARA Oceans dataset, we did not perform all the analyses that were run for the Malaspina dataset. Specifically, short V9 18S rRNA-gene reads or 16S rRNA-gene miTags [89] from TARA Oceans precluded robust phylogenetic reconstructions, which instead were possible with the longer reads produced for Malaspina. We used data from TARA Oceans surface (~5 m depth) stations only, including 41 samples (40 stations) for pico-nano eukaryotes (0.22–3 m [1 sample] and 0.8–5 m [40 samples]; 18S-V9 rRNA
gene amplicon data) [32] as well as 63 stations for prokaryotes (picoplankton, 0.22–3 m [45 samples] and 0.22–1.6 m [18 samples]; 16S rRNA genes, miTags) [50].

General analyses and phylogenetic inferences
Tables including OTUs_{99\%} were sub-sampled to 4,060 reads per sample using rrarefy in Vegan [90], resulting in sub-sampled tables containing 18,775 picoeukaryotic and 7,025 prokaryotic OTUs. OTUs_{99\%} with mean relative abundances >0.1% or <0.001% were defined as regionally abundant or rare respectively [91]. Phylogenetic trees were constructed by aligning 16S or 18S OTUs_{99\%} representative sequences against an aligned SILVA [86] template using mothur [92]. Afterwards, poorly aligned regions or sequences were removed using trimAl [93]. A phylogenetic tree was inferred using FastTree v2.1.9 [94]. Most analyses were performed in the R statistical environment [95] using APE [96], ggplot2 [97], gUniFrac [98], Maps, Mapplots, Picante [99] and Vegan. The Vegan function adonis and adonis2 were used to investigate the amount of variance in community composition explained by environmental or geographic variables. Variance partitioning analyses were run with varpart in Vegan and tested for significance with ANOVA. Distance decay, which refers to the decrease in microbial community similarity as geographic distance between communities increases was investigated in R using Mantel correlograms between geographic distance and -diversity, considering distance classes of 1,000 km. Local Contributions to Beta Diversity (LCBD) [36], which indicates the degree of uniqueness of each community in terms of its species composition, was measured with adespatial [100]. See Supplementary Methods, Additional file 6.

Quantification of selection, dispersal and drift
These processes were quantified using an approach that relies on null models, consisting of two main sequential steps: the first uses OTU phylogenetic turnover to infer the action of selection and the second uses OTU compositional turnover to infer the action of dispersal and drift [21]. The action of selection, dispersal and drift was quantified using both OTUs_{99\%} and OTUs_{ASVs}. In order to determine the action of selection using phylogenetic turnover, we first checked whether habitat preferences of phylogenetically closely related taxa (according to the 16S and 18S rRNA-genes) were more similar to each other than to those of more distantly related taxa, what is known as phylogenetic signal [101,
We tested for phylogenetic signal using temperature and fluorescence, which were the two variables that explained the highest fraction of community variance. We detected phylogenetic signal at relatively short phylogenetic distances (Figure S10, Additional file 19; Figure S11, Additional file 20), which is coherent with previous work [21, 103, 104]. We measured phylogenetic turnover using the abundance-weighted Mean Nearest Taxon Distance (MNTD) metric [17, 21], which quantifies the mean phylogenetic distances between the evolutionary-closest OTUs in two communities. MNTD values can be larger, smaller or equal to the values expected when selection is not affecting community turnover (that is, expected by chance). MNTD values higher than expected by chance indicate that communities experience heterogeneous selection [17]. In contrast, MNTD values which are lower than expected by chance indicate that communities experience homogeneous selection. Null models included 999 randomizations [21]. Differences between the observed MNTD and the mean of the null distribution are denoted as Nearest Taxon Index (NTI), with |NTI|>2 being considered as significant departures from random phylogenetic turnover, pointing to the action of selection.

The second step uses OTU turnover to calculate whether the -diversity of communities not structured by selection could be generated by drift (i.e. chance) or dispersal. We calculated the Raup-Crick metric [105] using Bray-Curtis dissimilarities (hereafter RC_{bray}) [21]. RC_{bray} compares the measured -diversity against the -diversity that would be obtained under random community assembly (drift); randomizations were run 9,999 times. RC_{bray} values between −0.95 and +0.95 point to a community assembly governed by drift. On the contrary, RC_{bray} values >+0.95 or <+−0.95 indicate that community turnover is driven by dispersal limitation or homogenizing dispersal respectively [105]. See Supplementary Methods, Additional file 6.

Estimation of interaction-adjusted indices
Taxa INteraction-Adjusted (TINA) and Phylogenetic INteraction Adjusted (PINA) indices were estimated following Schmidt et al. [24]. TINA is based on taxa co-occurrences while PINA considers phylogenetic similarities. TINA quantifies -diversity as the average association strength between all taxa in different samples. Thus, communities which are identical or include taxa that are perfectly associated will give
a TINA value of 1. TINA values will approach 0.5 in communities sharing no taxa or having neutral associations, and approach 0 if taxa display high avoidance. Dissimilarity matrices were generated as 1-TINA and used in downstream analyses (e.g. Figure 3). Full picoeukaryotic and prokaryotic subsampled OTU\textsubscript{.99} tables were used to calculate the abundance-weighted TINA\textsubscript{w} and PINA\textsubscript{w}. TINA\textsubscript{w} was calculated using picoeukaryotic and prokaryotic data from 119 Malaspina surface stations (most stations in Figure 1A). In addition, TINA\textsubscript{w} was calculated using data from TARA Oceans, including 63 surface stations for prokaryotes and 40 surface station for small unicellular eukaryotes (Figure 3).

**Associations between taxa and environmental parameters**

We analysed whether OTUs\textsubscript{.99} displayed associations with environmental variables and between themselves. Firstly, we used the Maximum Information Coefficient (MIC) which captures diverse relationships between two pairs of variables [106]. The Malaspina dataset consisted of 119 stations and 17 environmental variables. In the TARA Oceans dataset, prokaryotes were analysed across 63 surface stations (including 8 environmental variables), while microbial eukaryotes were analysed across 40 surface stations (including 6 environmental variables) [see Supplementary Methods, Additional file 6]. In both datasets, MIC analyses were run using CV = 0.5, B = 0.6, and statistically significant relationships with MIC $\geq$0.4 (Malaspina) or MIC $\geq$0.5 (TARA Oceans) were considered (MIC thresholds were adjusted to the characteristics of the datasets). MIC significance was assessed using precomputed p-values [106]. Secondly, we constructed association networks with the Malaspina dataset considering OTUs\textsubscript{.99} with $>100$ reads using SparCC [107] as implemented in FastSpar108. To determine correlations, FastSpar was run with 1,000 iterations, including 1,000 bootstraps to infer p-values. We used OTUs\textsubscript{.99} associations with absolute correlation scores $>0.3$ and $p<0.01$. Networks were visualized and analysed with Cytoscape [109] and igraph [110].

**Declarations**

**Ethics approval and consent to participate**

Not applicable

**Consent for publication**

Not applicable

**Availability of data and materials**
DNA sequences and metadata from the *Malaspina* expedition are publicly available at the European Nucleotide Archive (http://www.ebi.ac.uk/ena; accession numbers PRJEB23913 [18S rRNA genes] & PRJEB25224 [16S rRNA genes]). The data used from TARA Oceans is publicly available through Pangaea (https://doi.org/10.1594/PANGAEA.873275) as well as in (http://ocean-microbiome.embl.de/companion.html) [32, 50]. The code for generating OTUs, 99% tables is available in: https://doi.org/10.5281/zenodo.259579. R-Scripts for calculating the -Nearest Taxon Index and the Raup-Crick metric are available in https://github.com/stegen/Stegen_etal_ISME_2013. The code for calculating the TINA and PINA indices is available in https://github.com/defleury/Schmidt_et_al_2016_community_similarity, while the code for calculating MIC is available at http://www.exploredata.net. All used R packages as well as other software are cited in Methods.

Competing interests
The authors declare that they have no competing interests

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Authors’ contributions
RL designed the study. SGA, CMD, JMG and RM organized the sampling expedition. CR-Giner, GS, and MS collected samples, extracted the DNA and organized its sequencing. RL, IMD, AKK, PCJ analysed the DNA data, while CR-Giner, TSBS, LRR, MM, GS, CR-González, MS, CdV, RM, and JMG provided contextual ecological or environmental pre-processed data. RL, IMD, AKK, PCJ, TSBS, GS, CdV and RM interpreted the results. RL wrote the manuscript. All authors contributed substantially to manuscript
revisions. All authors read and approved the final manuscript.

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Additional Files

**Additional file 1: Figure S1**

(additional_file_1_FigureS1.png)

Position of the 120 analysed *Malaspina-2010* stations in the context of the Longhurst biogeographic provinces [35].

**Additional file 2: Figure S2**

(additional_file_2_FigureS2.pdf)

Bray Curtis and gUniFrac distances between picoeukaryotes and prokaryotes from the *Malaspina* dataset. Regression (blue) and 0:1 (red) lines are indicated.
Additional file 3: Table S1
(additional_file_3_TableS1.docx)

Additional file 4: Figure S3
(additional_file_4_FigureS3.png)

OTUs_{99\%} mean relative abundance (i.e. regional abundance) vs. occurrence (i.e. number of samples in which each OTUs_{99\%} is present) for the *Malaspina* dataset. The red and black horizontal lines indicate percentages of occurrences of 80\% and 20\% respectively. Cosmopolitan OTUs were considered as those with a percentage of occurrence >80\%, while restricted OTUs were those with a percentage of occurrence <20\% (see Table S2, Additional file 5). Blue and green vertical lines indicate regional abundances above and below which OTUs are considered regionally abundant (>0.1\%) or rare (<0.001\%) respectively.

Additional file 5: Table S2
(additional_file_5_TableS2.docx)

Additional file 6: Supplementary Methods
(additional_file_6_Supplementary_Methods.docx)

Additional file 7: Figure S4
(additional_file_7_FigureS4.pdf)

The 57 *Malaspina* stations for which 17 environmental parameters were available (*Meta-57* dataset).

Additional file 8: Figure S5
(additional_file_8_FigureS5.pdf)

Percentage of variance in Picoeukaryotic and Prokaryotic community composition (ADONIS $R^2$)
explained by water temperature and fluorescence when using different β-diversity metrics. Figure based on the *Malaspina Meta-57* dataset.

**Additional file 9: Figure S6**

(additional_file_9_FigureS6.pdf)

Species association networks for the tropical and subtropical surface-ocean microbiota as inferred from the *Malaspina* dataset. Left-hand side: Association networks of picoeukaryotes and prokaryotes considering positive (red) and negative (blue) correlations in panels A) [Eukaryotic Network (+e)] and B) [Prokaryotic Network (+e)], and only positive correlations in C) [Eukaryotic Network (+e)] and D) [Prokaryotic Network (+e)]. On the right-hand side, we present an alternative visualization of the network as well as the following network characteristics: number of nodes (n), number of edges with positive correlation (+e) and negative correlation (-e), average degree (avg. d), average path length (avg. l), global transitivity (t), number of modules with at least 3 nodes (m) and the number of nodes in each of those modules (sizes: n). The smaller network visualization on the right-hand side groups the nodes according to the modules. The colors of nodes in Left- and Right-hand side networks indicate the modules to which they belong (NB: colors in panels A, B, C & D are independent of each other).

**Additional file 10: Table S3**

(additional_file_10_TableS3.docx)

**Additional file 11: Table S4**

(additional_file_11_TableS4.docx)

**Additional file 12: Table S5**

(additional_file_12_TableS5.docx)
Percentage of OTUs, 99%, significantly associated to different environmental variables (MIC > 0.4) and their corresponding contribution to total sequence abundance (i.e. percentage of reads) in the *Malaspina* dataset. NB: Temperature, Oxygen, Conductivity and Salinity are correlated. OTUs can be associated to more than one variable.

Bray-Curtis dissimilarities and gUniFrac distances in Prokaryotes and Picoeukaryotes from the *Malaspina* dataset. In both cases, mean differences were significant (Wilcoxon test, p<0.05).

Prokaryotes (Bray Curtis mean=0.61, SD=0.19; gUniFrac mean=0.30, SD=0.07); Picoeukaryotes (Bray Curtis mean=0.74, SD=0.08; gUniFrac mean=0.50, SD=0.06).

Stations (total 36) from the *Malaspina* dataset featuring a comparatively large contribution to the overall b-diversity (LCBD = Local Contributions to Beta Diversity [36]; p<0.05)
Additional file 19: Figure S10
(additional_file_19_FigureS10.pdf)

Phylogenetic signal was detected across short phylogenetic distances for both the 16S and 18S rRNA-gene markers as indicated by phylogenetic mantel correlograms (*Malaspina* dataset). Phylogenetic signal was tested using temperature and fluorescence, the two variables that explain the highest fraction of community variance. Solid and open squares indicate significant and nonsignificant (using p=0.05) correlations respectively between environmental similarity (in terms of temperature and fluorescence) and phylogenetic relatedness. Correlations that are significantly positive indicate that the phylogenetic distance between OTUs-99% increases as environmental similarity decreases for the phylogenetic range being analysed. Phylogenetic distances were measured as abundance-weighted b-Mean Nearest Taxon Distances (bMNTD).

Additional file 20: Figure S11
(additional_file_20_FigureS11.pdf)

Same as Figure S10, Additional file 19 but using OTUs-ASVs. Solid and open squares indicate significant and nonsignificant (using p=0.05) correlations respectively between environmental similarity (in terms of temperature and fluorescence) and phylogenetic relatedness. Correlations that are significantly positive indicate that the phylogenetic distance between OTUs-ASVs increases as environmental similarity decreases for the phylogenetic range being analysed. Phylogenetic distances were measured as abundance-weighted b-Mean Nearest Taxon Distances (bMNTD).

Figures
Ecological mechanisms shaping the tropical and subtropical surface-ocean picoplankton.

Panel A) Position of the 120 stations included in this work that were sampled as part of the
Malaspina-2010 expedition (green dots) in the tropical and subtropical ocean. A snapshot of the global sea surface temperature, a main environmental driver affecting microbial distributions, is shown as a general representation of the temperature gradients in the surface ocean (as inferred using the ‘optimum interpolation sea surface temperature’ dataset from the NOAA corresponding to the 17 of March of 2018). Note that temperatures measured in situ were used in all analyses, not the ones displayed here. Panel B) Percentage of the community turnover associated to different ecological processes in prokaryotes and picoeukaryotes in the tropical and subtropical ocean as calculated using OTUs-99% and OTUs-ASVs. Note that percentage refers to the percentage of pairs of communities that appear to be driven by a given process.
Main variables influencing the structure of the surface-ocean microbiota as captured by different $\beta$-diversity metrics. Percentage of variance in picoeukaryotic and prokaryotic community composition (ADONIS R2) explained by Water Temperature and Longhurst Provinces when using different $\beta$-diversity metrics. Figure based on the Malaspina Meta-119 dataset (see Methods). TINA$_w$: TINA weighted, gUniFrac: Generalized Unifrac, PINA$_w$: PINA weighted. N.S. = Non-Significant. Note that TINA$_w$, which considers species association networks, captures a significantly higher proportion of community variance associated to temperature than Bray-Curtis, a compositional index, in prokaryotes.
Temperature-driven selection seems to affect species association networks in prokaryotes but not in pico-/nano-eukaryotes. Differences in community composition (as $1-\text{TINA-weighted} = \text{TINAw dissimilarities}$) vs. temperature differences (as Euclidean distances)
based on dimensionless z-scores) for both small unicellular eukaryotes and prokaryotes sampled during the Malaspina and TARA Oceans expeditions. Note that, in contrast to other indices, TINAw considers species-association patterns (i.e. co-occurrences and co-exclusions) when estimating $\bar{\alpha}$-diversity [24]. NB: While only picoeukaryotes were contemplated in Malaspina (cell sizes <3 µm), TARA Oceans data included pico- and nano-eukaryotes (cell sizes <5 µm). Pico- and nanoeukaryotes from both expeditions (left panels) displayed low or no correlations between TINAw distances and temperature differences (Mantel test results included in the panels). On the contrary, prokaryotes (right panels) displayed high to moderate correlations between TINAw distances and temperature differences. These differences in the correlations are likely due to the wider temperature ranges covered by TARA Oceans compared to Malaspina (see Results). The regression line is shown in red (Malaspina microbial eukaryotes N.S., Malaspina Prokaryotes R2=0.3, TARA Oceans microbial eukaryotes R2=0.1, TARA Oceans Prokaryotes R2=0.7; p<0.05). The maps at the bottom indicate the surface stations from the expeditions Malaspina (119 stations for both prokaryotes and picoeukaryotes) and TARA Oceans (63 stations for prokaryotes and 40 stations for small unicellular eukaryotes) that were used to calculate TINAw.
Picoeukaryotic communities display a higher spatial differentiation than prokaryotic counterparts in the tropical-subtropical surface-ocean. Panels A-C: Sequential change in community composition across space (sequential \( \beta \)-diversity). Communities were sampled along the Malaspina expedition (Panels A and B, black arrows), and the composition of each community was compared against its immediate predecessor. In Panels A and B, the size of each bubble represents the Bray-Curtis dissimilarity between a given community and the community sampled previously. Blue squares in Panels A and B represent the stations where \( \beta \)-diversity displayed abrupt changes (Bray Curtis values >0.8 for picoeukaryotes and >0.7 for prokaryotes). Abrupt changes coincided in a total of 11 out of 14 stations for both picoeukaryotes and prokaryotes, while one station displayed marked changes only for picoeukaryotes and two only for prokaryotes. Panel C summarizes the sequential Bray-Curtis values for prokaryotes and picoeukaryotes (Means were significantly different between domains [Wilcoxon text, \( p<0.05 \)]). Panel D indicates the differences in distance-decay
between prokaryotes and picoeukaryotes in the tropical and subtropical surface-ocean. Mantel correlograms between geographic distance and $\theta$-diversity featuring distance classes of 1,000 km for both picoeukaryotes and prokaryotes are shown. Coloured squares indicate statistically significant correlations ($p<0.05$). Note that $\theta$-diversity in picoeukaryotes displayed positive correlations with increasing distances up to $\sim$3,000 km, while prokaryotes had positive correlations with distances up to $\sim$2,000 km. Correlations tended to be smaller in prokaryotes than in picoeukaryotes, indicating smaller distance decay in the former compared to the latter.

Supplementary Files
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- additional_file_2_FigureS2.pdf
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