



Anthropogenic CO₂, air-sea CO₂ fluxes and acidification in the Southern 1 2 Ocean: results from a time-series analysis at station OISO-KERFIX (51°S-68°E). 3 4 Nicolas Metzl¹, Claire Lo Monaco¹, Coraline Leseurre^{1,2}, Céline Ridame¹, Gilles Reverdin¹, 5 Thi Tuyet Trang Chau³, Frédéric Chevallier³, Marion Gehlen³ 6 7 8 ¹ Laboratoire LOCEAN/IPSL, Sorbonne Université-CNRS-IRD-MNHN, Paris, 75005, France 9 ² Flanders Marine Institute (VLIZ), 8400 Ostend, Belgium 10 ³ Laboratoire LSCE/IPSL, CEA-CNRS-UVSQ, Université Paris-Saclay Gif-sur-Yvette, 91191, France 11 12 Correspondence to: Nicolas Metzl (nicolas.metzl@locean.ipsl.fr) 13 14 Abstract: The temporal variation of the carbonate system, air-sea CO₂ fluxes and pH is analyzed in the Southern 15 Indian Ocean, south of the Polar Front, based on in-situ data obtained from 1985 to 2021 at a fixed station 16 $(50^{\circ}40'S-68^{\circ}25'E)$ and results from a neural network model that reconstructs the fugacity of CO₂ (fCO₂) and 17 fluxes at monthly scale. Anthropogenic CO2 (Cant) was estimated in the water column and detected down to the 18 bottom (1600m) in 1985 resulting in an aragonite saturation horizon at 600m that migrated up to 400m in 2021 19 due to the accumulation of C_{ant} . In subsurface, the trend of C_{ant} is estimated at +0.53 (±0.01) µmol.kg⁻¹.yr⁻¹ with a 20 detectable increase in recent years. At the surface during austral winter the oceanic fCO₂ increased at a rate close 21 or slightly lower than in the atmosphere. To the contrary, in summer, we observed contrasting fCO_2 and 22 dissolved inorganic carbon (C_T) trends depending on the decade and emphasizing the role of biological drivers 23 on air-sea CO₂ fluxes and pH inter-annual variability. The region moved from an annual source of 0.8 molC.m 24 2 .yr⁻¹ in 1985 to a sink of -0.5 molC.m⁻².yr⁻¹ in 2020. In 1985-2020, the annual pH trend in surface of -0.0165 (\pm 25 0.0040).decade⁻¹ was mainly controlled by anthropogenic CO₂ but the trend was modulated by natural processes. 26 Using historical data from November 1962 we estimated the long-term trend for fCO₂, C_T and pH confirming 27 that the progressive acidification was driven by atmospheric CO₂ increase. In 59 years this leads to a diminution 28 of 11% for both aragonite and calcite saturation state. As atmospheric CO₂ will desperately continue rising in the 29 future, the pH and carbonate saturation state will decrease at a faster rate than observed in recent years. A 30 projection of future C_T concentrations for a high emission scenario (SSP5-8.5) indicates that the surface pH in 31 2100 would decrease to 7.32 in winter. This is up to -0.86 lower than pre-industrial pH and -0.71 lower than pH 32 observed in 2020. The aragonite under-saturation in surface waters would be reached as soon as 2050 (scenario 33 SSP5-8.5) and 20 years later for a stabilization scenario (SSP2-4.5) with potential impacts on phytoplankton 34 species and higher trophic levels in the rich ecosystems of the Kerguelen Island area. 35 36 Keywords: Ocean Carbonate System, Ocean acidification, anthropogenic CO₂, air-sea CO₂ fluxes, Southern

- 37 Ocean. Time-series station
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40 1 Introduction

41 The ocean plays an important role in mitigating climate change by taking up since decades a large part 42 of the excess of heat (Cheng et al., 2020; Fox-Kemper et al., 2021) and of CO₂ released by human activities 43 (Sabine et al., 2004; Gruber et al., 2019a; Canadell et al., 2021). Since 1750, the global ocean has captured 185 44 (± 35) PgC (Petagramm of Carbon) from a total of 700 (± 75) PgC of anthropogenic carbon emissions from 45 fossils fuels and land-used changes (Friedlingstein et al., 2022). From year to year, the ocean anthropogenic CO₂ sink increased progressively from 1.1 (\pm 0.4) PgC.yr⁻¹ in the 1960s to 2.3 (\pm 0.4) PgC.yr⁻¹ in the 2000s. Over the 46 47 decade 2012-2021, the partitioning of the anthropogenic CO₂ sinks was roughly equal between the ocean (2.9 \pm 48 0.4 PgC.yr⁻¹) and the land $(3.1 \pm 0.6 \text{ PgC.yr}^{-1})$ (Friedlingstein et al., 2022).

49 Ocean observations indicate that since the 1990s the Southern Ocean (SO) south of 45°S has been accumulating each year about 0.5 PgC.yr⁻¹ (e.g. Takahashi et al., 2009; Lenton et al., 2013; Rödenbeck et al., 50 51 2013; Long et al., 2021; Fay et al, 2023; Gray, 2024). Results based on BGC-Argo floats (Southern Ocean 52 Carbon and Climate Observations and Modeling project, SOCCOM) suggest that the CO₂ sink in the SO might 53 be much lower (0.16 PgC.yr⁻¹ south of 44°S for the period 2015-2017, Gray et al. 2018; Bushinsky et al., 2019) 54 but there is an ongoing debate (Long et al., 2021; Sutton et al., 2021; Gray, 2024). It is also well established that 55 the CO₂ sink in the SO undergoes substantial decadal variability first documented for the 1990s (Le Quéré et al., 56 2007; Metzl, 2009; Lenton et al., 2013) and subsequently identified for the period 1982-2018 (Landschützer et 57 al., 2015; Keppler and Landschützer, 2019; Mackay et al., 2022; Hauck et al., 2023). A recent extension of the 58 period to 1957-2020 suggests that the inter-annual to decadal variability of the SO CO₂ sink was most pronounced after the 1980s (Rödenbeck et al., 2022; Bennington et al., 2022). Whatever the variability of the SO 59 60 CO_2 sink since the 1960s, the ocean continuously absorbs atmospheric CO_2 and the distribution of anthropogenic 61 CO2 (Cant) in the SO is now relatively well documented (e.g. Pardo et al., 2014; Gruber et al., 2019a) thanks to 62 the GLODAP data synthesis effort for the global ocean (Global Ocean Data Analysis Project, Olsen et al., 2016, 63 2019, 2020). The SO takes up about 40% of the total anthropogenic carbon that enters the ocean (Khatiwala et 64 al., 2013; Gruber et al., 2019a).

65 The anthropogenic CO₂ uptake in the ocean results is lowering carbonate ion concentrations and pH, a 66 chemical process termed "ocean acidification" (OA) (Caldeira and Wickett 2003; Doney et al., 2009). This 67 decreases the saturation state with respect to carbonate minerals (aragonite, Ω ar and calcite, Ω ca), a process most pronounced in the cold and naturally at a low saturation state waters in high latitudes (Orr et al., 2005; Takahashi 68 69 et al., 2014; Jiang et al., 2015). The first estimate of Cant distribution in the global ocean (for a nominal year 70 1994, Sabine et al., 2004) shows that C_{ant} uptake led to an upward migration of the Ωar and Ωca saturation 71 horizon in all ocean basins (Feely et al., 2004). This change is particularly pronounced south of the Polar Front 72 (PF) in the SO linked to both Cant uptake and the upwelling of dissolved inorganic carbon (C_T) C_T-rich deep 73 waters (e.g. Hauck et al., 2010; Pardo et al., 2017). It has been suggested, through numerical studies, that 74 depending on future CO₂ emission levels, surface waters could reach under-saturation state for aragonite by 75 2030-2050 in the SO (Orr et al., 2005; Gangstø et al., 2008; McNeil and Matear, 2008; Negrete-Garcia et al., 76 2019). Such a change would have multiple and detrimental impacts on marine ecosystems (Fabry et al., 2008; 77 Doney et al., 2012; Bopp et al., 2013), in particular calcifying marine organisms especially aragonite producers 78 such as pteropods (Hunt et al., 2008: Gardner et al., 2023), but also calcite producing planktonic foraminifera (Moy et al., 2009), coccolithophorids (Beaufort et al., 2011), and non-calcifying species such as the abundant SO 79





diatoms (e.g. Benoiston et al., 2017; Petrou et al., 2019; Weir et al., 2020; Duncan et al., 2022) and krill
(Kawaguchi et al., 2013).

82 Hindcast simulations with Global Ocean Biogeochemical Models (GOBM), as well as projections with 83 Earth System Models (ESM) have been used to evaluate the ocean carbon cycle over the past decades and future 84 changes in Cant storage, ocean acidification or impacts of global change on marine ecosystems. However, current 85 model-based estimates of the contemporary SO CO₂ sink are subject to relatively large uncertainties (e.g. Long 86 et al., 2013; Hauck et al., 2020; Gooya et al., 2023; Hauck et al., 2023; Mayot et al., 2023; DeVries et al, 2023). Difference between GOBM models can reach 0.7 PgC.yr⁻¹ in the SO (Hauck et al., 2020), which is roughly 87 equivalent to the mean climatological flux of 0.5 PgC.yr⁻¹ (McNeil et al., 2007; Takahashi et al., 2009; Lenton et 88 89 al., 2013). In the high latitudes of the SO (> 50°S) for the 2010s, ESM from the Coupled Model Intercomparison 90 Project Phase 6 (CMIP6) simulated either a large sink or a modest source of CO₂ (McKinley et al, 2023). This is 91 mainly due to incorrect or missing physical and/or biological processes in the models (e.g. Pilcher et al., 2015; 92 Kessler and Tjiputra, 2016; Mongwe et al., 2018; Lerner et al., 2021) leading to biases in the seasonality of 93 temperature, C_T , partial pressure of CO₂ (pCO₂), air-sea CO₂ fluxes, pH or Ω (e.g. McNeil and Sasse 2016; 94 Rodgers et al., 2023; Rustogi et al., 2023; Joos et al., 2023). Such model imperfections should be resolved to 95 have future projections of CO_2 uptake, OA, productivity and the responses of the marine ecosystems, gain in 96 reliability (Frölicher et al., 2015; Hauck et al., 2015; Sasse et al., 2015; Kessler and Tjiputra, 2016; McNeil and 97 Sasse 2016; Kwiatkowski and Orr, 2018; Negrete-Garcia et al., 2019; Burger et al., 2020; Terharr et al., 2021; 98 Krumhardt et al., 2022; Jiang et al., 2023; Mongwe et al., 2023). In this context, as often concluded in modeling 99 studies (e.g. Kessler and Tjiputra, 2016; Gooya et al., 2023; Wright et al., 2023; Hauck et al., 2023; Mayot et al., 100 2023; Rodgers et al., 2023), long-term biogeochemical observations are particularly valuable to quantify and 101 understand recent past and current changes, and ultimately evaluate model simulations.

102 Although the SO south of the Polar Front remains much less observed than other oceanic regions, 103 several observations based studies have allowed to estimate the decrease in pH in the surface waters in response 104 to the increase in oceanic CO₂ fugacity, fCO₂ (Mirodikwa et al., 2012; Takahashi et al., 2014; Lauvset et al., 105 2015; Munro et al., 2015; Xue et al., 2018; Iida et al., 2021; Leseurre et al., 2022; Brandon et al., 2022). Results showed a large range of the pH trends from -0.008.decade⁻¹ to -0.035.decade⁻¹ depending on the period and 106 107 regions of interest. Most of these analyses were based on summer observations (Table 1) and some studies highlighted contrasting pH trends on a 5-10 year time probably linked to large scale climate variability such as 108 109 the Southern Annular Mode (SAM) (e.g. Xue et al., 2018). Given such variability, it is important to continue 110 monitoring fCO₂ and pH trend and, if possible, at different seasons as future change in CO₂ uptake and potential 111 tipping points of that carbonate saturation state also depends on seasonality (Sasse et al., 2015). The above 112 observational studies were dedicated to pH changes in surface waters. In contrast to Northern high latitudes (e.g. Olafsson et al., 2009, 2010; Franco et al., 2021; Skjelvan et al., 2022) few studies in the SO attempted to 113 114 evaluate decadal changes of carbonate system properties and acidification in the water column based on time-115 series stations. These changes in the SO water column were investigated from data collected during cruises 116 generally 3 to 15 years apart (Hauck et al., 2010; Van Heuven et al., 2011; Pardo et al., 2017; Tanhua et al., 117 2017; Carter et al., 2019).





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Table 1: Trends of oceanic fCO₂ (µatm.yr⁻¹) and pH (decade⁻¹) in the Southern Ocean south of the Polar Front
 based on observations and from this study. IO: Indian Ocean sector. PO: Pacific Ocean sector. AO: Atlantic
 Ocean sector. SO SPSS: Southern Ocean SubPolar Seasonally Stratified biome (around 50-60°S). PZ: Polar
 Zone. NR: Not Reported. Standard-deviations when available are in bracket.

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126	Period	Season	Zone	Trend fCO ₂	Trend pH	Reference
127				µatm.yr ⁻¹	decade ⁻¹	
128						
129	1991-2000	Summer	IO PZ 55-60°S	2.93	-0.035	Xue et al (2018)
130	2001-2011	Summer	IO PZ 55-60°S	1.41	-0.016	Xue et al (2018)
131						
132	2005-2019	Summer	IO PZ 54-64°S	NR	-0.026(0.003)	Brandon et al (2022)
133						
134	1998-2019	Summer	IO 50°S-68°E	1.9 (0.3)	-0.019 (0.004)	Leseurre et al (2022)
135	1998-2019	Summer	IO 55°S-63°E	2.1 (0.3)	-0.022 (0.003)	Leseurre et al (2022)
136	1998-2007	Summer	IO 55°S-63°E	5.3 (0.4)	-0.050 (0.016)	Leseurre et al (2022)
137	2006-2019	Summer	IO 55°S-63°E	0.3 (0.2)	no trend	Leseurre et al (2022)
138		_				
139	1969-2003	Summer	PO 55-62°S	1.7 (0.2)	-0.020 (0.003)	Midorikawa (2012)
140	2002 2012		D I M J	2 21 (0 55)	0.000 (0.007)	T 1 1 1: (2014)
141	2002-2012	Annual	Drake North	2.21 (0.55)	-0.023 (0.007)	Takahashi (2014)
142	2002-2012	Annual	Drake South	1.50 (0.65)	-0.015 (0.008)	Takahashi (2014)
143	2002 2015	C	DINA	1.05 (0.55)	0.001 (0.00()	M (1(2015)
144	2002-2015	Summer	Drake North	1.95 (0.55)	-0.021 (0.006)	Munro et al (2015)
145	2002-2015	winter	Drake North	1.92 (0.24)	-0.018 (0.003)	Munro et al (2015)
140	2002-2015	Summer	Drake South	1.30 (0.85)	-0.017 (0.010)	Munro et al (2015)
147	2002-2015	winter	Drake South	0.67(0.39)	-0.008 (0.004)	Munro et al (2015)
140	2002-2015	Annual	Drake North	1.74 (0.15)	-0.019 (0.002)	Munro et al (2015)
149	2002-2015	Annual	Drake South	1.10(0.27)	-0.015 (0.005)	Munro et al (2015)
150	1081 2011	Appual	50 5055	1 44 (0 10)	0.020 (0.002)	Louve at al (2015)
152	1981-2011	Annual	SO SPSS	1.44(0.10) 1.46(0.11)	-0.020(0.002) 0.021(0.002)	Lauvset et al (2015)
152	1991-2011	Allinual	50 51 55	1.40 (0.11)	-0.021 (0.002)	Lauvset et al (2015)
154	1993-2018	Δnnual	SO 44-75°S	NR	-0.0165 (0.0001)	Iida et al (2021)
155	1775-2010	7 tinituai	50		-0.0105 (0.0001)	fida et al (2021)
156						
157	1962-2016	November	IO 50°S-68°E	1 31 (0 20)	-0.014 (0.002)	This study Obs
158	1991-2021	Summer	IO 50°S-68°E	2 10 (0 22)	-0.014(0.002)	This study, Obs.
159	1991-2001	Summer	IO 50°S-68°E	0.76 (0.90)	-0.009 (0.010)	This study, Obs
160	2001-2010	Summer	IO 50°S-68°E	323(107)	-0.035(0.011)	This study, Obs
161	2010-2020	Summer	IO 50°S-68°E	0.84 (0.77)	-0.008 (0.008)	This study, Obs
162	2010 2020	Summer	10 0 0 0 00 2	0.01 (0.77)	0.000 (0.000)	11115 Stady, 005.
163	1985-2020	Summer	IO 50°S-68°E	1.71 (0.08)	-0.018 (0.001)	This study, FFNN
164	1991-2020	Summer	IO 50°S-68°E	1.85 (0.11)	-0.020 (0.001)	This study, FFNN
165	1991-2001	Summer	IO 50°S-68°E	1.18 (0.26)	-0.013 (0.004)	This study, FFNN
166	2001-2010	Summer	IO 50°S-68°E	2.87 (0.25)	-0.030 (0.003)	This study, FFNN
167	2010-2020	Summer	IO 50°S-68°E	0.98 (0.40)	-0.010 (0.004)	This study, FFNN
168	1991-2001	Winter	IO 50°S-68°E	0.98 (0.09)	-0.010 (0.001)	This study, FFNN
169	2001-2010	Winter	IO 50°S-68°E	1.99 (0.10)	-0.021 (0.001)	This study, FFNN
170	2010-2020	Winter	IO 50°S-68°E	2.21 (0.17)	-0.022 (0.002)	This study, FFNN
171	1985-2020	Annual	IO 50°S-68°E	1.57 (0.03)	-0.0165(0.0004)	This study, FFNN
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The present study complements in time, seasons, and in the water column, the surface fCO_2 and pH trends investigated by Leseurre et al., (2022) in different regions of the Southern Indian Ocean for the period 1998-2019 during austral summer. South of the PF around 50°S, Leseurre et al. (2022) showed that in summer the surface fCO_2 increase and pH decrease over 20 years were mainly driven by the increase in anthropogenic CO₂ sequestration about +0.6 (± 0.2) µmol.kg⁻¹.yr⁻¹ and by a small warming of +0.03 (± 0.02) °C.yr⁻¹. In addition Leseurre et al. (2022) showed that in the recent decade, 2007-2019, the fCO_2 trend was low +0.3 (± 0.2) µatm yr⁻¹ compared to +5.3 (± 0.4) µatm yr⁻¹ in 1998-2007, highlighting the sensitivity of the fCO_2 and pH trends to the





181	selected time period (especially during summer). In particular, they observed relatively stable pH values over
182	2010-2019 (i.e. no decrease in pH) with no clear explanation on the origin of the slow-down of the fCO_2 and pH
183	trends in surface waters south of the PF in recent years. To complement the analysis by Leseurre et al. (2022)
184	based on summer observations in 1998-2019 this study focusses on one location regularly visited south of the
185	Polar Front (around $50^{\circ}S-68^{\circ}E$ south-west of Kerguelen Island, Figure 1). The analysis period is extended back
186	to 1985 and forward to 2021 to investigate the recent status of fCO_2 and pH. We also evaluate the trends for
187	different seasons using sparse spring/winter data. The combination of in situ observations and monthly estimates
188	from a neural network model over the period 1985-2020 (Chau et al., 2022) enables to assess potential changes
189	in seasonality of the surface ocean carbonates system (including fCO ₂ , C_T , pH, Ω) as suggested in recent decades
190	or in future scenarios (Gallego et al., 2018; Landschützer et al., 2018; Kwiatkowski and Orr, 2018; Kwiatkowski
191	et al., 2020; Lerner et al., 2021; Fassbender et al., 2022; Yun et al., 2022; Rodgers et al., 2023; Joos et al., 2023).
192	The variability in surface waters will be related to changes in C_{ant} concentrations observed in the water column
193	and will be complemented by an analysis of OA at depth between 1985 and 2021. Finally we will explore the
194	long-term variability of fCO_2 and pH since 1960s and potential future changes of the carbonate system at this
195	time series site.

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- 197 2 Data selection, methods and quality control
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199 2.1 Study area and data selection

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201 This study focused on a High Nutrients Low Chlorophyll area (HNLC, Minas and Minas, 1992) of the 202 Indian sector of the Southern ocean (SO) in the Permanent Open Ocean Zone (POOZ) south of the Polar Front 203 (PF) and south-west of Kerguelen Islands (around 50°S-68°E, Figure 1). The Kerguelen Plateau is an extended 204 topographic feature that controls part of the Antarctic Circumpolar Current (ACC), generates eddies (Daniault 205 and Ménard, 1985) and the northward deflection of the PF near the Island (Pauthenet et al., 2018). The Plateau is 206 also a region of relatively high Chl-a (Moore and Abbott, 2000; Mongin et al., 2008) and strong CO₂ uptake 207 during austral spring-summer that contrasts with the weaker sink over the POOZ/HNLC (Metzl et al., 2006; 208 Jouandet et al., 2008, 2011; Lo Monaco et al., 2014; Leseurre et al., 2022). The POOZ/HNLC region west 209 (upstream) of the Kerguelen Plateau is characterized by rather stable water mass properties (temperature, 210 salinity, oxygen or nutrients) over time and low eddy activity compared to the Plateau (Daniault and Ménard, 211 1985; Chapman et al., 2015; Dove et al., 2022). In this region, located in the deep Enderby Basin, the flow is not 212 constrained by topography and there is no local upwelling that would import C_T-rich water to surface layers as 213 observed on the eastern side of the Kerguelen Plateau (Brady et al., 2021).

214 The Indian austral sector is also recognized to host the strongest winds in the SO leading to year-round high gas transfer coefficients (Wanninkhof and Trinanes 2017). As a result, and in contrast to the Atlantic sectors 215 216 of the SO, the Indian region south of 45°S was a periodic CO₂ source, especially in the 1960s to the 1980s 217 (Rödenbeck et al., 2022; Bennington et al., 2022; Prend et al., 2022; Gray, 2024). In the POOZ-HNLC region, high winter wind speed (monthly average up to 16 m.s⁻¹) and associated heat loss drive deep mixing. Deep 218 219 winter mixing entrains subsurface properties to the surface layer, increases C_T concentrations leading to 220 wintertime outgassing of CO₂ (Metzl et al., 2006). This combination of characteristics makes the region an ideal 221 test-bed for 1-D modeling studies investigating the temporal dynamics and drivers of biogeochemical processes







Figure 1: Left: Annual air-sea CO₂ flux (molC.m⁻².yr⁻¹) in the South Indian Ocean for year 2020 from the FFNN model (negative flux for ocean sink, positive flux for ocean source). The black box identified the location of the study south-west of Kerguelen. Right: Track of cruises with underway fCO₂ data South-West of Kerguelen Island. The station at 50°40'S-68°25'E occupied in 1985, 1992-1993 and 1998-2021 is indicated by a yellow circle. The yellow square is the region selected to calculate the mean values from the underway surface observations and from the FFNN model. Figures produced with ODV (Schlitzer, 2018).

245 Here we used surface and water-column observations around location 50°40'S-68°25'E (Figure 1, Table 246 S1), historically called KERFIX station (KERguelen FIXed station) sampled in 1990-1995 in the framework of 247 the WOCE/JGOFS programs (Jeandel et al., 1998). The station was first occupied in March 1985 during the INDIGO-1 cruise (Indian Ocean Geochemistry, Poisson, 1985; Poisson et al., 1988) and since 1998 it is 248 249 regularly visited during the OISO cruises (Océan Indien Service d'Observations, Metzl and Lo Monaco, 1998, 250 https://doi.org/10.18142/228). The regular occupation from 1985 to 2021 makes it the longest time-series station 251 in the Southern Ocean POOZ/HNLC area allowing investigating the inter-annual to decadal trends of carbonate 252 properties in surface waters and across the water-column (0-1600m). Despite the occasional variability in surface 253 waters properties (e.g. lower surface Salinity in 2011-2013) we consider all observations selected for this study 254 both in surface waters and the water-column to be representative of the water masses in this POOZ/HNLC region 255 upstream of the Kerguelen Plateau.

Data for 1985-2011 were extracted from the GLODAP data-product, version V2.2021 (Lauvset et al.,
2021 a, b; Table S1a). Observations collected during OISO cruises in 2012-2021 (Lo Monaco, 2020; Lo Monaco
et al., 2021) will be included in GLODAP-V3. For the surface water properties, all available underway fCO₂ data
were selected (Figure 1). This includes one cruise in November 1962 (Keeling and Waterman, 1968) and 41
cruises in 1991-2021 (Table S1b). All surface temperature, salinity and fCO₂ data were extracted from the
SOCAT data-product version v2022 (Surface Ocean CO₂ Atlas, Bakker et al., 2016, 2022).





263264 2.2 Methods

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The methods for surface underway fCO_2 and biogeochemical properties (Oxygen, A_T - C_T , nutrients) in the water-column for the INDIGO-1, KERFIX and OISO cruises were described in previous studies (e.g. Poisson et al., 1993; Louanchi et al., 2001; Metzl et al., 2006; Metzl, 2009; Mahieu et al., 2020; Leseurre et al., 2022). Here we briefly recall the methods for underway fCO_2 and water-column observations.

271 2.2.1 Surface fCO₂ data

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273 For fCO₂ measurements in 1991-2021, sea-surface water was continuously equilibrated with a "thin 274 film" type equilibrator thermostated with surface seawater (Poisson et al., 1993). The xCO₂ in the dried gas was 275 measured with a non-dispersive infrared analyser (NDIR, Siemens Ultramat 5F or 6F). Standard gases for 276 calibration (around 270, 350 and 490 ppm) were measured every 6 hours. To correct xCO₂ dry measurements to fCO₂ in situ data, we used polynomials from Weiss and Price (1980) for vapour pressure and from Copin-277 278 Montégut (1988, 1989) for temperature. Note that when incorporated in the SOCAT data-base, the original fCO_2 279 data are recomputed (Pfeil et al., 2013) using temperature correction from Takahashi et al. (1993). Given the 280 small difference between equilibrium temperature and sea surface temperature ($+0.56 \pm 0.30$ °C on average for 281 the cruises in 1998-2021), the fCO₂ data from SOCAT used in this analysis (Bakker et al., 2022) are almost 282 identical (within 1 µatm) to the original fCO2 values from our cruises (www.ncei.noaa.gov/access/ocean-carbon-283 data-system/oceans/VOS_Program/OISO.html).

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285 2.2.2 Water column data

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287 In 1990-1995, water samples were collected during the KERFIX program on the ship La Curieuse at 288 standard depths using 8 L Niskin bottles mounted on a stainless steel cable and equipped with reversing SIS pressure and temperature probes. Methods and accuracy for the geochemical measurements used in this analysis 289 290 (A_T, C_T, oxygen, nutrients) are detailed by Jeandel et al. (1998) and by Louanchi et al. (2001). From 1998 291 onwards, the station was occupied within the framework of the OISO long-term monitoring program onboard the 292 R.V. Marion-Dufresne. We used Conductivity-Temperature-Depth (CTD) sensors mounted on a 24 bottles 293 rosette equipped with 12 L Niskin bottles. Temperature and salinity measurements have an accuracy of 0.002 °C 294 and 0.005 respectively (Mahieu et al., 2020). Samples for A_T and C_T were filled in 500 mL glass bottles and poisoned with 100 μ L of saturated mercuric chloride solution to halt biological activity. Discrete C_T and A_T 295 296 samples were analyzed onboard by potentiometric titration derived from the method developed by Edmond 297 (1970) using a closed cell. Based on replicate samples from the surface or depth, the repeatability for A_T and C_T varies from 1 to 3.5 μ mol.kg⁻¹ depending on the cruise. The accuracy of ±3 μ mol.kg⁻¹ was ensured by daily 298 analyses of Certified Reference Materials (CRMs) provided by Andrew Dickson's laboratory (Scripps Institute 299 300 of Oceanography).

301 Dissolved oxygen (O_2) concentration was determined by a sensor fixed on the rosette and values were 302 adjusted based on discrete measurements (Winkler method, Carpenter, 1965) using a potentiometric titration 303 system. Accuracy for O_2 is $\pm 2 \mu \text{mol.kg}^{-1}$ (Mahieu et al., 2020). Although long-term deoxygenation in the





304 Southern ocean has been suggested (Ito et al., 2017; Schmidtko et al., 2017; Oschlies et al., 2018), no significant 305 trend in O2 was identified over 1985-2021 at this station around 50°S in both surface or in subsurface waters 306 (e.g. in the layer of the temperature minimum representing winter water for Cant calculations as described later). 307 However, in the station data a small O₂ decrease was detected around 800m in the O₂ minimum layer over 36 308 years (-0.22 \pm 0.07 μ mol.kg⁻¹.yr⁻¹). As this has no impact on the interpretation for pH and Ω trends for this 309 analysis, the observed change of O_2 at depth will be not discussed further. Here the O_2 data are mainly used for 310 the calculation of anthropogenic CO₂ concentrations and the observed O₂ change at depth is too small to have an 311 impact on temporal variations of Cant concentrations.

312 Nitrate (NO₃) and silicate (DSi) were analyzed on board or at LOCEAN/Paris by colorimetry following the methods described by Tréguer and Le Corre (1975) for 1998-2008 or from Coverly et al. (2009) for 2009-313 2021. The uncertainty of NO₃ and DSi measurements is $\pm 0.1 \ \mu mol.kg^{-1}$. Based on replicate measurements for 314 315 deep samples, we estimate an error of about 0.3 % for both nutrients. Phosphate (PO₄) samples were analyzed in samples from a few cruises following the method of Murphy and Riley (1962) revised by Strickland and Parsons 316 317 (1972) with uncertainty of $\pm 0.02 \ \mu mol.kg^{-1}$. When nutrient data are not available for a cruise, we used climatological values based on seasonal nutrients cycles inferred from data from 1990 to 2021. This has a very 318 319 small impact on the carbonate system calculations and the trend analysis as we did not detect any significant 320 trends in nutrients in surface or at depth since 1985 (not shown) as opposed to what has been observed at higher 321 latitude of the SO (Iida et al., 2013; Hoppema et al., 2015). However, we will see that the inter-annual variability 322 of nutrients (especially DSi in the HNLC region) might inform on potential changes in biological processes.

For Chlorophyll-a (Chl-a), samples were taken in the top layers (0-150m). One to two liters of seawater
were filtered onto 0.7 μm glass microfiber filters (GF/F, Whatman) and filters were stored at -80°C onboard.
Back at the LOCEAN/Paris laboratory, samples were extracted in 90% acetone (Strickland and Parsons, 1972)
and fluorescence of Chl-a was measured on a Turner Type 450 fluorometer in 1998-2007 and since 2009 at 670
nm on a Hitaschi F-4500 spectrofluorometer (Neveux and Lantoine, 1993).

328

329 2.2.3 Data quality-control and data consistency

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331 When exploring the trends of ocean properties based on different cruises more than 35 years apart, it is 332 important to first verify the consistency of the data and if there is any bias or drift. The INDIGO data from 1985 333 (i.e. prior to CRM available for A_T and C_T) were first controlled prior to their incorporation into the original 334 GLODAP product (Sabine et al., 1999; Key et al., 2004) and corrections for A_T and C_T were revisited within the 335 framework of the CARINA project (CARbon IN the Atlantic, Lo Monaco et al., 2010). A secondary quality 336 control was performed on the data from the OISO cruises collected between 1998 and 2011 within the CARINA 337 and GLODAP-v2 initiatives (Lo Monaco et al., 2010; Olsen et al., 2016). Significant off-sets were identified for A_T-C_T in samples from the KERFIX cruises (1990-1993) compared to INDIGO and OISO data and it was 338 proposed to correct the original values by -35 μ mol.kg⁻¹ for C_T and -49 μ mol.kg⁻¹ for A_T (Metzl et al., 2006). 339 340 These corrections were applied in GLODAP version v2.2019 (Olsen et al., 2019) and resulted in coherent AT and 341 C_T concentrations for KERFIX in the deep layers compared to other cruises (Supp Mat., Table S2, Figure S1). 342 The same data quality control protocol as for GLODAP-v2 was applied to data from OISO cruises for the years 2012-2021 (Mahieu et al., 2020). Given the accuracy of the data no systematic bias (excepted in 2014) was 343 344 found for the properties measured in 2012-2021. The time-series of A_T and C_T at depth below 1450 m for all





cruises in 1985-2021 show some variability but no trend over 36 years as expected in the bottom waters in this region (Supp. Mat, Figure S1). However, we identified a small bias for C_T in 2014 (cruise OISO-23) where C_T concentrations in the deep water appeared slightly lower (2228-2234 µmol.kg⁻¹ in 2014 compared to the mean value of 2240.7 (± 3.7) µmol.kg⁻¹, Table S2, Figure S1). When compared to fCO₂ in surface waters, we also suspect the C_T data in the mixed-layer in 2014 to be too low by about 10 µmol.kg⁻¹ (Figures S2, S3). Therefore we applied a WOCE/GLODAP flag 3 for C_T data of this cruise and will not use the station data in 2014 for the C_{ant} calculations and the trend analysis described in this study.

353 2.2.4 CMEMS-LSCE-FFNN model

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355 As most of the cruises took place during austral summer and data are not available each year, we 356 completed the observations with the results from an ensemble of feed-forward neural network model (CMEMS-357 LSCE-FFNN or FFNN for simplicity here, Chau et al., 2022). The FFNN model allows mapping at global scale 358 monthly surface fCO₂ given SOCAT gridded datasets and ancillary variables. The reconstructed fCO₂ is then 359 used to derive monthly surface pH fields as well as air-sea CO₂ fluxes. This data product enables to investigate 360 the trends for different seasons and to derive estimates of annual air-sea CO₂ fluxes to interpret the change in 361 CO_2 uptake, if any. For a full description of the model, access to the data and a statistical evaluation of fCO_2 362 reconstructions please refer to Chau et al. (2022). Within this study, we compared the FFNN fCO₂ with 363 observations from 35 cruises for the years between 1991 and 2020 (Table S3, Figure S2a). Excepted for a few periods (January 1993 and January 2002), model-data differences are generally within \pm 10 µatm with a mean 364 365 difference of 2.1 (\pm 7) µatm for the 35 co-located periods. Note that, as opposed to sea surface fCO₂, no temporal 366 trend was identified for the differences between the observed and reconstructed fCO_2 (Figure S2b), i.e. the trends 367 of sea surface fCO₂ derived from the observations and from the FFNN model should be the same. Aside from the fCO₂ reconstructions, surface ocean alkalinity (A_T) fields are also provided by using the multivariate linear 368 regression model LIAR (Carter et al., 2016; 2018) based on sea surface temperature, salinity, and nutrient 369 370 concentration.

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372 2.2.5 Calculations of carbonate properties

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374 Based on the data available for each cruise (fCO₂, or A_T and C_T) or from the FFNN model (fCO₂ and 375 A_T), other carbonate system properties (pH, [H⁺], [CO₃²⁻] and Ω) were calculated using the CO2sys program 376 (version CO2sys_v2.5, Orr et al., 2018) developed by Lewis and Wallace (1998) and adapted by Pierrot et al. 377 (2006) with K1 and K2 dissociation constants from Lueker et al. (2000) as recommended (Dickson et al., 2007; 378 Orr et al., 2015; Wanninkhof et al., 2015). The total boron concentration was calculated according to Uppström 379 (1974) and KSO₄ from Dickson (1990). To calculate the properties with the underway surface fCO_2 dataset, we 380 used the A_T/S relationship based on A_T-C_T data from OISO cruises in 1998-2019 in the South Indian sector as 381 described by Leseurre et al. (2022):

382

383 $A_T = 64.341 \text{ x S} + 106.764 \text{ (rmse} = 7.485 \text{ } \mu\text{mol.kg}^{-1}, \text{ n} = 4775 \text{) (Eq. 1)}$





385 The use of other A_T/S relationships (e.g. Millero et al., 1998; Jabaud-Jan et al., 2004; Lee et al., 2006; 386 Carter et al., 2018) would change slightly the A_T concentrations but neither the A_T trend nor the interpretation of 387 the C_T , pH or Ω trends. However, as salinity is an important predictor in the calculation of A_T , C_T or pH from 388 fCO₂ data, we have assessed the original underway salinity data and found biases for few cruises in 1992, 1993 389 and 1995 (Table S1b). For these cruises or when salinity was not measured we used the salinity from the World 390 Ocean Atlas, WOA (Antonov et al., 2006) in the SOCAT data-sets (Pfeil et al., 2013, identified "WOA" in Table 391 S1b). Monthly fCO2 and AT data extracted from the CMEMS-LSCE-FFNN datasets at the station location 392 (50.5°S-68.5°E) over 1985-2020 were used to calculate the carbonate properties in the same way as from 393 observations.

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395 2.2.6 Comparisons of different datasets and the FFNN model

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397 To validate the properties calculated using the fCO2 data for 1991-2021 or from the FFNN model over 1985-2020 we compared the calculated values (A_T, C_T, pH, [H⁺], [CO₃²⁻], Ω) with those from A_T-C_T data 398 measured in the mixed-layer at stations occupied in 1985 and in 1993-2021. For this comparison, we averaged 399 400 the continuous underway fCO₂ data selected in a box around the station location ($50^{\circ}S-51.5^{\circ}S/67.5-69^{\circ}E$, yellow 401 box in Figure 1). Results of the comparisons between various datasets are detailed in the Supplementary Material 402 (Tables S3 and S4). During the period 1993-2021, there are 22 stations with co-located fCO₂ data for different 403 seasons (but mainly in summer). Since we found a close agreement between measured fCO_2 and the FFNN 404 model (Table S3, Figure S2), mismatches in all calculated carbonate system properties between the underway 405 fCO2 dataset and the FFNN model are small, falling within the range of the errors associated with the 406 calculations (Orr et al., 2018). For example, for 35 co-located periods, the mean differences in calculated C_T of 1.5 (\pm 5) µmol.kg⁻¹ or pH of -0.002 (\pm 0.008) are in the range of the theoretical error of about 5 µmol.kg⁻¹ and 407 408 0.007 respectively when taking into account measurements errors on salinity, temperature, nutrients, fCO₂ and 409 A_T (Orr et al., 2018). On the other hand, compared to the station data in the mixed-layer (Table S4), bias for calculated A_T using Equation 1 is slightly higher by about 5 μ mol.kg⁻¹. This explains the relatively high 410 411 differences for C_T (mean difference around 8 µmol.kg⁻¹) and for pH (mean difference around 0.008) calculated 412 with fCO2 and the AT/S relationship. The differences of calculated values with observations in 1991-2021 are, on 413 average, in the range of uncertainties of the carbonate system calculations using A_T - C_T pairs (error for fCO₂) 414 around 13 µatm and for pH around 0.0144). Importantly, there is no temporal trend for the differences between 415 calculated and observed properties (Figure S3b). We are thus confident using the selected fCO2 data for the trend 416 analysis presented in this study. The independent comparison with A_T-C_T data at stations also indicates that the FFNN model results for A_T, C_T, are close to the observations (Table S4, Table S5, Figure S4) as well as for 417 calculated pH, [H⁺], [CO₃²⁻], Ω_{Ca} and Ω_{Ar} . This somehow validates the use of the FFNN data for the trend 418 419 analysis over the period 1985-2020 and for different seasons, although the FFNN model was not constrained by 420 in-situ fCO₂ before 1991 or Chl-a satellite data before 1998. Interestingly, in 1985 the atmospheric fCO₂ was 421 around 335-339 µatm (Dlugokencky and Tans, 2022) and the oceanic fCO2 from the FFNN model was higher 422 than in the atmosphere from March to October (Figure S4) resulting in an annual CO_2 source of +0.8 mol.m⁻².yr⁻¹ 423 in 1985.





425	3	Results	hne	discussion
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427 3.1 Variability and trend of sea surface fCO₂ and air-sea CO₂ fluxes: 1985-2021

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429 The fCO₂ observations around 50°S-68°E and their mean values for each cruise are shown in Figure 2a. 430 The fCO₂ data in 1991-2021 were available for different seasons but the sampling locations were mainly 431 reoccupied in austral summer (January-February). During austral summer, the ocean fCO2 was generally lower 432 than in the atmosphere (i.e. the ocean was a CO_2 sink) whereas in March to October it was near equilibrium. The same distribution is obtained from the FFNN model for 1991-2020 (Figure 2a). The model also indicates that in 433 434 1985-1998 the fCO2 during austral winter (May-September) was always higher than the atmospheric fCO2 435 leading to a CO₂ source during this period (Figure 3). The model estimates a decrease of the annual CO₂ source 436 in 1985-2001 followed by an increase of the source in 2001-2010 and an increase of the sink in 2010-2020.





461 Figure 2: (a): Time-series of sea surface fCO_2 observations (µatm) South-West of Kerguelen Island in 1985-2021 (insert map 462 shows the location of observations selected around station at 50°40'S-68°25'E). The color dots correspond to 5 seasons 463 (January-February, March-April, July-August, October and December) and triangles the average for each period. The 464 monthly sea surface fCO_2 from the FFNN model is presented for the period 1985-2020 (purple line) and the atmospheric 465 fCO_2 represented by red line. In March 1985 there were no underway fCO_2 observations and the triangle corresponds to fCO_2 466 calculated with A_T - C_T data in the mixed-layer. (b): Trends of atmospheric and oceanic fCO_2 (µatm.yr⁻¹) for different season 467 and periods based on observations (January) and the FFNN model (January or August).









Figure 3: Time series of the SAM index (in grey) in the Southern Ocean, wind-speed (in black, m.s⁻¹) and air-sea CO₂ flux (molC.m².yr⁻¹) from the FFNN model (in red) at location 50.5°S-68.5°E. Positive (negative) flux represents CO₂ source (sink). Wind-speed and SAM are presented for respectively 12-months and 24-months running mean based on monthly values. Note the positive SAM in 1998-2003 and 2010-2020. SAM data from Marshall (2003), http://www.nerc-bas.ac.uk/icd/gjma/sam.html, last access 14/8/2021. Wind speed data from ERA5 (Hersbach et al., 2020).

For the last cruise in February 2021, the average fCO₂ was 394.9 (\pm 1.5) µatm (Figure 2a), about 10 µatm lower than in the atmosphere (a small CO₂ sink). This is +50.5 µatm higher than fCO₂ observed during the first cruise in February 1991 (fCO₂ = 344.4 \pm 5.2 µatm). During the same period, the atmospheric CO₂ increased from 354 ppm in 1991 to 411 ppm in 2021 in this region (recorded at Crozet Island, Dlugokencky and Tans, 2022). This first comparison of two cruises 30 years apart indicates that the ocean fCO₂ increased at a rate (+1.7 µatm.yr⁻¹) close to that of the atmosphere (+1.9 µatm.yr⁻¹). During the same period, we observed some variations in A_T (average A_T = 2276.5 \pm 4.5 µmol.kg⁻¹) and a clear increase in C_T (Figure 4a and S5).

497 The C_T concentration in the mixed-layer in summer 2021 was 2134.0 (± 1.8) µmol.kg⁻¹, much higher 498 than in summer 1993 ($C_T = 2115.8 \pm 2.6 \mu \text{mol.kg}^{-1}$). The difference over 28 years of +22.1 $\mu \text{mol.kg}^{-1}$ 499 corresponds to an annual C_T increase of +0.8 µmol.kg⁻¹.yr⁻¹. At constant temperature and A_T, this would translate 500 in an increase of oceanic fCO₂ of $+1.9 \mu$ atm, yr⁻¹, i.e. equal to the atmospheric rate. The same comparison for 501 October shows that fCO₂ in 2016 was +43.8 µatm higher compared to 1995 (Figure 2a), i.e. a rate of +2.1 502 µatm.yr⁻¹. The C_T concentrations in October 2016 were also much higher than in 1993 (Figure 4a and S5). Over 23 years the observed C_T increase in October (+22.6 µmol.kg⁻¹) corresponds to a rate of +0.98 µmol.kg⁻¹.yr⁻¹ that 503 is faster than the rate of $+0.8 \ \mu\text{mol.kg}^{-1}$. yr⁻¹ derived from summer data in 2021 and 1993. At constant A_T this 504 505 would translate in an increase of oceanic fCO₂ of $+2.5 \mu atm.yr^{-1}$ in October, higher than the trend of +2.1µatm.yr⁻¹ computed from fCO₂ data. Part of the difference may be explained by A_T that was slightly higher (+6 506 507 umol.kg⁻¹) in October 2016 compared to 1993 (Figure S5).

Given the temporal variability of observed C_T in summer and the evolution of the annual air-sea CO_2 flux (Figure 3), decadal f CO_2 and pH trends as well as associated drivers need to be analyzed for different seasons and periods. This approach allows exploring links with the variability of primary production and/or the Southern Annual Mode (SAM). Shifts from a positive to a negative SAM index (Figure 3) will strengthen upwelling and impact ocean properties throughout the water column including C_T , nutrients, primary production or pH (e.g. Lovenduski and Gruber, 2005; Lenton et al., 2009; Hauck et al., 2013; Hoppema et al., 2015; Pardo et al., 2017).







Figure 4: (a): Time-series of surface C_T (µmol.kg⁻¹) around station at 50°40'S-68°25'E calculated from fCO₂ data (Figure 2) using the A_T/S relation (see text). The color dots correspond to 5 seasons (January-February, March-April, July-August, October and December) and triangles the average for each cruise. The monthly sea surface C_T from the FFNN model is presented for the period 1985-2020 (purple line). The annual C_T trend of +0.58 ±0.05 µmol.kg⁻¹.yr⁻¹ (dashed line) is derived from the FFNN monthly data. In March 1985 the triangle corresponds to the observed C_T in the mixed-layer. (b): Trends of sea surface C_T (µmol.kg⁻¹.yr⁻¹) for different season and periods based on observations (for January) and the FFNN model (for January or August). The trend for C_{ant} (µmol.kg⁻¹.yr⁻¹) is also shown (red bars) based on estimates in the winter water.

550 551 552 553 Summer data are characterized by a strong inter-annual variability between 1991-2021 (Figures 2a and 4a) with the ocean being a CO₂ source in January 2002, but a strong sink in January 1993, 1998, 2014, 2016 and 554 555 2019. In January 1998, when the surface ocean experienced a warm anomaly (Jabaud-Jan et al., 2004), the low 556 fCO₂ of 337 µatm and the low C_T of 2110 µmol.kg⁻¹ (Figure 4a and S5) co-occurred with intense primary production, probably supported by diatoms as suggested by very low DSi concentrations (< 2 µmol.kg⁻¹ down to 557 558 100m, Figure S6). In January 2014 and 2016, mixed-layer DSi concentrations were also remarkably small (< 5 559 µmol.kg⁻¹ down to 75m, Figure S6). In 2014 low DSi coincided with Chl-a levels that started to increase in mid-560 November 2013 and stayed at high level until February 2014 (Surface Chl-a > 0.3 mg.m⁻³, Figures 5 and S7). The intense primary production contributed to the low fCO₂ of 365 µatm reached by mid-January 2014, a value 561 as low as in 2004 (Figure 2a). To the contrary, in 2002 relatively low Chl-a (mean Chl-a $< 0.2 \text{ mg.m}^{-3}$, Figure 5) 562 563 was associated with higher levels of fCO2 (373 µatm), CT (2128 µmol.kg⁻¹, Figure 4a, Figure S5a) and DSi 564 (Figure S6). This was also associated with higher salinity indicative of entrainment that might be related to storm



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well below the atmospheric level (Figure 2a).



565 events that would occurred few days before the measurements leading to brief positive fCO₂ anomaly as recently 566 observed from Glider data in the subpolar South Atlantic (Nicholson et al., 2022). As opposed to the other 567 periods the ocean was a source of CO₂ in summer 2002 (this particular year was not well reconstructed by the FFNN model, Figure 2a and Figure S2b). The important inter-annual variability observed in summer indicates 568 569 that in this region historically referred to as HNLC (Minas and Minas, 1992), primary production could significantly impact fCO₂ level in summer (Jabaud-Jan et al., 2004; Pasquer et al., 2015; Gregor et al., 2018), a 570 571 result that needs to be taken into account when evaluating drivers of inter-annual variability (Rustogi et al., 572 2023) and the decadal trends of fCO₂ or pH.

The Chl-a time-series derived from MODIS suggests higher concentrations in recent years with Chl-a

peaks identified in 2014, 2016, 2018, 2019 and 2021 (Figure 5 and S7) when the oceanic fCO₂ in summer was

576 0,6 577 Chl-a Dec-Feb 0,5 578 📥 Chl-a mean Ā Δ 579 Chl-a (mg.m⁻³) Δ 580 0,3 581 0,2 582 583 0,1 584 0 585 1996 1998 2000 2002 2004 2006 2008 2010 2012 2014 2016 2018 2020 2022 Year 586

Figure 5: Time-series (1998-2021) of sea surface Chl-a (mg.m⁻³) in summer (December-February) from weekly satellite data (SeaWIFS and MODIS, triangles) and associated mean (green triangles). The trends in 1998-2010 and 2010-2021 of respectively -0.0099 (± 0.0041) and +0.0078 (± 0.0032) mg.m⁻³.yr⁻¹ (dashed orange) indicate a decrease or increase of the primary production that drives part of the fCO₂ and C_T stability observed in the recent period (Figure 2, Figure 4). The full Chl-a record is shown in Supp. Mat. Figure S7.

593 The primary production lowers C_T concentrations and fCO₂, i.e. opposite to the C_T increase from 594 anthropogenic CO₂ uptake. These counteracting processes might explain the relatively stable fCO₂ previously 595 observed in the Indian POOZ in summer 2007-2019 with an annual fCO₂ rate of increase of only $\pm 0.3 (\pm 0.2)$ μ atm.yr⁻¹ (Leseurre et al., 2022). This low rate is confirmed here with the new data obtained in 2020-2021 596 597 (Figure 2b and Figure S8). For the period 2010-2021, the oceanic fCO₂ trend in summer derived from observations and the FFNN model is lower than +1 µatm.yr⁻¹ (Table 1), i.e. much lower than the atmospheric 598 fCO₂ rate of +2.4 μ atm.yr⁻¹ and the oceanic fCO₂ trend of +2.21 (± 0.17) μ atm.yr⁻¹ in winter (Figure 2b). This 599 600 rate is also lower compared to the change observed in October (+2.9 µatm.yr⁻¹) albeit being only based on 2 601 cruises in 2011 and 2016 (Figure 2a). As the low fCO₂ trend in recent years is detected for summer only this is 602 likely linked to an increase in primary production, as suggested by Chl-a records (Figure 5). In 1998-2010 the summer Chl-a concentrations decreased at a rate of $-0.099 (\pm 0.041)$ mg.m⁻³.decade⁻¹ whereas in 2010-2021 Chl-603 604 a increased by $\pm 0.078 (\pm 0.032) \text{ mg.m}^3$. decade⁻¹ (Figure 5). These trends are coherent with previous studies, e.g. 605 the reduced net primary productivity reported in the Indian Antarctic zone in 1997-2007 (e.g. Arrigo et al., 2008; 606 Takao et al., 2012) and the shift of the Chl-a trend at large scale in the HNLC region of the Southern Ocean in 607 2010 (Basterretxea et al., 2023). As a consequence, after 2010 the difference between oceanic and atmospheric





 fCO_2 , ΔfCO_2 (where $\Delta fCO_2 = fCO_2^{\text{oce}} - fCO_2^{\text{atm}}$) decreased in summer (-1.4 $\mu atm.yr^{-1}$) and the annual CO₂ flux 608 progressively varied from a source of +0.45 molC.m⁻².yr⁻¹ in 2010 to a sink of -0.63 molC.m⁻².yr⁻¹ in 2020 609 (Figure 3). In addition, because the wind speed was stable during this period (12.0 \pm 0.9 m.s⁻¹ on average in 610 611 2010-2020, Figure 3), the variation of the air-sea CO₂ flux was mainly controlled by Δ fCO₂ (e.g. Gu et al., 2023) and the decadal variation of primary production imprinted a significant change on the fCO2 trend and air-sea 612 613 CO₂ flux in this HNLC region. In the region investigated here, increasing Chl-a levels co-occurred with shifts of 614 the SAM index to a positive state (Figure 3), a link previously suggested south of the Polar Front in the SO but 615 for a short period 1997-2004 (Lovenduski and Gruber, 2005).

Another process to take into account for interpreting fCO_2 trends is the change in temperature in surface waters. Previous analysis suggested a progressive warming in the region investigated here (Auger et al., 2021 for summer 1993-2017). For 1998-2019 Leseurre et al. (2022) estimated a warming of Indian POOZ surface waters of +0.03 (± 0.02) °C.yr⁻¹. Extending the time-series for the period 1991-2021 (Figure S9a) we note that the surface temperature presents sub-decadal variability and that the ocean cooled after 2018 with a trend of -0.474 (± 0.164) °C.yr⁻¹ in 2018-2021 based on the monthly sea surface temperature (SST, Figure S9b). The trend derived from our in-situ observations in summer 2018-2021 was -0.253 (± 0.092) °C.yr⁻¹.

In 2019, the lower temperature and relatively high Chl-a lead to low fCO₂ (380 µatm, Figure 2a) and 623 624 low C_T (2128 µmol.kg⁻¹) compared to 2018 (fCO₂ = 386 µatm; C_T = 2137 µmol.kg⁻¹, Figure 4a). The decrease in observed fCO2 from summer 2018 to 2019, also reconstructed by the FFNN model (Figure 2a), is contrary to the 625 626 expected fCO_2 and C_T increase due to anthropogenic uptake. In 2020, although the temperature was also lower, 627 the oceanic fCO₂ was higher (392 µatm) probably due to lower primary production as suggested by higher DSi (Figure S6), as well as from C_T (2135 μ mol.kg⁻¹, Figure 4a) and Chl-a records (Figure 5). In January 2021 the 628 629 temperature was close to that in January 2020, and both fCO2 and CT were slightly higher (395 µatm, 2139 630 μ mol.kg⁻¹). A_T concentrations were stable between 2018 and 2021 (2278.9 ± 1.8 μ mol.kg⁻¹, Figure S5) indicating 631 no effect of A_T on the observed fCO₂ change in this region as opposed to the areas north of the Polar Front in the Indian Ocean where A_T variations are often linked to coccolithophores blooms (Balch et al., 2016; Smith et al., 632 633 2017).

The inter-annual variability observed in 1991-2021 highlights the competitive processes that drive C_T , fCO₂ or pH temporal variations. In summer 2018-2019, cooling and increased primary production both lead to low fCO₂ counteracting the effect of anthropogenic CO₂ uptake. Given the changes of Chl-a, SST and air-sea CO₂ flux, trends will be evaluated for three periods, 1991-2001, 2001-2010 and 2010-2020. In order to separate natural and anthropogenic contributions, the anthropogenic CO₂ signal is estimated in the following section.

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640 3.2 Anthropogenic CO₂

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642 **3.2.1** Anthropogenic CO₂ in the water column

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To calculate anthropogenic CO_2 concentrations (C_{ant}), we used the TrOCA method developed by Touratier et al. (2007) and previously applied in the southern Indian Ocean (Mahieu et al., 2020; Leseurre et al., 2022). Such an indirect method is not suitable for evaluating C_{ant} concentrations in surface waters due to biological activity and gas exchange and we restrict the C_{ant} calculations below the productive layer around







648 150m. In the region south of the Polar Front, a well-defined subsurface temperature minimum is observed each

Figure 6: Hovmoller section (Depth-Time) of potential temperature (°C) and anthropogenic CO₂ (C_{ant}, µmol.kg⁻¹) in 19852021 at station OISO-KERFIX (50°40'S-68°25'E). The section for temperature is presented in the layer 0-500m and for
summer to highlight the temperature minimum around 200m (winter water, WW). The section for C_{ant} is limited below 200m.
Section produced with ODV (Schlitzer, 2018).

673 The CT and Cant concentrations increased over time in the water column, a signal that is most 674 pronounced in the top layers (0-400m, Figure 6b). In the deep layer, the presence of the Indo-Pacific Deep Water (IPDW) around 600-800m is identified by a maximum of C_T ($C_T > 2250 \mu mol.kg^{-1}$) and a minimum of O_2 (O_2 675 close to or < 180 µmol.kg⁻¹, Figure S10) (Talley, 2013; Chen et al., 2022). In the IPDW layer restricted to the 676 677 neutral density (ND) range 27.75-27.85 kg.m⁻³ there is no significant change in C_T over time (Figure S10). In that layer the C_{ant} concentrations in 1985 (17.3 μ mol.kg⁻¹) were almost identical to those evaluated in 2021 (21.2 678 μ mol.kg⁻¹), considering the uncertainty in the C_{ant} calculations (± 6.5 μ mol.kg⁻¹, Touratier et al., 2007). As 679 680 discussed above (section 2.2.3) the C_T and A_T concentrations in the bottom layer (>1450m) were stable in 1985-681 2021 (Table S2, Figure S1). Below 800m, the Cant concentrations were small but not null (Figure 6b). The average C_{ant} concentration below 800m for all years and seasons is 7.97 (± 5.31) µmol.kg⁻¹ (n=123) with a very 682 small change detected over time ($C_{ant} = 7.73 \pm 1.27 \ \mu mol.kg^{-1}$ in 1985 and $C_{ant} = 10.45 \pm 0.62 \ \mu mol.kg^{-1}$ in 2021). 683

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3.2.2 Anthropogenic CO₂ trend in the subsurface

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687To separate the natural and anthropogenic signals in surface waters for the driver analysis we assume688that C_{ant} in the WW is representative of C_{ant} in the mixed-layer (ML). This is confirmed with few stations689occupied during winter showing that C_{ant} concentrations in the WW in summer are almost equal to C_{ant} in the





690 ML during the preceding winter (Figure S11). The variation of C_{ant} in the WW for 1985-2021 is presented in 691 Figure 7a for all seasons. In 1985 the C_{ant} concentration in the WW was 47.1 µmol.kg⁻¹ and C_{ant} reached a 692 maximum of 71.7 µmol.kg⁻¹ in 2021. The data selected at 200m present some inter-annual variability like the 693 relatively low C_{ant} in 1998, 2005 or 2020 probably related to natural variability. In 1998 and in 2020 the O₂ 694 concentrations were slightly lower in the WW (< 300 µmol.kg⁻¹) explaining the lower C_{ant} concentration (44.8 695 µmol.kg⁻¹ in 1998 and 53.8 µmol.kg⁻¹ in 2020). In 2005 anomalies of C_T , O₂ and temperature concur to explain 696 the lower C_{ant} (43.9 µmol.kg⁻¹).



Figure 7: (a): Time-series of anthropogenic CO₂ ($C_{ant} \mu mol.kg^{-1}$) estimated in the winter water layer (WW around 200m, see figure 6) in 1985-2021 at station OISO-KERFIX (50°40'S-68°25'E). Black dots are the individual data in the WW and the grey squares the average for the 1990s, 2000s and 2010s (anomalies in 1998, 2005 and 2020 filtered). The C_{ant} trend of +0.53 (\pm 0.01) µmol.kg⁻¹.yr⁻¹ is represented (dashed line). (b): same data for C_{ant} versus atmospheric fCO₂ (the slope is +0.263 \pm 0.042 µmol.kg⁻¹.µatm⁻¹).

From 1985 to 2021, we estimate a Cant trend in WW of +0.49 (± 0.09) µmol.kg⁻¹.yr⁻¹. When the Cant 720 anomalies in 1998, 2005 and 2020 are discarded, this C_{ant} trend is +0.53 (± 0.01) µmol.kg⁻¹.yr⁻¹ (Figure 7a). As 721 722 expected, the C_{ant} concentrations in the ocean are positively related to atmospheric CO₂ (slope +0.263 ± 0.042 µmol.kg⁻¹.µatm⁻¹, Figure 7b). Interestingly the slope observed south of the PF in the Indian Ocean is close to that 723 observed in the Antarctic Intermediate waters (AAIW) in the South Atlantic ($+0.23 \pm 0.05 \mu mol.kg^{-1}.\mu atm^{-1}$, 724 725 Fontela et al., 2021). At large scale, Gruber et al. (2019 a, b) evaluated Cant changes between 1994 and 2007 in the global ocean. In the South Indian sector, they estimated a mean C_{ant} accumulation in the surface of +6.0 (\pm 726 1.1) µmol kg⁻¹ in the band 50-55°S south of the PF. At our station location (50-52°S/68°E) in the layer 0-250m, 727 728 the C_{ant} accumulated from 1994 to 2007 was +5.67 (± 1.47) µmol kg⁻¹. In 13 years, this corresponds to a trend of 729 +0.44 (± 0.11) µmol.kg⁻¹.yr⁻¹. Gruber et al. (2019 a, b) did not use the data presented here allowing for an 730 independent comparison to the present study. Estimates of Cant accumulation by Gruber et al. (2019 a, b) are in 731 agreement with ours for 1991-2008 (+0.46 µmol.kg⁻¹.yr⁻¹) but lower than reported here between 2008 and 2021 732 (+0.61 µmol.kg⁻¹.yr⁻¹).





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734 3.2.3 Anthropogenic and surface C_T seasonal trends

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The C_{ant} trend in WW over 1985-2021 (+0.53 ± 0.01 µmol.kg⁻¹.yr⁻¹) is slightly lower than the annual C_T 736 737 trend in surface derived from the FFNN model for 1985-2020 (C_T trend = +0.58 ± 0.05 µmol.kg⁻¹.yr⁻¹ Figure 4a) suggesting that anthropogenic CO_2 uptake explains 86% of the C_T increase in surface. In 1991-2020 the surface 738 C_T trend appears slightly higher in January (+0.68 ± 0.07 µmol.kg⁻¹.yr⁻¹) than in August (+0.56 ± 0.04 µmol.kg⁻¹.yr⁻¹) 739 ¹.yr⁻¹, Figure 4b). This suggests that in addition to the increase of C_T due to anthropogenic CO₂ other processes 740 741 count such as the variability of the biological activity, vertical mixing or upwelling. Indeed, as for fCO₂ (Figures 742 2b), the C_T growth rate also depends on seasons and decades (Figure 4b). In 1991-2001 the C_T trend from the observations ($+0.05 \pm 0.64 \mu$ mol.kg⁻¹.yr⁻¹) is highly uncertain due to few data and the large variability (Figures 743 744 4a, b). The FFNN model showed that the C_T trend in summer was faster than in winter and the winter C_T trend 745 lower than the Cant trend in subsurface (Figure 4b). This is because during that decade, the higher primary 746 production in 1998 created a negative C_T anomaly (Figure 4a) not compensated by the accumulation of C_{ant}.

In 2001-2010 the C_T trends were much faster than in 1991-2001 and they were the same for both 747 748 seasons (around 1 μ mol.kg⁻¹, yr⁻¹, Figure 4b). For this decade the summer C_T trends from the observations and the FFNN model are coherent. They were also twice the Cant rate in WW that could be explained by enhanced 749 750 upwelling of C_T-rich deep waters during this period after the SAM reached a high positive index (Figure 3; 751 Lenton and Matear, 2007; Le Quéré et al., 2007; Hauck et al., 2013). However, in 2001-2010 we did not detect 752 any clear change at depth for ocean properties (except for C_T and C_{anl}) that would support this assumption 753 (enhanced upwelling). The rapid C_T (and fCO₂) trend for this decade is probably due to processes at the surface 754 rather than changes in the water column (vertical mixing or upwelling). In 2010-2020 C_T trends are lower than in 755 2001-2010 (Figure 4b). For summer, this is identified from both observations and the FFNN model. In winter the 756 $C_{\rm T}$ trend is close to $C_{\rm ant}$ indicative of the anthropogenic accumulation. The low $C_{\rm T}$ trend at the surface in 757 summer, about half the Cant trend for the FFNN model, is likely due to the increase of primary production after 758 2010 as described above (Figure 5). It appears thus that the impact of biological activity and its variability in 759 summer could counteract that of anthropogenic CO₂ and explain the temporal change of the carbonate system in 760 surface.

761 Given the differences of the fCO₂ and C_T trends in summer and winter (Figures 2b and 4b) we explored 762 the temporal variations of the seasonality. For each year we estimated the differences between August and 763 January (Figure 8a). The seasonal amplitude for C_T was on average 26.1 (± 3.4) µmol.kg⁻¹ and for fCO₂ 15.1 (± 764 5.6). Interestingly, the fCO₂ seasonal amplitude reached a minimum around 2008-2010 and increased over 2010-765 2020. This signal appears correlated with surface Chl-a in summer (Figure 8b). The inter-annual variability of 766 the seasonality is clearly identified when comparing C_T with C_T calculated due only to C_{ant} accumulation after 2010 (Figure S12c). This supports the conclusion that in addition to the C_{ant} accumulation, the variations of 767 768 phytoplanktonic biomass imprinted inter-annual variability on C_T and fCO₂ in summer. This holds for the 769 seasonal amplitude as the results for winter follows the Cant trend (Figure 4b, Figure S12a). The same is true for 770 pH for which reduced seasonal amplitude was found when the production was low (not shown). However, over 771 36 years (1985-2020) we did not identify a long-term trend of the seasonal amplitude for C_T or for fCO₂ as 772 suggested by other studies (Landschützer et al., 2018; Rodgers et al., 2023; Shadwick et al 2023). Our results 773 highlight a variability over 5-10 years (Figure 8a) and suggest a potential change in seasonality and annual CO2





g

chl-a



sink if primary production changes in the future (e.g. Bopp et al., 2013; Leung et al., 2015; Fu et al., 2016;

775 Kwiatkowski et al., 2020; Krumhardt et al., 2022; Seifert et al., 2023).

Figure 8: (a): Time-series of the seasonal amplitude (August minus January) for surface C_T (black, µmol.kg⁻¹) and fCO₂
(grey, µam) from the FFNN model at station OISO-KERFIX (50°40'S-68°25'E). Also shown are the mean surface Chl-a
(green, mg.m⁻³) in summer in 1998-2021. (b): Seasonal amplitude of fCO₂ and C_T versus summer Chl-a for 1998-2020. The
dashed lines indicate that the seasonal amplitude (August-January) increases when Chl-a is higher.

800 3.3 Anthropogenic CO₂ drives acidification in surface and the water column

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802 3.3.1 Surface pH trend

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804 To explore the temporal change of pH in surface water we used the fCO₂ observations and the monthly 805 results from the FFNN model. For both data-sets pH was calculated from fCO_2 and A_T reconstructed as described in section 2.2.5. Figure 9a presents the time-series of pH in the surface (the same time-series for [H⁺] 806 concentrations is shown in Figure S13). For the full period, 1985-2020, the annual pH trend derived from the 807 FFNN model is -0.0165.decade⁻¹ (± 0.0004) exactly the same as derived at large scale in the Southern Ocean 808 809 (south of 44°S) for the period 1993-2018 (Iida et al., 2021, Table 1) but when restricted to this period, 1993-810 2018, the trend from the FFNN model appears slightly faster of -0.0182.decade⁻¹ (± 0.0006). This is less than the 811 pH trend of -0.020 (\pm 0.002).decade⁻¹ derived from pCO₂ data in the SO SubPolar Seasonally Stratified biome 812 around 40-50°S (SO-SPSS) for 1981-2011 (Table 1, Lauvset et al., 2015) and close to the pH trend of -0.0189 (\pm







Figure 9: (a): Time-series of surface pH (TS) around station OISO-KERFIX ($50^{\circ}40^{\circ}S-68^{\circ}25^{\circ}E$) calculated from fCO₂ data (Figure 2) using the A_T/S relation (see text). The color dots correspond to 5 seasons (January-February, March-April, July-August, October and December) and triangles the average for each cruise. The monthly sea surface pH from the FFNN model is presented for the period 1985-2020 (purple line). The annual pH trend in 1985-2020 of -0.0165.decade⁻¹ (\pm 0.0004) (dashed line purple) is derived from the FFNN monthly data (the same figure for [H⁺] concentrations is presented in Supp. Mat. Figure S13). (b): Trends of pH (TS.decade⁻¹) for different seasons and periods based on observations (January) and the FFNN model (January or August).

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848 The winter pH decreased was faster in recent years, mirroring the winter fCO₂ trend (Figure 2b). On the 849 opposite, in summer, the pH trend presents a large variability at decadal scale and was lower in 2010-2020. In summer 2001-2010, the pH trend from the FFNN model was -0.0304.decade⁻¹ (± 0.0026) whereas in 2010-2020, 850 851 it was -0.0098.decade⁻¹ (\pm 0.0042) (Figure 9b, Table 1). Although the trends based on the observations are less 852 robust because the cruises were not conducted each year the reduced pH trend in summer after 2010 is confirmed from in-situ data (-0.0351 \pm 0.0114 .decade⁻¹ in 2001-2010 against -0.0078 \pm 0.0079 .decade⁻¹ in 2010-2020, 853 854 Figure 9b, Table 1). These results show that the pH trend varied significantly from decade to decade and that the 855 decrease of pH since 1985 was mainly driven by anthropogenic CO2. This is revealed in the winter water when 856 comparing pH and pre-industrial pH (Figure 10a). Here, the pre-industrial pH (pH-PI) was calculated after subtracting C_{ant} values from the observed C_T concentrations for each sample in the WW layer. Interestingly the 857 pH trend in the WW of -0.0161 (\pm 0.0033).decade⁻¹ (here deduced from the station A_T-C_T data in 1985-2021) is 858 859 very close to the long-term trend in surface from the FFNN model in 1985-2020 (-0.0165.decade⁻¹ \pm 0.0004). This trend is slightly faster than the pH trends of -0.0134 (\pm 0.001).decade⁻¹ recently estimated in subsurface 860







Figure 10: (a): Time-series of pH (red dots) and pre-industrial pH (pH-PI, black dots) estimated in the winter water layer
(WW around 200m, see figure 6) in 1985-2021 at station OISO-KERFIX (50°40'S-68°25'E). pH-PI for each sample was
calculated after subtracting C_{ant} to C_T. The pH trend from the present days is -0.0161 (± 0.0033).decade⁻¹ (red dashed line).
No trend is observed for pH-PI (black dashed). The mean pH-PI in the WW is 8.173 (± 0.020, n =45). (b): Profiles of pH and
pH-PI evaluated from March 1985 (black symbols) and February 2021 data (red symbols). The profiles for pH-PI are shown
below 150m only as C_{ant} estimates are not available in surface layer. Note that the pH-PI profiles are the same either using
1985 or 2021 data.

899 A for other properties (A_T, O₂, temperature, salinity and nutrients), the pre-industrial pH (pH-PI) does 900 not change over time in the WW (mean pH-PI = 8.173 ± 0.020 , n=45, Figure 10a). The pH-PI in the WW is in the range of the pre-industrial surface pH value in the Southern Ocean (8.2 for year 1750 and 8.18 for year 1850) 901 902 derived from Earth system Models (Jiang et al., 2023, their Table S9). In the WW at our location the modern pH (1985-2021) was on average -0.147 (± 0.021) lower than pre-industrial pH. In 1985 pH in the WW was -0.119 903 904 lower than pH-PI and in 2021 it was -0.184 lower than pH-PI (Figure 10a). The progressive decrease of pH was 905 clearly linked to Cant concentrations in the WW layer and the pH decrease identified below that layer in the water 906 column (Figure 10b).





909 3.3.2 Temporal change in the water column

From 1985 to 2021, signals of decreasing pH and increasing C_T in surface waters are propagated in the water column down to about 500m. As mentioned above the data in 1985 (first occupation of the station) reveal significant C_{ant} levels across the water column (Figure 6b). Therefore the pH down to 1400m was already lower in 1985 than at pre-industrial times (Figure 10b). However, the largest C_{ant} increases were found in the top layers and changes in pH from 1985 to 2021 were small below 500m (Figure 10b, Figure S14). While observations for all years fall on a common linear relationship between C_{ant} and pH_{ant} for depths greater than 500 m, the change in pH for a given level of C_{ant} increases with time for layers shallower than 500 m (Figure 11).



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932Figure 11: Anthropogenic pH (pHant) and anthropogenic Ωar (Ωarant) versus anthropogenic CO2 concentrations (Cant,933µmol.kg⁻¹) at station OISO-KERFIX (50°40'S-68°25'E). The data are selected in the layer 150-1600m for the periods 1985934(blue), 1993-2020 (black) and 2021 (red). The arrow identifies the data in the layer 150-500m (for Cant > 30 µmol.kg⁻¹).935Below 500m no change of Cant was observed from 1985 to 2021 and thus for pHant and Ωarant.936

937 The increase in C_{ant} concentrations over time (Figure 6b) also leads to a decrease of carbonate ion 938 concentrations $[CO_3^{2-}]$ and of Ω ar and Ω ca (Figure S14, S15). These decreases are well identified since the pre-939 industrial era in the whole water column but in the last 36 years, observations do not show any appreciable 940 changes below 500m (Figure 11). The aragonite saturation state (Ω ar=1) was found around 600m in 1985 and 941 around 400m in recent years (2015-2021, Figures S14, S15). Moreover, during the period covered by 942 observations (1985-2021), we did not detect abrupt change of the aragonite saturation horizon from one year to 943 the next (including from season to season, Figure S16). This contrasts with previous regional studies in the SO 944 and most notably with results from the layers close to the deep minimum of carbonate ion concentrations (Hauri 945 et al., 2015; Negrete-Garcia et al., 2019). At our station the [CO3²⁻] minimum lies around 500-600m (Figure S14, 946 S15) and, along with the superimposed Cant accumulation, explains the upward shift of the aragonite and calcite 947 saturation between the pre-industrial and modern periods (Figure S15). At pre-industrial time under-saturation 948 with regard to aragonite (Ω ar<1) was found at the bottom only (1600m) whereas in 1985-2021 it was found in 949 the water column below 600 m or 400 m (Figure S15). The subsurface pre-industrial Ω ar value was around 1.9-2 950 (Figure S15) and in the range of Ω ar value in the Southern Ocean at pre-industrial time from ESM models (Jiang 951 et al., 2023, their Figure 4).





952The aragonite under-saturation already occurred in 1985 at 500-600m (Figure S15) and a small increase953of C_T (via C_{ant} accumulation) close to the $[CO_3^{2^-}]$ minimum would rapidly shift the aragonite saturation horizon954in layers above 500m. This might have already occurred and explains that Ω ar value was 1.02 at 350m in 2021955(Figure S15). These results suggest that for pelagic calcifiers living in subsurface (150m or deeper) such as956pteropods and/or foraminifera (e.g. Hunt et al., 2008; Meilland et al., 2018) the impact of acidification might957occur sooner than in surface.

958For the interpretation of the trend analysis based on observations, only data below 150m could be used959as C_{ant} was not evaluated in the surface layer. At 200m, based on A_T - C_T data, pH and Ωar decreased from 1985960to 2021 by -0.059 for pH (Figure 10b) and -0.16 for Ωar (Figure S15). In 36 years, this represents about 30% of961the total change since the pre-industrial era (-0.184 for pH and -0.6 for Ωar at 200m). This is mainly linked to the962 C_{ant} change that represents also 30% increase in 36 years (+24.6 µmol.kg⁻¹ from 1985 to 2021 for a total of +71.7963µmol.kg⁻¹ CO₂ accumulated at 200m in 2021, Figure 7). We conclude that anthropogenic CO₂ drives the change964of the carbonate system in subsurface and probably also in surface waters.

965 In order to quantify the propagation of surface trends to depth, the temporal variations of carbonate 966 properties in the surface for both summer and winter derived from the FFNN model are compared to the changes 967 observed across the water column (Figure 12). The comparison shows that the seasonal amplitude of surface 968 waters properties was of a similar magnitude to the observed changes in the mixed layer between 1985 and 2021. For example, the C_T and Ω ar seasonality, respectively around 20 µmol.kg⁻¹ and 0.2, corresponds to the C_T 969 increase and Ωar decrease from 1985 to 2021. The comparisons also highlight that in summer the FFNN results 970 were close to observations in the mixed-layer (e.g. C_T was 2120 µmol.kg⁻¹ in 1985 and 2140 µmol.kg⁻¹ in 2021). 971 In winter the surface properties are different (C_T was higher, and pH, $[CO_3^2]$, Ω ar were lower) and intercept the 972 973 observations at depth close to the winter water (150-200m). This is true in 1985 and 2020/2021. Specifically, surface C_T from the FFNN model in winter 1985 (2145.5 $\mu mol.kg^{\text{-}1})$ equaled the C_T measured at 150 m in 974 975 March 1985 (2148 μ mol.kg⁻¹). In 2020, the winter C_T at the surface (2168.3 μ mol.kg⁻¹) is equal to C_T concentrations observed at 150-180 m in January 2020 or in 2021. For Ω ar, the surface value derived from the 976 977 FFNN model in winter 1985 (1.6) equal to the Ωar observed at 125 m in March 1985. In 2020, the surface winter 978 estimate of Ω ar (1.42) was equal to Ω ar observed at 100-150 m in January 2020 or 2021. The same 979 correspondences between winter surface and WW data were identified for pH and [CO3²⁻] (Figure 12). This 980 supports the use of winter and summer surface data from the FFNN model to investigate the seasonal Ω ar trends 981 and their projection in the future.

The surface water Ωar (Ωca) trend from the FFNN model in summer of -0.0059.yr⁻¹ (-0.0094.yr⁻¹) was 982 stronger than in winter -0.0050.yr⁻¹ (-0.0079.yr⁻¹) and also higher than derived from observations in the WW (-983 984 $0.0043.yr^{-1}$ for Ω ar and $-0.0069.yr^{-1}$ for Ω ca). The results indicate that the change of carbonate properties in the years 1985-2021 were mainly driven by Cant accumulation in surface waters and across the water column. 985 986 However, potential changes in primary productivity after 2010 mitigated the effects of increasing Cant 987 accumulation in response to increasing atmospheric CO_2 leading to relatively stable summer C_T and fCO₂ and to 988 a stronger CO₂ sink (Figure 3). Consequently, when restricted to the period 2010-2020, the trend of Ωar in surface waters in summer was much smaller, -0.024.decade⁻¹ (\pm 0.027) than during the preceding period. This 989 was much smaller than derived from the all data in 1985-2021 (-0.048.decade⁻¹) or estimated from reconstructed 990 991 fields in the SO-SPSS in 1982-2021 (-0.0616.decade⁻¹, Ma et al., 2023). It underscores the uncertainty in 992 extrapolating long-term time-series depending on the selection of data and periods.







1017Figure 12: Profiles (0-400m left axis) of observed and calculated properties (C_T , pH, Ω-ar, [CO3²]) at station OISO-KERFIX1018(50°40'S-68°25'E) in Mach 1985, January 2020 and January 2021 along with surface time-series in 1985-2020 (right axis) of1019the same properties in January (yellow line) and August (blue line) from the FFNN model. The FFNN values in January 20201020are coherent with January 2020 observations in the mixed-layer and in January 1985 are close to the observations in March10211985. Note that the differences of properties between 2020-21 and 1985 have a similar magnitude as the seasonal amplitude1023(illustrated by the FFNN values for January and August).

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1026 The data described above allowed evaluating the temporal variations of the properties of the carbonate 1027 system and C_{ant} over 1985-2021 along with a comparison to the pre-industrial state in the water column 1028 excluding the surface layer. The results over 36 years informed on the recent changes, inter-annual variations 1029 and trends, but the time-series appears somehow short to extrapolate the trends over time. What was the change 1030 of the carbonate system in surface water before 1985 and what will be its future evolution ?

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1032 3.4.1 Back to the sixties: observed trends since 1962.

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1034To explore the long-term change, we start by comparing our recent data with the observations from the1035LUSIAD cruise conducted in 1962-1963 (Keeling and Waterman, 1968). Some data from this cruise were1036obtained mid-November 1962 south of the Polar Front, in the region south-west off Kerguelen. Because of the1037seasonality, we compared the November 1962 data with our observations obtained in October-November in

^{1024 3.4} Long-term change in surface water, from the sixties to the future.







1062Figure 13: Observed (black dots) sea surface temperature (°C), fCO2 (µatm), C_T (µmol.kg⁻¹), pH (TS), Ω-ar and Ω-ca around1063station OISO-KERFIX at 50°40'S-68°25'E for October-November. Also shown are the results for the FFNN model for1064November in 1985-2020 (Purple). The C_T concentrations, pH, Ω-ar and Ω-ca were calculated from fCO2 data using the A_T/S1065relation (Eq. 1). The red line is the atmospheric fCO2 and red dashed-lines in each plot are the evolution of properties since10661960 corrected to C_{ant} where fCO2, pH, Ω-ar and Ω-ca were recalculated using C_T+C_{ant}, A_T constant at 2290 µmol.kg⁻¹ andSST at 2°C. Grey triangles identified the mean values for C_T and pH.

1069 First, we note that measured SST in November 1962 (1.7°C) was slightly lower (on average about -1070 0.6°C) compared to recent years, but SST as low as 1.8°C for this season were also recorded in other periods (e.g. November 1995, 2014). The change in SST is unlikely to explain the long-term increase in fCO2 or 1071 1072 decrease in pH since 1962 (Figure 13). In 1962, the ocean fCO₂ was 324 µatm slightly higher than in the atmosphere (ΔfCO_2 =+8 µatm, a small source), whereas in November 1985-2020 the ocean was a small CO₂ sink 1073 1074 on average ($\Delta fCO_2 = -3.3 \pm 4.5 \mu atm$). The C_T concentration in 1962 (2135 $\mu mol.kg^{-1}$) was much lower than 1075 observed in the 90s and the pH (8.115) much higher than in recent years (Figure 13). Compared to 1962, pH in 1076 2016 was -0.078 lower, i.e. representing 70% of the pH decrease of -0.11 in the global ocean since the beginning of the industrial era (Jiang et al, 2019). In November 1962, surface C_T was lower by -15.1 μ mol.kg⁻¹ compared to 1077 the data in October 1995, i.e. a trend of $+0.46 \ \mu mol.kg^{-1}.yr^{-1}$ in 33 years close to the C_{ant} trend observed in the 1078 1079 WW in 1985-2021 as described above (+0.53 \pm 0.01 μ mol.kg⁻¹.yr⁻¹). Having the C_T value in 1962, we can 1080 project the C_T in time by adding the C_{ant} concentration based on the relationship observed between C_{ant} and 1081 atmospheric CO₂ (Figure 7b) assuming that the anthropogenic CO₂ uptake since the sixties is representative of 1082 the C_T change (i.e. the change of C_T due to natural variability is small). This projection is shown for all 1083 properties (red dashed-lines in Figure 13) and confirms that the progressive C_{ant} accumulation explained most of





1084 the C_T and fCO₂ increase in surface since 1962. We note that the C_T derived from the FFNN model suggests 1085 slightly lower C_T compared to the C_{ant} projection especially before 2004. The difference of projected C_T and the 1086 FFNN model (on average $-2.2 \pm 2.7 \ \mu mol.kg^{-1}$) is within the uncertainty of C_T calculations (error is $\pm 5 \ \mu mol.kg^{-1}$) when using the A_T/fCO_2 pairs) and the trend of the difference over 1985-2020 (-0.15 μ mol.kg⁻¹.yr⁻¹) is too small 1087 1088 to be related with confidence to changes associated with natural processes. On the other hand, the ocean fCO_2 1089 recalculated with the projected Cant trend suggested that for this season (November) the ocean moved from a CO2 source in 1962-1985 ($\Delta fCO_2 > 0$) to a sink in 1986-2021 ($\Delta fCO_2 < 0$) in line with results from the FFNN model. 1090 1091 The recalculated fCO₂ with C_{ant} (dashed red line in Figure 13) was close to that observed in 1995 or from the 1092 FFNN model in 1985-2014 (mean difference over 1985-2014 is -1.2 \pm 5.2 µatm). After 2016, the recalculated 1093 fCO2 suggest a stronger sink and the difference with observations in 2011 and 2016 or the FFNN model is 1094 slightly higher (mean difference over 2016-2020 is $-8.8 \pm 1.5 \mu$ atm). Although the differences are in the range of 1095 the error in fCO₂ calculation using A_T - C_T pairs (± 13 µatm), this might indicate that after 2016 a process could 1096 contribute to increase fCO_2 faster than the effect of C_{ant} only. This difference could be due to the warming that 1097 occurred after 2016 when SST was higher than 2°C and up to 3°C in November 2017 (Figure 13 and Figure S9). 1098 The same could be applied for pH that was slightly lower than the pH recalculated from Cant trend after 2015 (the 1099 mean difference between recalculated pH and FFNN-pH over 1985-2020 is only 0.002 ± 0.006). Therefore, we 1100 conclude that for November the pH decrease since 1962 was mainly driven by anthropogenic CO2. Aragonite 1101 and calcite saturation states also show a clear decrease since 1962 (Figure 13), a diminution of 11% in 59 years 1102 for both Ω ar and Ω ca. Based on these results over almost 60 years that confirm the conclusions from the 1103 observations in 1985-2021, we now evaluate the long-term change of the carbonate system in surface water in 1104 the future.

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1106 3.4.2 Projecting the observed trends in the future

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1108 The trends of the properties based on observations in 1962-2021 and the FFNN model in 1985-2020 1109 indicate relatively linear trends linked to Cant uptake albeit with some decadal variability in summer (Figure 4). A 1110 simple linear extrapolation of the trends in the future suggests that aragonite saturation in surface water would be reached in year 2110 for the winter season and 2120 for summer (Figure S17) whereas the trend in subsurface 1111 suggests under-saturation in 2090. In year 2100, surface pH and $[H^+]$ would be around 7.9 and 12 nmol.kg⁻¹ 1112 1113 (Figure S17). However, ESM CMIP6 models suggest that under a high emission scenario (SSP5-8.5), pH in 1114 2100 in the Southern Ocean near 50°S would be around 7.65 and $[H^+]$ around 22 nmol.kg⁻¹ (Jiang et al., 2023, their figure 4). This suggests that the simple linear extrapolation based on recent observed trends (Figure S17) 1115 1116 underestimated the future change of the carbonate system for a high emission scenario as previously shown in 1117 the South-Eastern Indian Ocean based on summer trends derived from observations in 1969-2003 (Midorikawa 1118 et al., 2012, their figure 4).

1119 To better investigate the changes in the next decades, we assumed that the C_{ant} trend for the modern 1120 period (Figure 7) that experienced a "business as usual" scenario after the sixties is representative of the future 1121 changes in the surface ocean carbonate system. For this analysis, we use two emissions scenarios (Shared 1122 Socioeconomic Pathways, SSP, Meinshausen et al., 2020) with atmospheric xCO₂ reaching 1135 ppm in 2100 (a 1123 "high" emission scenario SSP5-8.5) or xCO₂ reaching 603 ppm in 2100 after a stabilization around 2080 1124 (scenario SSP2-4.5). This enables to simulate future C_T concentrations for summer or winter (Figure 14) and to





1125 calculate other carbonate properties using C_T and A_T (Figure 15, Table 2) in response to approximated future 1126 changes in physical and geochemical properties excluding impacts of changes in atmospheric and oceanic 1127 circulation. As the calculated properties are sensitive to A_T values, we used a fixed A_T of 2280 μ mol.kg⁻¹ or 1128 applied a correction based on the long-term change of sea surface salinity observed in the last 6 decades (1960-1129 2017), i.e. a freshening in the Southern Ocean of around -0.01 to -0.02.decade⁻¹ (Durack and Wijffels, 2010; 1130 Cheng et al., 2020b). The decrease in salinity in the South Indian Ocean (-0.02.decade⁻¹ \pm 0.01) was recently 1131 analyzed by Akhoudas et al. (2023) who showed that in the years 1993-2021 the freshening was mainly due to 1132 an increase in the precipitation linked to the acceleration of the atmospheric hydrological cycle. From our data in the mixed-layer over 1985-2021, we estimated a trend in salinity of -0.0207.decade⁻¹ (\pm 0.0041). For the A_T 1133 1134 sensitivity test we thus select a salinity trend of -0.01.decade⁻¹ in 1962-1985 and -0.02.decade⁻¹ after 1985 and 1135 apply these trends to simulate A_T over 1960-2100 using the A_T /Salinity relationship (Equation 1). This leads to a 1136 salinity of 33.650 and A_T of 2272 µmol.kg⁻¹ in year 2100, about 8 µmol.kg⁻¹ lower than observed in 2021 (2280 μ mol.kg⁻¹). Compared to the C_T change from 2021 to 2100 (+50 and +193 μ mol.kg⁻¹ for the "low" and "high" 1137 1138 emissions scenario, Figure 14), the impact of A_T decrease has a minor effect on the future change for pH, $[CO_3^{2^-}]$ 1139 or Ω (Table 2). For example, in winter for the SSP5-8.5 scenario, when the A_T decrease is taken into account, pH 1140 in 2100 is 7.316 and Ω_{Ar} is 0.33 against 7.372 and 0.34 when A_T is constant (Table 2). In both cases, the 1141 aragonite saturation ($\Omega_{Ai}=1$) in winter occurred in 2055, whereas in summer it is identified in 2070. The effect of 1142 lower A_T in the future appeared also small compared to the seasonal differences of pH and Ω in 2100.

1143 As noted above, the Southern Ocean experienced a warming in recent decades (e.g. Auger et al., 2021) 1144 and it is projected that warming will continue in the future (IPCC, 2022). Therefore, to test the sensitivity of 1145 calculated properties to warming we applied a correction of $+0.0125^{\circ}$ C.yr⁻¹ in 1985-2020 and $+0.025^{\circ}$ C.yr⁻¹ after 1146 2020 (Azarian et al, 2023). As for A_T, these results are compared for winter using constant SST (Table 2). The 1147 effect of the long-term warming does mainly impact the projection of [H⁺] and pH (Table 2).

These sensitivity tests for temperature and A_T showed that as for the observed period 1962-2021 (Figure 1148 1149 13), the projection in the future depends mainly on the anthropogenic CO_2 accumulation. Here, the C_T 1150 concentrations were calculated using the Cant versus atmospheric CO2 relationship (Figure 7b). We thus tested 1151 the results for winter based on the error associated with this relationship (Figure S18). This leads to either higher 1152 or lower C_T compared to original calculation (Figure 14). For the SSP5-8.5 scenario, the winter C_T 1153 concentrations in 2100 range between 2328 and 2378 µmol.kg⁻¹, higher than simulated in the ESM CMIP6 models around 50°S (2300 µmol.kg-1, Jiang et al., 2023). As in the ESM models for the SSP2-4.5 scenario, the 1154 projected C_T concentration in 2100 at our location is much lower 2217 μ mol.kg⁻¹ (Figure 14). The future change 1155 of the carbonate system is not significantly different using low or high Cant accumulation (Figure S18) but this 1156 1157 test gives a range of years to reach aragonite and calcite under-saturation. In winter (SSP5-8.5 scenario), 1158 aragonite would reach under-saturation between year 2050 and 2060 and between year 2070 and 2080 for 1159 calcite. Note that for summer we derived under-saturation for Ω_{Ar} in year 2065 and for Ω_{Ca} in year 2085. For the SSP2-4.5 scenario, where C_T is 143 µmol.kg⁻¹ lower in 2100 compared to SSP5-8.5, aragonite under-saturation 1160 1161 would not be reached before 2070 (Figure 15).





Method	Year	Atm-CO ppm	2 fCO ₂ μatm	C _T µmol	A _T kg ⁻¹	pH TS ni	[H ⁺] nol.kg ⁻¹	[CO ₃ ²⁻] µmol.k	Ωca g ⁻¹	Ωar
Obs Jan	2020	410.6	391.9	2142.2	2281.8	8.044	9.04	105.2	2.53	1.59
Std obs.			(2.0)	(0.7)	(0.3)	(0.002)	(0.04)	(0.5)	(0.01)	(0.01)
FFNN Jan	2020	410.6	385.1	2138.5	2280.1	8.051	8.90	106.3	2.55	1.61
SSP Summer	2020	414.9	375.4	2137.5	2282.1	8.061	8.70	108.0	2.60	1.63
FFNN Aug	2020	410.6	410.0	2168.3	2289.8	8.024	9.45	94.2	2.27	1.42
SSP Winter	2020	414.9	434.5	2167.3	2282.1	8.001	9.98	90.4	2.18	1.37
SSP85 Summer	2050	562.8	526.5	2177.2	2278.3	7.928	11.79	84.2	2.02	1.28
SSP85 Winter	2050	562.8	624.7	2207.0	2278.3	7.857	13.91	68.5	1.65	1.04
SSP85 W-A-T	2050	562.8	585.7	2207.0	2280.0	7.880	13.17	69.0	1.66	1.04
SSP85 W-T	2050	562.8	592.7	2207.0	2278.3	7.875	13.32	68.1	1.64	1.03
SSP45 Winter	2050	506.9	554.8	2192.0	2278.3	7.905	12.46	75.8	1.92	1.15
SSP85 Summer	2100	1135.2	1986.9	2330.6	2271.8	7.394	41.31	26.9	0.65	0.41
SSP85 Winter	2100	1135.2	2306.3	2360.4	2271.8	7.316	48.26	21.8	0.52	0.33
SSP85 W-A-T	2100	1135.2	1993.1	2360.4	2280.0	7.372	42.44	22.6	0.54	0.34
SSP85 W-T	2100	1135.2	2097.0	2360.4	2271.8	7.349	44.74	21.3	0.51	0.32
SSP45 Winter	2100	602.8	7539	22177	2271.8	7 782	16 51	60.9	1 47	0.92



Figure 14: Evolution of atmospheric CO₂ (ppm) and sea surface C_T (µmol.kg⁻¹) in 1960-2110 evaluated for 2 scenarios (SSP2-4.5 black dashed and SSP5-8.5 red dashed), for summer (red line for SSP5-8.5) and winter (blue line for SSP5-8.5 and green line for SSP2-4.5). Grey lines are the high and low C_T for winter SSP5-8.5 based on the error in the C_{ant}/fCO_2 relationship (figure 7b). Also shown are the results for the FFNN model in 1985-2020 for summer (red diamonds) and winter (blue diamonds) and C_T in 1962 (black diamonds). The C_T values for different seasons and scenarios were used to calculate the carbonate properties in the future (Figure 15).







1246Figure 15: Evolution of sea surface fCO2 (μ atm), pH (TS), [CO3²⁻](μ mol.kg⁻¹), [H⁺] (nmol.kg⁻¹), Ω -Ar and Ω -Ca in 1960-12472110 evaluated for the SSP5-8.5 scenario for winter (blue line) and summer (red line) taking into account both A_T and SST1248future trends. For winter results are also presented using the SSP2-4.5 scenario (green line). Also shown are the results for the1249FFNN model in 1985-2020 for summer (red diamonds) and winter (blue diamonds). Atmospheric fCO2 is also shown for1250SSP5-8.5 (red dashed) and SSP2-4.5 (green dashed). Values in 2020, 2050 and 2100 for different sensitivity tests are listed in
Table 2.1251Table 2.

1254 4 Summary and concluding remarks

1256 The times-series of high quality observations in 1985-2021 and the results from the FFNN model at one 1257 location, south of the Polar Front in the Southern Indian Ocean ($50^{\circ}S-68^{\circ}E$) presented in this analysis offered 1258 new results on the inter-annual variability, decadal to long-term trends of the carbonate system in surface waters, 1259 air-sea CO₂ fluxes and associated drivers. The evaluation of anthropogenic CO₂ concentrations in the water 1260 column indicates that the trends of the carbonate species are mainly driven by the CO₂ uptake leading to a 1261 progressive acidification in surface waters and at depth.

1262 In 1985, the C_{ant} concentrations were approaching 50 µmol.kg⁻¹ at 200 m and C_{ant} was detected in the 1263 water column down to the bottom (1600m). This explains why aragonite under-saturation was observed at 1264 around 600m in 1985, where $[CO_3^{2^-}]$ concentration was at minimum, whereas at pre-industrial era the water 1265 column was super-saturated (this study Figure S15; Lauvset et al., 2020, their Figure S15). 36 years later because 1266 of the anthropogenic CO₂ accumulation, we observed an upward migration of the aragonite saturation horizon 1267 that was found around 400 m in 2021 (a shoaling rate of around -6 m.yr⁻¹).

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1268 At subsurface, in the winter water layer, the C_{ant} trend is estimated at +0.53 (± 0.01) µmol.kg⁻¹.yr⁻¹ in 1269 1985-2021 with a detectable increase in recent years (up to 72 µmol.kg⁻¹ in 2021 compared to 47 µmol.kg⁻¹ in 1270 1985). The C_{ant} concentrations in the ocean are closely related to the atmospheric CO₂ concentrations and the 1271 slope we observed south of the PF in the Indian sector of +0.263 ± 0.042 µmol.kg⁻¹.µatm⁻¹ is close to that 1272 observed in the AAIW in the South Atlantic (+0.23 ± 0.05 µmol.kg⁻¹.µatm⁻¹, Fontela et al., 2021). This suggests 1273 that local observations in the South Indian POOZ captured the link between C_{ant} and atmospheric CO₂ at larger 1274 scale.

1275 In surface waters, over 1991-2020 the oceanic fCO_2 increased at a rate close or slightly lower than in 1276 the atmosphere (Figure 2b) and the C_T trend followed the C_{ant} accumulation (Figure 4b, S12a). However in the 1277 last decade both observations and the FFNN model showed low fCO2 trends in summer (less than 1 µatm.yr⁻¹). 1278 The change in summer trend appears related to primary production as revealed by a decrease of Chl-a in 1998-1279 2010 followed by an increase after 2010. Biological activity counteracts the C_T increase due to C_{ant}, resulting in rather stable C_T and fCO_2 in summer 2010-2020 (+0.38± 0.26 μ mol.kg⁻¹.yr⁻¹ and +0.98 ± 0.40 μ atm.yr⁻¹). As a 1280 result, the region moved from an annual source of +0.8 molC.m⁻².yr⁻¹ in 1985 to a sink of -0.5 molC.m⁻².yr⁻¹ in 1281 1282 2020. The increase of the ocean CO_2 sink was particularly pronounced after 2011 (Figure 3) when phytoplankton 1283 biomass was stronger in this HNLC region and occurred when the SAM index was in a positive state.

1284 In 1959-1963, the SAM was also positive on average and moved to a negative phase in 1964 (Marshall, 2003; King et al., 2023). Historical data from 1962 suggest that in November the region was a small CO₂ source 1285 $(\Delta fCO_2 = +8 \mu atm)$. Assuming the seasonality was the same as in the 80s, we estimate that in 1962 the annual flux 1286 would be around 2.2 molC.m⁻².yr⁻¹. Extrapolating to the entire South Indian POOZ (50-58°S/20-120°E, 6.5 1287 Mkm²), this region was a CO₂ source of 0.17 PgC.yr⁻¹ in 1962, reduced to 0.06 PgC.yr⁻¹ in 1985 and a CO₂ sink 1288 1289 of -0.04 PgC.yr⁻¹ in 2020. This could be compared with reconstructed fluxes from a data-based model that 1290 produced a CO₂ source in 1960-1990 and a sink in 2020 in the south Indian sector (Rödenbeck et al., 2022, their 1291 Figure 6).

1292 For November 1962, the estimated C_T concentration in surface (2135 μ mol.kg⁻¹) is 21 μ mol.kg⁻¹ lower 1293 than observed mid-October 2016 in the mixed-layer (2156 µmol.kg⁻¹). This is almost equal to the increase of C_{ant} 1294 in 54 years (+22.3 µmol.kg⁻¹). As a result, surface ocean pH dropped from 8.11 in 1962 to 8.044 in 2020. Over 1295 multi-decadal time scale (30 years or more), acidification in the South Indian POOZ has been mainly controlled 1296 by uptake of anthropogenic CO₂. However, our data also indicate a modulation of the summer pH trend by 1297 natural processes. After 2010, a very small pH trend was estimated in summer (-0.0098.decade⁻¹ ± 0.0042) when the region experienced higher primary productivity. On the opposite, in winter, the pH trends continuously 1298 1299 increased with time, -0.010.decade⁻¹ (± 0.001) in 1991-2001 and -0.021.decade⁻¹ (± 0.002) in 2010-2020. In the 1300 subsurface (winter water layer), the trend of pH based on A_T - C_T data in 1985-2021 of -0.0161 (± 0.0033).decade 1301 ¹ is also almost equal to the annual surface trend from the FFNN model. A simple extrapolation of the trends in 1302 the WW indicated that under-saturation (Ω <1) would be reached at year 2090 for aragonite and year 2180 for 1303 calcite. However, as atmospheric CO₂ will desperately continue to rise and ocean C_T will increase in the future, 1304 the pH and Ω will decrease at a faster rate than observed in recent years. A projection of future C_T concentrations 1305 based on emissions scenario, excluding changes in ocean circulation, indicated that the winter surface pH in 1306 2100 would decrease to 7.32 for a high emission scenario (SSP5-8.5) or to 7.782 for a low emission scenario 1307 (SSP2-4.5). This is up to -0.86 lower than pre-industrial pH and -0.71 lower than pH observed in 2020. For the





winter season the aragonite saturation in surface would be reached around 2050 for a high emissions scenarioand 2070 for a low emission scenario.

1310 The time-series presented here for the Southern Ocean, along with other historical time-series of A_T-C_T in the water-column (BATS, HOT, ESTOC, KNOT, Iceland or Irminger seas; Bates et al., 2014; Lange et al., 1311 1312 2023) or the recent BG-Argo floats in the Southern Ocean (Mazloff et al., 2023) offer useful data for the 1313 evaluation of biogeochemical and Earth system models, especially the coupling of fCO2, CT, AT, and pH not well 1314 represented in current models at seasonal to decadal scale in the Southern Ocean (e.g. Hauck et al., 2023; 1315 Rodgers et al., 2023; Joos et al., 2023). Observing the decadal changes of the carbonate system in the water 1316 column is also an important step to extend the evaluation of biogeochemical and ESM models below the surface 1317 (Jiang et al., 2023). It is important to maintain such time-series for monitoring the future evolution of the ocean 1318 CO₂ sink, of the acidification and its impact on phytoplankton species and higher trophic levels. This is 1319 especially the case in Marine Protected Area such as the French Sub-Antarctic islands including the Kerguelen 1320 Archipelago which was listed as a UNESCO World Heritage site in 2019.

1322 Data availability:

Data used in this study are available in SOCAT (<u>www.socat.info</u>) for fCO₂ surface data, in GLODAP
 (<u>www.glodap.info</u>) for water-column data and at NCEI/OCADS (<u>www.ncei.noaa.gov/access/ocean-carbon-data-</u>
 <u>system/oceans/VOS Program/OISO.html</u>). The CMEMS-LSCE-FFNN model data are available at E.U.
 Copernicus Marine Service Information (https://resources.marine.copernicus.eu/products).

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1328 Authors contributions:

1329 CLM and NM are co-I of the ongoing OISO project. CLM, NM, CL and CR participated to OISO cruises. 1330 Underway fCO₂ was measured by CLM, NM, CL, and qualified by CLM and NM. Nutrients data were measured 1331 and qualified by CLM and CL. Chl-a data were measured and qualified by CR. Water column data were 1332 qualified by CLM, NM, CL, CR and GR. MG, FC and TTTC developed the CMEMS-LSCE-FFNN model and 1333 provided the model results. NM started the analysis, wrote the draft of the manuscript and prepared the figures 1334 All authors contributed to revising the draft manuscript.

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1336 Competing interest: The authors declare that they have no conflict of interest.

1337

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1351	and Ecosystem Research program (IMBER), to deliver a uniformly quality-controlled surface ocean CO2
1352	database.
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