



Large to submesoscale surface circulation and its implications on biogeochemical/biological horizontal distributions during the OUTPACE cruise (South West Pacific)

Louise Rousselet¹, Alain de Verneil¹, Andrea M. Doglioli¹, Anne A. Petrenko¹, Solange Duhamel², Christophe Maes³, and Bruno Blanke³

¹Aix Marseille Univ, Université de Toulon, CNRS, IRD, OSU PYTHEAS, Mediterranean Institute of Oceanography MIO, UM 110, 13288, Marseille, Cedex 09, France

²Lamont-Doherty Earth Observatory, Division of Biology and Paleo Environment, PO Box 1000, 61 Route 9W, Palisades, NY 10964, USA

³Laboratoire d'Océanographie Physique et Spatiale, CNRS, Ifremer, IRD, UBO, Brest, France

Correspondence to: Rousselet Louise (louise.rousselet@mio.osupytheas.fr)

Abstract. The patterns of the large-scale, meso- and submesoscale surface circulation on biogeochemical and biological distributions are examined in the Western Tropical South Pacific (WTSP) in the context of the OUTPACE cruise (Feb-April 2015). Multi-disciplinary original *in situ* observations were achieved along a zonal transect through the WTSP and their analysis was coupled with satellite data. The use of Lagrangian diagnostics allows for the identification of water mass pathways, mesoscale structures, and submesoscale features such as fronts. In particular, we confirmed the existence of a global wind-driven southward circulation of surface waters in the entire WTSP, using a new high-resolution altimetry-derived product, validated by *in situ* drifters, that includes cyclogeostrophy and Ekman components with geostrophy. Two subregions show counter-intuitive water mass trajectories due to mesoscale circulation: i) the Coral Sea with surface exchanges between the North Vanuatu Jet and the North Caledonian Jet; and ii) the zonal band between 180°W and 170°W with an eastward propagation whereas a westward general direction dominates. Fronts and small-scale features, detected with Finite-Size Lyapunov Exponents (FSLE), are correlated with 25% of surface tracer gradients which reveals the significance of such structures in the generation of submesoscale surface gradients. Additionally, two high-frequency sampling transects of biogeochemical parameters and micro-organism abundances demonstrate the influence of fronts in controlling the spatial distribution of bacteria and phytoplankton, and as a consequence the microbial community structure. All circulation scales play an important role that has to be taken into account when analysing the data from OUTPACE but also, more generally, to understand the global distribution of biogeochemical components.

1 Introduction

The zonal trophic gradient of the Western Tropical South Pacific (WTSP) represents a remarkable opportunity to study the interactions between marine biogeochemical Carbon (C), Nitrogen (N), Phosphorus (P), Silica (Si), and Iron (Fe) cycles between different trophic regimes. One of the OUTPACE cruise's goals is to understand how N₂ fixation controls production, miner-



alisation and export of organic matter (Moutin and Bonnet, 2015; Moutin et al., 2017). The ocean's circulation, at different time/space scales, can play a key role in biological variability and dynamics. In particular the meso- and submesoscale, which occurs at scales typical of phytoplankton blooms (Dickey, 2003), enhance carbon export through vertical motion (Guidi et al., 2012; Lévy et al., 2012) and thus strongly impact the biological pump.

5 Indeed, mesoscale dynamics (features with time/space scales on the order of months/100 km such as eddies) can affect biological and biogeochemical cycling through transport processes such as horizontal advection, lateral stirring and eddy trapping, as well as through processes that modify nutrients and/or light availability such as eddy pumping, eddy-wind interaction or frontal instabilities (Williams and Follows, 1998; McGillicuddy Jr, 2016). Some observations, mostly collected during the JGOFS program, have shown the influence of eddy circulation in sustaining primary production in oligotrophic regions (Jenkins, 1988; McGillicuddy and Robinson, 1997). Numerically modelled eddy fields also show an enhancement of biological productivity in most provinces of the North Atlantic Ocean (Oschlies and Garçon, 1998; Garçon et al., 2001). Then additional studies also discuss the structuring effect of mesoscale features, such as vortices, on ecological niche composition and distribution depending on eddy characteristics and eddy stirring (Sweeney et al., 2003; d'Ovidio et al., 2010; Perruche et al., 2011; d'Ovidio et al., 2013). Thus mesoscale features can have strong ecological impacts through the enhancement of biological production and the creation of favourable conditions for less competitive species, with implications for higher trophic levels.

Smaller scales may also have a significant role in the distribution of biological variability. The submesoscale represents the ocean processes characterized by horizontal scales 1-10 km whose origins might be linked with the stirring induced by mesoscale interactions and frontogenesis (Capet et al., 2008). At this typical scale, the flow can be characterized by strong stretching lines or vortex boundaries creating physical barriers such as fronts or filaments that are associated with sharp gradients. These structures can contribute to the separation or mixing of water masses and thus impact the horizontal distribution of tracers, biogeochemical and biological matter such as biomass of phytoplankton cells at a front boundary. Indeed microorganisms are buoyant material and their distribution, as well as the biogeochemical components, can be driven by submesoscale activity, whether through direct horizontal advection (Dandonneau et al., 2003), or indirectly following the biogeochemical dynamics. Nitrogen fixing organisms such as *Trichodesmium spp.*, which contribute in sustaining high primary productivity in the Pacific ocean, are known to concentrate around small-scale features in the North Pacific (Fong et al., 2008; Church et al., 2009; Guidi et al., 2012). In the WTSP, Bonnet et al. (2015) argued that *Trichodesmium spp.* abundances might follow gradient distributions. Besides regulating the spatial distribution of micro-organisms, these features can participate in biological dynamics as they can induce vertical movements of nutrient supplies and chlorophyll (Martin et al., 2001). Lévy et al. (2015) showed that the flow field brings populations into contact in frontal areas which can be characterized by a larger diversity of micro-organisms and more fast-growing species. Consequently, submesoscale circulation can influence the planktonic community structure. However, due to their typical scales, mesoscale and submesoscale features, require substantial means to be adequately observed (Mahadevan and Tandon, 2006) and their interactions with biogeochemistry and biology are also hard to elucidate due to their ephemeral nature (McGillicuddy Jr, 2016).

In the WTSP the large scale circulation is dominated by the anticyclonic South Pacific Gyre. The South Equatorial Current (SEC) flows in the equatorial band (0°S - 6°S) from East to West and is divided in multiple branches when approaching the



Coral Sea (Webb, 2000; Sokolov and Rintoul, 2000) (Figure 1). On the western boundary, the East Australian Current (EAC) feeds, through the Tasman Sea, the southern branch of the gyre which then flows east and reaches the Peru/Chile Current (PCC) near the western South American coast (Tomczak and Godfrey, 2013). Superimposed on these large scale patterns, several studies indicate a strong mesoscale variability due to barotropic instabilities and the interactions of the major currents and jets with the numerous islands of the region (Qiu et al., 2009; Hristova et al., 2014). As displayed in Figure 1, the OUTPACE cruise was conducted in a zonal band of relatively high eddy kinetic energy (Qiu et al., 2009). The influence of this intense variability, which results in mostly westward propagating eddies (Chelton et al., 2007; Rogé et al., 2015), has not been fully explored yet (Kessler and Cravatte, 2013), as well as its implications on biogeochemical/biological variations in the region. Recent studies have underlined the role of mesoscale activity as a conveyor of water masses, leading to the discovery of a potential water mass pathway in the Coral Sea (Maes et al., 2007; Ganachaud et al., 2008; Rousselet et al., 2016). This intense mesoscale activity is strongly linked to submesoscale fronts that might be responsible for surface small-scale features in temperature and salinity as shown by Maes et al. (2013) within the Coral Sea. Submesoscale dynamics are also thought to be responsible for ~20% of new production in oligotrophic regions as suggested by Lévy et al. (2014b) using an idealized model. Since the frequency of oceanic fronts and eddy kinetic energy should increase in oligotrophic regions with climate change (Matear et al., 2013; Hogg et al., 2015) the OUTPACE cruise offers an unprecedented opportunity to study large, meso- and submesoscale influences along a zonal gradient crossing the oligotrophic to ultra-oligotrophic WTSP with coupled physical and biogeochemical measurements.

In this study we investigate the large, meso- and submesoscale patterns using *in situ* observations obtained during the OUTPACE cruise, coupled with satellite data. Remote sensing provides daily physical and biological information over the entire WTSP for a time period covering the cruise duration and beyond (from June, 1 2014 to May, 31 2015). The inter-comparison between physical lagrangian diagnostics and biogeochemical/biological measurements explores the potential influence of small-scale ocean circulations on horizontal dispersal of tracers such as temperature, salinity or chlorophyll, as well as on biological dynamics. The use of multidisciplinary approaches, including *in situ* observations, remote sensing and numerical simulations is the key aspect of this study to investigate the surface circulation at different scales and try to understand their potential influence on biogeochemical-biological distributions during the OUTPACE cruise.

2 Materials and Methods

2.1 *In situ* observations

The OUTPACE (Oligotrophy to UItra-oligotrophy PACific Experiment) cruise performed a zonal transect across the WTSP aboard R/V Atalante from February 18, 2015 to April 3, 2015 (Moutin and Bonnet, 2015). The main objectives of the cruise were to study the interactions between planktonic organisms and the cycling of biogenic elements across trophic and N₂ fixation gradients (Moutin et al., 2017). A total of 15 hydrological stations were sampled along the transect as well as three long-duration (LD) stations named LDA, LDB and LDC (Fig. 1). LD station sampling lasted for almost 8 days each, and aimed to study the total export of carbon in 3 biogeochemically different regions. More details about the sampling strategy are



available in Moutin et al. (2017). The multi-disciplinary measurements used in this study are described hereinafter.

Of particular interest to understand the surface dynamics, SVP (*Surface Velocity Program*) floats were launched during each LD station to investigate the dispersion and the surface circulation relative to 15 m during the sampling period and beyond
5 (Lumpkin and Pazos, 2007). Three SVPs were launched during LDA, 6 during LDB and 4 during LDC. The *in situ* trajectories of the floats would be used to validate altimetry-derived surface velocities (see Section 2.2 and Fig.A1).

Continuous measurements of temperature and salinity were achieved using a ThermoSalinoGraph (TSG) that pumped sea water at 5 m depth. TSG data have been corrected and calibrated using independent measurements of salinity from water bottle
10 samples collected daily onboard RV *L'Atalante* (following the procedures described by Alory et al. (2015)) and are binned into minutes. In the following, temperature and salinity will refer to absolute salinity and conservative temperature, respectively, according to TEOS-10 standards (McDougall et al., 2012). A Wetstar SeaBird fluorimeter was deployed on the underway water flow. The fluorimeter provides measurements proportional to the chlorophyll a concentration with a time step of 10-15 min. Discrete samples were taken during the transit to calibrate the fluorimeter using the Aminot and K erouel (2004) method:

15
$$\text{Chla [mg m}^{-3}] = 1.99 \times \text{FluorescenceValue} - 0.083 \text{ (R}^2=0.87, n=55)$$

Due to technical issues the underway sampling of chlorophyll concentration started on March 7. Each of these data sets have been interpolated on a regular grid of 0.5 km resolution in order to keep the high resolution, but equally distributed along the travelled distance.

20 A high frequency sampling (every 20 min) was performed upon leaving LDA until arriving at LDB, in order to assess variability in 15 different biogeochemical parameters, in particular the Dissolved Inorganic Phosphate (DIP) turnover times and the abundances of bacteria (including the low and high nucleic acid content bacterial groups, LNA and HNA, respectively), *Prochlorococcus*, *Synechococcus* and picophytoeukaryotes (PPE). To enumerate cell abundances of these different microbial groups, water samples were collected directly from the underway pump, fixed with 0.25% (w/v) paraformaldehyde, flash frozen
25 and preserved at -80 C until analysis by flow cytometry following the protocol described in Bock et al. (2017). Briefly, bacteria were discriminated in a sample aliquot stained with SYBR Green I DNA dye (1:10,000 final) while pigmented groups were discriminated in an unstained sample aliquot. Reference beads (Fluoresbrite, YG, 1  m) were added to each sample. Particles were excited at 488 nm (plus 457 nm for unstained samples) and bacteria were discriminated based on their green fluorescence and forward scatter (FSC) characteristics, while *Prochlorococcus*, *Synechococcus* and PPE were discriminated based on their
30 chlorophyll (red) fluorescence and FSC characteristics. LNA and HNA groups were further distinguished based on their relatively low and high SYBR Green fluorescence, respectively, in a green fluorescence vs side scatter plot. *Prochlorococcus* were further distinguished from *Synechococcus* by their relative lack of a phycoerythrin signal (orange fluorescence). Using a FSC detector with small particle option and focusing a 488 plus a 457 nm (200 and 300 mW solid state, respectively) laser into the same pinhole greatly improved the resolution of dim surface *Prochlorococcus* population from background noise in unstained



samples. Because the *Prochlorococcus* population cannot be uniquely distinguished in the SYBR stained surface samples, bacteria were determined as the difference between the total cell numbers of the SYBR stained sample and *Prochlorococcus* enumerated in unstained samples. Cytograms were analyzed using FCS Express 6 Flow Cytometry Software (De Novo Software, CA, US). These data are used to investigate the small-scale distribution of the different community groups and its relation
5 with the concomitant dynamics at submesoscale.

2.2 Satellite data

Several satellite datasets were exploited during the campaign to guide the cruise through an adaptive sampling strategy using the SPASSO software package (<http://www.mio.univ-amu.fr/SPASSO/>) following the same approach of previous cruises such as LATEX (Doglioli, 2013; Petrenko et al., 2017) and KEOPS2 (d'Ovidio et al., 2015). SPASSO was also used after the cruise
10 in order to extend the spatial and temporal vision of the *in situ* observations.

Four different altimetry-derived velocity products were tested in this study to choose the product that best represents the surface circulation during the cruise. First the daily Ssalto/Duacs product (Ducet et al., 2000), from AVISO (Archiving, Validation and Interpretation of Satellite Oceanographic 3) data base, for the period from 1 January 2004 to 31 December 2015,
15 was used to extract daily delayed-time maps of absolute geostrophic velocities ($1/4^\circ \times 1/4^\circ$ on a Mercator grid, since 15 April 2014). Three other altimetry products were specifically produced, for the first time, at $1/8^\circ$ resolution for the WTSP region by Ssalto/Duacs and CLS (Collecte Localisation Satellites), with support from CNES (Centre National d'Études Spatiales), from June 2014 to June 2015. In particular, they provided daily maps of absolute geostrophic velocities, daily maps of the sum of absolute geostrophic velocities and Ekman components, and the same product as the latter that also includes a cyclogeostrophy
20 correction (Penven et al., 2014). Ekman surface currents refer to the wind-induced circulation relative to 15 m and are computed from ECMWF ERA INTERIM windstress with an Ekman model fitted onto drifting buoys (Rio et al., 2014).

A preliminary comparison between Lagrangian numerical particle trajectories (see Section 2.3.1), computed with each of the above products, and the observed trajectories of the different floats launched during OUTPACE (section 2.1) allowed us to identify the product including geostrophic and Ekman components with the cyclogeostrophy correction (hereafter total
25 altimetry-derived velocity field) as the most accurate *in situ* surface currents (see figure A1 for a comparison between float trajectories and numerical particle derived streamfunction).

Daily near-real-time maps of sea surface temperature (SST) and ocean color were also specifically produced for the WTSP from December 2014 to May 2015 by CLS with support from CNES. They are constructed with a simple weighted data average over
30 the 5 previous days (giving more weight to the most recent data), and have a $1/50^\circ$ resolution (2 km at the Equator) in latitude and longitude. The temperature product corresponds to maps of SST deduced from a combination of several intercalibrated infrared sensors (AQUA/MODIS, TERRA/MODIS, METOP-A/AVHRR, METOP-B/AVHRR). The ocean color product corresponds to maps of chlorophyll concentration issued from the Suomi/NPP/VIIRS sensor (<http://npp.gsfc.nasa.gov/viirs.html>). These satellite data are compared with *in situ* data from the underway survey. A correlation of 0.8 between *in situ* mea-



surements and co-located satellite data validates the satellite-derived SST and chlorophyll concentration. A supplementary correlation with the daily High-Resolution SST blended from NCDC/NOAA (Reynolds et al., 2007) and *in situ* SST showed a similar correlation. These results corroborate the accuracy of the CLS products in our region of interest.

2.3 Lagrangian diagnostics

5 2.3.1 Surface water mass pathways detection

To investigate the water mass movements at large and meso-scale, we used the Lagrangian diagnostic tool Ariane that can trace water mass movements from the trajectories of numerical particles that enter and exit a predefined domain (Blanke and Raynaud, 1997; Blanke et al., 1999). In this study the numerical particle trajectories are computed with altimetry-derived surface currents from the products listed above (see section 2.2). As many Lagrangian particles as desired can be integrated in two different ways: backward in time to assess the origins of the water masses or forward in time to investigate their fate. Additionally, this Lagrangian tool allows for the computation of two different diagnostics : i) qualitative diagnostics that compute typically few particles with a steady recording of the positions along their trajectories; ii) quantitative diagnostics that compute thousands of particles with statistics available for initial and final positions, and with the diagnostic of the main pathways. In this study we use 3 different configurations of the Ariane tool depending on the objectives.

15 First, to identify the altimeter product that best fit the observed trajectories of the floats launched during OUTPACE, a comparison was done with the trajectories of numerical particles (typically considering release of several hundred of particles). They were initially positioned around the launching position of the floats with a resolution of 1-2 km. The particles were advected forward in time for 96 (LDA), 78 (LDB) and 70 (LDC) days, corresponding to the time lapse between the launch day of the floats and the last available day of satellite data. These qualitative experiments allows for the comparison of successive positions (every 6 hours) of numerical particles computed with the 4 different products and those of the floats. Thus the choice of the best surface velocity product relied on the best fit between observed and numerical trajectories. The total altimetry-derived velocity field (i.e. the product including geostrophy, cyclogeostrophy and Ekman components) will be used for all diagnostic computations, as it best fits *in situ* data (Fig.A1).

20

To study the large scale circulation in the WTSP, quantitative experiments were performed to find the main pathways of the waters entering and exiting the box contouring the WTSP (Fig. 2). Particles were launched along each section of the box (North, East, South, West) and advected forward with the total altimetry-derived surface currents. We simulated ten years of particle trajectories by repeating ten times the available dataset. We compare the results of this simulation with the ten years integration of available geostrophic AVISO surface currents over this time period. The comparison between these simulations ensures that the use of a one year time period looped several times does not significantly modify statistical outputs. It is clearly more interesting to use CLS data instead of commonly used AVISO geostrophic surface currents because they include the wind effect and cyclogeostrophy with higher resolution, as well as better represents *in situ* data.

30

Another objective is to identify the mesoscale trajectories of surface water masses sampled at each of the LD stations. Backward and forward quantitative experiments were performed to identify the main pathways of the surface water masses arriving



and leaving each LD station using total altimetry-derived surface currents (Figure 4). Almost one million numerical particles were initially distributed along a square box surrounding the position of the LD station. The calculations were stopped whenever the particles return to the initial box (hereafter called meanders) or were intercepted on one of the four remote sections located around the LD station. The boxes size are tuned in order to minimize meanders and the loss of particles in the domain.

5 Percentage of both quantities are reported in Table A1. Forward computation times are identical to the qualitative diagnostics. For backward experiments, the particles were advected for 183 (LDA), 201 (LDB) and 209 (LDC) days corresponding to the maximum time lapse allowed by CLS satellite data availability.

2.3.2 Eddies and filaments identification

To set up the mesoscale context during the OUTPACE cruise, we used the Lagrangian Averaged Vorticity Deviation method (Hadjighasem and Haller, 2016; Haller et al., 2016) that allows identification of coherent structures from altimetry-derived surface velocity fields (code available at <https://github.com/Hadjighasem/Lagrangian-Averaged-Vorticity-Deviation-LAVD>). The detected features are able to trap water masses for a certain period (defined by the integration time) and transport them along their route. In this study we computed the detection with the total altimetry-derived velocity field and chose an 8 day time integration with respect to the duration of LD stations. Indeed this time interval provides a confirmation or rebuttal of the assumption that LD stations have been performed in a coherent structure, as targeted during the cruise. This Lagrangian diagnostic is also compared with a hybrid Lagrangian and Eulerian approach combining the calculation of the Okubo-Weiss (OW) parameter and a retention parameter (RP), computed with the same velocity field. The OW parameter identifies structures such as eddies by separating the flow into a vorticity-dominated region and a strain-dominated region (Okubo, 1970; Weiss, 1981). The RP identifies the number of days a fluid parcel remains trapped within a structure core, defined by a negative OW parameter. Both parameter calculations are detailed in d'Ovidio et al. (2013). As for the LAVD detection method, the RP allows for the identification of potentially trapping coherent structures.

10
15
20

Submesoscale flow features in two-dimensional data are evaluated with altimetry-derived finite size Lyapunov exponents (FSLE), computed with the algorithm of d'Ovidio et al. (2004). This Lagrangian diagnostic detects frontal zone on which passive elements of the flow should theoretically align. Here we used the total altimetry-derived velocity field to compute the algorithm. The main parameter values for the algorithm are described in de Verneil et al. (2017b). The OUTPACE cruise occurred in the relatively open ocean and far enough from islands to trust the FSLE diagnostic calculated from altimetry.

25

We compare the horizontal positions of the fronts, detected with FSLE, with surface gradients measured both with the TSG (temperature and salinity) and the underway survey (chlorophyll). Indeed these high frequency samplings provide access to submesoscale gradients. A point by point correlation is calculated whenever a strong gradient of density (or chlorophyll) corresponds or not to a high FSLE value (i.e. $> 0.05 \text{ day}^{-1}$) indicative of a front. Sensitivity tests have been performed to choose the thresholds on density (chlorophyll) gradients to ensure the stability of the correlations calculated. These tests ended with the selection of gradient larger than $0.1 \text{ kg m}^{-3} \text{ km}^{-1}$ and a $0.2 \text{ mg m}^{-3} \text{ km}^{-1}$ thresholds for density and chlorophyll gradients, respectively.

30



3 Results and discussions

3.1 Large scale wind-driven pathways

The geostrophic large scale mean circulation and directions of the main currents in the WTSP are well established from the literature. However, the trajectories and pathways of surface waters may change and be more complex when the effect of the wind is added and resolution is increased especially in the context of inter-annual ENSO (El Niño Southern Oscillation) variability. We decided to use a Lagrangian integration of numerical particles to simulate the transport of surface fluid parcels at the scale of the WTSP region using altimetry-derived ocean currents including the wind effect. Figure 2 shows the transport calculated from the sum of almost 13 million numerical particles advected with the total surface altimetry-derived flow for 10 years (see Section 2.3.1). This figure highlights the westward transport of the SEC in the northwestern part of the domain, but also the eastward transport at 10°S due to the output of the South Equatorial Counter Current (SECC) from the Solomon Sea (Ganachaud et al., 2014). Both these pathways follow the well-known circulation of the SEC and SECC in this region. Additionally, a clear surface meridional transport is noticeable from 10°S to 25°S. In the very southeastern part of the domain some surface waters seem to recirculate to the east from 170°W, which is probably an indicator of the gyre circulation or an artefact of mesoscale dynamics due to the short averaging period considered by the present study. The meridional transport does not correspond to any surface current but is mainly due to the south-easterly trade winds. This meridional component appears due to the addition of the Ekman component in altimetry surface velocities (Fig.B1, Supplementary Material). The large scale transport of surface waters in the OUTPACE area is thus a combination of the transport by general well-known surface currents and wind-driven circulations. Surface waters globally travel the WTSP from northeast to southwest. This meridional transport seems to separate the gyre surface waters that will recirculate towards the east near 175°W and waters that will continue to be advected in the WTSP towards the Coral Sea. At the scale of the WTSP, the individual transport calculated from each initial section (see Section 2.3.1) reveals that 80% of the surface waters crossed during the OUTPACE cruise originate from the «North» section, 8-15% from the «East» section and very few from the «South» and «West» sections (Fig. C1). We can thus identify a general wind-driven surface transport in the WTSP as follows: the surface waters enter the WTSP from the northeast with the SEC and are gradually advected to the south with a part (east of 170°W) that directly recirculates within the gyre and another part (west of 170°W) that follows a southwestern propagation through the different WTSP archipelagos. From a biogeochemical point of view, two types of waters can be differentiated: the relatively oligotrophic but richer Melanesian waters (from 160°E to 170°W) and the ultra-oligotrophic gyre waters (west of 170°W) (Fumenia, personal communication). The wind-driven surface transport highlighted here could participate in the biogeochemical variations between western and eastern waters. Indeed the path through the Melanesian area may enrich these waters due to the contact with multiple islands whereas waters that directly recirculate within the gyre keep their ultra-oligotrophic characteristics. This pathway induces complex recirculation through the Melanesian archipelago, not visible on the large scale transport.

The WTSP circulation is also strongly impacted by ENSO conditions, responsible for SST variability on inter-annual to decadal timescales (Sarmiento and Gruber, 2006). A negative Southern Oscillation Index (SOI), El Niño phase, is characterized



by a decrease or even an overturn of trade winds whereas during La Niña phase (positive SOI) trade winds are strengthened. The mean wind velocity measured during OUTPACE is shown to be close to mean velocities during El Niño (data not shown) and Moutin et al. (2017) clearly showed that the OUTPACE cruise took place during an El Niño phase but they determined that climatological effects, upon the results of the cruise, were minimized. Otherwise, as this region is characterized by an intense mesoscale circulation, we can expect that it participates in the biogeochemical variations in the region. Thus we investigate the mesoscale feature trajectories on the entire WTSP and in particular the mesoscale circulation around three biogeochemically different locations: i) LDA located in the Coral Sea, at the end of the surface waters' journey across the WTSP; ii) LDB, in the Melanesian waters, west of the transition zone with gyre waters; iii) LDC, in the gyre waters.

3.2 Mesoscale activity and trajectories of surface waters

The major goal in this section is to identify whether mesoscale activity, and in particular trapping features, actively participate in the transport of different water mass properties across the WTSP ocean. The LAVD method is chosen in order to track the coherent structures for a time period of 8 days (see Section 2.3.2). Figure 3 shows the total altimetry-derived velocity field and the mesoscale structures identified for the first day of each LD station (LDA, LDB and LDC). It reveals that many mesoscale structures are found in the entire WTSP with no specific region with higher abundances of these features. A comparison with the eulerian Okubo-Weiss (OW) parameter method shows a good agreement with the LAVD detection method. Indeed, a mean OW parameter value of -0.24 day^{-2} is calculated inside the contour of coherent structures detected with the LAVD method. It indicates that wherever a coherent structure is identified, the OW parameter also identifies a mesoscale feature. Most of the mesoscale structures detected with the LAVD method are identified as retention areas, with the RP, ensuring the trapping characteristics of these features (Fig. C2). A tracking of these coherent structures highlights that they all show a general zonal propagation as expected from the mean transport in this area. If the major part propagates westward, the meridional band between 180°W and 170°W is identified as a region with mostly eastward propagation of mesoscale structures. Several studies have discussed the eastward flow south of Fiji, detectable on Figure 1, named the South Tropical Countercurrent (STCC) (Qiu and Chen, 2004). This flow, when bordering the Fijian coast, could lead to the eastward transport of mesoscale structures bounded east of Fiji. The location of this band is also interesting as it is found in the transition area between Melanesian waters and gyre waters. Unfortunately, the zonal equidistant biogeochemical sampling during OUTPACE did not sample both mesoscale features and surrounding waters and consequently no differentiation is possible between potentially trapped waters and surrounding waters. The small differences between water mass properties in this region may also make it difficult to confidently notice a biogeochemical marker of different water masses. However, the particular trajectories of these features could induce enriched-fluid transport into nutrient-poor gyre waters. Even if in this case the influence of mesoscale activity on biogeochemical variations is not directly visible on *in situ* data, through eddy trapping and transport, the role of mesoscale dynamics on the trajectories of surface waters can document the possible exchanges between biogeochemically differentiated regions.

Here we dynamically explore the origins and fates of surface waters sampled during each LD station located in three different



environments. LDA is situated on the path of the westward North Caledonian Jet (NCJ), that flows between New Caledonia and Vanuatu, in relatively highly productive waters (Fig.5). Far to the west, LDB lies in the eastward eddy-propagation band inside a phytoplankton bloom whereas LDC is located in nutrient-poor waters in the South Pacific (SP) gyre (Fig.5). Figure 4 shows the streamfunctions calculated from numerical particle advection using total altimetry-derived surface velocities (see Section 2.3.1). They represent the origin (Fig. 4 top) and the fate (Fig.4 bottom) of each LD stations' waters, respectively the backward and forward computations. One would expect LDA surface waters to come from the East as it is located on the path of the westward NCJ. However they seem to have multiple origins: i) easterly, directly from the NCJ transport; ii) northerly, directly from waters that have circulated between the Vanuatu islands before heading south to LDA; and iii) from a meridional tortuous recirculation path ($\sim 162^{\circ}\text{E}$) within the Coral Sea. Both i) and ii) origins agree with the integrated transport entering the Coral Sea induced by the complex topography (Kessler and Cravatte, 2013). The meridional recirculation determined here suggests that eddy-eddy interactions might be responsible for the emergence of complex paths between the NVJ and the NCJ. Indeed it matches the area described by Rousselet et al. (2016) as the region of exchange between NCJ and NVJ waters through eddy trapping and transport. We identify a probable water mass mixing area, in the Coral Sea center, through complex mesoscale stirring. This stirring may also create surface gradients as depicted by Maes et al. (2013). After LDA sampling, an intense signature of the NCJ is detected at the surface. Another portion of surface waters directly crash on New Caledonia's northern coast but this pathway might not be so relevant due to the satellite's lack of resolution near coastal areas. The westward circulation scheme is consistent with previous studies focusing on the NCJ (Ganachaud et al., 2008; Gasparin et al., 2011) and the results reported by Barbot et al. (2017).

In the eastern WTSP, LDB surface waters seem to follow the same general path: they flow from northeast towards southwest before they reach a group of islands and then recirculate to the east towards LDB. After LDB sampling, they continue their way to the east before heading back to the south or to the south-west. The eastward propagation of mesoscale structures, mentioned above, is sustained by the eastward circulation of surface waters around the LDB site (Fig. 4). de Verneil et al. (2017b) also pointed out a possible eastward transport to explain the origin of the bloom sampled at LDB. Indeed the surface waters might be iron-enriched through contact with the islands and thus create favorable conditions for a phytoplankton bloom. At this site, adjacent to the nutrient-poor SP gyre, the biological dynamics could be specially enhanced by the mesoscale eastward transport of essential chemical supplies for phytoplankton development. If this eastward transport is revealed to be quasi-permanent, it could be associated with recurrent bloom events in this area but this assumption requires further analyses to be generalized.

Further west, as one would expect, the waters sampled during LDC travelled from the east and flow to the west after LDC (Fig.4). We notice a recirculation area east of LDC where the waters seemed to follow a looping trajectory before reaching LDC. Both lagrangian methods (LAVD detection and advection of numerical particles) agree that a mesoscale structure trapped water masses east of LDC (from 164°W) and transported them until the sampling region of LDC. Indeed, the high rates of meanders (70% and 44% for backward and forward integration, respectively) (Table A1), also suggest the waters were still trapped in the structure core during LDC sampling. This observation is in good agreement with de Verneil et al. (2017a) who conclude that LDC was performed in a stable coherent water mass for the entire LDC sampling.



3.3 Fine scale distribution of tracers

The surface tracers' distribution is mostly driven by the wind-induced circulation but also by the transport through mesoscale activity. In a more ephemeral way, tracers can be dispersed following small-scale perturbations such as frontal features. Here we try to detect and quantify the influence of such features on the density and chlorophyll surface gradients using both *in situ* and satellite observations.

As described in Section 2.2 and shown by Figures 5a and 5b, the comparison between *in situ* observations and satellite-derived data results in reasonable correlations. As both datasets are comparable, SST and chlorophyll concentrations from remote sensing are then used to investigate horizontal gradients in the WTSP (Fig. 5a and Fig. 5b). To assess the spatial scale of submesoscale gradients (typical of 1-10 km) in terms of ocean dynamics, we use a lagrangian methodology based on the calculation of FSLE (see Section 2.3.2) that allow for fronts detection. Figure 5c shows regions where fronts are frequently generated during the time period of the cruise. We notice that the gyre is a region less suitable for fronts to occur and persist. We also identify east of the Fiji islands a zonal band at 18°S where almost no fronts occur during the OUTPACE cruise. Southeast of LDB a mesoscale structure, that is also identified on Figure 3, creates a frontal barrier that lasted for more than 30 days. It seems that this structure matches with strong surface gradients in chlorophyll and in SST, consequently separating colder and relatively chlorophyll-rich waters to the south from warmer but chlorophyll-poor waters to the north of the front. Overall, as shown by Figure 5, the most frequent and long-lived fronts seem to help in structuring the spatial distribution of tracers such as SST and chlorophyll concentration by creating physical barriers, isolating areas with different biogeochemical characteristics. To try to quantify the influence of frontogenesis on the structuring effect of surface tracers, we decided to compare the surface sharp gradients measured by the TSG or with the underway fluorimeter with the presence of a front detected from satellite products.

The strong surface density gradients (as defined in Section 2.3.2) represent 9% of the data measured by the TSG during the OUTPACE cruise. The comparison with FSLE reveals that 25% of the strong surface density gradients match with a physical front. Through a bootstrapping re-sampling, the same method is applied to the 91% of TSG data identified as non-gradient, and demonstrates that only $14 \pm 1\%$ of homogeneous density areas match with a front. These latter results also exhibit that an FSLE existence does not necessarily create a gradient but probably needs pre-existing tracer gradients and a lifetime longer than few days. The same calculations have also been performed for temperature and salinity gradients independently and show similar results. The relatively better correlation between density gradients and FSLEs than between no-density gradients and FSLEs, attests that gradients are not randomly distributed with regard to FSLE structure and proves that FSLE detection can be a good candidate to explain the presence of *in situ* surface gradients. Despite the effort to increase the altimetry resolution to $1/8^\circ$, it is still not enough to fully resolve the submesoscale gradients. Consequently the correlation between surface gradients and FSLE can not increase much higher than the 25% calculated here. This result converges with Hernández-Carrasco et al. (2011) who exhibited that FSLEs would still give an accurate picture of Lagrangian small-scale features despite some missing dynamics. Additionally, we perform a comparison between absolute values of gradients and co-located FSLE. Due to the lack of resolution of satellite products, this point by point comparison may induce a few kilometer offset between the area identified



with FSLE and the gradient sampled with the TSG, and consequently the method applied here may not identify the match. Therefore the methodology could be improved by focusing on FSLE values around a certain radius from the position of the gradient to eliminate the uncertainty caused by the absolute point by point comparison. Another way to improve the method would be to only take into account cross-front gradients that are more likely to be induced by a physical front than along-
5 front gradients. However, considering that the SST distribution is also governed by other processes than advection, such as the diurnal cycle, for example, the 25% correlation is large enough to sustain the idea that the submesoscale circulation can participate actively in the spatial structuring of surface tracers such as SSS, SST or density.

The same approach was performed with a reactive tracer, the chlorophyll concentration sampled with high frequency. It shows that 35% of strong surface chlorophyll gradients, representative of 1% of the entire underway sampling, agree with the
10 presence of FSLE. Re-sampling over the 99% of non-gradient areas, indicates that $28 \pm 14\%$ of homogeneous chlorophyll areas match with an FSLE. The high percentage of FSLEs matching with a «no chlorophyll-gradient» area gives little confidence on the fact that 35% of chlorophyll gradients were actually caused by the presence of a physical barrier. As chlorophyll concentration is driven by many biological processes, it may be more accurate to associate gradients of phytoplankton abundances, responsible for chlorophyll gradients, with small-scale features. Hereinafter, using two case studies of plankton high
15 frequency sampling, we propose to compare microbial abundances with frontal features.

3.4 Physical barriers' influences on phytoplankton community

To test the hypothesis of Bonnet et al. (2015), that pointed out the use of FSLE to explain some correlations between *Trichodesmium* spp. abundances and gradients, we measured the abundances of microbial groups of plankton in samples collected during two high-frequency sampling transects (Fig. 6 and 7). LDB high frequency sampling crosses from North to South the
20 bloom patch described in de Verneil et al. (2017b). The spatial distribution of organisms presents relatively high concentration of bacteria at the center of the bloom but decreases when exiting this feature (Fig. 6). The tip of an FSLE barrier is visible near the center of the transect. This barrier coincides with a salinity and temperature gradient (data not shown) associated with a sharp decrease in surface chlorophyll concentration and in PPE, HNA and LNA (bacteria in general) abundances (Fig. 6b). There is a sharp increase of DIP turnover time when exiting the patch in the south, indicating that Phosphorus is quickly
25 consumed inside the bloom. The front thus seems to create a barrier for organisms which grow and accumulate on one side of the front as demonstrated by relatively high surface chlorophyll (peak at $0.8 \mu\text{g l}^{-1}$) and low Phosphorus. According to Mann and Lazier (2013) phytoplankton growth may be stimulated in aggregates where they can easily take up nutrients released by bacterial decomposition of organic matter. This phenomenon may also support the persistency of the bloom during LDB. Indeed the bloom may be sustained in time by submesoscale features creating an aggregation of micro-organisms that can
30 benefit from each other. Aggregates can thus influence the community structure by creating favorable growing conditions for a species at the expense of others. A good example is LDA high-frequency sampling (Fig. 7a). Indeed we can notice a region of high abundance of bacteria at $165^\circ\text{E } 15'$ that is to be bounded by two FSLE barriers. This trend is confirmed by figure 7b which shows an increase of the abundance of *Prochlorococcus*, bacteria, HNA and LNA associated with a spike of FSLE values at 45 km. At the same time the abundance of PPE seems to decrease where bacterial abundances are the highest indicating



that this group may not find an advantage in these features. Another FSLE peak at 80 km was characterized by the decrease of *Prochlorococcus*, bacteria, HNA and LNA while the PPE abundance increases. The surface chlorophyll follows the same pattern as that of *Prochlorococcus*, bacteria, HNA and LNA with a relative increase of chlorophyll concentration ($0.3 \mu\text{g l}^{-1}$) within the region bounded by the FSLEs. Another peak in chlorophyll concentration is noticeable around 30 km ($0.4 \mu\text{g l}^{-1}$) but may be associated with other organisms than those analysed with cytometry. The physical fronts thus not only structure the spatial distribution of organisms by creating barriers but seems to create border regions influencing the community structure, abundances and diversity. Indeed modelled fine-scale structures have already been shown to delimit niches of different phytoplankton types but also to modify phytoplankton assemblages and diversity (d'Ovidio et al., 2010; Lévy et al., 2014a). Consistent with these previous numerical results, we therefore find out with *in situ* observations that microbial growth seems to benefit from the conditions engendered by frontal structures. It remains important to determine how these two study cases could be generalized for the organisms implied in the N_2 fixation cycle.

4 Conclusions

We document here the surface circulation at different spatial scales (from 1000 km to 10 km) and its influence on horizontal dispersal of biogeochemical components in the WTSP during the OUTPACE cruise. This study is conducted thanks to the combined use of value-added high-resolution altimetry products, *in situ* observations and Lagrangian numerical simulations. The total altimetry-derived velocity field, combining geostrophy and wind components, revealed a wind-driven meridional pathway of surface waters in the WTSP. This surface trajectory can be linked to the biogeochemical differences between Melanesian waters and gyre waters: a part of the water masses directly recirculates into the gyre whereas the other part is driven across the multiple islands of the WTSP.

The mesoscale activity is confirmed to be intense and mostly westward except in the region of transition between Melanesian waters and gyre waters (180°W - 170°W). Most of these mesoscale structures demonstrated ability to trap waters, however no obvious biogeochemical variations were linked to eddy entrainment. We identify two areas where the mesoscale circulation might have a strong influence on water mass transport. First, the central Coral Sea appears as a region of exchange between distinct NVJ and NCJ waters through eddy transport. Second, the band between 180°W and 170°W could emerge as a recurrent bloom formation area due to the simultaneous effect of N_2 fixation, well-known to sustain summertime blooms in the WTSP, and the eastward mesoscale transport of island-enriched waters.

Associating the surface small-scale gradients with Lagrangian diagnostics of frontal features, we showed a correlation of at least 25% highlighting the role of submesoscale activity in governing the horizontal dispersal of surface tracers. The small-scale features also participated in the horizontal distribution and community structure of phytoplankton patches sampled during two original high-frequency sampling of surface phytoplankton abundances.

Future studies in the area will need to take into account the interactions between physical features of the flow at large and fine-scale to better understand the phenomenon that drives the distribution of buoyant matter. In particular, the region around station LDB should be investigated during other bloom events to confirm the possible role of enriched-water mesoscale



transport in instigating/driving the bloom. This study also revealed the necessity to perform high-frequency sampling during oceanographic cruises to fully resolve submesoscale impacts on biogeochemical distributions.

Acknowledgements. This is a contribution of the OUTPACE (Oligotrophy from Ultra-oligoTrophy PACific Experiment) project (Moutin and Bonnet, 2015) funded by the French national research agency (ANR-14-CE01-0007-01), the LEFE-CyBER program (CNRS-INSU), the
5 GOPS program (IRD) and CNES (BC T23, ZBC 4500048836). Solange Duhamel was funded by the National Science Foundation (OCE-1434916). The authors thank the crew of the RV L'Atalante for outstanding shipboard operations, Gilles Rougier, Dominique Lefevre and Francesco d'Ovidio for their valuable help on different dataset. This work is supported by CLS in the framework of CNES funding and the authors would like to thank M-I Pujol and G. Taburet for providing enhanced satellite data.



References

- Alory, G., Delcroix, T., Téchiné, P., Diverrès, D., Varillon, D., Cravatte, S., Gouriou, Y., Grelet, J., Jacquin, S., Kestenare, É., et al.: The French contribution to the voluntary observing ships network of sea surface salinity, *dsrI*, 105, 1–18, 2015.
- Aminot, A. and Kérouel, R.: *Hydrologie des écosystèmes marins: paramètres et analyses*, Editions Quae, 2004.
- 5 Barbot, S., Petrenko, A., and Maes, C.: Specific floats behaviors in the southern tropical Pacific, this issue, 2017.
- Blanke, B. and Raynaud, S.: Kinematics of the Pacific Equatorial Undercurrent: an Eulerian and Lagrangian approach from GCM results, *J. Phys. Oceanogr.*, 27, 1038–1053, 1997.
- Blanke, B., Arhan, M., Madec, G., and Roche, S.: Warm water paths in the equatorial Atlantic as diagnosed with a general circulation model, *J. Phys. Oceanogr.*, 29, 2753–2768, 1999.
- 10 Bock, N., Dion, M., Van Wambeke, F., and Duhamel, S.: Microbial distribution across the Western Tropical South Pacific Ocean, this issue, 2017.
- Bonnet, S., Rodier, M., Turk-Kubo, K., Germineaud, C., Menkes, C., Ganachaud, A., Cravatte, S., Raimbault, P., Campbell, E., Desnues, A., Maes, C., and Eldin, G.: Contrasted geographical distribution of N₂ fixation rates and *nit*/H phylotypes in the Coral and Solomon Seas (South-Western Pacific) during austral winter conditions, *Global. Biogeo. Cyc.*, 2015.
- 15 Capet, X., McWilliams, J. C., Molemaker, M. J., and Shchepetkin, A.: Mesoscale to submesoscale transition in the California Current System. Part I: Flow structure, eddy flux, and observational tests, *jpo*, 38, 29–43, 2008.
- Chelton, D. B., Schlax, M. G., Samelson, R. M., and de Szoeke, R. A.: Global observations of large oceanic eddies, *Geophys. Res. Lett.*, 34, 2007.
- Church, M. J., Mahaffey, C., Letelier, R. M., Lukas, R., Zehr, J. P., and Karl, D. M.: Physical forcing of nitrogen fixation and diazotroph community structure in the North Pacific subtropical gyre, *Global Biogeochemical Cycles*, 23, 2009.
- 20 Dandonneau, Y., Vega, A., Loisel, H., Du Penhoat, Y., and Menkes, C.: Oceanic Rossby waves acting as a "hay rake" for ecosystem floating by-products, *Science*, 302, 1548–1551, 2003.
- de Verneil, A., Rousselet, L., Doglioli, A. M., Petrenko, A. A., Maes, C., Bouruet-Aubertot, P., and Moutin, T.: OUTPACE long duration stations: physical variability, context of biogeochemical sampling, and evaluation of sampling strategy, *Biogeosciences Discuss.*, 2017a.
- 25 de Verneil, A., Rousselet, L., Doglioli, A. M., Petrenko, A. A., and Moutin, T.: The fate of a south-west Pacific bloom: gauging the impact of submesoscale vs. mesoscale circulation on biological gradients in the subtropics, *Biogeosciences*, 14, 3471–3486, doi:10.5194/bg-14-3471-2017, <https://www.biogeosciences.net/14/3471/2017/>, 2017b.
- Dickey, T. D.: Emerging ocean observations for interdisciplinary data assimilation systems, *Journal of Marine Systems*, 40, 5–48, 2003.
- 30 Doglioli, A.: Software Package for Adaptive Sampling Strategy for Oceanographic cruises (SPASSO, <http://www.mio.univ-amu.fr/SPASSO>), 2013.
- d'Ovidio, F., Fernández, V., Hernández-García, E., and López, C.: Mixing structures in the Mediterranean Sea from finite-size Lyapunov exponents, *Geophys. Res. Lett.*, 31, 2004.
- d'Ovidio, F., Della Penna, A., Trull, T. W., Nencioli, F., Pujol, M.-I., Rio, M.-H., Park, Y.-H., Cotté, C., Zhou, M., and Blain, S.: The biogeochemical structuring role of horizontal stirring: Lagrangian perspectives on iron delivery downstream of the Kerguelen plateau, *Biogeosciences*, 12, 5567–5581, 2015.
- 35



- Ducet, N., Le Traon, P.-Y., and Reverdin, G.: Global high-resolution mapping of ocean circulation from TOPEX/Poseidon and ERS-1 and-2, *J. Geophys. Res.-O.*, 105, 19 477–19 498, 2000.
- d’Ovidio, F., De Monte, S., Alvain, S., Dandonneau, Y., and Lévy, M.: Fluid dynamical niches of phytoplankton types, *Proceedings of the National Academy of Sciences*, 107, 18 366–18 370, 2010.
- 5 d’Ovidio, F., De Monte, S., Della Penna, A., Cotté, C., and Guinet, C.: Ecological implications of eddy retention in the open ocean: a Lagrangian approach, *Journal of Physics A: Mathematical and Theoretical*, 46, 254 023, 2013.
- Fong, A. A., Karl, D. M., Lukas, R., Letelier, R. M., Zehr, J. P., and Church, M. J.: Nitrogen fixation in an anticyclonic eddy in the oligotrophic North Pacific Ocean, *The ISME journal*, 2, 663, 2008.
- Ganachaud, A., Gourdeau, L., and Kessler, W.: Bifurcation of the Subtropical South Equatorial Current against New Caledonia in December
10 2004 from a Hydrographic Inverse Box Model*, *J. Phys. Oceanogr.*, 38, 2072–2084, 2008.
- Ganachaud, A., Cravatte, S., Melet, A., Schiller, A., Holbrook, N., Sloyan, B., Widlansky, M., Bowen, M., Verron, J., Wiles, P., et al.: The Southwest Pacific Ocean circulation and climate experiment (SPICE), *J. Geophys. Res.-O.*, 119, 7660–7686, 2014.
- Garçon, V. C., Oschlies, A., Doney, S. C., McGillicuddy, D., and Waniek, J.: The role of mesoscale variability on plankton dynamics in the North Atlantic, *Deep Sea Research Part II: Topical Studies in Oceanography*, 48, 2199–2226, 2001.
- 15 Gasparin, F., Ganachaud, A., and Maes, C.: A western boundary current east of New Caledonia: Observed characteristics, *dsrI*, 58, 956–969, 2011.
- Guidi, L., Calil, P. H., Duhamel, S., Björkman, K. M., Doney, S. C., Jackson, G. A., Li, B., Church, M. J., Tozzi, S., Kolber, Z. S., et al.: Does eddy-eddy interaction control surface phytoplankton distribution and carbon export in the North Pacific Subtropical Gyre?, *Journal of Geophysical Research: Biogeosciences*, 117, 2012.
- 20 Hadjighasem, A. and Haller, G.: Level set formulation of two-dimensional Lagrangian vortex detection methods, *Chaos: An Interdisciplinary Journal of Nonlinear Science*, 26, 103 102, 2016.
- Haller, G., Hadjighasem, A., Farazmand, M., and Huhn, F.: Defining coherent vortices objectively from the vorticity, *Journal of Fluid Mechanics*, 795, 136–173, 2016.
- Hernández-Carrasco, I., López, C., Hernández-García, E., and Turiel, A.: How reliable are finite-size Lyapunov exponents for the assessment
25 of ocean dynamics?, *Ocean Modelling*, 36, 208–218, 2011.
- Hogg, A. M., Meredith, M. P., Chambers, D. P., Abrahamsen, E. P., Hughes, C. W., and Morrison, A. K.: Recent trends in the Southern Ocean eddy field, *Journal of Geophysical Research: Oceans*, 120, 257–267, 2015.
- Hristova, H. G., Kessler, W. S., McWilliams, J. C., and Molemaker, M. J.: Mesoscale variability and its seasonality in the Solomon and Coral Seas, *J. Geophys. Res.-O.*, 119, 4669–4687, 2014.
- 30 Jenkins, W.: Nitrate flux into the euphotic zone near Bermuda, 1988.
- Kessler, W. and Cravatte, S.: Mean circulation of Coral Sea, *J. Geophys. Res.-O.*, 118, 1–26, doi:10.1002/2013JC009117, 2013.
- Lévy, M., Iovino, D., Resplandy, L., Klein, P., Madec, G., Tréguier, A.-M., Masson, S., and Takahashi, K.: Large-scale impacts of sub-mesoscale dynamics on phytoplankton: Local and remote effects, *Ocean Modelling*, 43, 77–93, 2012.
- Lévy, M., Jahn, O., Dutkiewicz, S., and Follows, M. J.: Phytoplankton diversity and community structure affected by oceanic dispersal and
35 mesoscale turbulence, *Limnology and Oceanography: Fluids and Environments*, 4, 67–84, 2014a.
- Lévy, M., Resplandy, L., and Lengaigne, M.: Oceanic mesoscale turbulence drives large biogeochemical interannual variability at middle and high latitudes, *Geophysical Research Letters*, 41, 2467–2474, 2014b.



- Lévy, M., Jahn, O., Dutkiewicz, S., Follows, M. J., and d'Ovidio, F.: The dynamical landscape of marine phytoplankton diversity, *Journal of The Royal Society Interface*, 12, 20150481, 2015.
- Lumpkin, R. and Pazos, M.: Measuring surface currents with Surface Velocity Program drifters: the instrument, its data, and some recent results, *Lagrangian analysis and prediction of coastal and ocean dynamics*, pp. 39–67, 2007.
- 5 Maes, C., Gourdeau, L., Couvelard, X., and Ganachaud, A.: What are the origins of the Antarctic Intermediate Waters transported by the North Caledonian Jet?, *Geophys. Res. Lett.*, 34, 2007.
- Maes, C., Dewitte, B., Sudre, J., Garçon, V., and Varillon, D.: Small-scale features of temperature and salinity surface fields in the Coral Sea, *J. Geophys. Res.-O.*, 118, 1–13, doi:10.1002/jgrc.20344, 2013.
- Mahadevan, A. and Tandon, A.: An analysis of mechanisms for submesoscale vertical motion at ocean fronts, *Ocean Modelling*, 14, 241–256,
10 2006.
- Mann, K. H. and Lazier, J. R.: *Dynamics of marine ecosystems: biological-physical interactions in the oceans*, John Wiley & Sons, 2013.
- Martin, A. P., Richards, K. J., and Fasham, M. J.: Phytoplankton production and community structure in an unstable frontal region, *Journal of Marine Systems*, 28, 65–89, 2001.
- Matear, R., Chamberlain, M., Sun, C., and Feng, M.: Climate change projection of the Tasman Sea from an Eddy-resolving Ocean Model,
15 *Journal of Geophysical Research: Oceans*, 118, 2961–2976, 2013.
- McDougall, T., Jackett, D., Millero, F., Pawlowicz, R., and Barker, P.: A global algorithm for estimating Absolute Salinity, *Ocean Sci.*, 8, 1123–1134, 2012.
- McGillicuddy, D. and Robinson, A.: Eddy-induced nutrient supply and new production in the Sargasso Sea, *Deep Sea Research Part I: Oceanographic Research Papers*, 44, 1427–1450, 1997.
- 20 McGillicuddy Jr, D. J.: Mechanisms of physical-biological-biogeochemical interaction at the oceanic mesoscale, 2016.
- Moutin, T. and Bonnet, S.: OUTPACE cruise, RV L'Atalante, doi:10.17600/15000900, 2015.
- Moutin, T., Doglioli, A., De Verneil, A., and Bonnet, S.: The Oligotrophy to the UTRa-oligotrophy PACific Experiment (OUTPACE cruise, Feb. 18 to Apr. 3, 2015), *Biogeosciences Discussions*, 2017, 1–23, doi:10.5194/bg-2017-50, <http://www.biogeosciences-discuss.net/bg-2017-50/>, 2017.
- 25 Okubo, A.: Horizontal dispersion of floatable particles in the vicinity of velocity singularities such as convergences, in: *Deep sea research and oceanographic abstracts*, vol. 17, pp. 445–454, Elsevier, 1970.
- Oschlies, A. and Garçon, V.: Eddy-induced enhancement of primary production in a model of the North Atlantic Ocean, *Nature*, 394, 266, 1998.
- Penven, P., Halo, I., Pous, S., and Marié, L.: Cyclogeostrophic balance in the Mozambique Channel, *Journal of Geophysical Research: Oceans*, 119, 1054–1067, 2014.
30
- Perruche, C., Rivière, P., Lapeyre, G., Carton, X., and Pondaven, P.: Effects of surface quasi-geostrophic turbulence on phytoplankton competition and coexistence, *Journal of marine research*, 69, 105–135, 2011.
- Petrenko, A. A., Doglioli, A. M., Nencioli, F., Kersalé, M., Hu, Z., and d'Ovidio, F.: A review of the LATEX project: mesoscale to submesoscale processes in a coastal environment, *Ocean Dynamics*, 67, 513–533, 2017.
- 35 Qiu, B. and Chen, S.: Seasonal modulations in the eddy field of the South Pacific Ocean, *Journal of Physical Oceanography*, 34, 1515–1527, 2004.
- Qiu, B., Chen, S., and Kessler, W. S.: Source of the 70-Day Mesoscale Eddy Variability in the Coral Sea and the North Fiji Basin*, *J. Phys. Oceanogr.*, 39, 404–420, 2009.



- Reynolds, R. W., Smith, T. M., Liu, C., Chelton, D. B., Casey, K. S., and Schlax, M. G.: Daily high-resolution-blended analyses for sea surface temperature, *Journal of Climate*, 20, 5473–5496, 2007.
- Rio, M.-H., Mulet, S., and Picot, N.: Beyond GOCE for the ocean circulation estimate: Synergetic use of altimetry, gravimetry, and in situ data provides new insight into geostrophic and Ekman currents, *Geophysical Research Letters*, 41, 8918–8925, 2014.
- 5 Rogé, M., Morrow, R. A., and Dencausse, G.: Altimetric Lagrangian advection to reconstruct Pacific Ocean fine-scale surface tracer fields, *Ocean Dynam.*, 65, 1249–1268, 2015.
- Rousselet, L., Doglioli, A., Maes, C., Blanke, B., and Petrenko, A.: Impacts of mesoscale activity on the water masses and circulation in the Coral Sea, *J. Geophys. Res.*, 121, 7277–7289, 2016.
- Sarmiento, J. and Gruber, N.: *Ocean Biogeochemical Dynamics*, 2006.
- 10 Sokolov, S. and Rintoul, S.: Circulation and water masses of the southwest Pacific: WOCE section P11, Papua New Guinea to Tasmania, *J. Mar. Res.*, 58, 223–268, 2000.
- Sweeney, E. N., McGillicuddy, D. J., and Buesseler, K. O.: Biogeochemical impacts due to mesoscale eddy activity in the Sargasso Sea as measured at the Bermuda Atlantic Time-series Study (BATS), *Deep Sea Research Part II: Topical Studies in Oceanography*, 50, 3017–3039, 2003.
- 15 Tomczak, M. and Godfrey, J. S.: *Regional oceanography: an introduction*, Elsevier, 2013.
- Webb, D.: Evidence for shallow zonal jets in the South Equatorial Current region of the southwest Pacific, *J. Phys. Oceanogr.*, 30(4), 2000.
- Weiss, J.: The Dynamics of enstrophy transfer in two-dimensional hydrodynamics, La Jolla Inst. La Jolla, CA, Tech. rep., LJI-TN-81-121, 1981.
- Williams, R. G. and Follows, M. J.: The Ekman transfer of nutrients and maintenance of new production over the North Atlantic, *Deep Sea Research Part I: Oceanographic Research Papers*, 45, 461–489, 1998.
- 20

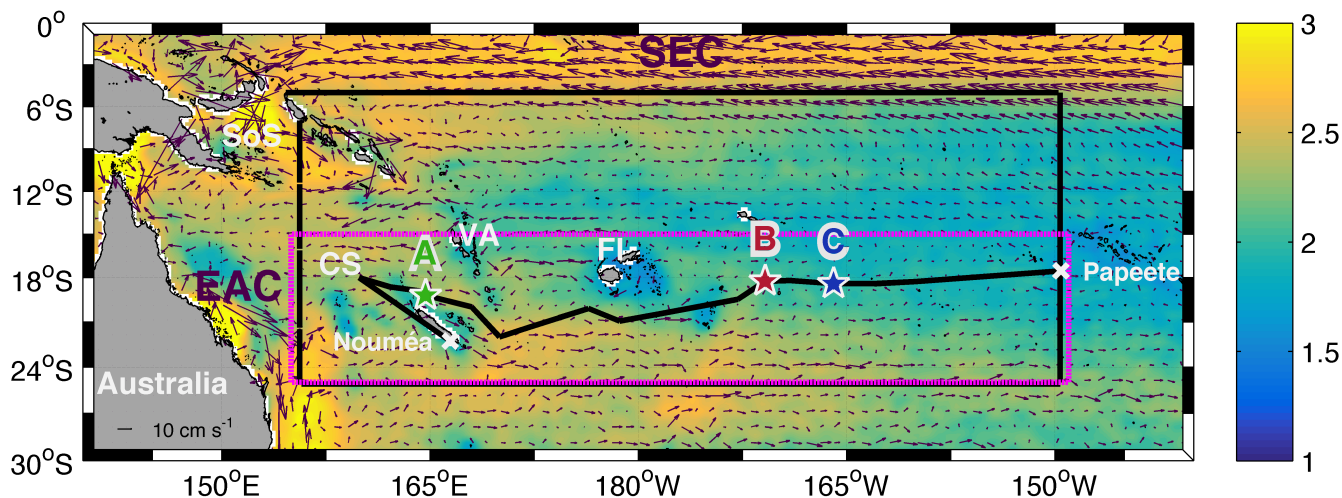


Figure 1. Mean eddy kinetic energy (log scale, colorbar) with surface velocity (arrows in cm s^{-1}) computed for 10 years (2005 - 2015) from satellite-derived altimetry. The South Equatorial Current (SEC) and the western boundary East Australian Current (EAC) are indicated. The black line shows the ship track during OUTPACE from Nouméa to Papeete. The positions of the three Long-Duration (LD) stations are drawn with green, red and blue stars for LD-A, LD-B and LD-C respectively. SoS: Salomon Sea; CS: Coral Sea; VA: Vanuatu, FI: Fiji.

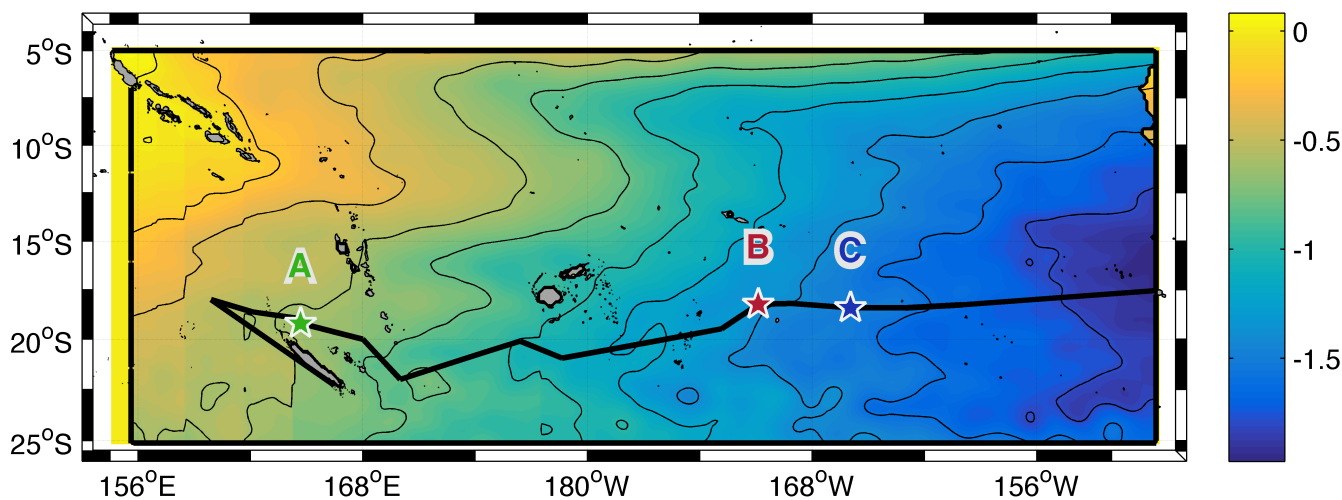


Figure 2. Total transport (Sv, colorbar) and streamfunction (black lines with contour interval of 0.25 Sv) computed for ten years with the velocity field combining geostrophy and Ekman components with correction for cyclogeostrophy. The ship track and locations of OUTPACE LD stations are indicated with the black line and colored stars as in Figure 1.

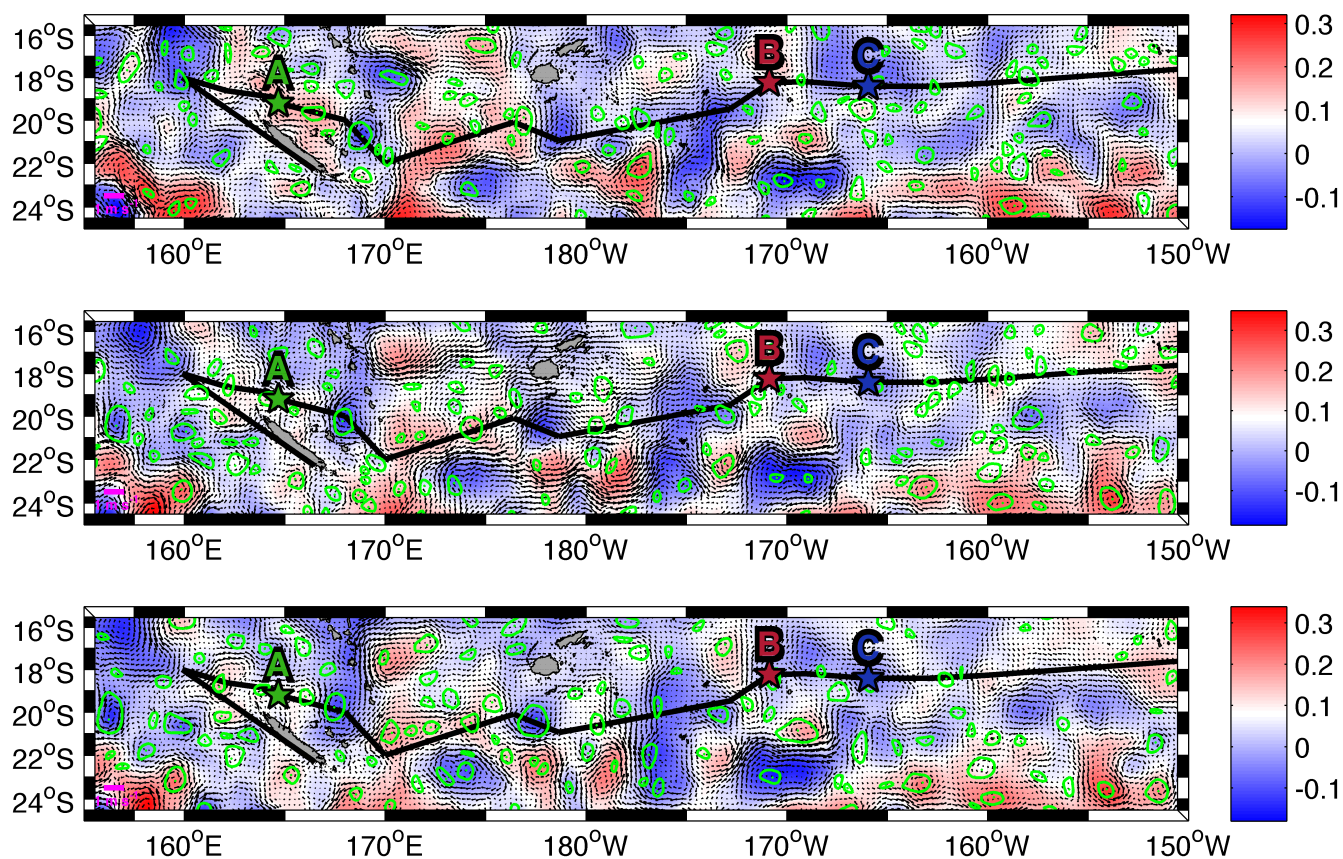


Figure 3. Daily sea surface level anomaly [m, colorbar] and velocity field [m s^{-1}] from the geostrophy, Ekman and cyclogeostrophy included product for the first day of LDA (February 25, top), LDB (March 15, center) and LDC (March 23, bottom). Contours of LAVD detected structures are drawn in green. The center of the structures is also indicated by a green point. The ship track and locations of OUTPACE LD stations are indicated with the black line and colored stars as in Figure 1.

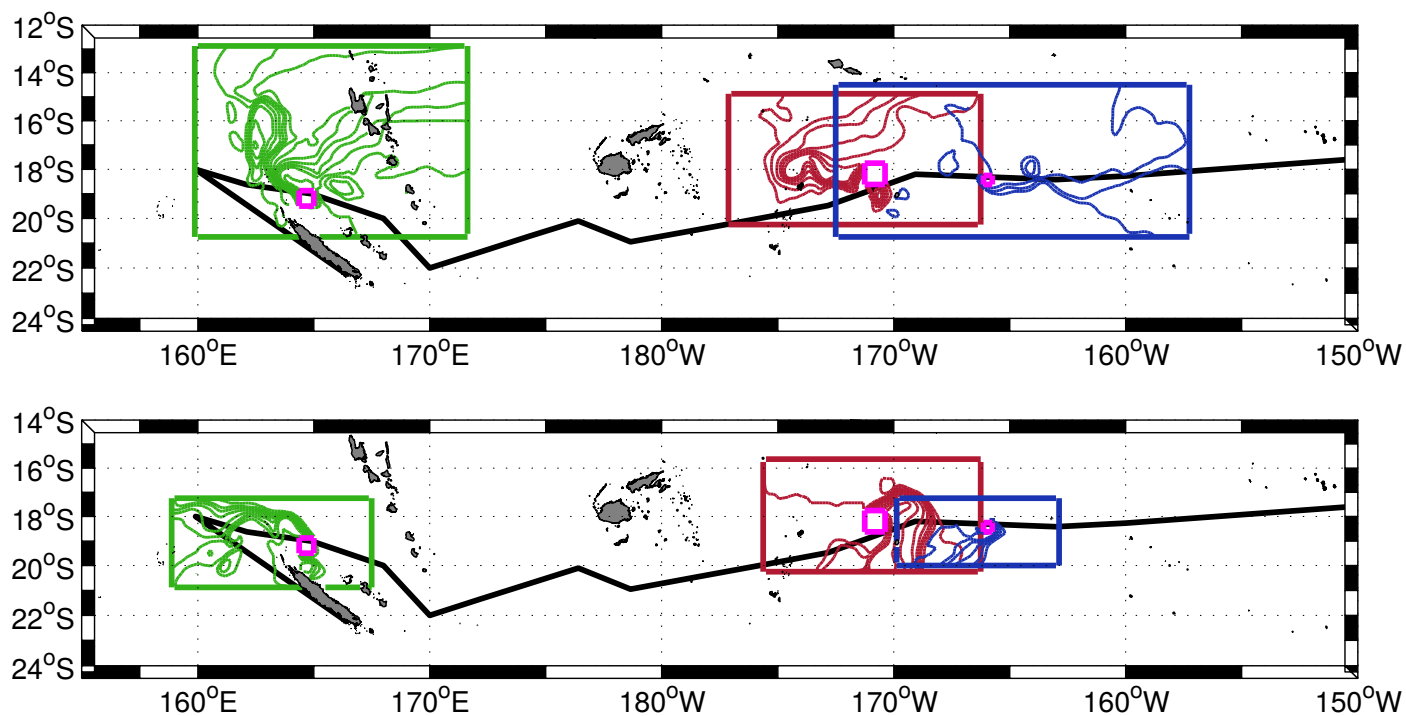


Figure 4. Backward (top) and forward (bottom) streamfunctions for LDA (green lines), LDB (red lines) and LDC (blue lines). Numerical particles are initially launched on the magenta boxes which represent the position of each LD station. The domain limit of each Ariane Lagrangian analysis are shown by the large green, red and blue boxes respectively. The ship track of OUTPACE is indicated with the black line.

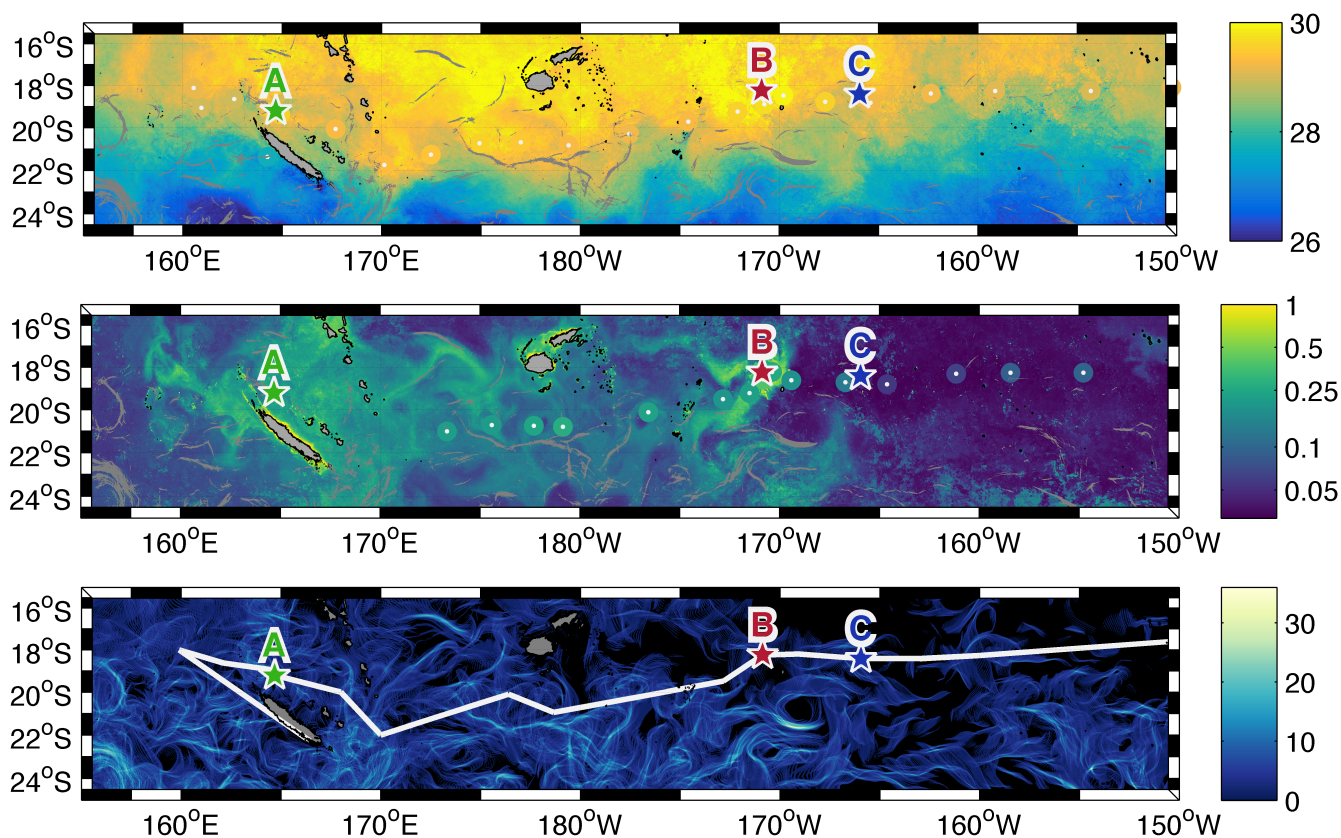


Figure 5. Top: Lagrangian satellite sea surface temperature ($^{\circ}\text{C}$) (for the time period of the OUTPACE cruise, adapted from de Verneil et al. (2017b)) from CLS superimposed with sea surface temperature ($^{\circ}\text{C}$) from TSG (colored circles with centres indicated in white). Center: Lagrangian satellite-derived surface chlorophyll concentration (mg m^{-3}) (for the time period of the OUTPACE cruise, adapted from de Verneil et al. (2017b)) from CLS superimposed with surface chlorophyll concentration (mg m^{-3}) measured onboard during OUTPACE (colored circles with centres indicated in white). Fronts present for at least 10 days during the OUTPACE cruise are indicated in gray in both figures. Bottom: Recurrence of FSLE structures (number of days of FSLE presence, colorbar).

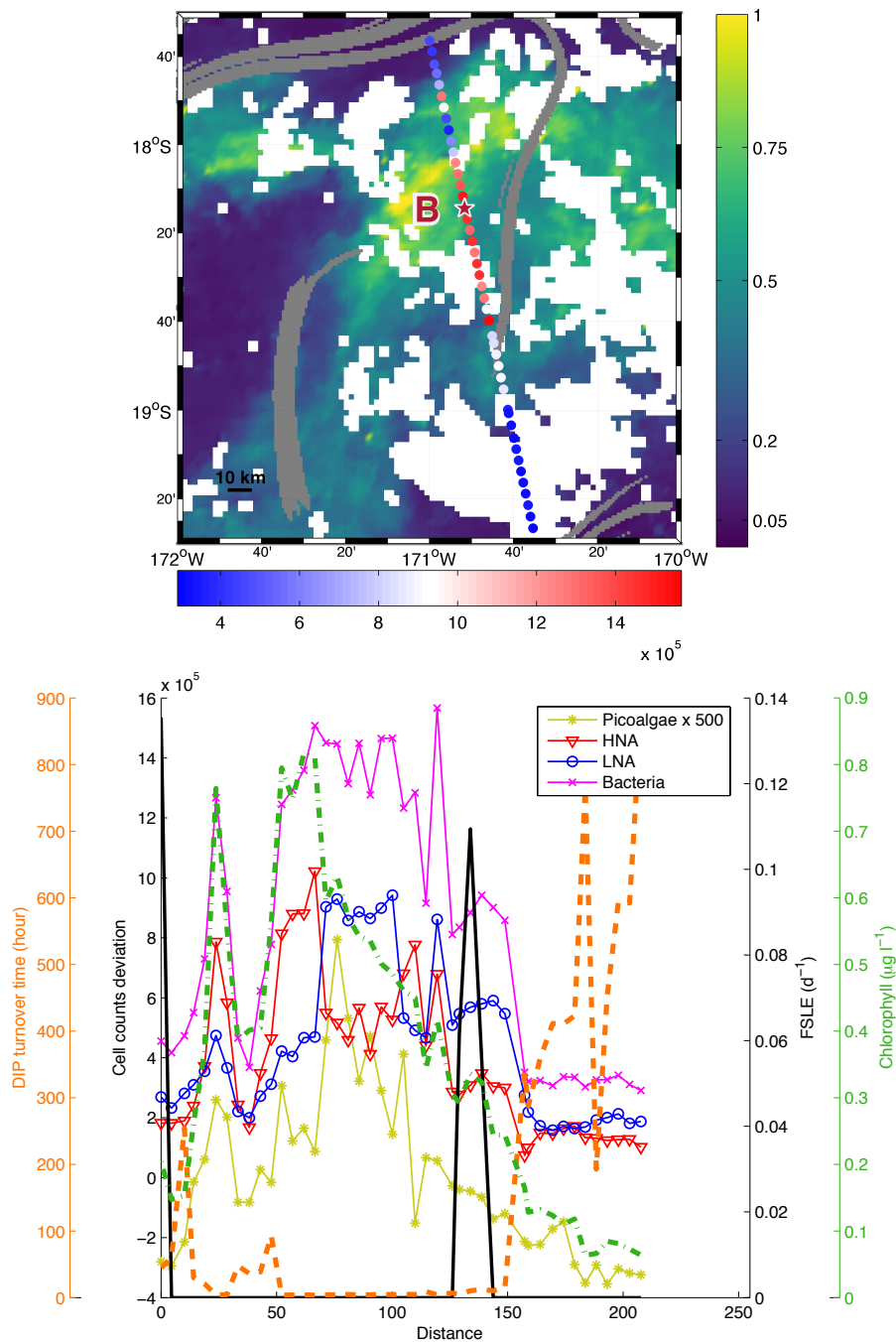


Figure 6. Top panel: chlorophyll concentration (mg m^{-3} , colorbar) from March 13, 2015 superimposed with bacteria counts (cell ml^{-1} , red-blue colorbar) sampled during LDB. FSLE fronts (values $> 0.05 \text{ day}^{-1}$) are shown in gray. The location of LDB is indicated with the red star. Bottom panel: Cell counts deviation from mean of PPE and bacteria (HNA and LNA) superimposed with FSLE (day^{-1} , black line), surface chlorophyll concentration ($\mu\text{g l}^{-1}$, dashed green line) and dissolved inorganic phosphate turnover time (hours, dashed orange line) along the high frequency transect of LDB.

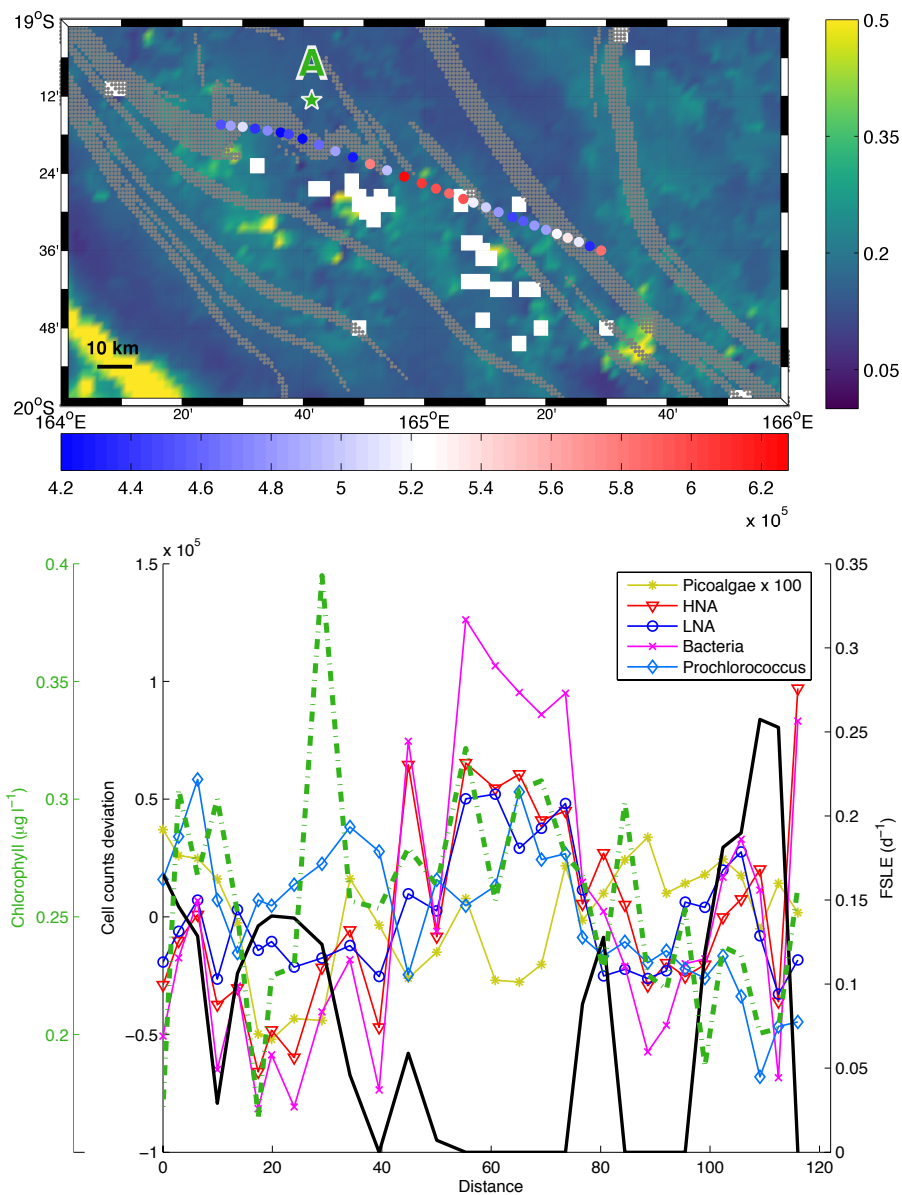


Figure 7. Top panel: chlorophyll concentration (mg m^{-3} , colorbar) from March 3, 2015 superimposed with bacteria counts (cell ml^{-1} , red-blue colorbar) sampled during LDA. FSLE fronts (values $> 0.05 \text{ day}^{-1}$) are shown in gray. The location of LDA is indicated with the green star. Bottom panel: Cell counts deviation of *Prochlorococcus*, PPE and bacteria (HNA and LNA) superimposed with FSLE values (day^{-1} , black line) and surface chlorophyll concentration ($\mu\text{g l}^{-1}$, dashed green line) along the high frequency transect of LDA.



Table A1. Statistics of meanders (see in the text for definition) and particles lost during each backward and forward lagrangian experiments near LD stations.

STATION	LDA		LDB		LDC	
	Backward	Forward	Backward	Forward	Backward	Forward
Meanders	39%	26%	65%	18%	70%	44%
Particle lost	22%	34%	2%	18%	3%	2%

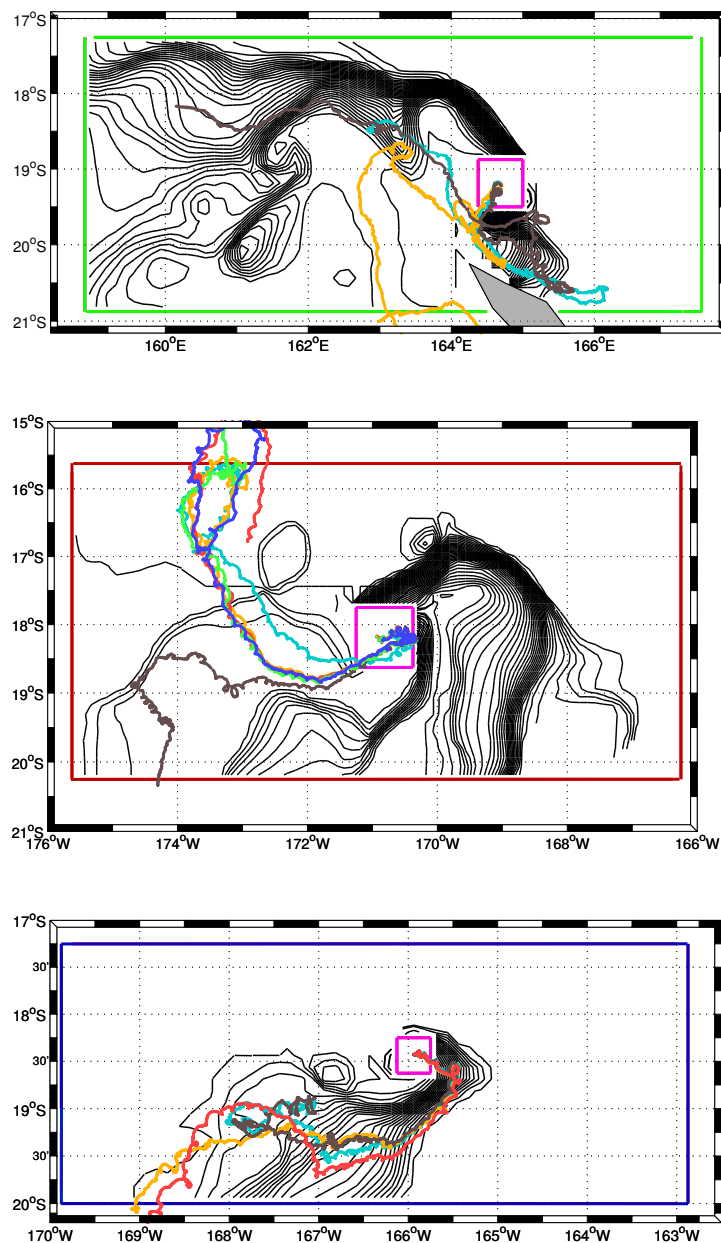


Figure A1. Forward streamfunctions for LDA (top panel), LDB (middle panel) and LDC (bottom panel) superimposed with SVP trajectories (colored lines) launched during each LD station. Numerical particles are initially launched on the magenta boxes (roughly in the center) which represent the position of each LD station. The domain limit of each Ariane Lagrangian analysis are shown by the large green (LDA), red (LDB) and blue (LDC) boxes, respectively.

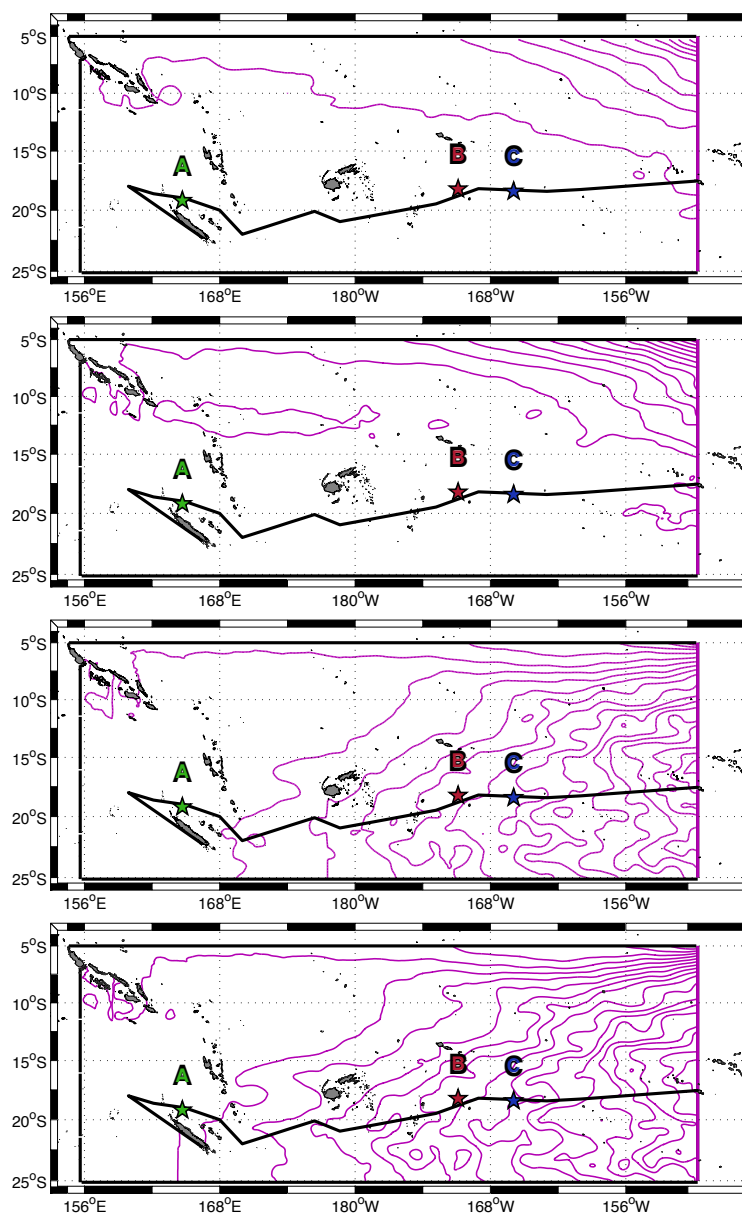


Figure B1. Forward streamfunctions computed for a ten year period with, from top to bottom: the low resolution geostrophic product of AVISO; the high resolution geostrophic product from CLS; the high resolution geostrophic and Ekman (at 15 m) product from CLS; and the high resolution geostrophic, Ekman (at 15 m) and cyclogeostrophic product from CLS (referred as the total altimetry-derived product in the text). The initial section of Ariane Lagrangian analysis is indicated with a purple vertical line to the east. The ship track and locations of OUTPACE LD stations are indicated with the black line and colored stars as referred to in Figure 1.

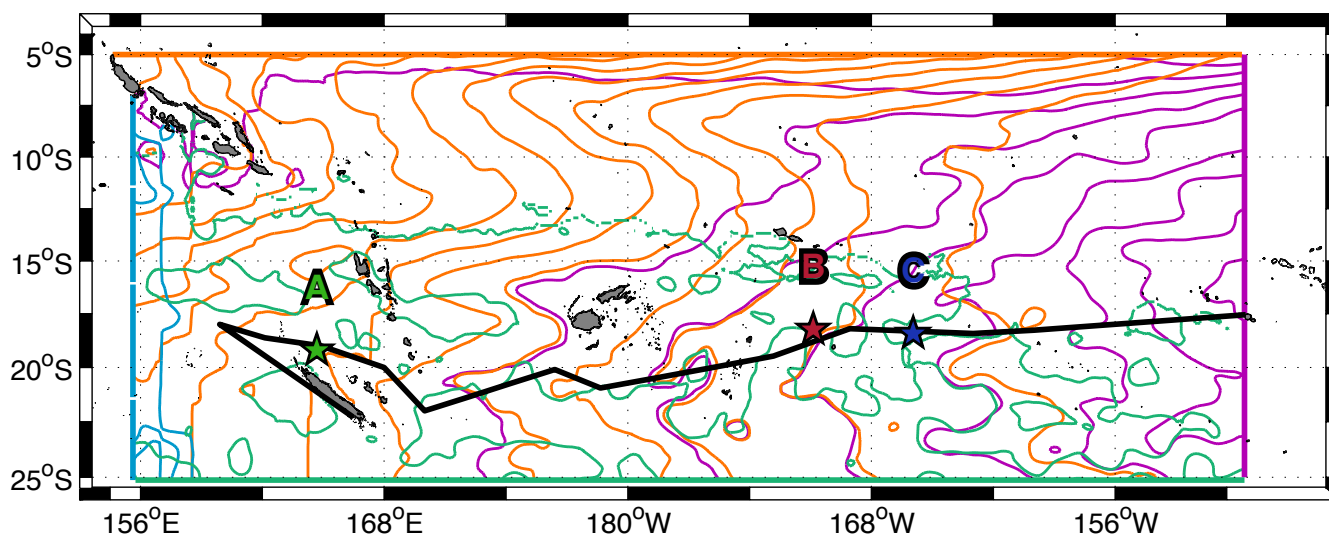


Figure C1. Forward streamfunctions computed for a ten year period with the total altimetry-derived product from CLS. Each streamline's color corresponds to the initial section of numerical particles: North section (orange lines), East section (purple lines), South section (green lines) and West section (magenta lines). The ship track and locations of OUTPACE LD stations are indicated with the black line and colored stars as in Figure 1.

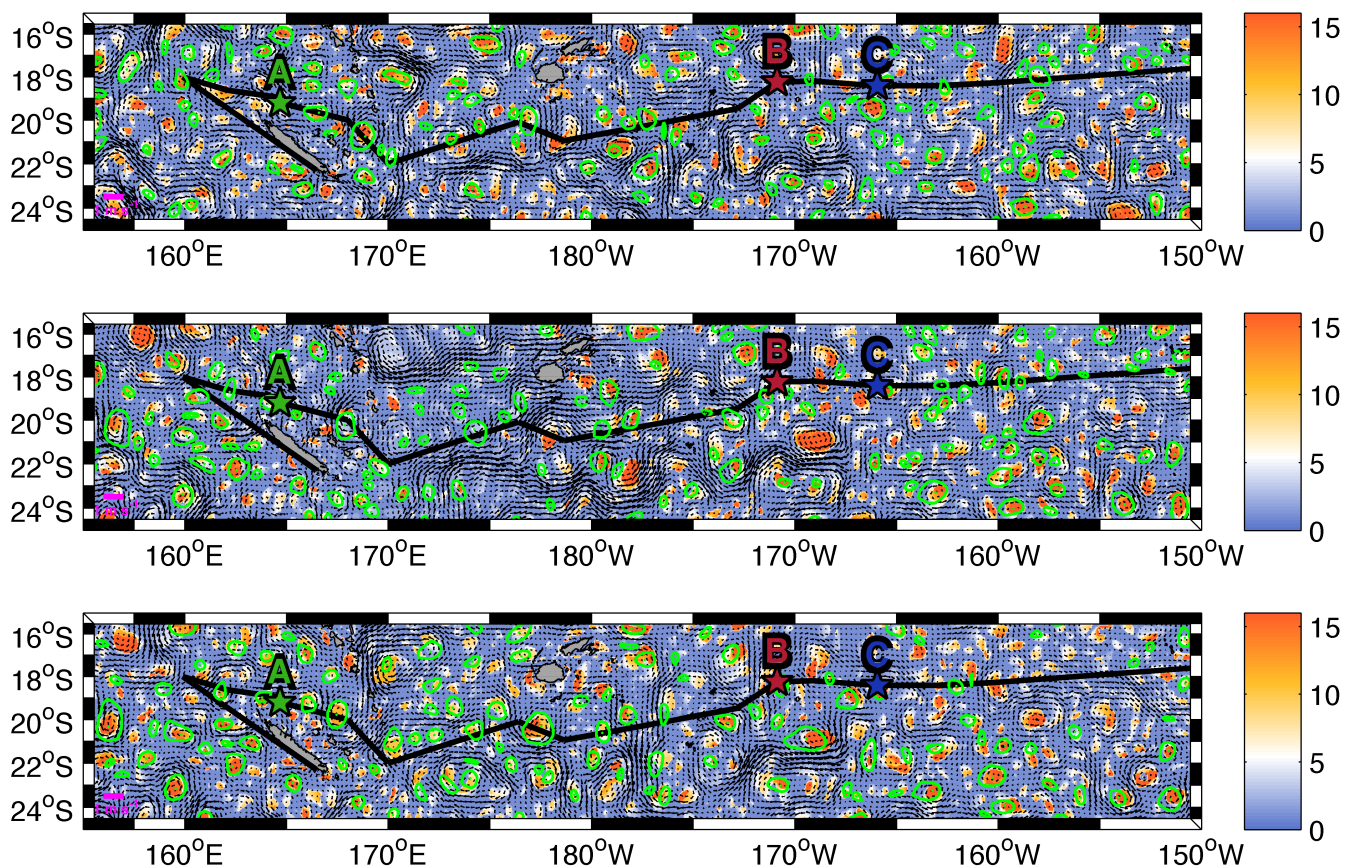


Figure C2. Particle retention time [days, colorbar] and velocity field [m s^{-1}] derived from the geostrophy, Ekman and cyclogeostrophy included product for the first day of LDA (February 25, top), LDB (March 15, center) and LDC (March 23, bottom). Contours of LAVD detected structures are drawn in green. The center of the each structure is marked with a green point. The ship track and locations of OUTPACE LD stations are indicated with the black line and colored stars as in Figure 1.