

# How Well Do We Understand the Land-Ocean-Atmosphere Carbon Cycle?

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## Abstract

Fossil fuel combustion, land use change and other human activities have increased the atmospheric carbon dioxide ( $\text{CO}_2$ ) abundance by almost 50% since the beginning of the industrial age. These changes would have been much larger if natural sinks in the land biosphere and ocean had not removed over half of this anthropogenic  $\text{CO}_2$ . Here, we review the current state of knowledge of the ocean, land and atmospheric carbon cycles, identify emerging measurement and modeling capabilities, and gaps that must be addressed to diagnose the processes driving the carbon cycle and predict their response to human activities and a changing climate. The anthropogenic  $\text{CO}_2$  uptake by the ocean has increased over this period, as the atmospheric  $\text{CO}_2$  partial pressure ( $\text{pCO}_2$ ) has increased. For the land carbon cycle, the emerging picture is more complicated. Over the past three decades, the uptake of  $\text{CO}_2$  by intact tropical humid forests appears to be declining, but these effects are offset by a net greening across mid- and high-latitudes associated with afforestation, agricultural, and longer growing seasons. These studies have also revealed measurement gaps and other limitations in our understanding of the evolving carbon cycle. They show that continued ship-based observations combined with expanded deployments of autonomous platforms are needed to quantify ocean-atmosphere fluxes on policy relevant spatial and temporal scales. They also reinforce the urgent need for more comprehensive measurements of stocks, fluxes and atmospheric  $\text{CO}_2$  in humid tropical forests and across the Arctic and boreal regions, which appear to be experiencing rapid change.

## How Well Do We Understand the Land-Ocean-Atmosphere Carbon Cycle?

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### Key Points:

- Carbon dioxide (CO<sub>2</sub>) emissions from fossil fuel use and other human activities have increased its atmospheric abundance by almost 50% during the industrial age.
- These increases would have been much larger if natural sinks in the ocean and on land had not removed over half of this CO<sub>2</sub>, maintaining an airborne fraction < 45%.
- New measurements and models are providing insights into the carbon cycles as the land and ocean sinks continue to respond to human activities and climate change.

## Abstract

Fossil fuel combustion, land use change and other human activities have increased the atmospheric carbon dioxide (CO<sub>2</sub>) abundance by almost 50% since the beginning of the industrial age. These changes would have been much larger if natural sinks in the land biosphere and ocean had not removed over half of this anthropogenic CO<sub>2</sub>. Here, we review the current state of knowledge of the ocean, land and atmospheric carbon cycles, identify emerging measurement and modeling capabilities, and gaps that must be addressed to diagnose the processes driving the carbon cycle and predict their response to human activities and a changing climate. The anthropogenic CO<sub>2</sub> uptake by the ocean has increased over this period, as the atmospheric CO<sub>2</sub> partial pressure (pCO<sub>2</sub>) has increased. For the land carbon cycle, the emerging picture is more complicated. Over the past three decades, the uptake of CO<sub>2</sub> by intact tropical humid forests appears to be declining, but these effects are offset by a net greening across mid- and high-latitudes associated with afforestation, agricultural, and longer growing seasons. These studies have also revealed measurement gaps and other limitations in our understanding of the evolving carbon cycle. They show that continued ship-based observations combined with expanded deployments of autonomous platforms are needed to quantify ocean-atmosphere fluxes on policy relevant spatial and temporal scales. They also reinforce the urgent need for more comprehensive measurements of stocks, fluxes and atmospheric CO<sub>2</sub> in humid tropical forests and across the Arctic and boreal regions, which appear to be experiencing rapid change.

## Plain Language Summary

Since the beginning of the industrial age in the mid-1700s, fossil fuel combustion, land use change and other human activities have increased the atmospheric carbon dioxide (CO<sub>2</sub>) concentration to levels never before seen in human history such that CO<sub>2</sub> is now the primary driver of climate change. The atmospheric CO<sub>2</sub> abundance and its effect on the climate would have been much larger if natural processes in the ocean and on land carbon cycle had not absorbed over half of the CO<sub>2</sub> emitted by these human activities. Here, we review our understanding of anthropogenic CO<sub>2</sub> emissions and the processes controlling the emissions and uptake of CO<sub>2</sub> by the natural carbon cycle as it responds to continuing human activities a changing climate.

## 1 Introduction

Since the beginning of the industrial age, human activities have increased the atmospheric concentrations of carbon dioxide (CO<sub>2</sub>) and other greenhouse gases (GHGs) to levels never before seen in human history. These large increases are driving climate change, because CO<sub>2</sub> is an efficient greenhouse gas with an atmospheric lifetime measured in centuries. Bottom-up statistical inventories indicate that fossil fuel combustion, industry, agriculture, forestry, and other human activities are now adding more than 11.5 petagrams of carbon (Pg C) to the atmosphere each year (Friedlingstein et al., 2019; 2020). Direct measurements of CO<sub>2</sub> in the atmosphere and in air bubbles in ice cores indicate that these and other human activities have increased the globally averaged atmospheric CO<sub>2</sub> dry air mole fraction from less than 277 parts per million (ppm) in 1750 to more than 410 ppm today. Measurements from Mauna Loa

64 Observatory show that over half of this increase has occurred since 1985 and over a quarter has  
65 happened within the past 15 years (Dlugokencky et al., 2018).

66 These increases would be much larger if processes operating in the natural carbon cycle  
67 had not removed over half of these anthropogenic CO<sub>2</sub> emissions. Carbon cycle measurements  
68 and modeling studies show that these anthropogenic CO<sub>2</sub> emissions are superimposed on an  
69 active natural carbon cycle that regulates CO<sub>2</sub> through photosynthesis and respiration on land  
70 and in the ocean (Beer et al., 2010), as well as temperature-driven solubility coupled with the  
71 ocean circulation (Takahashi et al., 2002; 2009). The land biosphere and ocean emit almost 20  
72 times as much CO<sub>2</sub> into the atmosphere as human activities each year, but then reabsorb a  
73 comparable amount along with about half of the anthropogenic CO<sub>2</sub> emissions (Jones and Cox,  
74 2005; Canadell et al., 2007; Raupach et al., 2008; Knorr 2009; Bennedsen et al., 2019,  
75 Friedlingstein et al., 2020). While the fraction of the anthropogenic CO<sub>2</sub> that stays in the  
76 atmosphere (the “airborne fraction”) is remarkably constant, at about 0.45 for the multi-year  
77 average (e.g. Ballantyne et al., 2012; Raupach et al., 2008; 2014), it can change substantially  
78 from year to year. In some years, the airborne fraction can be as high as 80%, while in others, it  
79 can be as low as 30% (Raupach et al., 2008; 2014). Some of the largest changes in this airborne  
80 fraction appear to be associated with changes in uptake of CO<sub>2</sub> by the land biosphere (the land  
81 sink) in response to large-scale temperature and precipitation anomalies, like those associated  
82 with major El Niño events or large volcanic aerosol injections into the stratosphere (Frölicher et  
83 al., 2013). The ocean uptake also responds to El Niño events and large volcanic eruptions  
84 (Eddebbar et al., 2019; McKinley et al., 2004; 2020), but has a smaller impact on the amplitude  
85 of variability in the airborne fraction (Chatterjee et al., 2017). The relative roles of these and  
86 other processes that link the land, ocean and atmospheric carbon cycles with the climate are less  
87 well understood, compromising our ability to predict how the atmospheric CO<sub>2</sub> growth rate  
88 might change as the carbon cycle responds to climate change (Ballantyne et al., 2012, McKinley  
89 et al., 2017).

90 Over the past two decades, our understanding of the natural and anthropogenic  
91 contributions to the carbon cycle has grown steadily with the deployment of progressively more  
92 sophisticated ground-based, oceanic, airborne, and space-based carbon cycle measurement  
93 systems. These advances have been accompanied by the development of far more comprehensive  
94 diagnostic and prognostic carbon cycle modeling tools. On time scales of a decade or longer,  
95 “bottom-up” measurements of changes in carbon stocks on land and in the ocean provide a  
96 reliable indicator of the stability of these reservoirs and a useful integral constraint on their  
97 fluxes of carbon to the atmosphere. On shorter time scales, stock measurements are less reliable  
98 for estimating atmospheric emissions because land and ocean stock changes that are too small to  
99 be accurately quantified can produce large changes in the atmospheric CO<sub>2</sub> concentration. On  
100 these scales, direct measurements of CO<sub>2</sub> fluxes at the surface, fluxes derived from vertical  
101 gradients in pCO<sub>2</sub> across the air-sea interface or “top-down” fluxes estimated from spatial and  
102 temporal gradients in atmospheric CO<sub>2</sub> concentrations can provide more reliable results.

103 Both bottom-up stock and flux estimates and “top-down” atmospheric estimates are  
104 providing key insights into the carbon cycle. Bottom-up methods use empirical or process-based  
105 models of the land biosphere and ocean to estimate fluxes, or to upscale in situ measurements of  
106 the time change of stocks or of direct flux observations (e.g. Sabine et al., 2004; Doney et al.,  
107 2004; Gruber et al., 2019a; Pan et al., 2011; Sitch et al., 2015; Hubau et al., 2020; Piao et al.,  
108 2020; Jung et al., 2020; Long et al., 2013; Landschützer, et al., 2013; Rödenbeck et al., 2014;

2015; Gregor et al., 2019; Watson et al., 2020; Hauck et al., 2020; Carroll et al., 2020). “Top-down” models use inverse methods to estimate the surface CO<sub>2</sub> fluxes from the land or ocean needed to maintain the observed atmospheric or ocean CO<sub>2</sub> concentrations in the presence of the prevailing winds and ocean circulation (e.g. Enting et al., 1995; Mikaloff-Fletcher et al., 2006; Jacobson et al., 2007; Khatiwala et al., 2009; Chevallier et al., 2010; 2019; DeVries 2014; Crowell et al., 2019).

Both bottom-up and top-down methods benefit from remote sensing as well as in situ data. For example, a bottom-up forest stock inventory might use in situ measurements to estimate the above ground biomass from an ensemble of specific plots and then use remote sensing measurements to upscale those measurements to larger areas. Similarly, a top-down approach might combine in situ and remote sensing observations of atmospheric CO<sub>2</sub> along with models of atmospheric transport constrained by both in situ and satellite meteorological measurements to estimate regional-scale fluxes. In practice, top-down and bottom-up methods can be combined. For example, top-down methods for estimating net biospheric exchange (NBE) may use fossil fuel emissions derived from bottom-up inventories as a static (i.e. not optimized) input (i.e. Crowell et al., 2019).

As the world embarks on efforts to monitor and control CO<sub>2</sub> emissions, there is growing evidence that the natural carbon cycle is evolving in response to human activities, severe weather, disturbances and climate change. In this context, an improved understanding of both the anthropogenic and natural processes in the land biosphere and ocean that control the emissions and uptake of CO<sub>2</sub> is critical to our ability to predict its rate of increase in the CO<sub>2</sub> in the atmosphere and its impact on the climate. To address this need, advanced measurement capabilities are being deployed and their results are being analyzed with more comprehensive diagnostic and prognostic modeling tools. While these advances do not yet provide the full capabilities of the policy-relevant carbon observing system advocated by Ciais et al. (2014) or envisioned by the WMO Integrated Global Greenhouse Gas Information System (IG3IS), they are providing substantial new insights with respect to the anthropogenic and natural processes operating in the carbon cycle.

Here, we review the current state of knowledge of the anthropogenic and natural processes driving the land, ocean and atmospheric carbon cycles. We summarize advances in each domain and identify ongoing efforts to exploit this new information to create a more accurate and complete baseline and to identify trends. We also identify critical measurement and modeling gaps that must be addressed to produce an effective carbon monitoring system.

## 2 A Note on Units

Because the bottom-up and top-down land, ocean and atmospheric carbon communities focus on different aspects of the carbon cycle, they have developed a sometimes confusing array of units to quantify stocks and fluxes of carbon and CO<sub>2</sub>. For example, the land carbon community typically quantifies the mass of stocks and fluxes carbon, the atmospheric community typically measures and reports the CO<sub>2</sub> dry air mole fraction,  $X_{\text{CO}_2}$  and the ocean

Table 1: Quantities and Units Describing the Carbon Cycle

Quantity	Acron ym	Typical units	Comment
carbon dioxide dry air mole fraction	CO <sub>2</sub> , xCO <sub>2</sub> or $X_{\text{CO}_2}$	parts per million by volume (ppm)	Number of CO <sub>2</sub> molecules relative to each million (10 <sup>6</sup> ) molecules of dry air. If CO <sub>2</sub> is assumed to be an ideal gas and its dry air mole fraction is increased by 1 ppm at constant temperature, the CO <sub>2</sub> partial pressure will increase by one micro atmosphere (μatm).
partial pressure of carbon dioxide	pCO <sub>2</sub>	μatm	At sea level, $p\text{CO}_2 = (P - p\text{H}_2\text{O}) \times X_{\text{CO}_2}$ , where P is the total atmospheric pressure and pH <sub>2</sub> O is the water vapor saturation vapor pressure (see Woolf et al., 2016). 1 μatm = 10 <sup>-6</sup> atmospheres = 0.10325 Pascals.
carbon dioxide fugacity	fCO <sub>2</sub>	μatm	Effective ideal gas partial pressure of CO <sub>2</sub> that has the same temperature and Gibbs free energy as the real gas. At the surface, $f\text{CO}_2 = X_{\text{CO}_2} \times \phi_{\text{CO}_2}$ , where $\phi_{\text{CO}_2} \approx 0.0002/\text{K}$ is the fugacity coefficient for CO <sub>2</sub> and K is the temperature in Kelvin.
column averaged carbon dioxide dry air mole fraction	XCO <sub>2</sub>	ppm	This vertically-averaged quantity used by the atmospheric remote sensing community is derived from the ratio of the CO <sub>2</sub> column abundance and the dry air column abundance. The dry air column abundance is estimated from the measured molecular oxygen (O <sub>2</sub> ) column abundance, assuming an O <sub>2</sub> dry air mole fraction of 0.20955.
carbon stock or stock change		petagrams of carbon per year (Pg C yr <sup>-1</sup> )	1 Pg C = 10 <sup>15</sup> g C. 1 PgC = 10 <sup>12</sup> kg C = 10 <sup>9</sup> tons of carbon = 1 Gt C. When oxidized to form CO <sub>2</sub> , 1 Pg C generates 3.664 Pg CO <sub>2</sub> .
Gross Primary Production	GPP	Pg C yr <sup>-1</sup>	Total flux of carbon fixed through photosynthetic reduction of CO <sub>2</sub> by plants in an ecosystem.
Net Primary Production	NPP	Pg C yr <sup>-1</sup>	Net flux of organic carbon produced by plants in an ecosystem. NPP equals GPP minus autotrophic respiration by plants, R <sub>a</sub> .
Net Ecosystem Exchange or Net Ecosystem Production	NEE or NEP	Pg C yr <sup>-1</sup>	Net primary production, NPP, minus carbon losses by heterotrophic (non plant) respiration, R <sub>h</sub> . NEE and NEP are generally interchangeable, with NEE used more often to refer to fluxes measured in the atmosphere, while NEP is more often used for fluxes inferred from measurements of carbon stock changes.
Net Biospheric Exchange or Net Biome Exchange	NBE	Pg C yr <sup>-1</sup>	Change in mass of carbon stocks after episodic carbon losses due to natural or anthropogenic disturbance.

community uses the partial pressure, pCO<sub>2</sub>, fugacity, fCO<sub>2</sub>, and the air-sea carbon flux. To relate these units, it is useful to note that one petagram of carbon (1 Pg C) in the atmospheric CO<sub>2</sub> is equivalent to a concentration of ~ 2.124 ppm (c.f. Ballantyne et al., 2012; Friedlingstein et al., 2020). Table 1 summarizes these and other commonly used quantities and units used by the carbon cycle community and describes their relationships.

### 3 Anthropogenic Contributions to the Carbon Cycle

Atmospheric CO<sub>2</sub> emissions from fossil fuel combustion, cement production, land use change and other human activities are currently being tracked by the regulatory, commercial and scientific communities (e.g. Andrew, 2020). International organizations such as the International Energy Agency (IEA) originally compiled fossil fuel statistics to avoid disruptions in the world's oil supplies, but now provide annual reports on a range of technologies to support sustainable energy development (IEA 2020). Those from national organizations, such as the U.S. Energy Information Administration (EIA) serve a similar purpose, tracking short-term and long-term trends in supply and demand around the world to support the energy industry, while those from national regulatory organizations such as the U.S. Environmental Protection Agency (EPA) compile statistics for regulating and reporting national emissions to other government agencies or organizations such as the United Nations Framework Convention on Climate Change (UNFCCC). Commercial organizations, such as British Petroleum, produce inventories to track trends in energy markets (BP 2020).

Similarly, to track emissions from land use change, international organizations such as the United Nations Food and Agriculture Organization (FAO) collect and disseminate global information on agriculture, forestry and other land use (AFOLU). Two methods are typically used to track emissions from land use change. In the first, data collected by FAO are used in so-called bookkeeping models that prescribe carbon loss per pool over time (Hansis et al., 2015, Houghton and Nassikas, 2017). In the second, satellite remote sensing data are used to determine the amount of land cover change and associate emission losses with the change by emission factors or more detailed biogeochemical models (van der Werf et al., 2017). For tracking historical land use changes, a map of historical land use is required such as LUH2-GCB2020 (Hurrt et al., 2020; see also Friedlingstein et al., 2020). Using this information, it is possible to project land use change impact on the carbon cycle using the new generation of dynamic global vegetation models (DGVMs). At the national level, land use change emissions are compiled and delivered to the UNFCCC by country level organizations such as the U.S. EPA, Japan's Ministry of the Environment and the European Union's European Environment Agency.

Scientific inventories, such as those compiled by the Carbon Dioxide Information Analysis Center (CDIAC; Boden et al., 2017) and the annual reports compiled by the Global Carbon Project (GCP), combine information from all of these sources to support scientific investigations of the energy and carbon cycles as well as other applications. The science community has also produced high resolution gridded inventories such as EDGAR (Janssen-Maehout et al., 2019), ODIAC (Oda et al., 2018), and Hestia (Gurney et al., 2019). These inventories use other data (population, night lights, etc.) to disaggregate national scale emissions to support carbon cycle investigations on spatial scales spanning individual forest plots or urban areas to full biomes. These gridded inventories also provide more actionable information on anthropogenic CO<sub>2</sub> emissions for policy makers working on urban to sub-national scales.

#### 3.1 *Regulating Greenhouse Gas Emissions - the Paris Agreement*

The primary international regulatory framework for tracking anthropogenic CO<sub>2</sub> emissions is the UNFCCC. This treaty was established in 1994 to stabilize "greenhouse gas concentrations in the atmosphere at a level that would prevent dangerous anthropogenic interference in the climate system." To implement this treaty, parties to the 2015 UNFCCC Paris Agreement resolved to define Nationally Determined Contributions (NDCs) to a global GHG

reduction effort and to report progress toward these NDCs, at 5-year intervals through Global Stocktakes, the first of which is scheduled for 2023. The NDC reports are to be compiled based on the Intergovernmental Panel on Climate Change (IPCC) Guidelines for National Greenhouse Gas Inventories (IPCC 2006; 2019) and consist of bottom-up emission inventories, compiled from a statistical analysis of emissions reported from sources in specific sectors, including Energy, Industrial Processes Product use, Agriculture, Forestry, and Other Land Use (AFOLU), and Waste. Each sector is subdivided into a series of categories. For example, AFOLU is divided into six categories, including Forest Land, Cropland, Grassland, Wetlands, Settlements, and Other Land categories.

While CO<sub>2</sub> emissions from fossil fuel combustion in the energy sector constitute the largest direct anthropogenic contribution to the global carbon cycle, emissions of CO<sub>2</sub> and other greenhouse gases from managed lands tracked under the AFOLU sector are responsible for almost one quarter of all anthropogenic GHG emissions (Smith et al., 2014). These emissions originate primarily from deforestation and forest degradation, but also include contributions from agricultural land and livestock management. The IPCC Guidelines only require reporting of emissions on managed lands, but they acknowledge that natural disturbances (wildfires, insects, severe weather) contribute a significant source of interannual variability and uncertainty in this sector. The concept of natural and anthropogenic changes are becoming more diffuse as climate change itself affects “natural disturbances”. A comprehensive understanding of the processes operating in the natural carbon cycle is therefore essential for tracking and attributing observed interannual changes in atmospheric CO<sub>2</sub> concentrations, and for linking bottom-up inventories with top-down monitoring, recording, and verification (MRV) strategies (Janssens-Maenhout et al., 2020).

### 3.2 Carbon Cycle Insights from Scientific Inventories

The Global Carbon Budget (GCB) compiled annually by the Global Carbon Project document the globally-averaged budget of anthropogenic carbon fluxes for five key components: atmosphere, fossil fuel emissions, land use change, uptake by the terrestrial biosphere (“land sink”) and uptake by the ocean (“ocean sink”). From 2000-2006 to 2019, emissions from fossil fuel use and cement production increased from 7.6 to 9.7 Pg C yr<sup>-1</sup>, with a peak of 10 Pg C yr<sup>-1</sup> in 2018. The ocean and land sinks increased during the same time from 2.2 to 2.6 and 2.8 to 3.6 Pg C yr<sup>-1</sup> respectively (Friedlingstein et al., 2020). The anthropogenic sinks are defined as their perturbation caused by direct effects of increasing atmospheric CO<sub>2</sub> and indirect effects by climate change.

In addition to these flux estimates, the GCBs document uncertainties, expressed as one standard deviation around the mean. Figure 1 shows the relative error of these estimates (uncertainty/mean) as they progress along the years for the 2006-2019 budgets. The estimates refer to each individual year for which the budget was prepared. As such, they indicate the progression in understanding of the uncertainties in the budget at that time (as opposed to an *a posteriori* analysis of the uncertainties of all years in a similar manner). The relatively low, stable uncertainties associated with both the fossil fuel emissions and atmospheric CO<sub>2</sub> concentrations result from two factors (Ballantyne et al., 2012). The first is the precision of the atmospheric in situ CO<sub>2</sub> measurements and efficient mixing of CO<sub>2</sub> throughout the atmosphere, although analytical errors and sampling bias do play a role. Second, while fossil fuel combustion is the primary source of anthropogenic CO<sub>2</sub> emissions, the relative error on this contribution is



relatively small because the fossil fuel industry provides reliable numbers on their sales, which are well correlated with the amount of fossil fuel burned. The highest relative errors are associated with land use change emissions. These estimates are based on either bookkeeping methods (Houghton, 2003; Houghton and Nassikas, 2017), and since 2018, also on DGVMs (Friedlingstein et al., 2019, 2020). Compared to the early period, 2000-2006, the relative error has not substantially decreased, nor has the mean value of the land use change emissions of 1.5 to 1.8 Pg C yr<sup>-1</sup>, respectively.

In the GCB published in 2016 and before, the land sink was calculated as a residual: land sink = emissions - atmosphere - ocean sink. Since 2017, the land sink in the Global Carbon Budget has been calculated directly from DGVMs with a corresponding increase in relative error due to the divergence of the individual DGVM estimates. Since then, the ocean sink estimate from models is no longer normalized to a data-based estimate from the 1990s (Denman et al., 2007), which was previously applied to ensure the land sink estimate from the budget residual

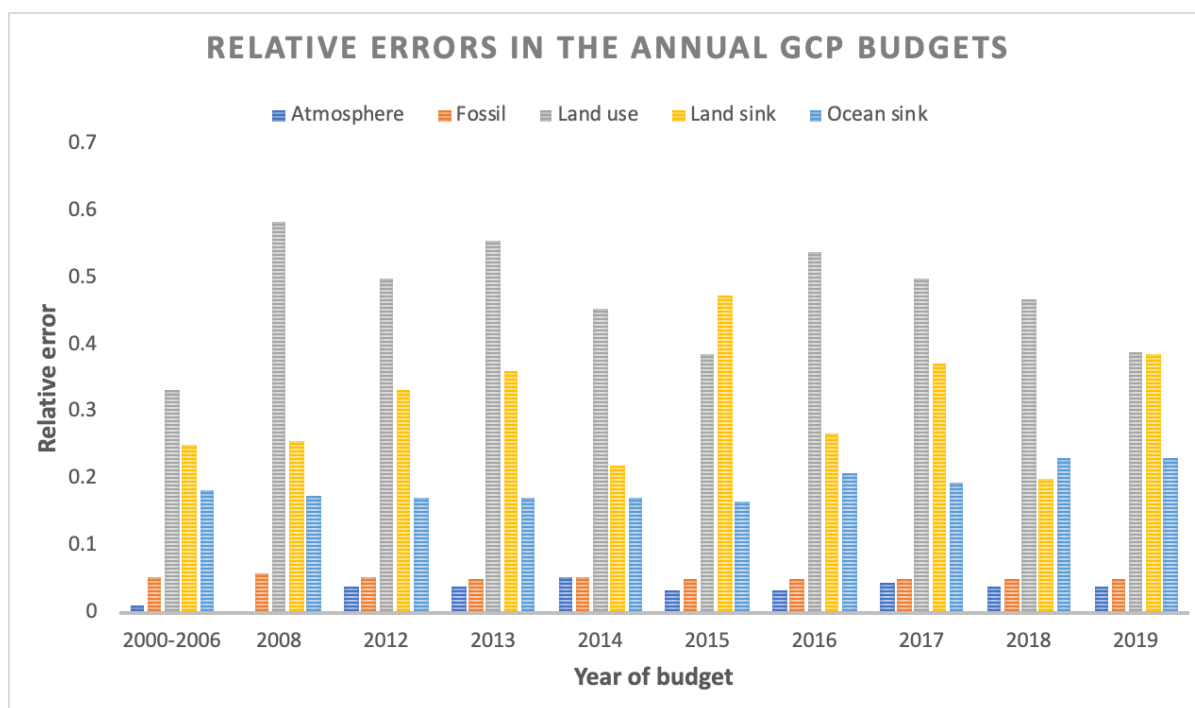


Figure 1. Relative error (1 standard deviation uncertainty / mean) for the Global Carbon Budget estimates since 2000. Numbers are taken for the individual year(s) reported each year from Canadell et al. (2007), LeQuéré et al. (2009) and LeQuéré et al. (2013-2018) and Friedlingstein et al. (2019-2020) and refer to the annual estimates.

had a realistic mean value. This change in methodology also affected the ocean sink uncertainty, which slightly increased from 17% in 2015 to 19% in 2016. The fact that the ocean sink was no longer scaled to the mean 1990s value led to a smaller ocean sink estimate that, together with the same absolute uncertainty, resulted in the slightly higher relative uncertainty. The ocean sink uncertainty had also varied between 17 and 19% for the years 2006 to 2015.

With the advent of a direct estimate of the land sink from DGVMs, it is now possible to assess the degree to which the overall global carbon budget can be closed, i.e., the difference

between the sum of the fluxes and the atmospheric accumulation. This budget imbalance was estimated at 0.3 Pg C for 2018 and 2019, or approximately 10% of the magnitude of the land and ocean sinks (Friedlingstein et al., 2019, 2020). The budget imbalance indicates that there remains substantial uncertainty in global annual mean fluxes.

#### 4 The Ocean Carbon Cycle

The ocean holds a large natural reservoir of carbon that exchanges with the atmosphere on short (non-geological) time-scales. Superimposed upon the cycling of this natural reservoir, the increasing atmospheric CO<sub>2</sub> concentration is causing the ocean to absorb a significant fraction of anthropogenic carbon emissions. Due to the natural carbon cycle of the ocean, 39,000 Pg C is stored in the ocean, which amounts to ~90% of the carbon contained in the combined land, ocean and atmosphere domains (Bolin 1983; Sundquist 1993; Sabine and Tanhua, 2010). The natural carbon cycle is driven by ocean circulation, seasonal heating and cooling, and biological processes (Figure 2, left).

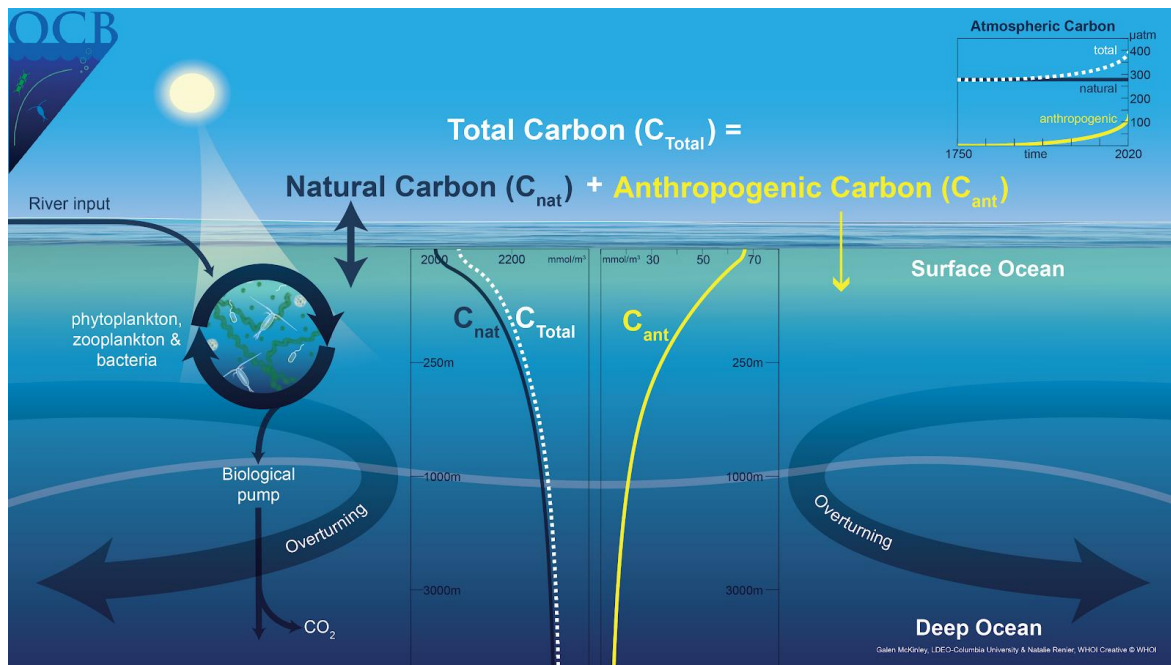


Figure 2: The total carbon cycle in the ocean ( $C_{\text{Total}}$ ) is the sum of the natural carbon cycle ( $C_{\text{nat}}$ ) and the anthropogenic carbon cycle ( $C_{\text{ant}}$ ). The natural carbon cycle is quantitatively dominant, as shown in the observed data (GLODAPv2, Olsen et al., 2016) plotted in the center, and includes contributions from biological activity and the large-scale circulation of the ocean. Overlain is the uptake of additional carbon due to anthropogenic emissions to the atmosphere that occurs in the present ocean as atmospheric pCO<sub>2</sub> continues to rise.

The ocean carbon budget can be quantified as the storage of inorganic and organic carbon in the ocean, the fluxes of carbon across the air-sea interface, river input, and a small term for sedimentation. The natural carbon inventory is very large compared to the anthropogenic component and is believed to have been in an approximate long-term steady state in preindustrial times, such that there was zero global mean air-sea flux of natural carbon ( $F_{\text{nat}}$ ). The

anthropogenic uptake flux ( $F_{\text{ant}}$ ) is the additional ocean uptake due to the direct effect of increasing atmospheric  $\text{CO}_2$  concentration and occurs as a perturbation to the vigorous natural cycle (Figure 2, right), with the column inventory of anthropogenic carbon ( $C_{\text{ant}}$ ) from the latest data-based estimates mapped in Figure 3 (bottom).

The downward increase in natural carbon ( $C_{\text{nat}}$ ) from surface to depth (Figure 2) is largely due to the biological carbon pump (BCP) (Sarmiento & Gruber, 2006). If the BCP did not operate, the atmospheric  $\text{CO}_2$  concentration would be around 200 ppm higher (Maier-Reimer et al., 1996). During the last glacial maximum, changes in the efficiency of the BCP may have played an important role in lowering atmospheric  $\text{CO}_2$  (Galbraith and Skinner, 2020; Sigman et al., 2010). Biological feedbacks may accompany anthropogenic climate change (Sabine & Tanhua, 2010; Hauck et al., 2015; Moore et al., 2018), but there is significant spread in model projections (Laufkötter et al., 2015, 2016; Frölicher et al., 2016). To date, observed timeseries are too short to provide evidence for long-term biologically-driven trends in the ocean carbon cycle (Henson et al., 2016). Thus, the ocean carbon sink for anthropogenic carbon over the industrial era is currently understood as a physical and chemical process (Figure 2, right). The contemporary (or ‘net’) air-sea  $\text{CO}_2$  flux ( $F_{\text{net}}$ ) is the sum of  $F_{\text{nat}}$  and  $F_{\text{ant}}$ . The surface ocean partial pressure of  $\text{CO}_2$ ,  $p\text{CO}_2$ ,  $\text{CO}_2$  flux and interior ocean inventory of anthropogenic carbon ( $C_{\text{ant}}$ ) are shown in Figure 3.

The ocean surface layer carbon content equilibrates with the atmosphere on time-scales of months. The ocean continually removes  $C_{\text{ant}}$  from the atmosphere because the ocean circulation transports  $C_{\text{ant}}$ -laden waters away from the surface layer and into the ocean interior, while the water that returns to the surface tends to have low  $C_{\text{ant}}$  content. Thus, the ocean circulation is essential to continued  $\text{CO}_2$  uptake. At the global scale, the ocean overturns relatively slowly, on timescales of 1000 years. Because of this, 75% of all anthropogenic carbon attributable to the industrial age remains in the upper 1000m (Gruber et al., 2019a). Because carbon is highly soluble in ocean water, the fundamental limit on the rate of anthropogenic carbon uptake by the ocean is the rate of overturning, as it determines how fast waters with  $C_{\text{ant}}$  uptake capacity are exposed to the surface.

Since the beginning of the industrial era, the uptake of  $C_{\text{ant}}$  by the land carbon sink has been largely offset by emissions associated with land use change. The ocean has therefore been the primary cumulative net  $C_{\text{ant}}$  sink over this period (Friedlingstein et al., 2019; 2020). Looking forward, the behavior of the ocean carbon sink is expected to continue playing a critical role in determining how much anthropogenic carbon remains in the atmosphere (Randerson et al., 2015; Zickfeld et al., 2016; Schwinger and Tjiputra, 2018).

313           The following sections describe the approaches used to study the ocean carbon sink. A  
314 mechanistic understanding of this sink is essential for diagnosing its state and for making reliable  
315 future predictions. This requires quantification of carbon stocks and fluxes at higher spatio-  
316 temporal resolution than is available from interior data alone. Air-sea fluxes on monthly to  
317 decadal timescales are quantified using surface ocean observations and ocean models of varying

318 complexity. Agreement between independent estimates for mean fluxes and temporal variability  
 319 indicates growing confidence in global-scale mechanistic understanding. Yet, key uncertainties  
 320 remain and must be resolved to support better predictions for future ocean carbon sink and to  
 321 allow for reduced diagnostic uncertainty for the global carbon cycle as it evolves. Substantial  
 322 advances in observing systems, quantification of land-to-ocean fluxes of carbon, and models of

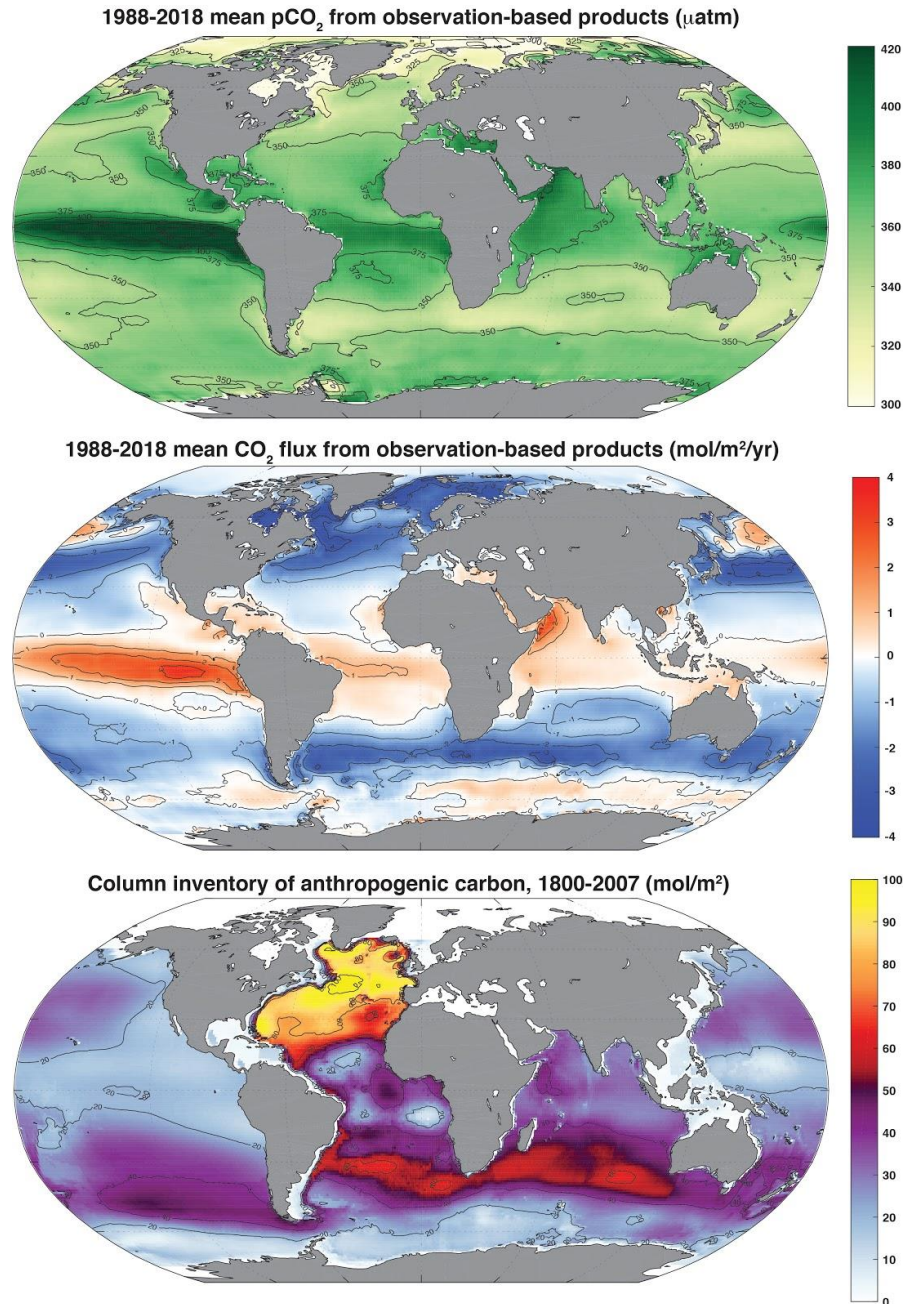


Figure 3: Surface ocean  $p\text{CO}_2$  (top); and air-sea  $\text{CO}_2$  flux ( $F_{\text{net}}$ ), positive flux to the atmosphere (middle), 1988-2018, mean of 6 observation-based products (Fay et al., in prep); column inventory of anthropogenic carbon ( $C_{\text{ant}}$ , bottom), 1800-2007 (Sabine et al., 2004, Gruber et al., 2019).

ocean circulation are needed to reduce these uncertainties. In addition, as nations implement substantial reductions in carbon emissions, we need to assess the near-term response of the ocean carbon sink to reduced atmospheric  $p\text{CO}_2$  growth rates.

#### **4.1 Bottom-up Estimates of Anthropogenic Carbon Accumulation In The Ocean From Interior Observations**

Based on a bottom-up accounting using interior ocean data, Gruber et al. (2019a) find a total ocean  $C_{\text{ant}}$  accumulation of  $152 \pm 20$  Pg C for the industrial era through 2007. By combining evidence from top-down and bottom-up approaches, Khatiwala et al. (2013) find an inventory of  $160 \pm 26$  Pg C in 2010. Consistent with previous inventories (Sabine et al., 2004), these studies find that the ocean has cumulatively absorbed excess carbon equivalent to 45% of industrial-era fossil fuel emissions until 2010, or 30% of the total emissions including LUC. The column inventory of ocean  $C_{\text{ant}}$  accumulation from Gruber et al. (2019a) is shown in Figure 3 (bottom).

The amount of  $C_{\text{ant}}$  estimated for 2010 ( $160 \pm 26$  Pg C) represents only about  $\sim 0.4\%$  of the ocean carbon stock, indicating the significant challenge of directly observing the temporal change in carbon stock over time. Direct measurements are only possible in areas with rapid change in dissolved inorganic carbon (DIC; e.g. Tanhua and Keeling, 2012). Instead, it is more practical to infer ocean storage of  $C_{\text{ant}}$  against the large natural background, and then to calculate the change in storage over time. A few different methods have been used to estimate the storage of  $C_{\text{ant}}$ , either based on observations of biogeochemistry variables, or by transient tracers (see Sabine and Tanhua (2010) for a review).

On a global scale, different methods converge within the uncertainties, but significant differences persist regionally (e.g. Waugh et al., 2006, Khatiwala et al., 2009). Multivariate techniques (e.g. Friis et al., 2005, Clement and Gruber, 2018) can be used to disentangle variability and calculate decadal-scale trends. A global estimate of the storage of anthropogenic carbon finds an increase of  $34 \pm 4$  Pg C between 1994 and 2007 (Gruber et al., 2019a), indicating a mean  $F_{\text{ant}}$  uptake of  $-2.6 \pm 0.3$  Pg C (negative flux into the ocean) annually over this time frame. This relatively accurate ( $\sim 12\%$ ) estimate provides an important benchmark for the ocean's role in sequestering anthropogenic carbon, and acts as a direct constraint on the net magnitude of the land flux given low uncertainty on fossil fuel emissions and atmospheric carbon accumulation. The magnitude of the uptake implies that the ocean is continuing to take up anthropogenic carbon at a rate proportional to anthropogenic carbon emissions.

A critical element to the success of global estimates of anthropogenic carbon stocks and changes in carbon storage is the GLODAP data product (Key et al., 2004), which collects all carbon-relevant interior ocean data into a product that has undergone the extensive quality control (Tanhua et al., 2010) required to quantify small changes over a large background. This data product is now being released on an annual basis and the GLODAPv2.2020 version contains data from over 1.2 million water samples collected during 946 cruises (Olsen et al., 2020).



## 4.2 *Bottom-up Estimates of Ocean-Atmosphere CO<sub>2</sub> Fluxes from Observations of Surface Ocean pCO<sub>2</sub>*

In order to understand the ocean carbon sink on annual to interannual timescales relevant to climate change policy, more frequent estimates of the sink are required than those produced from decadal timescale interior ocean observations. These data come from observations of the surface ocean partial pressure of CO<sub>2</sub> (pCO<sub>2</sub>), and are used to directly estimate net air-sea CO<sub>2</sub> fluxes (F<sub>net</sub>). The reported variable is surface ocean fugacity of CO<sub>2</sub> (fCO<sub>2</sub>) which equals the partial pressure of CO<sub>2</sub> corrected for the non-ideal behavior of the gas (Pfeil et al., 2013). The fugacity of CO<sub>2</sub> is 0.3-0.4% smaller than the partial pressure of CO<sub>2</sub> (Zeebe and Wolf-Gladrow, 2001). For simplicity, we use the terminology pCO<sub>2</sub> to refer to these data for the remainder of this paper. Over the past decade, the number of publicly available observations of pCO<sub>2</sub> has increased rapidly from 6 million in the first release of the Surface Ocean CO<sub>2</sub> Atlas (SOCAT) database (Pfeil et al., 2013, Bakker et al., 2014, 2016, 2020) in 2011 to 28 million in 2020 (www.socat.info). These observations and their automated organization into a consistent database have enabled scientists to create a variety of new observationally-based estimates of the ocean carbon sink that use correlated data (sea surface temperature, salinity and height, chlorophyll, mixed layer depth) to drive upper ocean extrapolation techniques and machine-learning algorithms so as to fill the observational gaps (Rödenbeck et al., 2014; 2015, Landschützer et al., 2013; 2014; 2020; Denvil-Sommer et al., 2019, Gregor et al., 2019).

As the SOCAT database currently offers pCO<sub>2</sub> data for only ~2% of all 1° x 1° locations across the surface ocean, the extrapolation is quite significant. Nonetheless, comparisons of the extrapolated, observationally-based products to independent data indicate relatively low bias and convergence of the independent estimates (Gregor et al., 2019). Root mean square errors (RMSE) range from 10 to 35 µatm. The fact that bias and RMSE comparisons are largely consistent across the variety of approaches suggests that it is data sparsity rather than extrapolation methodology that is now a fundamental limitation on further error reduction (Gregor et al., 2019). Additional tests of the machine-learning based extrapolation approaches using an Earth System Model testbed indicate that they generally have low bias and are skillful in representing the amplitude and timing of seasonality across the global ocean. However, higher and lower frequency variations are more poorly represented because of inadequate sampling on these timescales (Gloege et al., 2020, Stamell et al., 2020). Several challenges remain in using these data, including the uneven distribution of data over time, methodological differences in the calculation of air-sea flux from pCO<sub>2</sub> (Fay et al., 2021, Woolf et al., 2019, Zavarisky and Marandino, 2019), and the potential need for adjustments to pCO<sub>2</sub> data to account for near-surface temperature and salinity gradients (Watson et al., 2020).

Despite the significant extrapolation and remaining uncertainties, it is a major advance for ocean carbon cycle science to have spatially-resolved, data-based estimates of air-sea CO<sub>2</sub> fluxes on monthly timescales. This allows for new investigation into the magnitudes and mechanisms of interannual and decadal variability in the ocean carbon sink, and a key point of comparison to ocean models that were previously the only basis for this analysis. Models are discussed in the next section, and results are compared in the following.

## 4.3 *Bottom-Up Estimates of Ocean-Atmosphere CO<sub>2</sub> Fluxes from Ocean Models*

Global ocean biogeochemical hindcast models estimate interior ocean carbon cycling and, from this, air-sea CO<sub>2</sub> fluxes. Models simulate the carbon distribution in the ocean with currents, water mass formation and mixing, and for biological carbon turn-over. The bottle-neck for ocean carbon uptake in the models, as in the real world, is the carbon transport across the mixed layer depth. As a result, the models' carbon uptake is sensitive to simulated physics (Doney et al., 2004; Goris et al., 2018; Huber and Zanna, 2017). Models can also provide air-sea flux estimates prior to the 1990s when surface pCO<sub>2</sub> observations were rare.

Models are routinely evaluated against observations or observation-derived estimates that characterize the physical and biogeochemical state of the ocean for the last several decades (Doney et al., 2004; Schourup-Kristensen et al., 2014; Aumont et al., 2015; Schwinger et al., 2016; Stock et al., 2020). For the suite of models used in the Global Carbon Budget, comparison of pCO<sub>2</sub> at observed locations from SOCAT reveals the models' ability to capture variability and trends on annual (RMSE <10 µatm) and decadal timescales (RMSE <10 µatm). However, large model-data mismatches on the seasonal timescale also exist (RMSE of 20–80 µatm; Hauck et al., 2020). Despite concurrence on annual and decadal timescales, ocean carbon sink estimates diverge (Hauck et al., 2020), indicating the available data do not fully constrain the models' emergent carbon sink.

Global ocean biogeochemical models are routinely used to quantify the ocean sink in the GCB. For example, for 2019, they find that the ocean sink accounted for 22% of 2019 anthropogenic CO<sub>2</sub> emissions (Friedlingstein et al., 2020). Models have also shed light on processes behind observed variability such as the weakening of the Southern Ocean carbon sink in response to increased westerlies (LeQuéré et al., 2007), and to explore the role of stationary Rossby waves in subduction of anthropogenic carbon (Langlais et al., 2017). As a component of Earth System Models, ocean models are the single tool for future projections. In the future, on timescales from decadal to centennial, models project a decreased carbon sink efficiency due to changes in the air-sea pCO<sub>2</sub> gradient, ocean circulation and reduced buffer capacity (Schwinger et al., 2014, Randerson et al., 2015, Zickfeld et al., 2016, Schwinger and Tjiputra, 2018, Ridge and McKinley, 2020b).

#### **4.4 Reconciling Air-Sea Flux Estimates from Different Methods**

We must accurately quantify the ocean sink and understand its mechanisms so that the sink can be monitored as it evolves, and so that reliable projections can be developed. The best window into our current understanding is the degree to which the above-mentioned independent estimates of the present-day sink's magnitude agree. We discuss the degree of agreement in this section, where a negative flux refers to a flux from atmosphere to ocean, and we discuss mechanistic understanding in the next section.

Surface ocean carbon observations indicate the net air-sea flux of carbon into the ocean ( $F_{\text{net}} \sim -1.6 \text{ Pg C yr}^{-1}$ ), while interior measurements offer estimates of the anthropogenic uptake and storage ( $F_{\text{ant}} \sim -2.6 \text{ Pg C yr}^{-1}$ ). Dynamic hindcast models as used in the GCB estimate the total of anthropogenic perturbations, that is the sum of anthropogenic uptake ( $F_{\text{ant}}$ ) and anthropogenic climate change induced natural carbon fluxes ( $F_{\text{nat, ns}}$ ). Closure terms of significant net magnitude ( $\sim 1 \text{ Pg C yr}^{-1}$ ) are required to bridge the gap between  $F_{\text{net}}$  and  $F_{\text{ant}}$ .

To reconcile flux estimates from pCO<sub>2</sub>-based data products with ocean models and estimates from interior data, it is well established that an adjustment due to the riverine input of



natural carbon that outgasses from the ocean ( $F_{\text{nat},\text{riv}}$ ) must be applied (Sarmiento and Sundquist, 1992; Aumont et al., 2001; Lacroix et al., 2020). Unfortunately, high quality direct estimates of  $F_{\text{nat},\text{riv}}$  do not exist, and so the closure between surface flux estimates of  $F_{\text{net}}$  and  $F_{\text{ant}}$  remains a significant uncertainty. Lacking better evidence, values typically used are between 0.45 and 0.78  $\text{Pg C yr}^{-1}$  (Jacobson et al., 2007, Resplandy et al., 2018), with uncertainty estimates that are approximately 100% of these mean values. Anthropogenic changes to the riverine input of carbon are an additional closure term not usually considered with no temporally-resolved estimates available and one estimate for 2000-2010 suggesting it to be small (0.1  $\text{Pg C yr}^{-1}$ , Regnier et al., 2013). No estimates on anthropogenic changes to the outgassing of the riverine carbon in the ocean are available.

It has also been proposed that climate change is having an effect on the natural carbon cycle ( $F_{\text{nat},\text{ns}}$ , Le Quéré et al., 2010; Gruber et al., 2019a). The magnitude of this non-steady state component of the natural carbon cycle is still highly uncertain.  $F_{\text{nat},\text{ns}}$  has been estimated with one model for the period 1981-2007 (Le Quéré et al., 2010) and with a back-of-the-envelope calculation for the period 1994-2007 (Gruber et al., 2019a), suggesting a reduction of  $F_{\text{ant}}$  by 10 to 15%. Gruber et al. (2019a) estimate  $F_{\text{nat},\text{ns}}$  by assuming that the accumulation of anthropogenic carbon in the ocean follows a strictly linear scaling with the atmospheric load. However, this assumption is known to hold only when the atmospheric growth is strictly exponential, which has not been the case (Raupach et al., 2014, Ridge and McKinley 2020b), and thus the resulting estimate of +0.38  $\text{Pg C yr}^{-1}$  is likely an upper-bound. Another approach for estimating  $F_{\text{nat},\text{ns}}$  is to use ocean models that represent the natural carbon cycle, and to make a reasonable assumption that the total carbon cycle response to climate variability is dominated by the natural component. With this assumption, models indicate  $F_{\text{nat},\text{ns}}$  for 1994-2007 of +0.06 to +0.31  $\text{Pg C yr}^{-1}$  (DeVries et al., 2019; McKinley et al., 2020). Better quantification of this term is clearly needed.

Numbers for the ocean sink efficiency relative to emissions vary between 22% and 45% in the literature (Friedlingstein et al., 2020; Khatiwala et al., 2013; Sabine et al., 2004). These seemingly contradicting numbers are explained by different components of the ocean sink compared to different components of the emissions. Quantitatively the most important choice is the denominator used. For studies of the interior ocean cumulative ocean sink, efficiency is calculated relative to anthropogenic fossil emissions: 45% for the industrial era through 2010 (Khatiwala et al., 2013), and 41% for the industrial era through 2007 (Sabine et al., 2004). GCB estimates compare the ocean sink to total anthropogenic  $\text{CO}_2$  emissions, which includes emissions to the atmosphere from land-use change. From the GCB, a lower number with an efficiency of 22% compared to total anthropogenic emissions is estimated for 2010-2019. The GCB's approach also includes climate perturbation effects ( $F_{\text{nat},\text{ns}}$ ), which reduces the magnitude of the ocean sink.

The choice to compare studies of interior ocean accumulation to fossil fuel emissions is motivated by the fact that these numbers are cumulative over the industrial era, and over this time, the land use source and land sink have been in approximate balance. Thus, this approach also circumvents the higher uncertainties associated with land-use changes emissions and the land sink. The GCB's approach, on the other hand, acknowledges that fossil fuel and land-use change emissions increase the atmospheric  $\text{CO}_2$  concentration indistinguishably, and this determines the magnitudes of the ocean and land carbon sink.

#### 4.5 *Recent Evidence for Decadal Variability of the Ocean Carbon Sink*

In the mid-2000s, studies using ocean hindcast models suggested a slowing of the ocean carbon sink from the mid-1990s and attributed this change to processes in the Southern Ocean (Lovenduski et al., 2007; 2008; Le Quéré et al., 2007). In the following decade, the release of both the LDEO pCO<sub>2</sub> database and the development of the international SOCAT database allowed for new analyses of trends in air-sea CO<sub>2</sub> fluxes directly from observations (Le Quéré et al., 2009; McKinley et al., 2011; Fay and McKinley 2013). Additionally, a variety of extrapolations of these data to global monthly coverage were developed (Rödenbeck et al., 2015), and a recovery of the ocean carbon sink following the low near the year 2000 was noted (Fay and McKinley, 2013; Landschützer et al., 2015; DeVries et al., 2017; Gruber et al., 2019b). The Southern Ocean was generally identified as a key regional driver of these trends. A number of studies agreed that the stagnation of the Southern Ocean carbon sink in the 90s was related to a trend towards a more positive Southern Annular Mode (SAM) index associated with stronger westerly winds leading to more upwelling of natural carbon and hence dampened net air-to sea CO<sub>2</sub> flux (Le Quéré et al., 2007; Lovenduski et al., 2007; Lenton and Matear, 2007; Hauck et al., 2013).

Increasing nutrient concentrations in surface waters of all sectors of the Southern Ocean provide further evidence of strengthened upwelling during the late 1990s (Iida et al., 2013; Ayers and Strutton, 2013; Hoppema et al., 2015; Pardo et al., 2017; Panassa et al., 2018). However, the same driving mechanisms cannot explain the reinvigoration of the sink in the 2000s, as the trends towards a more positive SAM and stronger winds in the 2000s continued. Asymmetric changes in atmospheric circulation (Landschützer et al., 2015), a weaker upper ocean overturning circulation (DeVries et al., 2017) and regional wind variability (Keppler and Landschützer, 2019) were proposed as possible explanations, but no consensus was reached on the driving mechanisms of the reinvigoration. Several studies concluded that ocean models were substantially underestimating the magnitude of decadal variability in the ocean carbon sink (De Vries et al., 2019; Gruber et al., 2019b).

In the last few years, more observation-based estimates have become available (Denvil-Sommer et al., 2019; Gregor et al., 2019), and now the size of the ensemble of observation-based estimates and of hindcast models is more comparable. With similar size ensembles for both observation-based and hindcast models, estimates of decadal variability are more similar in magnitude and phase, and not as large as the initial observation-based products had suggested (McKinley et al., 2020; Hauck et al., 2020). Both the ensemble of hindcast models and observation-based products indicate a larger ocean carbon sink in the early 1990s, then a slowing of the sink through about 2000, and then a strong and steady recovery through 2018 (Figure 4). In both the products and models, flux variability is largely homogenous across the globe (McKinley et al., 2020). The mechanism driving this variability is still unknown.

By representing the surface ocean as a single abiotic box that exchanges water with the deep ocean at a constant rate, McKinley et al. (2020) are able to reproduce the variability of the ocean carbon sink with two external forcings (Figure 4). The two external forcings are the observed atmospheric pCO<sub>2</sub> and surface ocean temperature anomalies caused by eruptions of large volcanoes (El Chichon, 1982; Mt Pinatubo, 1991). This result emerges because the globally-averaged air to sea pCO<sub>2</sub> gradient - the fundamental driver of the flux - is only 6-10  $\mu$ atm, and thus anomalies in the atmospheric growth rate of a few  $\mu$ atm over several years can rapidly modify the global air-sea gradient. Large volcanic eruptions, such as Mt Pinatubo in

1991, cause a rapid surface ocean cooling, which increases solubility and creates an uptake pulse (Eddebbbar et al., 2019). Then, as the ocean warms from this rapid cooling, solubility is lowered and the sink is damped for 5-7 years beyond the eruption (Figure 4).

This model of McKinley et al. (2020) is arguably simple. Yet, it can reproduce the ocean carbon uptake that occurs in the ensemble mean of much more complex models and observation-based products. What does this mean? It can be interpreted simply as Henry's Law operating at the global scale, wherein the partial pressure in the water is moving toward equilibration with the partial pressure in the air. Since the atmospheric  $p\text{CO}_2$  continues to increase, the ocean continues to adjust toward equilibrium. In other words, McKinley et al. (2020) demonstrate that the ocean carbon sink temporal variability today is likely dominated by the external forcing from an ever-growing atmospheric  $p\text{CO}_2$  concentration. This perspective is consistent with recent analysis that shows heat uptake and interior redistribution in the ocean is far more sensitive to the details of the ocean circulation than is the pattern and magnitude of carbon uptake and storage (Bronse laer and Zanna, 2020).

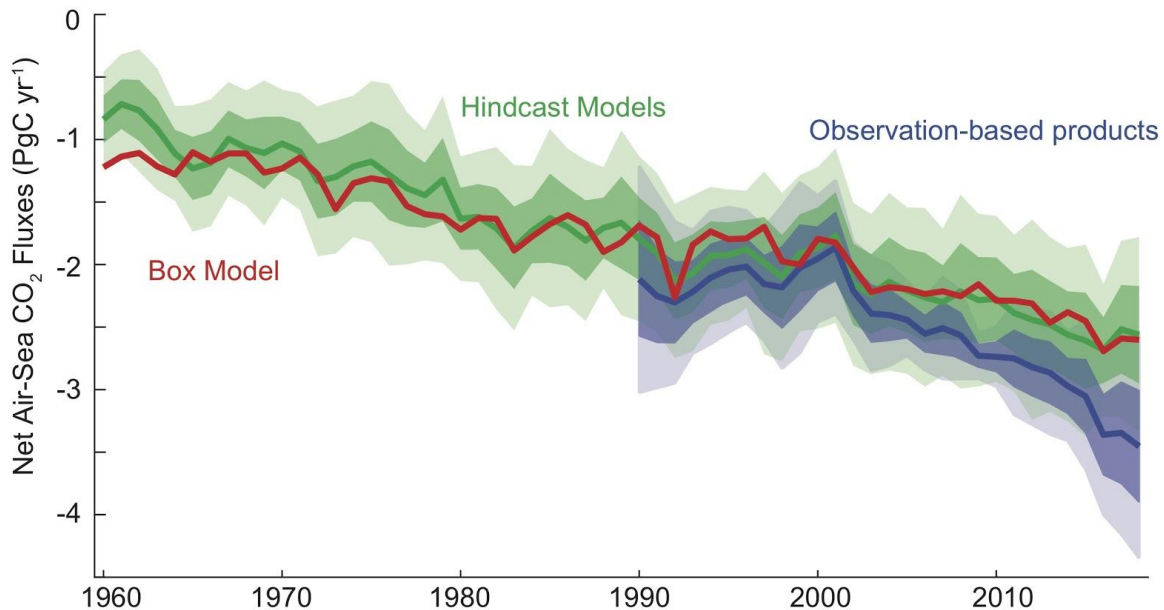


Figure 4. Air-sea  $\text{CO}_2$  flux of carbon ( $F_{\text{ant}} + F_{\text{nat,ns}}$ ) from observationally-based products (blue), hindcast models (green) and upper ocean diagnostic box model (red); negative flux into the ocean. Global ensemble means (bold), with 1 sigma and 2 sigma of individual members (shading). Hindcast ocean models from Global Carbon Budget 2020 (Friedlingstein et al., 2020). Observationally-based product  $p\text{CO}_2$  fields have missing ocean areas filled with a full-coverage climatology (Landschützer et al., 2020) and air-sea flux calculated as average of 3 wind reanalyses (CCMP, ERA5, JRA55) with a quadratic parameterization (Wanninkhof 2014, Fay et al., 2021); to this  $F_{\text{net}}$  estimate,  $F_{\text{riv,nat}} = 0.62 \text{ Pg C yr}^{-1}$  (Jacobson et al., 2007, Resplandy et al., 2018) is added. The upper ocean diagnostic box model (McKinley et al., 2020) is forced with observed atmospheric  $p\text{CO}_2$  and surface ocean temperature changes associated with the eruptions of three large volcanoes (Agung 1963, El Chichon 1982, and Mt. Pinatubo 1991; Eddebbbar et al., 2019).

551

552 **4.6 Advancing Understanding of the Current and Future Ocean Carbon Sink**

553 In order to continue to quantify the global carbon cycle, the constraint provided by the  
 554 relatively low-uncertainty estimates for decadal anthropogenic carbon accumulation must be  
 555 maintained. To better quantify fluxes on monthly to decadal timescales, increased observational  
 556 capacity at the surface and higher fidelity models are needed. In order to be prepared to support  
 557 climate management efforts in the near-term, the likely behavior of the ocean sink under  
 558 emissions mitigation must receive increased attention.

559 Observations of ocean interior carbon require measurements with high accuracy and  
 560 precision due to the small perturbations on a large background signal. For example, in 2010, the  
 561  $C_{\text{ant}}$  content was  $\sim 160$  Pg C out of a total inorganic carbon content of  $\sim 39,000$  Pg C. For the  
 562 surface ocean flux estimates, the high spatiotemporal variability in  $p\text{CO}_2$  and a low average  
 563 deviation from air-sea equilibrium concentration needed to drive the observed net flux, i.e. a net  
 564 flux of  $\sim 2.5$  Pg C  $\text{yr}^{-1}$  over a total flux of  $\sim 90$  Pg C  $\text{yr}^{-1}$ , indicates that accuracy and data  
 565 coverage are possibly the most important components of the observing system. There is a  
 566 seasonal bias in the observing system, with fewer observations being made in winter at high  
 567 latitudes. This is particularly important for observations of surface fluxes, which tend to be high  
 568 in winter, but less so for the interior ocean observations where seasonality tends to be low below  
 569 the winter mixed layer.

570 **4.6.1 Expanding autonomous observations:**

571 Although ship-based observations remain a central resource for the carbon observing system,  
 572 these are expensive and tend to be seasonally biased. Driven by these demands, there is a  
 573 continuous development of sensors for inorganic carbon system measurements with at least some  
 574 of these attributes; increased precision and accuracy, lower power consumption and lower  
 575 instrument drift (Johnson et al., 2016; Sabine et al., 2020; Seelmann et al., 2019; Sutton et al.,  
 576 2014). Similarly, there is a continuous development of autonomous platforms capable of carrying  
 577 sensors for ocean carbon. These include moorings (Sutton et al., 2014), profiling floats (e.g.  
 578 BGC Argo, Claustre et al., 2020), underwater gliders (Rudnick, 2016), and autonomous surface  
 579 vehicles powered by wind or waves (Sabine et al., 2020). These developments are rapidly  
 580 changing the capability to monitor ocean carbon with higher spatial and temporal resolution. For  
 581 instance, observations from Biogeochemical (BGC) Argos floats enable the calculation of  
 582 surface  $p\text{CO}_2$  (from pH and alkalinity estimates) with reasonable accuracy and precision,  $\sim 11$   
 583  $\mu\text{atm}$  (Takeshita et al., 2018; Williams et al., 2017). Although not as good as the  $2 \mu\text{atm}$  target  
 584 for the ship-based observations, this system has shown potential to fill spatiotemporal gaps in the  
 585 observations, with important implications for the carbon flux estimates. For example, Bushinsky  
 586 et al. (2019) report on significantly lower uptake of carbon in the Southern Ocean by including  
 587 winter time  $p\text{CO}_2$  from BGC-Argo floats using a neural network interpolation.

588 **4.6.2 Improved constraints on carbonate chemistry:**

589 Although individual components of the ocean carbon observing system have high technical  
 590 readiness levels, the new capabilities have not yet been integrated with existing, well-tested  
 591 technologies to provide an observing system that can quantify ocean carbon uptake to within  
 592 10%. One key need is an improved understanding of the ocean inorganic carbonate system.  
 593 There are four measurable inorganic carbon variables in the ocean - total alkalinity (TA), total  
 594 dissolved inorganic carbon (DIC), pH and  $f\text{CO}_2$ . By measuring two out of those, the complete

inorganic carbon system can, in theory, be calculated. Small errors in the dissociation constants, the boron-salinity ratio, and small contribution to the total alkalinity from unknown bases, can cause significant discrepancies in directly measured and calculated carbon variables (Fong and Dickson, 2019, Takeshita et al., 2020). A recent study by Álvarez et al. (2020) shows that inconsistencies between calculated and measured pH has decreased during the last decade, and they conclude that improved standard operating procedures for measurements and calculation of pH are urgently needed. An improved understanding of these issues is essential to fully utilize data from, for instance, BGC Argo floats equipped with pH sensors.

#### 4.6.3 *High-quality, timely data:*

As noted above, the anthropogenic perturbation in the global ocean is more than an order of magnitude smaller than the background natural state. Thus, if we are to track the changing anthropogenic carbon uptake by the ocean, we must maintain very high standards for accuracy and precision of carbon system data. New autonomous technologies offer great promise for expanding the observing system, but cannot be incorporated into the observing system if this substantially increases overall uncertainties. For the foreseeable future, ship-based measurements will continue to be required to calibrate and validate autonomous observations. Cross-over evaluations should occur both with deployment and post-deployment (Fay et al., 2018). At the same time, ocean carbon data should be ingested into publicly-released databases (SOCAT, GLODAP) in a timely manner that supports at least annual diagnoses of the ocean carbon sink. It is essential that these data be carefully quality controlled. As the timescales at which the user community requires these diagnoses become shorter, these data will need to be available more quickly.

#### 4.6.4 *Improved quantification of closure terms to link estimates of surface flux and interior $C_{ant}$ accumulation:*

In order to reduce uncertainties in the global and regional ocean carbon cycle, we need to understand how interior-based estimates of  $F_{ant}$  and surface flux estimates of  $F_{net}$  are quantitatively linked. A key barrier to this is the significant magnitude and high uncertainty in current estimates for natural fluxes of carbon in rivers ( $F_{nat,riv}$ ) and interannual variability in the natural carbon cycle ( $F_{nat,ns}$ ).

#### 4.6.5 *Constraining mechanisms of surface flux variability:*

Though recent work has identified the important role of external forcing from atmospheric  $pCO_2$  and volcanoes in driving ensemble-mean estimates of recent variability of the ocean carbon sink, individual models and individual observation-based products deviate from the mean of the ensembles (Hauck et al., 2020, McKinley et al., 2020). These deviations are due to internal variability of the ocean circulation and biology in each individual ensemble member. We do not currently have understanding of which of these individual estimates best represent the real ocean. To understand the actual total variability of the real ocean carbon sink (total = forced + internal), we need to select the observation-based products and models of highest fidelity. More stringent application of observational constraints can allow for focused analysis of the mechanisms driving variability in the highest-fidelity models and help improve others.

Another approach for combining observations and models is through data-assimilation that constrains the model ocean state and fluxes using observations, and closes data gaps by model dynamics rather than extrapolation. While assimilation applications so far have not provided annually updated global ocean sink estimates with full spatial and temporal

resolution (e.g., Mikaloff Fletcher et al., 2006; DeVries 2014; Verdy and Mazloff, 2017; DeVries et al., 2019), the first spatially and temporally resolved global data-assimilated models are starting to become available (Carroll et al., 2020).

#### 4.6.6 *Potential change in the natural carbon cycle:*

Climate change induced modifications of the ocean, such as ocean acidification, warming and ecosystem composition could significantly influence the transport of particulate and dissolved organic carbon from the surface to the interior ocean, i.e. the “biological pump”. The efficiency of this transport is a key factor regulating the atmospheric CO<sub>2</sub> concentration and is thought to play a role in regulating glacial / deglacial atmospheric CO<sub>2</sub> (e.g. Galbraith and Skinner, 2020). For instance, Marsay et al. (2015) suggest that a warmer ocean might lead to reduced sequestration of CO<sub>2</sub> by the biological pump. Complex interactions in the marine ecosystem will affect carbon export in a changing climate in ways that are difficult to predict and currently inadequately quantified (Laufkötter et al., 2015, 2016, Frölicher et al., 2016).

#### 4.6.7 *The future ocean sink under scenarios of emission mitigation:*

On centennial timescales under high emissions scenarios, slowing of the overturning circulation and reduced buffer capacity will significantly slow the rate of ocean carbon uptake (Randerson et al., 2015, Ridge and McKinley 2020b). But how will the ocean sink evolve under the increasingly more likely scenario of emission mitigation (Hausfather and Peters, 2020)? Given that the long-term growth and interannual variability of the ocean sink observed to date is driven by the exponential growth of atmospheric pCO<sub>2</sub> (Joos, 1996, Raupach et al., 2014, McKinley et al., 2020, Ridge and McKinley, 2020b), the ocean sink can be expected to slow in parallel to slowing in the growth rate of atmospheric pCO<sub>2</sub>. In effect, the anthropogenic carbon stuck in the near-surface ocean will begin to equilibrate with the atmosphere and the sink will be significantly reduced in response to the mitigation of emissions. This will occur simply due a change in the growth of atmospheric pCO<sub>2</sub> - no change in the ocean circulation or buffer capacity is required (Ridge and McKinley, 2020b). Slowing of the ocean sink will feedback on the atmosphere, reducing the effectiveness of mitigation actions and increasing climate warming.

Though a series of idealized studies have established the general fact that the ocean sink will be reduced with mitigation (Joos et al., 1996, Raupach et al., 2014, Zickfeld et al., 2016, Schwinger and Tjiputra, 2018, MacDougall et al., 2020, Ridge and McKinley, 2020b), the spatially and temporally resolved response of the ocean sink to emission mitigation has received little attention. Thus, we do not know how rapidly the ocean sink will slow, nor where surface flux changes will be most substantial. We do not know what will be required from our monitoring systems to detect these changes.

Current uncertainties in ocean models suggest despite the fact that the current ensemble of models largely agrees as to the recent evolution of the sink (Figure 4), there may be substantial divergence in feedback strength and ocean sink response to emission mitigation. Since the majority of the anthropogenic carbon is held in the ocean's thermocline (Gruber et al., 2019a), the circulation here is critical to the ocean sink's near-term response to mitigation (Iudicone et al., 2016, Rodgers et al., 2020, Ridge and McKinley 2020b). There is substantial spread in the regional distribution of ocean carbon uptake in current models (McKinley et al., 2016, Hauck et al., 2020), and major differences in representations of seasonality (Mongwe et al., 2018), which illustrates knowledge gaps with respect to physical and biological processes and their representations in models. In addition, circulation in these critical upper-ocean regions

is not consistently represented in state-of-the-art models (Bronselaer and Zanna, 2020). Uncertainties in the ocean sink under mitigation need first to be assessed, and then they need to be reduced by model development efforts so that robust projections can be made. Especially in these first decades of climate management via emission mitigation, there will be great public interest in how emission cuts are changing atmospheric CO<sub>2</sub>. Scientists need to be prepared to explain ocean carbon sink changes as they occur.

## 5 The Terrestrial Carbon Cycle

The terrestrial carbon cycle is characterized by large, spatially diffuse fluxes that should generally balance on multi-year time scales. Its primary stocks and fluxes are summarized in Figure 5. The Gross Primary Production (GPP) was estimated to be 123 Pg C yr<sup>-1</sup> from an analysis of direct flux observation made by a network of eddy covariance towers (Beer et al., 2010). Roughly one third of this (40.8 Pg C yr<sup>-1</sup>) is produced in the tropical forests, and one quarter (31.3 Pg C yr<sup>-1</sup>) in the tropical savannas, making the tropics by far the largest contributor to global GPP. Temperate and boreal forest are estimated to have a GPP of only 9.9 Pg C yr<sup>-1</sup> and 8.3 Pg C yr<sup>-1</sup>. When integrated over the globe, the GPP of croplands contributes an estimated 14.8 Pg C yr<sup>-1</sup>.

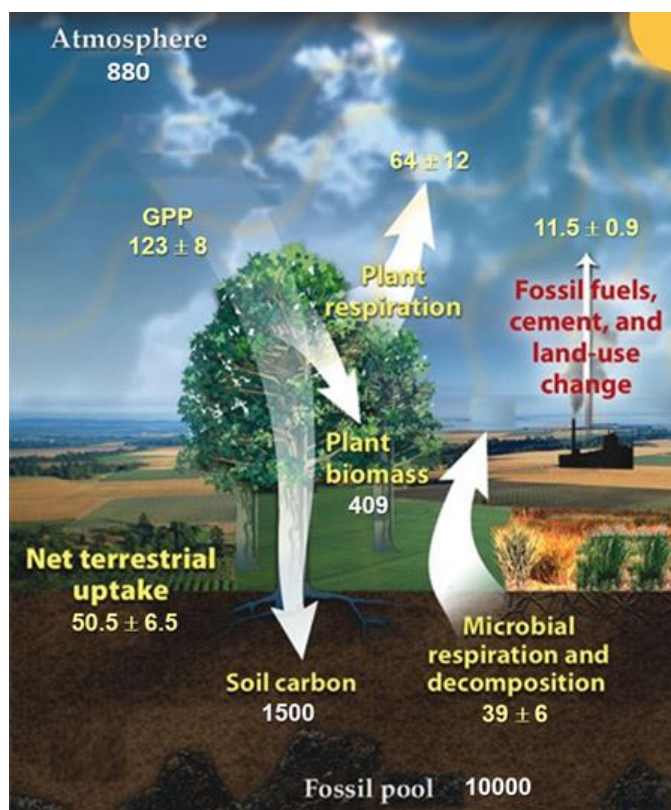


Figure 5: the land carbon cycle, showing the primary fluxes and reservoirs. The sources for the fluxes, shown in yellow, are listed in Table 2. Stock estimates are shown in white. Atmospheric carbon stocks were calculated assuming 410 ppm CO<sub>2</sub> and 1.884 ppm methane. Plant biomass stocks were taken from Spawn et al. (2020). Soil carbon stocks are from Scharlemann et al. (2014). Fossil pool stocks are from Ajani et al. (2013). (Adapted from U.S. Department of Energy Genomic Science program - <https://genomicscience.energy.gov>).



Welp et al. (2011), using oxygen isotopes, suggest that this value of Global GPP may be too low and would be closer to 150 -175 Pg C yr<sup>-1</sup>. However, Anav et al. (2015) argue that Welp et al. used a limited number of observations and a simple model that included gross photosynthesis, but neglected photorespiration by land plants. They note that plants immediately respire away 20-40% of the carbon fixed by photosynthesis. When photorespiration is included, they note that these GPP values are more in line with those obtained from other methods.

Global Autotrophic Respiration is estimated at 64 (±12) Pg C yr<sup>-1</sup> (Ito, 2020). This is the term that is also called maintenance respiration and consists mainly of dark respiration. Precise determination is difficult as it also involves a substantial below ground component. It is expected to vary with biome and climate (Ito, 2020). Estimates of Net Primary Production (NPP), GPP minus autotrophic respiration, are generally assumed to be of the order of 50% of GPP (i.e. Ito, 2020), but Ciais et al. (2020) suggest values in the range of 44-57 Pg C yr<sup>-1</sup>. They suggest that decomposition of organic matter or soil (heterotrophic) respiration (SHR) produces 39 Pg C yr<sup>-1</sup> within an interquartile range of 33-46 Pg C yr<sup>-1</sup>. This estimate is lower than conventionally assumed but agrees with recent large-scale estimates based on site soil respiration measurements (e.g. Jian et al., 2020). An additional flux to the atmosphere of 8.3 ± 9 Pg C yr<sup>-1</sup> is due to a series of processes listed Table 2. The remainder of GPP-NPP-SHR and these additional losses constitutes how much carbon is currently taken up by the land, and yields a value of the Net Ecosystem Exchange (NEE) of 2.2 ± 0.6 Pg C yr<sup>-1</sup>. This value is derived from a combination of methods such as close analysis of stock changes, flux measurements and inventories at the continental scale obtained during the Regional Carbon Cycle Assessment and Processes (RECCAP) Project. It is clear from the above description that the terrestrial carbon cycle is complex, composed of many, sometimes interacting elements, that often are not easy to quantify.

Table 2: Land Carbon Fluxes (Note: numbers without uncertainties are assumed to have uncertainties comparable to their stated values.)

Quantity	Flux (Pg C yr <sup>-1</sup> )	Reference
Gross Primary Production (GPP)	123 ± 8	1
Net Primary Production	~50.5 (44 - 57)	2
Autotrophic Respiration	~64 ± 12	3
Soil Heterotrophic Respiration	39 (33 - 46)	2
Outgassing by Rivers, Lakes and Estuaries	0.8 to 2.3	2
Fires	1.6	2
Consumption of Harvested Crops	1.5	2
Land Use Change	1.1	2
Grazing	1.0	2
Biogenic Reduced Carbon	0.8	2
Decay and Burning of Wood Products	0.7	2
<sup>1</sup> Beer et al. (2010); <sup>2</sup> Ciais et al. (2020); <sup>3</sup> Ito (2020)		



### 5.1 Bottom-up Estimates of CO<sub>2</sub> Emissions from Land

Fluxes of carbon between the land biosphere and atmosphere have traditionally been inferred from changes in land carbon stocks. To create a "bottom-up" estimate of the CO<sub>2</sub> emissions or uptake due to deforestation, reforestation, disturbance, or land use change requires information about the area affected, the corresponding carbon stock per unit area, and the fraction of carbon exchanged with the atmosphere due to the observed change. In practice, all three of these properties are challenging to quantify accurately (e.g., Saatchi et al., 2011; Ramankutty et al., 2007; Pearson et al., 2017), but all have benefited from new in situ and remote sensing measurement techniques and more advanced bottom-up modeling techniques.

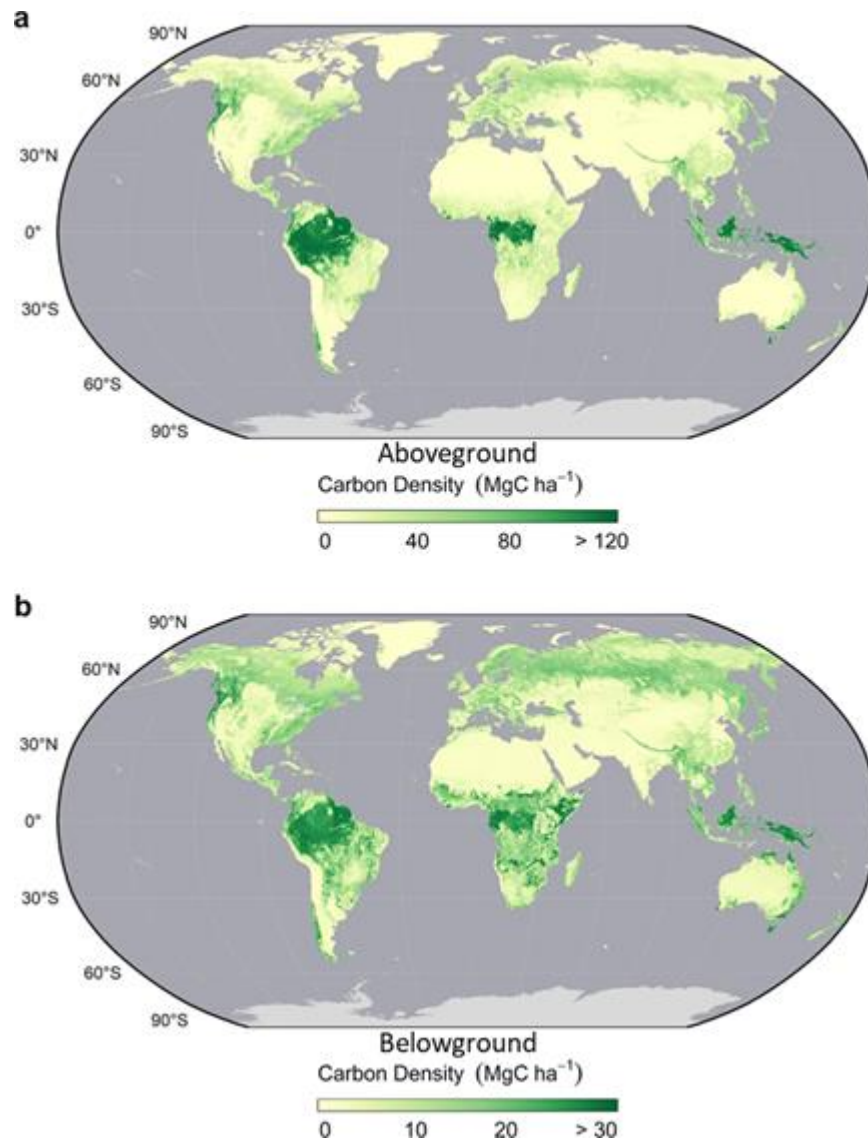


Figure 6. Maps of above and belowground living biomass carbon densities. (a) Aboveground biomass carbon density (AGBC) and (b) belowground biomass carbon density (BGBC). Maps have been aggregated at 5 km spatial resolution (Spawn et al., 2020).

Improved estimates of the area affected by the change have been facilitated by the availability of high-resolution space-based remote sensing observations from sensors such as LandSat, Moderate Resolution Spectroradiometer (MODIS) and Sentinel-2. Until recently, estimates of the carbon stocks per unit area associated with above-ground and below ground biomass still relied primarily on in situ measurements collected from a limited number of dedicated research plots at regular intervals (e.g. Pan et al., 2011). However, the increasing availability of above-ground biomass estimates derived from light detection and ranging (Lidar) and radio detection and ranging (radar) sensors on airborne and space-based platforms is providing improved spatial coverage and temporal sampling frequency. Soil carbon inventories still rely exclusively on in situ measurements, that are often characterized by limited spatial coverage and infrequent (decadal) repeat intervals (Ciais et al., 2014).

Figure 6 shows a map of global biomass (Spawn et al., 2020). This map was produced by combining and harmonizing several satellite products and site information. This involved using the satellite products of biomass with land cover with machine learning techniques to produce aerial estimates, and link this to below ground carbon density information (see Spawn et al., 2020 for more detail). These estimates yield a total living biomass of 409 Pg C, composed of an above ground biomass of 287 Pg C and a below ground biomass carbon density of 122 Pg C. Note that this is several orders of magnitude smaller than the ocean carbon reservoir, and in fact more comparable to the amount of total anthropogenic carbon stored in the ocean (160 Pg C).

CO<sub>2</sub> exchanged between the land and the atmosphere can be measured in situ using networks of eddy covariance flux towers, such as those deployed by FLUXNET (Baldocchi et al., 2001). The global network of eddy covariance sites have grown substantially over the past 25 years, with some records spanning that full period. These data provide unique constraints on the CO<sub>2</sub> fluxes from a broad range of vegetation types, climate regions and disturbance types. Globally, this network still has large gaps, particularly in the tropics, and each flux tower has a limited spatial footprint. Thus, early efforts to upscale results from local to regional scales often introduced large uncertainties (Baldocchi, 2003; Jung et al., 2009; Beer et al., 2010; Xiao et al., 2012; Keenan and Williams, 2018).

More recent methods that combine flux tower data with other remote sensing data in machine learning algorithms to produce upscaled fluxes (see Jung et al., 2020 for a review) yield global GPP estimates that agree well with those obtained from other methods, while providing insights into the processes controlling the carbon cycle of the land biosphere and their changes over time, particularly in the temperate Northern latitudes. Using radar derived estimates of biomass and soil carbon data from the harmonized world soil database and other sources combined with flux estimates of the global product of Beer et al. (2010), Carvalhais et al. (2014) calculated residence times of carbon. They found that the sensitivity of the residence time to soil moisture and temperature did not agree with the sensitivity of a set of DGVMs, while the overall pattern of increasing residence time at higher latitudes was reproduced. The following sections summarize recent results from bottom-up inventories that combine plot-based in situ measurements and remote sensing observations to constrain the carbon uptake and emissions from the land biosphere.

## **5.2 Indications of a Declining Tropical Land Sink from Bottom-up Inventories**

The carbon sink associated with intact tropical forests has been a major focus of bottom-up inventory methods like those described above. For example, Pan et al. (2011) estimated the intact tropical forest carbon sink to be approximately  $1.2 \text{ Pg C yr}^{-1}$  over 1990–2007 using an in situ inventory derived from up-scaled plot measurements. Hubau et al. (2020) present a more up-to-date assessment of the carbon sink in African and Amazon forest and conclude that while the African sink strength showed no trend ( $0.66 \text{ Mg C ha}^{-1} \text{ yr}^{-1}$ ), the Amazon forest sink slowed down  $-0.034 \text{ Mg C ha}^{-1} \text{ yr}^{-2}$  between 1983 and 2010, citing Brien et al. (2015). The results presented in Figure 7 show that this trend has persisted. Hubau et al. (2020) attribute the downward trend in the sink strength by intact forest primarily to increased mortality of live biomass. Disentangling the compound effects of  $\text{CO}_2$  fertilization (i.e. the increased rate of photosynthesis resulting from increased levels of  $\text{CO}_2$  in the atmosphere), increased temperature and drought, they imply that chronic long-term environmental change factors, temperature and  $\text{CO}_2$ , rather than simply the direct effects of drought, underlie longer-term trends in tropical forest tree mortality. Temperature increases enhance respiration and a reduce growth by increasing the vapor pressure deficit (VPD).

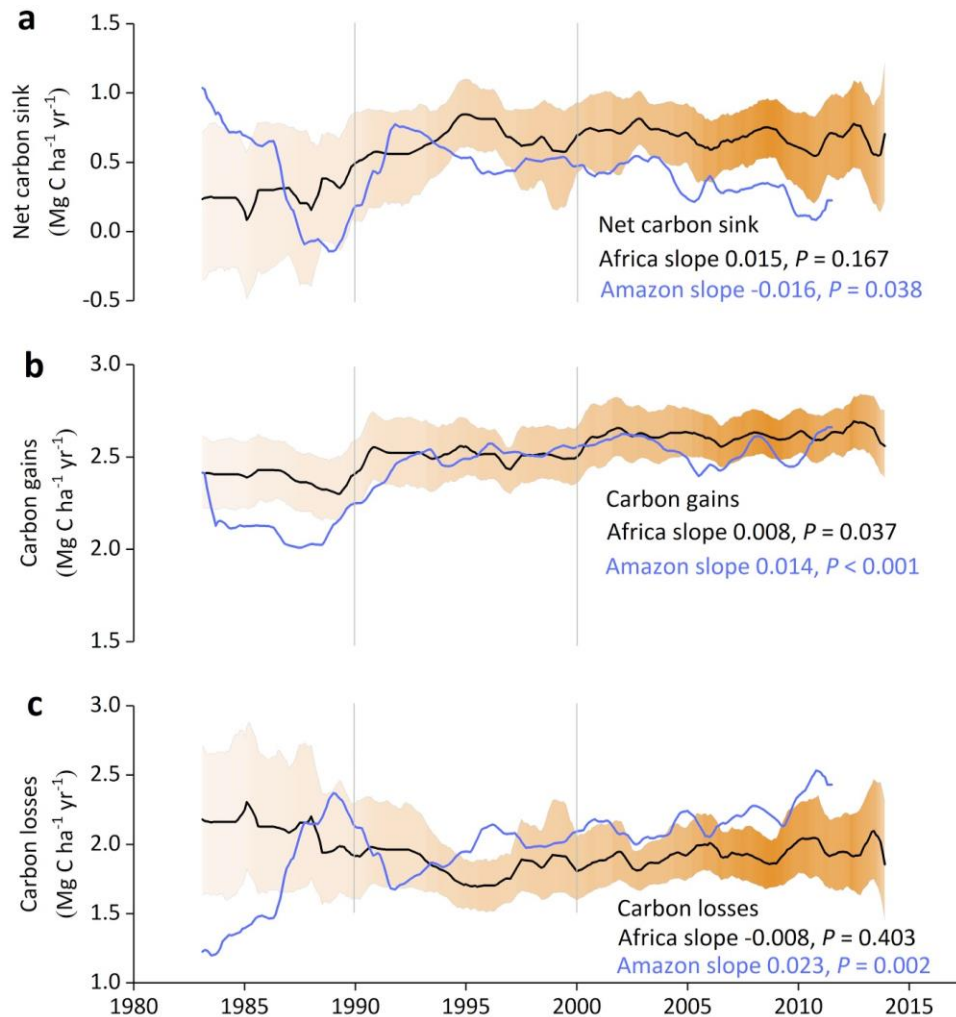


Figure 7. Time series of carbon dynamics from structurally-intact old-growth tropical forests in Africa and Amazonia from 1985 through 2015 (Data: Hubau et al., 2020).

While it is difficult to show the impact of climate extremes such as strong El Niños using in situ inventory data alone, bottom-up inventories of above ground biomass stocks compiled from microwave remote sensing observations provide a temporally denser record of such impacts. For example, contrary to Hubau et al. (2020), Wigneron et al. (2020) show that there was a strong legacy effect after the 2015-2016 El Niño event in both African and Amazon forest (0.9 and 0.5 Pg C loss in 2014-2017 respectively). For the overall tropics, Fan et al. (2019) use Vegetation Optical Depth (VOD) data from microwave sensors to show how changes in the above ground biomass of the forest of Tropical Africa and Tropical Asia contributed strongly to the interannual variability in CO<sub>2</sub> growth rates, while for semi-arid regions, those of Tropical America were the most important.

The net balance of tropical forest biomass remains the sum of the carbon gain from intact forest minus land use change, which is mostly deforestation. Estimates of deforestation in the Amazon fell from about 15000-20000 km<sup>2</sup> yr<sup>-1</sup> in the 1990s to about 6000 km<sup>2</sup> yr<sup>-1</sup> in 2016-2017, but have recently peaked again ([http://terrabrasilis.dpi.inpe.br/app/dashboard/deforestation/biomes/legal\\_amazon/rates](http://terrabrasilis.dpi.inpe.br/app/dashboard/deforestation/biomes/legal_amazon/rates)) to around 10000 km<sup>2</sup> yr<sup>-1</sup>. Preliminary analysis suggests that this increase would cause 12-14 teragrams of carbon (Tg C) to be released annually, compared to 8-10 Tg C over the period 2005-2015. The strong El Niño of 2015-2016 caused an estimated release of almost 19 Tg C, (<https://www.globalfiredata.org/regional.html#amazonas>) of which, 75% was due to forests.

Another possible cause of mortality could be a decrease in photosynthesis or GPP. Quantitative estimates of these quantities are now available from space-based observations of solar induced chlorophyll fluorescence (SIF). As plants absorb sunlight to perform

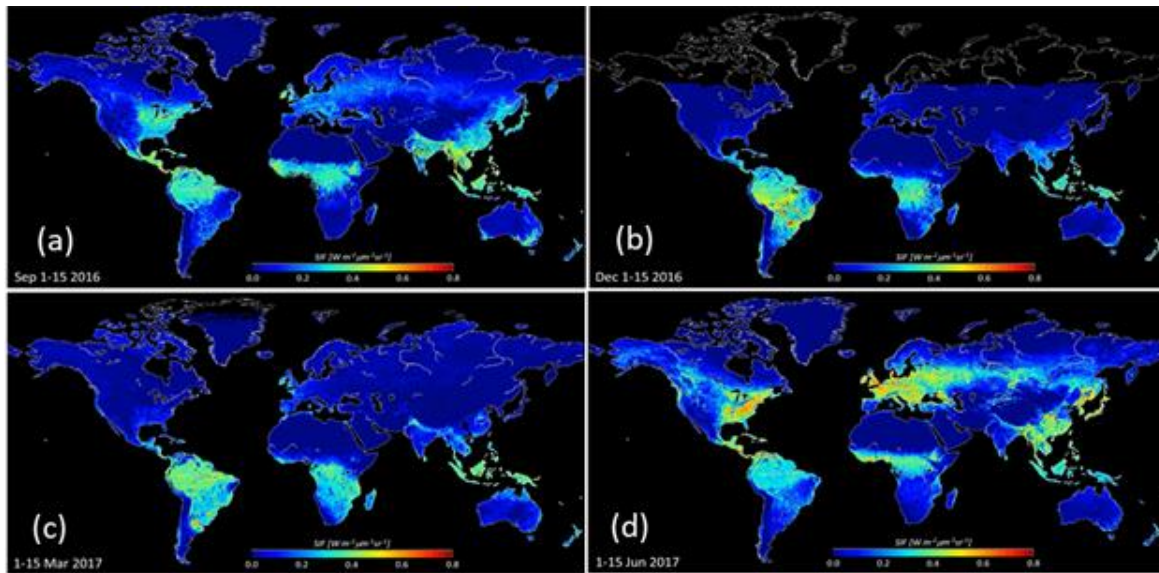


Figure 8. OCO-2 observations of SIF for (a) 1-15 September 2016; (b) 1-15 December 2016; (c) 1-15 March 2017, and 1-15 June 2017. Blue indicates low SIF and therefore low photosynthetic activity. The warmer colors indicate higher SIF and higher photosynthetic activity (Ying Sun, Personal communication, 2018).

photosynthesis, a fraction of that light (< 2%) is re-emitted at longer wavelengths (fluorescence), which can be detected in the cores of strong solar Fraunhofer lines by high resolution space-

based spectrometers (Frankenberg et al., 2014; Sun et al., 2018). SIF is a rapidly-responding indicator of light use efficiency that has been adopted as a functional proxy for GPP (Figure 8). Koren et al. (2018) find that SIF was strongly suppressed over areas with anomalously high temperatures and reduced availability of soil moisture. SIF fell below its climatological range starting from the end of the 2015 dry season (October), but returned to normal levels by February 2016 when atmospheric conditions returned to normal. Importantly, the impacts of the El Niño were not uniform across the Amazon basin.

The eastern part of the Amazon experienced a much larger (10–15%) reduction in SIF than the western part (2–5%). Koren et al. estimated the integrated loss of GPP across the Amazon basin relative to eight previous years to be 0.34 – 0.48 Pg C in the three-month period October-December 2015. Ciais et al. (2020) calculated a small sink of 0.06 Pg C yr<sup>-1</sup> using a combination of top down and bottom up inventories for the whole of South America, albeit with a large uncertainty of 0.29 Pg C yr<sup>-1</sup>. Thus, in contrast with the African sink, which shows no trend, the Amazon forest sink strength appears to be gradually declining, and that extreme events imposed on this trend cause additional losses.

### 5.3 Evidence for a Global Greening from Bottom-up Inventories

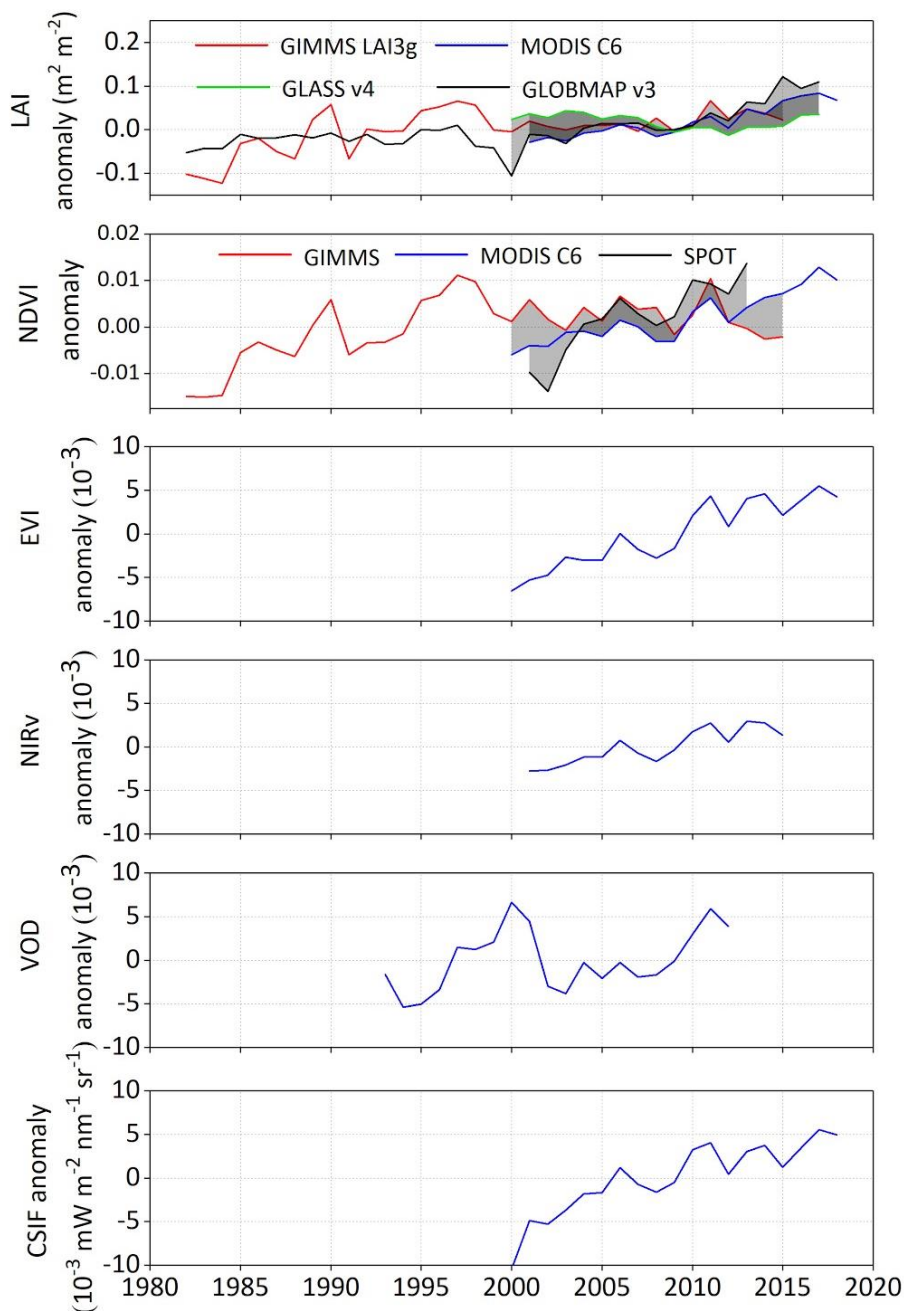
In a recent analysis of land use change over 1982-2016, Song et al. (2018) show that global tree cover has increased by 2.24 million km<sup>2</sup> (+7.1%) relative to the 1982 value. This overall net gain is the result of a net loss in the tropics being outweighed by a net gain in the extra tropics. Global bare ground cover has decreased by 1.16 million km<sup>2</sup> (-3.1%), most notably in agricultural regions of tropical and subtropical Asia. Of all land changes, 60% are associated with direct human activities and 40% with indirect drivers such as climate change.

Several factors are thought to impact vegetation greening, including rising atmospheric CO<sub>2</sub> concentrations, climate change, nitrogen deposition and land- use changes. Supporting the conclusions of Song et al., using long-term satellite records, Piao et al. (2020) find a significant global greening of vegetated areas since the 1980s, which they suggest has continued past 2010 (Figure 9). The most pronounced greening is observed in China and India and is due to afforestation and in particular agricultural intensification. They also analyzed a set of DGVMs that suggest CO<sub>2</sub> fertilization is the main driver of global vegetation greening. In addition, warming was the major cause of greening in boreal and Arctic biomes. However, in contrast to this result and in line with Hubau et al., they find that CO<sub>2</sub> fertilization had negative effects on greening in the tropics. Overall, they suggest that the greening contributed to mitigation of global warming through enhanced land carbon uptake and evaporative cooling, but could also lead to decreased albedo, which could potentially cause local warming.

847 Piao et al. (2020) find widespread greening occurred since the 1980's north of 50°N.  
848 Warming was considered the major cause of greening in boreal and Arctic biomes. However, a  
849 small, but persistent browning was seen at 3% of the high latitudes between 1982 and 2014. Such  
850 browning trends are caused by trends in disturbances such as fires, harvesting and insect



851 defoliation (Beck and Goetz, 2011). North American boreal forests exhibit browning areas nearly  
 852 20 times larger than the Eurasian boreal forests. Importantly, dramatic increases in fire



**Figure 9.** Changes in satellite-derived global vegetation indices, including anomalies in the normalized difference vegetation index (NDVI), Enhanced Vegetation Index (EVI), near-infrared reflectance of vegetation (NIRV), vegetation optical depth (VOD) and contiguous solar-induced fluorescence (CSIF) (Data: Piao et al., 2020.)

853 disturbances and insect infestation such as those from the bark beetle have also been seen in  
 854 browning areas, partially offsetting the decades long greening trend.

While the CO<sub>2</sub> emissions and uptake by agriculture and other land use practices are still highly uncertain, there is growing evidence that these activities are contributing to the global greening trend. Zeng et al. (2014) analyzed the impact of the Agricultural Green Revolution, which enhanced crop yields through hybridization, irrigation and fertilization, on the amplitude of the atmospheric CO<sub>2</sub> seasonal cycle. They use a DGVM to analyze CO<sub>2</sub> observations and atmospheric inversions for the 50-year period extending from 1961 to 2010. Between the decades 1961-1970 and 2001-2010, they attribute the observed the ~15% increase in the CO<sub>2</sub> seasonal amplitude to a peak growing season that now starts ~10 days earlier and lasts about two weeks longer, with the land biosphere absorbing ~0.5 Pg more carbon at its peak in July than it did at the beginning of the period. They find that the increased CO<sub>2</sub> seasonal amplitude originates primarily from two regions: the midlatitude cropland between 25° N and 60° N and the high-latitude natural vegetation between 50° N and 70° N. Zeng et al. note that the leading role of mid-latitude cropland in CO<sub>2</sub> uptake is reinforced by recent space-based observations of SIF, which show that at the peak of the growing season, cropland is substantially more productive than surrounding dense forests with similar climate conditions. These conclusions are reinforced by other studies of the relationship between SIF and crop productivity (Guan et al., 2016; 2017; He et al., 2020).

#### **5.4 Current Status and Next Steps in Bottom-up Land Carbon Observations and Analysis**

The overall picture that emerges from recent observations of aboveground biomass stocks is that the classical sinks in the tropical humid forests are slowly losing their sink strength, with these changes amplified by deforestation. In extra-tropical areas, greening has taken place due to afforestation and increased agriculture. At Arctic latitudes, browning, i.e. a loss of vegetation activity, is increasing. These trends provide the fragile background for a still slowly increasing land uptake. The underlying causes for these increases are complex and consist of interacting processes of CO<sub>2</sub> fertilization, nutrient and water availability compounded by climate variability. On top of this, the impact of human activities including deforestation and intensifying agriculture are additional complications.

This myriad of interacting processes complicates predictions of the future trajectory of the terrestrial sink. Until now, the sink has grown in harmony with the increase in fossil fuel emissions with the result that the atmospheric fraction has remained remarkably constant over the past 40 years or so. Theoretical and empirical evidence, such as that described here in this paper, suggest that this may stop at some point in the future. Theoretically it is unlikely that the global vegetation will continue to grow indefinitely. Water and nutrient shortages will impede increased growth (Wang et al., 2020). Similarly, the balance between GPP, as determined by plant photosynthetic properties, and the heterotrophic respiration is expected to shift. It has long been known that on short time scales, the efficiency of photosynthesis decreases beyond a critical temperature, while that of heterotrophic respiration continues to increase (Doughty et al., 2008; Mau et al., 2018). This behavior has recently been experimentally confirmed by Duffy et al. (2021) using FLUXNET data. How this works out at longer time scales, when heterotrophic carbon limitation on microbial decomposition may also start playing a role is virtually unknown (Soong et al., 2021). It is likely that the first signs of a deceleration of the terrestrial carbon sink, and a decoupling with the fossil fuel increase are already there, and that the atmospheric fraction will increase.



One factor that has impeded progress in the analysis of trends inferred from aboveground biomass stocks is they are not well represented in the current generation of DGVMs. For example, Sitch et al. (2015) use an ensemble of nine DGVMs to study global and regional processes and trends in the land sink for a period extending from 1990 - 2009. They conclude that for this period, the global land sink is increasing, led by CO<sub>2</sub> fertilization of plant production, with the largest increases seen in the natural ecosystems of the tropics. They find no significant trend in northern land regions. This is largely inconsistent with the observations presented above.

Another cause for concern is the recent finding that the effects of CO<sub>2</sub> fertilization appear to decline due to nutrient limitations and water availability (Wang et al., 2020). Like the results presented above, this conclusion is, largely based on satellite and ground-based data analysis of NDVI and SIF, and cannot be reproduced by the current generation of DGVMs. A decline in the effectiveness of CO<sub>2</sub> fertilization of the order as suggested by Wang et al. (2020) gives cause for concern of the efficiency of the terrestrial sink going forward. To date, both the land and ocean sink have increased steadily over time such that the airborne fraction has remained relatively stable since the 1960s (Raupach et al., 2014). If the land biospheric processes responsible for maintaining the airborne fraction are disrupted by human activities or climate variability and change, this could compromise our ability to predict the coupled evolution of the land carbon cycle and its interactions with an evolving climate.

Fortunately, advances in bottom-up observation capabilities and modeling tools are coming on line to facilitate more comprehensive and responsive monitoring and analysis of the land carbon cycle. Ground-based estimates of stocks and fluxes will continue to provide the most accurate and site-specific information. However, remote sensing observations from airborne and space-based active and passive sensors and modeling tools will play an increasingly important role for upscaling these results to yield useful constraints on regional to global scales.

Xiao et al. (2019) review the evolution of remote sensing observations of terrestrial carbon stocks over the past 50 years, spanning the electromagnetic spectrum from the visible, infrared, and microwave. They then review the methods being used to analyze the observations to yield quantitative estimates of carbon stocks and fluxes, including vegetation indices, SIF, light use efficiency models, DGVMs, as well as data driven (including machine learning) techniques. Xiao et al. discuss the use of these data and analysis techniques to quantify the impacts of disturbances and to quantify uncertainties in carbon stock estimates, noting advances achieved by integrating in situ and remote sensing observations into progressively more advanced, process-based carbon cycle models. Looking forward, they predict substantial improvements in our ability to track above ground biomass stocks, through the use of merged datasets, such as the NASA Harmonized LandSat and Sentinel 2 (HLS) products, ultra-high resolution imaging products from QuickBird, IKONOS, and UAVs, lidar measurements from GEDI, future active microwave products from NASA's NISAR (Rosen et al., 2016), TanDEM-L and BIOMASS missions (Quegan et al., 2019).

While in situ and space-based measurements of aboveground biomass play a critical role in efforts to monitor trends in managed and natural forest, they are not sufficient for monitoring the rapid turnover of carbon stocks in croplands and grasslands. Until recently, high resolution imaging observations and moderate resolution estimates of vegetation indices provided the primary tools for scaling up plot-based observations to national and continental scales. Recently, these capabilities have been augmented by space-based observations of SIF. SIF relates the

emission of excess radiative energy from the photosynthesis process of leaves at two wavelengths 685 nm and 740 nm to photosynthesis, or GPP. Estimates of SIF from GOME, GOME2, GOSAT, OCO-2 and TROPOMI are increasingly being used to monitor crop and grassland productivity and crop yield prediction (He et al., 2020; Peng et al., 2020; Parazoo et al., 2020; Qiu et al., 2020; Yin et al., 2020). Future SIF observations from the ESA FLuorescence EXplorer (FLEX), Japan's GOSAT-GW, NASA's GeoCarb, and the Copernicus CO2M missions promise substantial improvements in resolution.

To fully exploit these new measurements to describe long term trends in the terrestrial carbon cycle, the in situ and remote sensing measurements must be reconciled so that their climate data records can be combined to increase their spatial and temporal resolution and coverage. The protocol for cross-validating aboveground biomass products described by Duncanson et al. (2019) and the effort by the Forest Observation Initiative to develop a global in situ forest biomass databases for validating remote sensing observations (Schepaschenko et al., 2018) are positive steps in this direction. While the current generation of DGVMs and other terrestrial biosphere models are evolving rapidly and provided key insights into the processes driving the land carbon cycle, these modeling tools are still yielding widely diverging results the uptake of CO<sub>2</sub> by the land biosphere and its trends (i.e. Fisher et al., 2014; Sitch et al., 2015; Keenan and Williams, 2018; Parazoo et al., 2020). These limitations have raised concerns about their use in CO<sub>2</sub> emission inventory development activities (Grassi et al., 2018; Petrescu et al., 2020). Pioneering model intercomparison efforts such as the Carbon-Land Model Intercomparison Project (C-Lamp; Randerson et al., 2009) are being followed up by the International Land Model Benchmarking (ILAMB) project (see <https://www.ilamb.org/>) to address these concerns and accelerate the development of these critical tools.

## 6 The Atmospheric Carbon Cycle

Until recently, the concentrations of atmospheric CO<sub>2</sub> and other greenhouse gases were measured in situ by instruments deployed at surface stations, tall towers or on aircraft. Continuous measurements of atmospheric CO<sub>2</sub> were initiated in 1958 by Charles David Keeling of the Scripps Institution of Oceanography, when he established stations at Mauna Loa, Hawaii and the South Pole. These measurements are now being collected at a global network that includes the U.S. National Oceanic and Atmospheric Administration (NOAA) Global Greenhouse Gas Reference Network (GGGRN), the European Integrated Carbon Observation System (ICOS) network and other partners of the World Meteorological Organization Global Atmospheric Watch (WMO GAW) program. These in situ measurements provide the most

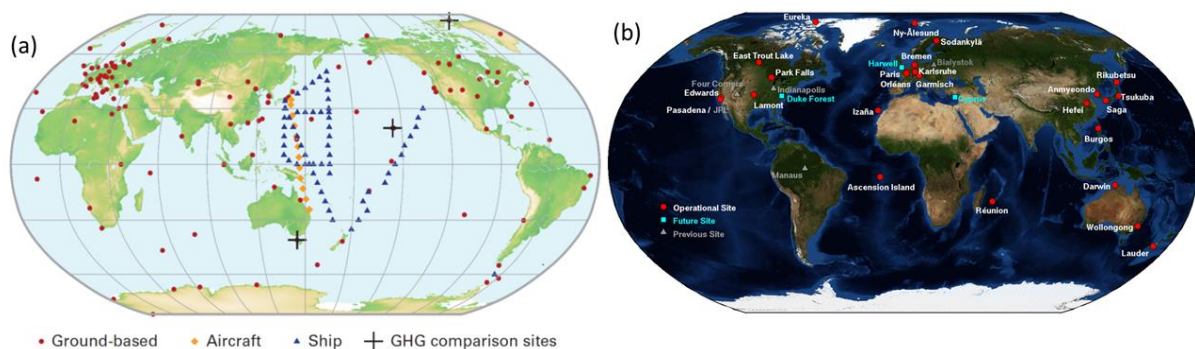


Figure 10: Spatial distribution of stations in the ground-based atmospheric CO<sub>2</sub> monitoring network. The vast majority of the stations are in North America and western Europe. (a) In situ CO<sub>2</sub> measurements are collected routinely at the WMO Global Atmospheric Watch Stations (from WMO Greenhouse Gas Bulletin, 25 Nov. 2019). (b) Solar-looking remote sensing observations of CO<sub>2</sub> are collected at Total Carbon Column Observing Network (TCCON) stations.

accurate estimates of the CO<sub>2</sub> concentrations and their trends on global scales. They also quantify chemical and isotopic tracers, such as carbon-14 (<sup>14</sup>C), which help to discriminate fossil fuel from biogenic contributions to the observed CO<sub>2</sub> trends. Perhaps the most important asset of the ground based in situ measurements is the length of their climate data records, which now extend over 70 years. While the ground-based networks have grown over time, the spatial resolution and coverage of in situ measurements are still far too limited to identify and quantify the natural and anthropogenic emission sources emitting CO<sub>2</sub> into the atmosphere and the natural “sinks” absorbing it at the surface on spatial scales ranging large urban areas to nations. The coverage is particularly sparse in Arctic, boreal and tropical land regions and over most ocean basins (Figure 10).

Recent advances in space based remote sensing capabilities are providing new insights into the current state of the atmospheric carbon cycle as well as the primary processes controlling its interactions with the land and ocean carbon reservoirs. With the launch of Japan’s Greenhouse gases Observing SATellite, GOSAT in 2009, and NASA’s Orbiting Carbon Observatory in 2014, space-based remote sensing measurements are complementing the ground-based and airborne CO<sub>2</sub> measurements with much greater spatial resolution and coverage, yielding tens of thousands of observations over the sunlit hemisphere each day. These two satellites have recently been joined by their sister missions, GOSAT-2 and OCO-3, providing additional coverage and resolution. These space-based sensors collect high-resolution spectra of reflected sunlight within molecular oxygen (O<sub>2</sub>) and CO<sub>2</sub> bands that can be analyzed to yield precise, spatially resolved estimates of the column-averaged CO<sub>2</sub> dry air mole fraction, XCO<sub>2</sub>. The principal challenge of this technique is the need for unprecedented levels of precision and accuracy for a space-based atmospheric trace gas measurement. While intense local sources, such as large coal-fired power plants or large urban areas can increase the near surface CO<sub>2</sub> concentrations by more than 10%, these variations decay rapidly with altitude, such that they rarely yield XCO<sub>2</sub> variations larger than 1-2 ppm (0.25 to 0.5%) on the spatial scale of a satellite footprint (1 to 100 km<sup>2</sup>). Natural sinks of CO<sub>2</sub>, such as forests or ocean basins, produce even smaller changes in XCO<sub>2</sub>.

In addition to XCO<sub>2</sub>, these sensors collect simultaneous observations of SIF, providing additional constraints on the CO<sub>2</sub> uptake by the land biosphere. The primary challenge of this approach is the need for unprecedented precision and accuracy to resolve the small (< 0.25% or 1 ppm) XCO<sub>2</sub> variations associated with surface sources and sinks of CO<sub>2</sub>. While these space-based measurements are not as accurate as those collected by ground-based and airborne systems, they complement those observations with improved coverage and spatial resolution.

Atmospheric CO<sub>2</sub> estimates collected by ground-based, airborne, and space-based sensors are now being assimilated into atmospheric inverse models, along with meteorological data and space based global measurements of the land biosphere to provide new insights into the carbon cycle. These models illustrate the spatial and temporal relationships between SIF and the seasonal CO<sub>2</sub> drawdown in the northern hemisphere spring (Eldering et al., 2017; Byrne et al., 2018; Yin et al., 2020; Qui et al., 2020). They have also been used to quantify the atmospheric signature of the land biosphere’s response to the record-setting 2015-2016 El Niño (Chatterjee et al., 2017; Liu et al., 2017; Palmer et al., 2019). The high spatial and temporal resolution of these atmospheric CO<sub>2</sub> observations, combined with their global coverage, should facilitate the development of improved diagnostic models for studying processes operating in the present-day carbon cycle. This evolving land-ocean-atmosphere carbon monitoring system also fosters the

development of more comprehensive and reliable prognostic models for predicting the evolution of the carbon cycle as it responds to climate change.

From a carbon monitoring perspective, these atmospheric measurements are now providing quantitative constraints on CO<sub>2</sub> emission and uptake at policy relevant spatial and temporal scales. CO<sub>2</sub> estimates retrieved from GOSAT and OCO-2 measurements clearly show persistent positive anomalies associated with the anthropogenic emissions over East Asia, Western Europe and eastern North America (Hakkarainen et al., 2016; 2019; Wang et al., 2018). On smaller scales, space-based XCO<sub>2</sub> estimates are being used to quantify CO<sub>2</sub> emissions from large urban areas (Hedelius et al., 2018; Wu et al., 2018; Wu et al., 2020) and individual power plants (Nassar et al., 2017; Reuter et al., 2019). Estimates of CO<sub>2</sub> emissions and uptake derived from atmospheric measurements of XCO<sub>2</sub> are not as source specific as traditional bottom-up statistical methods, which infer CO<sub>2</sub> emissions from fuel use, power generation statistics, etc. However, they complement those methods by providing an integral constraint on the total amount of CO<sub>2</sub> added to or removed from the atmosphere by all natural and anthropogenic processes.

The availability of high spatial resolution atmospheric emissions products within a few weeks of acquisition could help to identify and track rapidly-evolving emission hotspots that are often missed in the bottom-up statistical inventories. As these tools are integrated into a more comprehensive carbon management system, they could also help carbon managers to assess the effectiveness of their carbon management strategies, and help to identify emerging emission reduction opportunities.

## **6.1 The Tropical Land Carbon Cycle Did Not Recover as Expected from 2015–2016 El Niño**

As noted above, bottom-up inventories indicate that the strength of the tropical forest sink is gradually declining over time. This conclusion is reinforced by atmospheric CO<sub>2</sub> observations. Gatti et al. (2014), using in situ observations of atmospheric CO<sub>2</sub>, determined that the Amazon basin lost  $0.48 \pm 0.18$  Pg C yr<sup>-1</sup> during a dry year (2010), but was carbon neutral ( $0.06 \pm 0.1$  Pg C yr<sup>-1</sup>) when measurements were taken during a wet year (2011). Correcting for carbon losses, they derived a basin net biome exchange that was carbon neutral during the dry year. During the wet year, vegetation was a net carbon sink of  $0.25 \pm 0.14$  Pg C yr<sup>-1</sup>. The latter value compares well with the bottom-up estimates by Hubau et al. (2020).

The availability of time-resolved global measurements of atmospheric CO<sub>2</sub> at high spatial resolution is providing a more comprehensive description of impacts of severe weather and climate on the exchange of carbon between land and ocean reservoirs and the atmosphere on regional scales. These emerging capabilities were recently demonstrated in studies of the record-setting 2015-2016 El Niño. GOSAT and OCO-2 measurements collected over the central and eastern tropical Pacific basin were combined to quantify 0.5 ppm XCO<sub>2</sub> decreases associated with reductions in outgassing in the tropical Pacific Ocean during March through July of 2015 (Chatterjee et al., 2017). These reduced CO<sub>2</sub> values were then replaced by 0.5 to 2 ppm increases in XCO<sub>2</sub> associated with reduced uptake and increased emissions of CO<sub>2</sub> by tropical forests in South America, Africa and tropical Asia (Liu et al., 2017; Heymann et al., 2017; Palmer et al., 2019; Crowell et al., 2019; Figure 11).

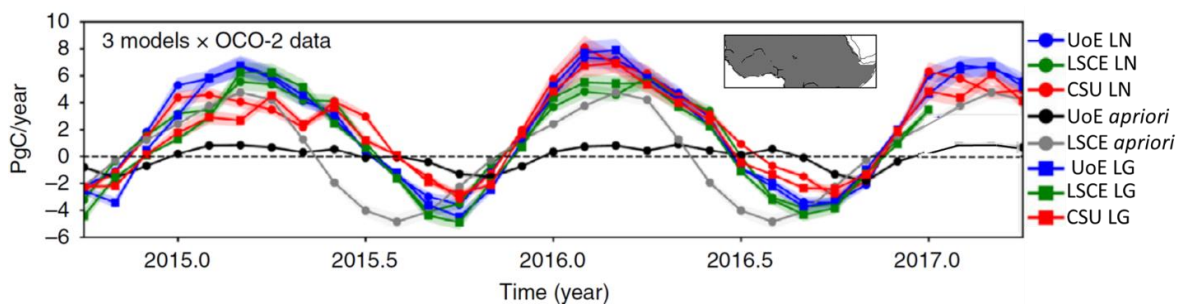


Figure 11. CO<sub>2</sub> fluxes from tropical northern Africa inferred from the University of Edinburgh (UoE), LSCE and Colorado State University (CSU) models constrained by in situ CO<sub>2</sub> measurements as well as XCO<sub>2</sub> data from GOSAT and OCO-2. Positive fluxes indicate CO<sub>2</sub> emissions from the land surface to the atmosphere. LN and LG denote OCO-2 XCO<sub>2</sub> measurements taken using nadir and glint observing modes, respectively. The geographical region is shown in the inset. Fluxes inferred from OCO-2 data have larger amplitudes and a larger seasonal cycle than those from in situ data. An extended mission will provide new opportunities to validate these results and track their changes (Adapted from Palmer et al., 2019).



Liu et al., (2017) find that the pan-tropical biosphere released an additional  $2.36 \pm 0.34$  Pg C into the atmosphere, or about 78% of the global total. Emissions originated throughout the tropics with  $0.91 \pm 0.24$ ,  $0.85 \pm 0.21$ , and  $0.60 \pm 0.31$  Pg C from tropical South America, tropical Africa, and tropical Asia, respectively. Although the enhanced emissions from these three regions were comparable, *different* processes appeared to dominate in each region. Fire emissions dominated over tropical Asia. Both increased respiration and fires associated with historically high temperatures dominated over tropical Africa. Increased CO<sub>2</sub> over the Amazon was attributed to GPP reductions associated with drought. These results support the hypothesis that El Niño related increases in CO<sub>2</sub> growth rates are primarily due to tropical land carbon fluxes, but they show that specific mechanisms can differ from continent to continent.

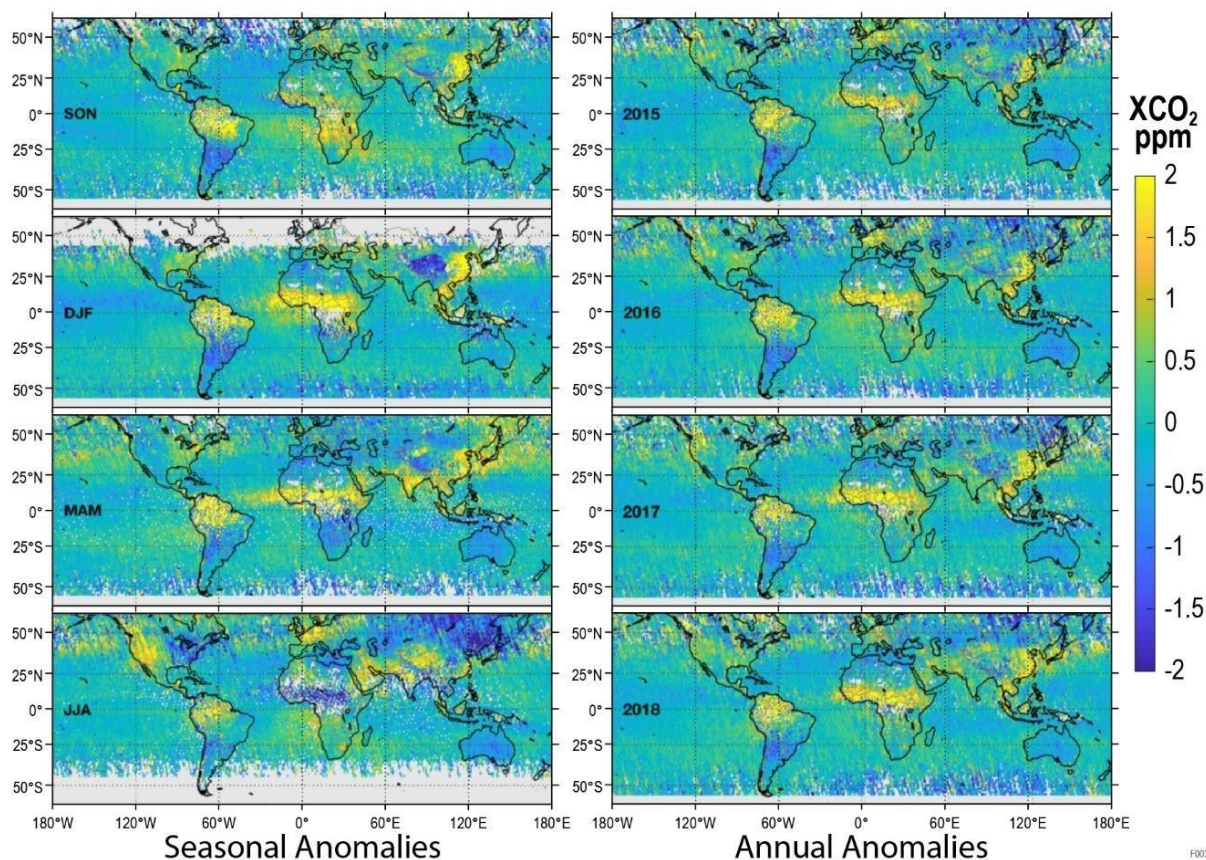


Figure 12. OCO-2 XCO<sub>2</sub> estimates indicate that some regions have persistently high (positive) or low (negative) spatial anomalies that persist from season to season (left, integrated over the four-year period) and from year to year (right). A longer data record is needed to determine whether the tropical anomalies reflect short-term climate variations or long-term climate trends. (Adapted from Hakkarainen et al., 2019).

Palmer et al., (2019) use three different models to analyze *in situ* CO<sub>2</sub> measurements along with XCO<sub>2</sub> and SIF observations from Japan's Greenhouse gases Observing SATellite (GOSAT) and OCO-2 (Figure 11). Like Liu et al., in 2015–2016, they find that the largest CO<sub>2</sub> emissions over western Ethiopia and western tropical Africa, where there are large soil organic carbon stores and substantial land use change. While the amplitude of the XCO<sub>2</sub> anomalies that

produced these sources may have been overestimated in the early OCO-2 XCO<sub>2</sub> products used in this investigation (version 7), they clearly reveal important components of the tropical carbon budget that is largely missing from in carbon flux inverse models constrained by *in situ* measurements alone.

As the 2015–2016 El Niño transitioned to a weak La Niña in 2017, and then to more neutral conditions in 2018, OCO-2 XCO<sub>2</sub> and SIF estimates indicate that tropical forests, once thought to be significant net sinks of CO<sub>2</sub> (Pan et al., 2011; Sellers et al., 2018) may now be persistent net sources (Hakkarainen et al., 2019; Palmer et al., 2019; Crowell et al., 2019). Hakkarainen et al. (2019) processed OCO-2 Version 9 (V9) XCO<sub>2</sub> data to remove the seasonal cycle and reveal spatially persistent anomalies. They find positive XCO<sub>2</sub> anomalies over tropical forests with amplitudes as high as 2 ppm above the background (Figure 12). The spatial extent of the Amazon anomaly was slightly greater during the 2015–2016 El Niño than in 2017, but the positive XCO<sub>2</sub> anomalies there have persisted from season to season and from year to year throughout the OCO-2 mission. The Amazon is now a persistent *source* of CO<sub>2</sub>, rather than a sink. Positive anomalies over topical Africa and Southeast Asia are seen on annual time scales. However, in Figure 11, tropical African fluxes are negative during June–July–August (JJA), indicating that this region becomes a weak sink during that season (Palmer et al., 2019). These conclusions are supported by some satellite-based aboveground biomass studies (Baccini et al., 2017; Wigneron et al., 2020), but are inconsistent with plot-based studies (Pan et al., 2011; Hubau et al., 2020), which conclude that tropical forests are absorbing less CO<sub>2</sub>, but are still a net sink of carbon.

Additional insight into the tropical land carbon cycle can be obtained by comparing XCO<sub>2</sub> anomalies to observations of SIF. Specifically, the largest positive CO<sub>2</sub> anomalies derived from the space-based XCO<sub>2</sub> estimates are seen in regions where SIF observations indicate the highest photosynthetic activity (Figure 8). This suggests that in spite of significant growth, tropical forests are now emitting more CO<sub>2</sub> than they absorb, when integrated over the annual cycle. This may have always been the case, but it may also be due to increased respiration, drought stress, fires, and other processes.

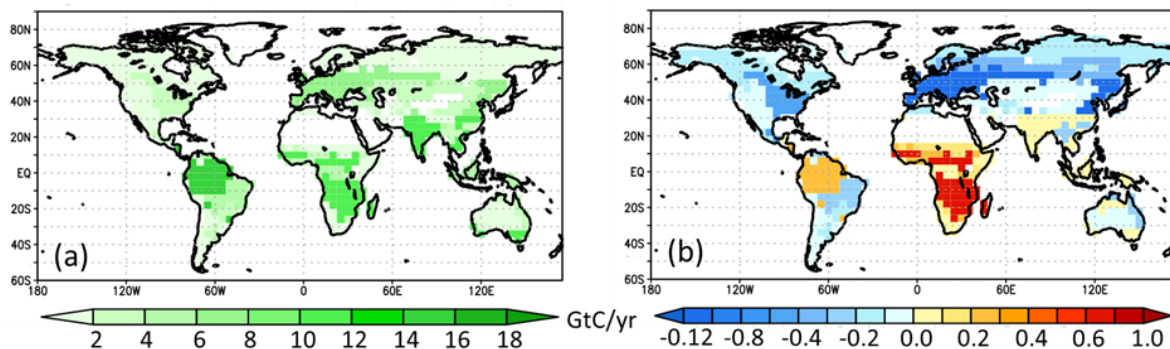


Figure 13. (a) GPP from OCO-2 SIF and (B) Net Biospheric Exchange (NBE) from XCO<sub>2</sub> and SIF, both expressed in gigatons of carbon per year (Pg C yr<sup>-1</sup>) for 2015–2018. Negative NBE indicates sinks while positive values indicate sources. NBE is typically < 5% of the GPP, but is positive in the tropics where we see the highest GPP, in sharp contrast to existing models (Junjie Liu, Personal communication, 2019).

These conclusions are reinforced by experiments that assimilate XCO<sub>2</sub> and SIF results into atmospheric inverse models (Liu et al., 2017; 2020; Palmer et al., 2019; Chevallier et al., 2019; Crowell et al., 2019). Even in the Amazon and tropical Africa, where the GPP exceeds 10 Pg C yr<sup>-1</sup> (Figure 13a), the net biospheric exchange (NBE) can still be positive (Figure 13b). This contradicts results from carbon flux inversion models constrained by in situ data, alone, which indicate that, at least until recently, these tropical forests were significant net sinks of CO<sub>2</sub> (Hubau et al., 2020).

## 6.2 New Insights into the Extratropical Land Biosphere from Space

Unlike for the tropics, the XCO<sub>2</sub> anomalies and retrieved CO<sub>2</sub> fluxes from northern temperate and boreal forests are generally more consistent with expectations. During the NH summer (JJA in Figure 12), we see negative XCO<sub>2</sub> anomalies. Annual average flux inversion experiments show moderately strong GPP and negative NBE (Figure 13). The OCO-2 XCO<sub>2</sub> and SIF measurement therefore indicate that these forests have continued to be significant net CO<sub>2</sub> sinks as the CO<sub>2</sub> seasonal cycle amplitude has grown in response to warming. While mid- and high-latitude CO<sub>2</sub> fluxes derived from inverse models constrained by OCO-2 XCO<sub>2</sub> and SIF estimates are generally consistent with those from earlier flux inversion experiments constrained by *in situ* CO<sub>2</sub> data alone (Lucht et al., 2002; Graven et al., 2013; Byrne et al., 2018), there are substantial differences in amplitude and phase in some regions (Reuter et al., 2017; Crowell et al., 2019; Palmer et al., 2019; Chevallier et al., 2019).

Space-based observations of XCO<sub>2</sub> and SIF provide unique opportunities to study the relationships between the land and atmospheric carbon cycles and the hydrological cycle. Yin et al. (2020) combine SIF with atmospheric CO<sub>2</sub> observations to quantify the effects of large-scale flooding on cropland carbon sequestration. Widespread flooding during spring and early summer of 2019 delayed crop planting across the U.S. Midwest. As a result, satellite observations of SIF from OCO-2 and the Tropospheric Monitoring Instrument (TROPOMI) reveal a shift of 16 days in the seasonal cycle of photosynthetic activity relative to 2018, along with a 15% lower peak photosynthesis. Yin et al. find that the 2019 anomaly produced an estimated GPP reduction of -0.21 Pg C in June and July that was partially compensated in August and September with a +0.14 Pg C increase. The growing season integral corresponds to a 4% reduction in cropland GPP for the Midwest, but a 3% increase for areas where cropland occupies less than 10% of the land. Using an atmospheric transport model, they show that a decline of ~0.1 Pg C in the net carbon uptake in June and July is consistent with observed ~10 ppm CO<sub>2</sub> enhancements in the midday boundary layer from the Atmospheric Carbon and Transport - America (ACT-America) aircraft and the ~1 ppm increases in XCO<sub>2</sub> seen by OCO-2.

In another study, Gonsamo et al. (2019) combined OCO-2 SIF observations with soil moisture (SM) observations from NASA's Soil Moisture Active Passive (SMAP) mission to study the impact of environmental limiting factors on terrestrial ecosystem productivity of drylands and croplands. For drylands (dry sub-humid, semi-arid, and arid zones) and the majority of croplands, soil water content is typically low and topsoil moisture is critical for plant growth. As expected, SMAP SM retrievals show positive daily relationships with OCO-2 SIF for drylands and croplands of the tropics and Australia, where SM is limiting plant growth and concurrent data records are sufficient to make statistical inferences. Negative relationships between SIF and SM were observed in forested areas of mid-latitude dry sub humid zones with high average annual SM. In these regions, SIF showed a positive relationship with air



temperature. They find strong evidence that the OCO-2 SIF is accurately capturing monthly SMAP SM dynamics, particularly for regions with distinct seasonality of rainfall such as Sub-Saharan North Africa, Indian subcontinent, and southern Africa.

### 6.3 *Lessons Learned from Space-based XCO<sub>2</sub> Estimates over the Ocean*

In general, XCO<sub>2</sub> anomalies over the ocean were expected to be much smaller than those over land, but they have still yielded some surprises in first-generation space-based observations of CO<sub>2</sub>. For example, the earliest version of the OCO-2 product, version 7 (v7) revealed large (2–3 ppm) positive XCO<sub>2</sub> anomalies over the midlatitude ocean (20 – 50 S) in the southern hemisphere during the Austral winter. These anomalies were initially suspicious because they were not seen in the XCO<sub>2</sub> observations from the Total Carbon Column Observing Network (TCCON) stations at Lauder, New Zealand or Wollongong, or Reunion Island (Wunch et al., 2017). However, they could not be dismissed immediately because there are few CO<sub>2</sub> measurements over the ocean at the latitudes where they had their largest amplitudes. Retrieval algorithm validation efforts subsequently traced these anomalies to a bias introduced by a thin layer of volcanic aerosol in the stratosphere that had been neglected in the retrieval algorithm (O'Dell et al., 2018). This bias was corrected in the version 8 (v8) and later versions of the OCO-2 products by explicitly retrieving the stratospheric aerosol optical depth.

While the v8 and version 9 (v9) OCO-2 XCO<sub>2</sub> products were generally more reliable over the ocean at mid latitudes (O'Dell et al. 2018; Kiel et al., 2019), they included a new anomaly that compromised their acceptance by the ocean carbon cycle community. In contrast to existing carbon cycle models, which assume that ocean-atmospheric CO<sub>2</sub> fluxes are correlated with ocean surface pCO<sub>2</sub> values, Figure 12 shows persistent negative XCO<sub>2</sub> anomalies over the tropical Pacific and Indian Oceans, where surface ocean pCO<sub>2</sub> is high (Figure 3, top). The lowest persistent spatial anomalies in XCO<sub>2</sub> are seen in regions that receive the highest rainfall (1000 – 3000 mm/year), suggesting a possible correlation. Comparisons of OCO-2 XCO<sub>2</sub> estimates with TCCON XCO<sub>2</sub> observations indicate that the OCO-2 estimates are systematically low over the tropical Pacific and Indian Oceans (S. Kulawik personal communication, 2019), but this does not explain the apparent spatial correlation of the deepest anomalies with the rain bands. These spatial correlations are not entirely surprising. Atmospheric CO<sub>2</sub> dissolves in cloud and raindrops, reducing the pH of rain to values between 5 and 6 even for regions well away from anthropogenic nitrate and sulfate acid rain sources (Willey et al., 2000; Bogan et al., 2009; Liu et al., 2010). Liu et al. (2010) estimate that precipitation transports ~0.2 Pg C yr<sup>-1</sup> (0.35 to 0.7 Pg CO<sub>2</sub>) from the atmosphere to the ocean. This process is not seen in carbon cycle models because most assume that rain contains no inorganic carbon, and falls with a pH of 7.0. If these OCO-2 XCO<sub>2</sub> anomalies can be validated, this assumption will have to be revisited.

### 6.4 *The Continuing Need for in situ Measurements of CO<sub>2</sub> and other Greenhouse Gases*

While this first generation of space-based measurements of CO<sub>2</sub> have provided new insights into land, ocean, and atmospheric carbon cycles, they have also revealed some key limitations of this approach, and the continuing need to maintain and expand the ground-based and airborne atmospheric greenhouse measurement networks. First, while CO<sub>2</sub> and CH<sub>4</sub> can now be measured from space with the accuracies need to quantify surface sources and sinks, other critical greenhouse gases including nitrous oxide (N<sub>2</sub>O), chlorofluorocarbons (CFCs), hydrochlorofluorocarbons (HCFCs), hydrofluorocarbons (HFCs), perfluorocarbons (PFCs), and

sulfur hexafluoride (SF<sub>6</sub>) cannot be measured to adequate accuracy. For these gases, as well as CO<sub>2</sub> and CH<sub>4</sub>, in situ measurements continue to provide the most reliable means for tracking their concentrations on hemispheric to global scales. These measurements therefore continue to be a critical element of any global stocktake of these gases, like that mandated by the UNFCCC Paris Agreement. Other species that are useful for discriminating fossil fuel from biospheric CO<sub>2</sub> emissions, such as carbon-14 (<sup>14</sup>C) can also only be measured in situ (Miller et al., 2012; 2020).

While spatial coverage is a critical asset of space-based atmospheric CO<sub>2</sub> measurements, there are situations where in situ measurements complement the coverage provided by these data. For example, remote sensing observations of near-surface CO<sub>2</sub> can only be collected in regions that are sufficiently free of clouds and aerosols. In situ measurements are therefore essential in persistently cloudy regions like those occupied by the intertropical convergence zone (ITCZ), and the Asian summer monsoon. In situ measurements are also needed at high latitudes during the winter, where clouds, low sun angles, and the low reflectivity of snow and ice covered surfaces at shortwave infrared wavelengths reduce the number and quality of remote sensing observations of CO<sub>2</sub>. In addition, in situ data from instruments on airborne platforms complement the vertically-integrated remote sensing results with information about the vertical profile of CO<sub>2</sub>, which can provide critical insight into net flux of CO<sub>2</sub> into the atmosphere. Finally, because the air-sea flux of CO<sub>2</sub> is determined mainly by the pCO<sub>2</sub> gradient between the ocean surface layer and the atmospheric surface boundary layer, in situ observations of near-surface atmospheric CO<sub>2</sub> concentrations are critical over the ocean.

Finally, because both surface in situ and remote sensing observations are more accurate than space-based remote sensing measurements, these data are critical for validating the remote sensing measurements. Currently, space-based XCO<sub>2</sub> estimates are validated against ground-based remote sensing observations collected by the Total Carbon Column Observing Network (TCCON), which, in turn, are validated against airborne in situ measurements, to provide a transfer standard to the WMO atmospheric greenhouse gas standard (Wunch et al., 2017). This validation approach has yielded valuable information about surface and atmospheric properties that introduce bias and scatter in the space-based remote sensing measurements. However, while this network now includes 27 stations in 14 countries, it still provides little coverage of the tropics and high latitudes, some of the most dramatic features are seen in the space-based remote sensing data.

While the number of ground-based and airborne in situ and remote sensing reference stations have grown slowly over the past decade, new measurement capabilities are coming on line that promise substantial increases in coverage. The up-looking remote sensing measurements being collected by the TCCON spectrometers are being complemented by measurements from smaller, less costly, and more portable Bruker EM27/SUN systems. These spectrometers are now being deployed as networks in urban settings (Hedelius et al., 2018) and in remote locations (Frey et al., 2019). In situ vertical profiles of CO<sub>2</sub>, CH<sub>4</sub> and other gases are now being collected at altitudes as high as 25 km by AirCore instruments deployed on low-cost weather balloons (Karion et al., 2010). Additional in situ profiles and upper tropospheric measurements are now being made by commercial aircraft in Japan's Comprehensive Observation Network for Trace gases by Airliner (CONTRAIL) and Europe's In-service Aircraft for a Global Observing System (IAGOS).

## 7 Discussion

When integrated over the industrial age, the land sink has roughly balanced the land use source. The ocean has therefore been the only *de facto* cumulative net sink of fossil carbon from the atmosphere (Friedlingstein et al., 2019; 2020). However, since 1958, when continuous atmospheric CO<sub>2</sub> measurements have been available, CO<sub>2</sub> emissions from fossil fuel combustion have increased by about a factor of four, from less than 2.5 Pg C yr<sup>-1</sup> to almost 10 Pg C yr<sup>-1</sup> in 2019. During this period, the land sink has taken up an increasing fraction of anthropogenic emissions. Together, sinks in ocean and on land have absorbed enough anthropogenic CO<sub>2</sub> to limit the fraction that has remained in the atmosphere to a remarkably constant value around 45% (Raupach et al., 2014). This implies that, to first order, the uptake capacity of the ocean and land sinks has increased in step with the emissions (Friedlingstein et al., 2020). There has been debate as to whether increases in the atmospheric fraction, i.e. declines in sink efficiency, are already observable (Canadell et al., 2007; Knorr 2009; Raupach et al., 2014). Even if an increasing atmospheric fraction is not yet detectable, process-level understanding and regional trends indicate that the atmospheric fraction should increase as climate change progresses. While the exact timing and magnitude of changes in the land and ocean sinks remains unclear, the likelihood is high that substantial climate-carbon feedbacks will occur during this century. Any upward change in the atmospheric fraction, or reduction in sink capacity, will decrease the allowable fossil carbon that can still be burned without violating the temperature targets specified in the Paris agreement.

For the ocean, despite remaining uncertainties and missing closure terms, distinct methodologies for quantifying the ocean sink agree that the sink has increased over the industrial era, including in recent decades. Since the uptake of atmospheric CO<sub>2</sub> on annual to decadal time scales is primarily controlled by the pCO<sub>2</sub> gradient at its surface, the carbon sink is expected to grow steadily as long as near-exponential growth of atmospheric pCO<sub>2</sub> continues. However, if anthropogenic emissions are reduced, atmospheric pCO<sub>2</sub> will grow more slowly, and thus there will be a reduced ocean carbon sink even if the ocean circulation and chemical buffer capacity do not change. To understand these likely changes, it is essential that ocean carbon studies start to focus more attention on the near-term response to emission mitigation scenarios (Hausfather and Peters, 2020). If emissions are not mitigated, current climate models suggest that by the middle to late 21st century, a slowing ocean overturning rate and reduced chemical capacity in the ocean will reduce the rate of growth in the ocean sink.

To develop an integrated ocean carbon observing system that can track the evolution of the ocean sink on the annual to interannual timescales most relevant to climate change policy, we need to sustain existing and continue to develop improved observation systems for the surface and interior ocean. Ocean carbon instruments deployed on autonomous platforms are revolutionizing ocean carbon measurement spatial and temporal resolution and coverage, but reduced uncertainties in the carbonate constants are needed to fully exploit these data. High-quality shipboard observations will continue to be required. We also need improved ocean hindcast models and better understanding of uncertainties in observation-based data products derived through statistical extrapolation of sparse surface ocean pCO<sub>2</sub> data.

For the land carbon cycle, the picture is more complicated and appears to already be changing. Classical sinks in the tropical humid forests are slowly losing their strength and these changes are amplified by the losses associated with deforestation, forest degradation and extreme climate events. In the extra tropics, bottom-up measurements show evidence for a mid-latitude

greening associated with afforestation and increased agriculture, accompanied by increased browning in some parts of the Arctic. Overall, these trends provide the fragile background for a slowly increasing land uptake of CO<sub>2</sub>. Over the next century, carbon-climate feedback is expected to play a larger role in the Arctic, although the magnitude and timing of those changes is subject to considerable debate (Schuur et al., 2015).

Space-based remote sensing observations are helping to revolutionize our ability to monitor the land and atmospheric carbon cycles to anthropogenic forcing and a changing climate. From a bottom-up perspective, microwave and lidar measurements are providing higher spatial and temporal resolution estimates of above ground biomass stocks. SIF measurements are providing a more responsive estimate of light use efficiency and CO<sub>2</sub> uptake by plants. From a top-down perspective, space-based remote sensing estimates of XCO<sub>2</sub> are complementing ground-based and aircraft in situ measurements with much greater spatial and temporal resolution and coverage. These space-based measurements sometimes reinforce, amplify or contradict the results inferred from ground-based in situ measurements, painting a somewhat controversial picture of the evolution of the land carbon cycle. For example, in the tropics, both space-based microwave estimates of above ground biomass (Wigneron et al., 2020) and top-down atmospheric inverse models constrained by space-based estimates of XCO<sub>2</sub> (Liu et al., 2017; 2020; Palmer et al., 2019; Crowell et al., 2019) indicate that the humid tropical forests never fully recovered from the 2015-2016 El Niño, and have transitioned from net sinks to net sources of CO<sub>2</sub>. Meanwhile, at mid- and high latitudes, bottom-up and top-down models constrained by space-based remote sensing measurements largely reinforce the in situ results, showing a long term increase in the amplitude of the seasonal cycle (Graven et al., 2013; Byrne et al., 2018; 2020; Liu et al., 2020) and that mid-latitude and boreal forests are strong net sinks of CO<sub>2</sub> when averaged over the seasonal cycle. More generally, the space based measurements are also providing more information about rapid changes in the land carbon cycle associated with severe weather, such as droughts (Gonsamo et al., 2019; Castro et al., 2020) and floods (Yin et al., 2020). They are also beginning to provide estimates of CO<sub>2</sub> emissions from fossil fuel combustion and other human activities (Hakkarainen et al., 2016; 2019; Wang et al., 2018; Hedelius et al., 2018; Wu et al., 2018; 2020; Reuter et al., 2019).

In spite of these advances, the reliability of the space-based remote sensing results are still a subject of substantial debate within the land carbon cycle community. This is especially true for the tropics, where CO<sub>2</sub> fluxes derived from the space-based XCO<sub>2</sub> estimates differ in both sign and magnitude from the results of earlier flux inversion experiments constrained by bottom-up stock or flux estimates or ground-based in situ measurements of atmospheric CO<sub>2</sub>. This apparent inconsistency suggests one of three possibilities. First, the space-based XCO<sub>2</sub> estimates might still include biases that compromise the accuracy of the top-down flux estimates. Recent efforts to validate the space-based XCO<sub>2</sub> estimates using measurements from TCCON and other standards (Wunch et al., 2017) indicate biases with amplitudes less than one third as large as the observed tropical XCO<sub>2</sub> anomalies. However, there are few TCCON stations or other validation capabilities in the tropics. Second, fluxes constrained by surface in situ measurements, alone, may tell an incomplete story of the land carbon cycle in sparsely sampled regions. The spatial resolution and coverage provided by surface in situ measurements of carbon stocks, fluxes, or atmospheric CO<sub>2</sub> are still very limited, especially in the tropics and boreal regions, where the largest flux differences are seen. Both top-down and bottom-up methods may yield unreliable results where there are few measurements. Third, flux estimates based on the much denser space-based XCO<sub>2</sub> measurements may be tracking changes in the natural carbon

cycle on time and space scales too short to be resolved by the in situ measurements of stocks or CO<sub>2</sub> concentrations. There is increasing evidence from atmospheric CO<sub>2</sub> measurements and modeling studies and other carbon cycle observations, that the tropical land carbon cycle is evolving rapidly in response to human activities (deforestation and degradation, biomass burning, land use change) and climate change (drought, heat stress, flooding). Satellite observations of aboveground biomass (Baccini et al., 2017) also support this conclusion.

All three of these possibilities may be valid to some extent. To address these questions and improve our understanding of the land carbon cycle in the tropics and at high latitudes, we need additional ground-based and aircraft validation measurements in these areas, advances in the space-based measurement calibration, retrieval algorithms, and validation techniques to further reduce regional scale biases and a longer, continuous space-based data record that clearly resolves the carbon cycle impacts of short-term climate anomalies from long-term secular climate trends.

## 8 Conclusions

Our understanding of the carbon cycle and its response to natural and anthropogenic forcing has grown steadily over the past two decades as more advanced carbon cycle measurement systems have been deployed and their results have been analyzed with more sophisticated diagnostic and prognostic carbon cycle models. These results reveal a strongly coupled, dynamic system that responds on daily, to seasonal, to interannual time scales across spatial scales spanning individual fields, forest plots or coal-fired power plants on land or individual eddies in the ocean to entire continents or ocean basins. While surface or ocean interior in situ measurements still provide the most precise and accurate results, these data are now being complemented by a broad range of ground and ocean-based, airborne and space based remote sensing observations that extend their spatial resolution and coverage. The growing international collaborations between the top-down and bottom-up carbon cycle measurement and modeling communities, with their continued focus on a rigorous peer review process is contributing to the transparency of the carbon emission inventory process, an increasingly urgent requirement of any carbon management system.

While these observations and models have provided new insights into this system, they have also revealed measurement gaps and modeling limitations that must be addressed to develop a true global carbon monitoring system that can track changes in both natural and anthropogenic sources and sinks of CO<sub>2</sub> on policy relevant time and space scales. For example, space-based remote sensing observations of atmospheric CO<sub>2</sub> and land and ocean surface properties can expand the coverage and resolution of surface-based in situ measurements. However, passive remote sensing observations are largely precluded in persistently cloudy regions such as tropical rain forests, or mid- and high-latitude forests during the fall, winter and spring. These regions are often centers of action in the carbon cycle, but are also among the most challenging to observe systematically with surface-based in situ measurement systems. Remote sensing observations provide little insight into the carbon budget of the interior ocean, but here networks of autonomous in situ sensors are providing new tools for gathering data. Like remote sensing observations, their data typically has larger uncertainties and biases than conventional shipboard in situ measurements. Thus, a robust ocean carbon observing system will require continued shipboard observations for calibration and validation.

Top-down atmospheric CO<sub>2</sub> inventories are now complementing bottom-up statistical inventory methods by providing an integral constraint on total CO<sub>2</sub> emitted into the atmosphere and absorbed at the surface on scales ranging from large urban areas to nations. However, the inventory community has been slow to adopt top-down methods because they are much less source specific, and thus provide less direct, actionable information to policy makers than the bottom-up inventories. Optimal methods for combining top-down and bottom up emissions estimates are essential to both improve the accuracy and ensure the transparency urban to national scale inventories should be a high priority in any carbon management system. An effort to integrate top-down and bottom-up methods could also yield significant scientific benefits by fostering the development of more reliable models for diagnosing the current state of the carbon cycle, and for more accurately predicting the carbon cycle response to a changing climate.

The world's space agencies are actively working to coordinate ambitious plans for an expanded space-based remote sensing capability that supports atmospheric CO<sub>2</sub> measurements, high resolution maps of land surface type and biomass and ocean biological productivity. These efforts are being led by the Committee on Earth Observation Satellites (CEOS) and Coordination Group on Meteorological Satellites (CGMS) through their Joint Working Group on Climate (WGClimate) Greenhouse Gas Task team. In parallel, national agencies such as the U.S. NOAA and Japan's National Institute for Environmental Studies (NIES) and European organizations, such as ICOS and IAGOS, are working with international organizations including WMO GAW and the Global Climate Observing System and the Global Ocean Observing System (GCOS, GOOS) to coordinate the deployment of ground-based, ocean and airborne in situ sensors. These programs are receiving less attention and much less resources than the space-based measurement systems, but are equally critical to a global carbon monitoring system. The modeling systems needed to ingest and analyze the data collected by these expanding measurement systems are also advancing, but efforts to organize carbon cycle modeling efforts are also receiving less attention from the carbon cycle science community and their stakeholders. Ambitious efforts to track changes in the emission and uptake of CO<sub>2</sub> associated with human activities and the carbon cycle's response to climate change, such as the global stocktakes mandated by the UNFCCC Paris agreement, will require both expanded capabilities and much more coordination among all of these groups.

## 9 Open Research

This is a review of other published work. No new data has been created or archived specifically for this manuscript. Original data are available through the citations listed here. Figures have been redrawn to avoid copyright conflicts.

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## 11 References

- Ajani, J. I., Keith, H., Blakers, M., Mackey, B. G. and King, H. P. (2013). Comprehensive carbon stock and flow accounting: A national framework to support climate change mitigation policy. *Ecological Economics*, **89**, 61–72. doi:10.1016/j.ecolecon.2013.01.010
- Álvarez, M., Fajar, N. M., Carter, B. R., Guallart, E. F., Pérez, F. F., Woosley, R. J. and Murata, A. (2020). Global Ocean Spectrophotometric pH Assessment: Consistent Inconsistencies. *Environ. Sci. Technol.*, **54**, 10977–10988. doi:10.1021/acs.est.9b06932
- Anav, A., Friedlingstein, P., Beer, C., Ciais, P., Harper, A., Jones, C., Murray-Tortarolo, G., Papale, D., Parazoo, N. C., Peylin, P., Piao, S., Sitch, S., Viovy, N., Wiltshire, A. and Zhao, M. (2015). Spatiotemporal patterns of terrestrial gross primary production: A review. *Rev. Geophys.*, **53**, 785–818. doi:10.1002/2015RG000483
- Andrew, R. M. (2020). A comparison of estimates of global carbon dioxide emissions from fossil carbon sources. *Earth Syst. Sci. Data*, **12**, 1437–1465. doi:10.5194/essd-12-1437-2020
- Aumont, O., Orr, J.C., Monfray, P., Ludwig, W., Amiotte-Suchet, P. and Probst, J.-L. (2001). Riverine-driven interhemispheric transport of carbon. *Global Biogeochemical Cycles*, **15**, 393–405. doi:10.1029/1999GB001238
- Aumont, O., Ethé, C., Tagliabue, A., Bopp, L. and Gehlen, M. (2015). PISCES-v2: An ocean biogeochemical model for carbon and ecosystem studies. *Geosci. Model Dev.*, **8**, 2465–2513. doi:10.5194/gmd-8-2465-2015
- Ayers, J.M. and Strutton, P.G. (2013). Nutrient variability in Subantarctic Mode Waters forced by the Southern Annular Mode and ENSO: Subantarctic mode water nutrients. *Geophysical Research Letters*, **40**, 3419–3423. doi:10.1002/grl.50638
- Baccini, A., Walker, W., Carvalho, L., Farini, M., Sulla-Menashe, D. and Houghton, R. A. (2017). Tropical forests are a net carbon source based on aboveground measurements of gain and loss. *Science*, **358**, 230–234. doi:10.1126/science.aam5962
- Bakker, D. C. E., Pfeil, B., Smith, K., Hankin, S., Olsen, A., Alin, S. R., Cosca, C., Harasawa, S., Kozyr, A., Nojiri, Y., O'Brien, K. M., Schuster, U., Telszewski, M., Tilbrook, B., Wada, C., Akl, J., Barbero, L., Bates, N. R., Boutin, J., Bozec, Y., Cai, W.-J., Castle, R. D., Chavez, F. P., Chen, L., Chierici, M., Currie, K., de Baar, H. J. W., Evans, W., Feely, R. A., Fransson, A., Gao, Z., Hales, B., Hardman-Mountford, N. J., Hoppema, M., Huang, W.-J., Hunt, C. W., Huss, B., Ichikawa, T., Johannessen, T., Jones, E. M., Jones, S. D., Jutterström, S., Kitidis, V., Körtzinger, A., Landschützer, P., Lauvset, S. K., Lefèvre, N., Manke, A. B., Mathis, J. T., Merlivat, L., Metzl, N., Murata, A., Newberger, T., Omar, A. M., Ono, T., Park, G.-H., Paterson, K., Pierrot, D., Ríos, A. F., Sabine, C. L., Saito, S.,



- 1454 Salisbury, J., Sarma, V. V. S. S., Schlitzer, R., Sieger, R., Skjelvan, I., Steinhoff, T.,  
 1455 Sullivan, K. F., Sun, H., Sutton, A. J., Suzuki, T., Sweeney, C., Takahashi, T., Tjiputra,  
 1456 J., Tsurushima, N., van Heuven, S. M. A. C., Vandemark, D., Vlahos, P., Wallace, D. W.  
 1457 R., Wanninkhof, R. and Watson, A. J. (2014). An update to the Surface Ocean CO<sub>2</sub> Atlas  
 1458 (SOCAT version 2). *Earth Syst. Sci. Data*, **6**, 69-90. doi:10.5194/essd-6-69-2014
- 1459 Bakker, D. C. E., Pfeil, B., Landa, C. S., Metzl, N., O'Brien, K. M., Olsen, A., Smith, K., Cosca,  
 1460 C., Harasawa, S., Jones, S. D., Nakaoka, S. I., Nojiri, Y., Schuster, U., Steinhoff, T.,  
 1461 Sweeney, C., Takahashi, T., Tilbrook, B., Wada, C., Wanninkhof, R., Alin, S. R.,  
 1462 Balestrini, C. F., Barbero, L., Bates, N. R., Bianchi, A. A., Bonou, F., Boutin, J., Bozec,  
 1463 Y., Burger, E. F., Cai, W. J., Castle, R. D., Chen, L., Chierici, M., Currie, K., Evans, W.,  
 1464 Featherstone, C., Feely, R. A., Fransson, A., Goyet, C., Greenwood, N., Gregor, L.,  
 1465 Hankin, S., Hardman-Mountford, N. J., Harlay, J., Hauck, J., Hoppema, M., Humphreys,  
 1466 M. P., Hunt, C. W., Huss, B., Ibanhez, J. S. P., Johannessen, T., Keeling, R., Kitidis, V.,  
 1467 Kortzinger, A., Kozyr, A., Krasakopoulou, E., Kuwata, A., Landschutzer, P., Lauvset, S.  
 1468 K., Lefevre, N., Lo Monaco, C., Manke, A., Mathis, J. T., Merlivat, L., Millero, F. J.,  
 1469 Monteiro, P. M. S., Munro, D. R., Murata, A., Newberger, T., Omar, A. M., Ono, T.,  
 1470 Paterson, K., Pearce, D., Pierrot, D., Robbins, L. L., Saito, S., Salisbury, J., Schlitzer, R.,  
 1471 Schneider, B., Schweitzer, R., Sieger, R., Skjelvan, I., Sullivan, K. F., Sutherland, S. C.,  
 1472 Sutton, A. J., Tadokoro, K., Telszewski, M., Tuma, M., van Heuven, S. M. A. C.,  
 1473 Vandemark, D., Ward, B., Watson, A. J. and Xu, S. (2016). A multi-decade record of  
 1474 high-quality fCO<sub>2</sub> data in version 3 of the Surface Ocean CO<sub>2</sub> Atlas (SOCAT). *Earth*  
 1475 *Syst. Sci. Data*, **8**, 383-413. doi:10.5194/essd-8-383-2016
- 1476 Bakker, D. C. E., Alin, S. R., Bates, N., Becker, M., Castaño-Primo, R., Cosca, C. E., Cronin,  
 1477 M., Kadono, K., Kozyr, A., Lauvset, S. K., Metzl, N., Munro, D. R., Nakaoka, S.,  
 1478 O'Brien, K. M., Ólafsson, J., Olsen, A., Pfeil, B., Pierrot, D., Smith, K., Sutton, A. J.,  
 1479 Takahashi, T., Tilbrook, B., Wanninkhof, R., Andersson, A., Atamanchuk, D., Benoit-  
 1480 Cattin, A., Bott, R., Burger, E. F., Cai, W.-J., Cantoni, C., Collins, A., Corredor, J. E.,  
 1481 Cronin, M. F., Cross, J. N., Currie, K. I., De Carlo, E. H., DeGrandpre, M. D., Dietrich,  
 1482 C., Emerson, S., Enright, M. P., Evans, W., Feely, R. A., García-Ibáñez, M. I., Gkritzalis,  
 1483 T., Glockzin, M., Hales, B., Hartman, S. E., Hashida, G., Herndon, J., Howden, S. D.,  
 1484 Humphreys, M. P., Hunt, C. W., Jones, S. D., Kim, S., Kitidis, V., Landa, C. S.,  
 1485 Landschützer, P., Lebon, G. T., Lefèvre, N., Lo Monaco, C., Luchetta, A., Maenner  
 1486 Jones, S., Manke, A. B., Manzello, D., Mears, P., Mickett, J., Monacci, N. M., Morell, J.  
 1487 M., Musielewicz, S., Newberger, T., Newton, J., Noakes, S., Noh, J.-H., Nojiri, Y.,  
 1488 Ohman, M., Ólafsdóttir, S., Omar, A. M., Ono, T., Osborne, J., Plueddemann, A. J.,  
 1489 Rehder, G., Sabine, C. L., Salisbury, J. E., Schlitzer, R., Send, U., Skjelvan, I.,  
 1490 Sparnocchia, S., Steinhoff, T., Sullivan, K. F., Sutherland, S. C., Sweeney, C., Tadokoro,  
 1491 K., Tanhua, T., Telszewski, M., Tomlinson, M., Tribollet, A., Trull, T., Vandemark, D.,  
 1492 Wada, C., Wallace, D. W. R., Weller, R. A., and Woosley, R. J. (2020). Surface Ocean  
 1493 CO<sub>2</sub> Atlas Database Version 2020 (SOCATv2020) (NCEI Accession 0210711), NOAA  
 1494 National Centers for Environmental Information. doi:10.25921/4xkx-ss49
- 1495 Baldocchi, D., Falge, E., Gu, L., Olson, R., Hollinger, D., Running, S. W., Anthoni, P.,  
 1496 Bernhofer, C., Davis, K. J., Evans, R., Fuentes, J., Goldstein, A., Katul, G., Law, B., Lee,  
 1497 X., Malhi, Y., Meyers, T., Munger, W., Oechel, W., Paw, U. K. T., Pilegaard, K.,  
 1498 Schmid, H. P., Valentini, R., Verma, S., Vesala, T., Wilson, K. and Wofsy, S. (2001).

- 1499 FLUXNET: A new tool to study the temporal and spatial variability of ecosystem-scale  
1500 carbon dioxide, water vapor, and energy flux densities. *Bull. Amer. Meteor. Soc.*, **82**,  
1501 2415–2434. doi:10.1175/1520-0477
- 1502 Baldocchi D. D. (2003). Assessing the eddy covariance technique for evaluating carbon dioxide  
1503 exchange rates of ecosystems: past, present and future. *Global Change Biology*, **9**, 479–  
1504 492. doi:10.1046/j.1365-2486.2003.00629.x
- 1505 Ballantyne, A. P., Alden, C. B., Miller, J. B., Tans, P. P. and White, J. W. C. (2012). Increase in  
1506 observed net carbon dioxide uptake by land and oceans during the past 50 years. *Nature*,  
1507 **488**, 70–72. doi:10.1038/nature11299
- 1508 Beck, S. A. and Goetz, S. J. (2011). Satellite observations of high northern latitude vegetation  
1509 productivity changes between 1982 and 2008: ecological variability and regional  
1510 differences. *Environ. Res. Lett.*, **6**, 045501. doi:10.1088/1748-9326/6/4/045501
- 1511 Beer, C., Reichstein, M., Tomelleri, E., Ciais, P., Jung, M., Carvalhais, N., Rödenbeck, C.,  
1512 Arain, M. A., Baldocchi, D., Bonan, G., B., Bondeau, A., Cescatti, A., Lasslop, G.,  
1513 Lindroth, A., Lomas, M., Luyssaert, S., Margolis, H., Oleson, K. W., Rouspard, O.,  
1514 Veenendaal, E., Viovy, N., Williams, C., Woodward, F., I. and Papale, D. (2010).  
1515 Terrestrial gross carbon dioxide uptake: global distribution and covariation with climate.  
1516 *Science*, **329**, 834–838. doi:10.1126/science.1184984
- 1517 Bennedsen, M., Hildebrand, E. and Koopman, S. (2019). Trend analysis of the airborne fraction  
1518 and sink rate of anthropogenically released CO<sub>2</sub>. *Biogeosciences*, **16**, 3651–3663.  
1519 doi:10.5194/bg-16-3651-2019
- 1520 Boden T. A., Marland G. and Andres R. J. (2017). Global, Regional, and National Fossil-Fuel  
1521 CO<sub>2</sub> Emissions. Carbon Dioxide Information Analysis Center, Oak Ridge National  
1522 Laboratory, U.S. Department of Energy, Oak Ridge, Tenn., U.S.A.  
1523 doi:10.3334/CDIAC/00001\_V2017
- 1524 Bogan, R. A. J., Ohde, S., Arakaki, T., Mori, I. and McLeod, C. W. (2009). Changes in rainwater  
1525 pH associated with atmospheric carbon dioxide after the Industrial Revolution. *Water*,  
1526 *Air, and Soil Pollution*, **196**, 263–271. doi:10.1007/s11270-008-9774-0
- 1527 Bolin, B., Björkström, A., Holmén, K. and Moore, B. (1983). The simultaneous use of tracers for  
1528 ocean circulation studies. *Tellus B*, **35B**, 206–236. doi:10.1111/j.1600-  
1529 0889.1983.tb00025.x
- 1530 BP Statistical Review of World Energy 2020. available at  
1531 [https://www.bp.com/en/global/corporate/energy-economics/statistical-review-of-world-](https://www.bp.com/en/global/corporate/energy-economics/statistical-review-of-world-energy/co2-emissions.html)  
1532 [energy/co2-emissions.html](https://www.bp.com/en/global/corporate/energy-economics/statistical-review-of-world-energy/co2-emissions.html), last access 1 February 2021.
- 1533 Brien, R. J.W., Phillips, O. L., Feldpausch, T. R., Gloor, E., Baker, T. R., Lloyd, J., Lopez-  
1534 Gonzalez, G., Montagudo-Mendoza, A., Malhi, Y., Lewis, S. L., Vásquez Martinez, R.,  
1535 Alexiades, M., Álvarez Dávila, E., Alvarez-Loayza, P., Andrade, A., Aragão, L. E. O. C.,  
1536 Araujo-Murakami, A., Arets, E. J. M. M., Arroyo, L., Aymard, C. A. G., Bánki, O.S.,  
1537 Baraloto, C., Barroso, J., Bonal, D., Boot, R. G. A. , Camargo, J. L. C., Castilho, C. V.,  
1538 Chama, V., Chao, K. J., Chave, J., Comiskey, J. A., Cornejo Valverde, F., da Costa, L.,  
1539 de Oliveira, E. A., Di Fiore, A., Erwin, T. L., Fauset, S., Forsthofer, M., Galbraith, D. R.,  
1540 Grahame, E. S., Groot, N., Hérault, B., Higuchi, N., Honorio Coronado, E. N., Keeling,

- 1541 H., Killeen, T. J., Laurance, W. F., Laurance, S., Licona, J., Magnussen, W. E., Marimon,  
1542 B. S., Marimon-Junior, B. H., Mendoza, C., Neill, D. A., Nogueira, E. M., Núñez, P.,  
1543 Pallqui Camacho, N. C., Parada, A., Pardo-Molina, G., Peacock, J., Peña-Claros, M.,  
1544 Pickavance, G. C., Pitman, N. C. A., Poorter, L., Prieto, A., Quesada C. A., Ramírez, F.,  
1545 Ramírez-Angulo, H., Restrepo, Z., Roopsind, A., Rudas, A., Salomão, R. P., Schwarz,  
1546 M., Silva, N., Silva-Espejo, J. E., Silveira, M., Stropp, J., Talbot, J., ter Steege, H., Teran-  
1547 Aguilar, J., Terborgh, J., Thomas-Caesar, R., Toledo, M., Torello-Raventos, M., Umetsu,  
1548 R. K., van der Heijden, G. M. F., van der Hout, P., Guimarães, Vieira I. C., Vieira, S. A.,  
1549 Vilanova, E., Vos, V. A. and Zagt, R. J. (2015). Long-term decline of the Amazon carbon  
1550 sink. *Nature*, **519**(7543), 344-348. doi:10.1038/nature14283
- 1551 Bronselaer, B., and Zanna, L. (2020), Heat and carbon coupling reveals ocean warming due to  
1552 circulation changes. *Nature*, **584**, 227–233. doi:10.1038/s41586-020-2573-5
- 1553 Bushinsky, S. M., Landschützer, P., Rödenbeck, C., Gray, A. R., Baker, D., Mazloff, M. R.,  
1554 Resplandy, L., Johnson, K. S., and Sarmiento, J. L. (2019). Reassessing Southern Ocean  
1555 air-sea CO<sub>2</sub> flux estimates with the addition of biogeochemical float observations. *Global*  
1556 *Biogeochemical Cycles*, **33**(11), 1370-1388, doi:10.1029/2019GB006176
- 1557 Byrne, B., Wunch, D., Jones, D. B. A., Strong, K., Deng, F., Baker, I., Köhler, P., Frankenberg,  
1558 C., Joiner, J., Arora, V. K., Badawy, B., Harper, A. B., Warneke, T., Petri, C., Kivi, R.  
1559 and Roehl, C. M. (2018). Evaluating GPP and respiration estimates over northern  
1560 midlatitude ecosystems using solar-induced fluorescence and atmospheric CO<sub>2</sub>  
1561 measurements. *Journal of Geophysical Research: Biogeosciences*, **123**(9), 2976–2997.  
1562 doi:10.1029/2018JG004472
- 1563 Byrne, B., Liu, J., Lee, M., Baker, I., Bowman, K. W., Deutscher, N. M., Feist, D. G., Griffith D.  
1564 W. T., Iraci, L. T., Kiel, M., Kimball, J. S., Miller, C. E., Morino, I., Parazoo, N. C.,  
1565 Petri, C., Roehl, C. M., Sha, M. K., Strong, K., Velazco, V. A., Wennberg, P. O. and  
1566 Wunch, D. (2020). Improved constraints on northern extratropical CO<sub>2</sub> fluxes obtained  
1567 by combining surface-based and space-based atmospheric CO<sub>2</sub> measurements. *Journal of*  
1568 *Geophysical Research: Atmospheres*, **125**, e2019JD032029. doi:10.1029/2019JD032029
- 1569 Canadell, J. G., Le Quéré, C., Raupach, M. R., Field, C. B., Buitenhuis, E., Ciais, P., Conway,  
1570 T. J., Gilett, N. P., Houghton, J. T. and Marland, G. (2007). Contributions to accelerating  
1571 atmospheric CO<sub>2</sub> growth from economic activity, carbon intensity, and efficiency of  
1572 natural sinks. *Proceedings of the National Academy of Sciences*, **104**, 18,866 – 18,870.  
1573 doi:10.1073/pnas.0702737104
- 1574 Carroll, D., Menemenlis, D., Adkins, J. F., Bowman, K. W., Brix, H., Dutkiewicz, S., Fenty, I.,  
1575 Gierach, M. M., Hill, C., Jahn, O., Landschützer, P., Lauderdale, J. M., Liu, J., Manizza,  
1576 M., Naviaux, J. D., Rodenbeck, C., Schimel, D. S., Van der Stocken, T. and Zhang, H.  
1577 (2020). The ECCO-Darwin data-assimilative global ocean biogeochemistry model:  
1578 Estimates of seasonal to multi-decadal surface ocean pCO<sub>2</sub> and air-sea CO<sub>2</sub> flux. *J. Adv.*  
1579 *Model. Earth Syst.*, **12**, e2019MS001888. doi:10.1029/2019MS001888
- 1580 Carvalhais, N., Forkel, M., Khomik, M., Bellarby, J., Jung, M., Migliavacca, M. Mu, M.,  
1581 Saatchi, S., Santoro, M., Thurner, M., Weber, U., Ahrens, B., Beer, C., Cescatti, A.,  
1582 Randerson, J. T. and Reichstein, M. (2014). Global covariation of carbon turnover times  
1583 with climate in terrestrial ecosystems. *Nature*, **514**(7521) 213-7. doi:10.1038/nature13731

- 1584 Castro, A. O., Chen, J., Zang, C. S., Shekhar, A., Jimenez, J. C., Bhattacharjee, S., Kindu, M.,  
1585 Morales, V. H. and Ramming, A. (2020). OCO-2 Solar-Induced Chlorophyll  
1586 Fluorescence Variability across Ecoregions of the Amazon Basin and the Extreme  
1587 Drought Effects of El Nino (2015-2016), *Remote Sensing*, **12**(7), 1202.  
1588 doi:10.3390/rs12071202
- 1589 Chatterjee, A., Gierach, M. M., Sutton, A. J., Feely, R. A., Crisp, D., Eldering, A., Gunson, M.  
1590 R., O'Dell, C. W., Stephens, B. B. and Schimel, D. S. (2017). Influence of El Niño on  
1591 atmospheric CO<sub>2</sub> over the tropical Pacific Ocean: Findings from NASA's OCO-2  
1592 mission. *Science*, **358**, eaam5776. doi:10.1126/science.aam5776
- 1593 Chevallier, F., Ciais, P., Conway, T. J., Aalto, T., Anderson, B. E., Bousquet, P., Brunke, E. G.,  
1594 Ciattaglia, L., Esaki, Y., Frohlich, M., Gomez, A. J., Gomez-Pelaez, A. J., Haszpra, L.,  
1595 Krummel, P., Langenfelds, R., Leuenberger, M., Machida, T., Maignan, F., Matsueda,  
1596 H., Morgui, J. A., Mukai, H., Nakazawa, T., Peylin, P., Ramonet, M., Rivier, L., Sawa,  
1597 Y., Schmidt, M., Steele, P., Vay, S. A., Vermeulen, A. T., Wofsy, S. and Worthy, D.  
1598 (2010). CO<sub>2</sub> surface fluxes at grid point scale estimated from a global 21-year reanalysis  
1599 of atmospheric measurements. *Journal of Geophysical Research: Atmospheres*, **115**,  
1600 D21307. doi:10.1029/2010JD013887
- 1601 Chevallier, F., Remaud, M. O'Dell, C. W., Baker, D., Peylin, P. and Cozic, A. (2019). Objective  
1602 evaluation of surface and satellite driven CO<sub>2</sub> atmospheric inversion. *Atmospheric*  
1603 *Chemistry and Physics*, **19**, 14233–14251. doi:10.5194/acp-19-14233-2019
- 1604 Ciais, P., Dolman, A. J., Bombelli, A., Duren, R., Peregon, A., Rayner, P. J., Miller, C., Gobron,  
1605 N., Kinderman, G., Marland, G., Gruber, N., Chevallier, F., Andres, R. J., Balsamo, G.,  
1606 Bopp, L., Bréon, F.-M., Broquet, G., Dargaville, R., Battin, T. J., Borges, A.,  
1607 Bovensmann, H., Buchwitz, M., Butler, J., Canadell, J. G., Cook, R. B., DeFries, R.,  
1608 Engelen, R., Gurney, K. R., Heinze, C., Heimann, M., Held, A., Henry, M., Law, B.,  
1609 Luyssaert, S., Miller, J., Moriyama, T., Moulin, C., Myneni, R. B., Nussli, C.,  
1610 Obersteiner, M., Ojima, D., Pan, Y., Paris, J.-D., Piao, S. L., Poulter, B., Plummer, S.,  
1611 Quegan, S., Raymond, P., Reichstein, M., Rivier, L., Sabine, C., Schimel, D., Tarasova,  
1612 O., Valentini, R., Wang, R., van der Werf, G., Wickland, D., Williams, M. and Zehner,  
1613 C. (2014). Current systematic carbon-cycle observations and the need for implementing a  
1614 policy-relevant carbon observing system, *Biogeosciences*, **11**, 3547–3602.  
1615 doi:10.5194/bg-11-3547-2014
- 1616 Ciais, P., Yao, Y., Gasser, T., Baccini, A., Wang, Y., Lauerwald, R., Peng, S., Bastos, A., Li, W.,  
1617 Raymond, P. A., Canadell, J. G., Peters, G. P., Andres, R. J., Chang, J., Yue, C., Dolman,  
1618 A. J., Haverd, V., Hartman, J., Laruelle, G., Konings, A. G., King, A. W., Liu, Y.,  
1619 Luyssaert, S., Maignan, F., Patra, P. K., Peregon, A., Regnier, P., Pongratz, J., Poulter,  
1620 B., Shvidenko, A., Valentini, R., Wang, R., Broquet, G., Yin, Y., Zscheischler, J.,  
1621 Guenet, B., Goll, D., S., Ballantyne, A.-P., Yang, H., Qiu, C. and Zhu, D. (2020).  
1622 Empirical estimates of regional carbon budgets imply reduced global soil heterotrophic  
1623 respiration. *National Science Review*, **0**, nwaa145,1-14. doi:10.1093/nsr/nwaa145
- 1624 Clement, D., and Gruber, N. (2018). The eMLR(C\*) Method to determine decadal changes in the  
1625 global ocean storage of anthropogenic CO<sub>2</sub>. *Global Biogeochemical Cycles*, **32**, 654-679.  
1626 doi:10.1002/2017gb005819

- 1627 Claustre, H., Johnson, K. S., and Takeshita, Y. (2020). Observing the global ocean with  
1628 Biogeochemical-Argo, *Annual Review of Marine Science*, **12**, 23-48.  
1629 doi:10.1146/annurev-marine-010419-010956
- 1630 Crowell, S., Baker, D., Schuh, A., Basu, S., Jacobson, A., Chevallier, F., Liu, J., Deng, F., Feng,  
1631 L., McKain, K., Chatterjee, A., Miller, J., Stephens, B., Eldering, A., Crisp, D., Schimel,  
1632 D., Nassar, R., O'Dell, C., Oda, T., Sweeney, C., Palmer, P., and Jones, D. (2019). The  
1633 2015-2016 carbon cycle as seen from OCO-2 and the global in situ network. *Atmospheric*  
1634 *Chemistry and Physics*, **19**, 7347–7376. doi:10.5194/acp-19-7347-2019
- 1635 Denman, K.L., G. Brasseur, A. Chidthaisong, P. Ciais, P.M. Cox, R.E. Dickinson, D.  
1636 Hauglustaine, C. Heinze, E. Holland, D. Jacob, U. Lohmann, S Ramachandran, P.L. da  
1637 Silva Dias, S.C. Wofsy and X. Zhang, (2007). Couplings between changes in the climate  
1638 system and biogeochemistry. In: Climate Change 2007: The Physical Science Basis.  
1639 Contribution of Working Group I to the Fourth Assessment Report of the  
1640 Intergovernmental Panel on Climate Change [Solomon, S., Qin, D., Manning, M., Chen,  
1641 Z., Marquis, M., Avery, K. B., Tignor, M. and Miller, H. L. (eds.)]. Cambridge  
1642 University Press, Cambridge, United Kingdom and New York, NY, USA.
- 1643 Denvil-Sommer, A., M. Gehlen, M. Vrac, and C. Mejia (2019). LSCE-FFNN-v1: A two-step  
1644 neural network model for the reconstruction of surface ocean pCO<sub>2</sub> over the global ocean.  
1645 *Geosci. Model Dev.*, **12**(5), 2091–2105. doi:10.5194/gmd-12-2091-2019
- 1646 DeVries, T. (2014). The oceanic anthropogenic CO<sub>2</sub> sink: Storage, air-sea fluxes, and transports  
1647 over the industrial era, *Global Biogeochemical Cycles*, **28**(7), 631–647.  
1648 doi:10.1002/2013GB004739
- 1649 DeVries, T., Holzer, M. and Primeau, F. (2017). Recent increase in oceanic carbon uptake driven  
1650 by weaker upper-ocean overturning. *Nature*, **542**, 215–218. doi:10.1038/nature21068
- 1651 DeVries, T., Le Quéré, C., Andrews, O., Berthet, S., Hauck, J., Ilyina, T., Landschützer, P.,  
1652 Lenton, A., Lima, I. D., Nowicki, M., Schwinger, J., Séférian, R. (2019). Decadal trends  
1653 in the ocean carbon sink. *Proceedings of the National Academy of Sciences*, 201900371.  
1654 doi:10.1073/pnas.1900371116
- 1655 Dlugokencky, E. J., Hall, B.D., Montzka, S.A., Dutton, G., Muhle, J. and Elkins, J.W. (2018).  
1656 Long-lived greenhouse gases [in "State of the Climate in 2017"]. *Bulletin of the American*  
1657 *Meteorological Society*, **99**(8), S46-S48. doi:10.1175/2018BAMSStateoftheClimate.1
- 1658 Doney S.C., Lindsay, K., Caldeira, K., Campin, J.-M., Drange, H., Dutay, J. C., Follows, M.,  
1659 Gao, Y., Gnanadesikan, A., Gruber, N., Ishida, A., Joos, F., Madec, G., Maier-Reimer,  
1660 E., Marshall, J. C., Matear, R. J., Monfray, P., Mouchet, A., Najjar, R., Orr, J. C.,  
1661 Plattner, G.-K., Sarmiento, J., Schlitzer, R., Slater, R., Totterdell, I. J., Weirig, M. F.,  
1662 Yamanaka, Y. and Yool, A. (2004). Evaluating global ocean carbon models: The  
1663 importance of realistic physics. *Global Biogeochemical Cycles*, **18** (3), GB3017.  
1664 doi:10.1029/2003GB002150
- 1665 Doughty, C. E. and Goulden, M. L. (2008). Are tropical forests near a high temperature  
1666 threshold? *Journal of Geophysical Research: Biogeosciences*, **113**, G00B07.  
1667 doi:10.1029/2007JG000632

- 1668 Duffy, K., A., Schwalm, C. R., Arcus, V. L., Koch, G. W., Liang, L. L. and Schipper, L. A.  
1669 (2021). How close are we to the temperature tipping point of the terrestrial biosphere?  
1670 *Science Advances*, **7**, eaay1052.
- 1671 Duncanson, L., Armston, J., Disney, M., Avitabile, V., Barbier, N., Calders, K., Carter, S.,  
1672 Chave, J., Herold, M., Crowther, T. W., Falkowski, M., Kellner, J. R., Labrière, N.,  
1673 Lucas, R., MacBean, N., McRoberts, R. E., Meyer, V., Naesset, E., Nickeson, J. E., Paul,  
1674 K. I., Phillips, O. L., Réjou-Méchain, M., Román, M., Roxburgh, S., Saatchi, S.,  
1675 Schepaschenko, D., Scipal, K., Siqueira, P. R., Whitehurst, A. and Williams, M.  
1676 (2019). The importance of consistent global forest aboveground biomass product  
1677 Validation. *Surveys in Geophysics volume*, **40**, 979–999. doi:10.1007/s10712-019-09538-  
1678 8
- 1679 Eddebbar, Y. A., Rodgers, K. B., Long, M. C., Subramanian, A. C., Xie S.-P. and Keeling, R. F.  
1680 (2019). El Niño-like physical and biogeochemical ocean response to tropical eruptions.  
1681 *Journal of Climate*, **32**(9), JCLI-D-18-0458.1, doi:10.1175/JCLI-D-18-0458.1
- 1682 Eldering, A., Wennberg, P. O., Crisp, D., Schimel, D., Gunson, M. R., Chatterjee, A., Liu J.,  
1683 Schwandner, Y. Sun, C.W. O'Dell, C. Frankenberg, T. Taylor, B. Fisher, G.B. Osterman,  
1684 D. Wunch, F., Hakkarainen, J. and Tamminen, J. (2017). The Orbiting Carbon  
1685 Observatory-2 early science investigations of regional carbon dioxide fluxes. *Science* ,  
1686 **358**, eaam5745. doi:10.1126/science.aam5745
- 1687 Enting, I.G., Trudinger, C.M. and Francey, R.J. (1995). A synthesis inversion of  
1688 the concentration and  $\delta^{13}\text{C}$  atmospheric  $\text{CO}_2$ . *Tellus*, **47B**, 35-52. doi:10.1034/j.1600-  
1689 0889.47.issue1.5.x
- 1690 Fan, L., Wigneron, J.-P., Ciais, P., Chave, J., Brandt, M., Fensholt, R., Saatchi, S. S. Bastos, A.,  
1691 Al-Yaari, A., Hufkens, K., Qin, Y., Xiao, X., Chen, C., Myneni, R. B., Rernandez-  
1692 Moran, R., Mialon, A., Rodriguez-Fernandez, N. J., Kerr, Y., Tian, F. and Peñuelas, J.  
1693 (2019). Satellite-observed pantropical carbon dynamics. *Nature Plants*, **5**, 944–951.  
1694 doi:10.1038/s41477-019-0478-9
- 1695 Fay, A. R., McKinley, G. A. (2013). Global trends in surface ocean  $\text{pCO}_2$  from in situ data.  
1696 *Global Biogeochemical Cycles*, **27**, 541–557. doi:10.1002/gbc.20051
- 1697 Fay, A.R., Gregor, L., Landschützer, P. McKinley, G.A., Gruber, N., Gehlen, M., Iida, Y.,  
1698 Laruelle, G. G., Rödenbeck, C. and Zeng J. (2021). Harmonization of global surface  
1699 ocean  $\text{pCO}_2$  mapped products and their flux calculations; an improved estimate of the  
1700 ocean carbon sink. *Earth Syst. Sci. Data*, submitted 2020.
- 1701 Fay, A. R., Lovenduski, N. S., McKinley, G. A., Munro, D. R., Sweeney, C., Gray, A. R.,  
1702 Landschützer, P., Stephens, B. B., Takahashi, T. and Williams, N. (2018). Utilizing the  
1703 Drake Passage Time-series to understand variability and change in subpolar Southern  
1704 Ocean  $\text{pCO}_2$ . *Biogeosciences*, **15**(12), 3841–3855. doi:10.5194/bg-15-3841-2018
- 1705 Fisher, J. B., Huntzinger, D. N., Schwalm, C. R. and Sitch, S. (2014). Modeling the terrestrial  
1706 biosphere. *Annual Review of Environment and Resources*, **39**, 91–123.  
1707 doi:10.1146/annurev-environ-012913-093456

- 1708 Fong, M. B. and Dickson, A. G. (2019). Insights from GO-SHIP hydrography data into the  
1709 thermodynamic consistency of CO<sub>2</sub> system measurements in seawater. *Marine*  
1710 *Chemistry*, **211**, 52-63. doi:10.1016/j.marchem.2019.03.006
- 1711 Frankenberg, C., O'Dell, C., Berry, J., Guanter, L., Joiner, J., Köhler, P., Pollock, R. and Taylor,  
1712 T. E. (2014). Prospects for chlorophyll fluorescence remote sensing from the orbiting  
1713 carbon observatory-2. *Remote Sensing of Environment*, **147**, 1–12.  
1714 doi:10.1016/j.rse.2014.02.007. ISSN 0034–4257
- 1715 Frey, M., Sha, M. K., Hase, F., Kiel, M., Blumenstock, T., Harig, R., Surawicz, G., Deutscher,  
1716 N. M., Shiomi, K., Franklin, J. E., Bösch, H., Chen, J., Grutter, M., Ohyama, H., Sun, Y.,  
1717 Butz, A., Mengistu Tsidu, G., Ene, D., Wunch, D., Cao, Z., Garcia, O., Ramonet, M.,  
1718 Vogel, F. and Orphal, J. (2019). Building the COllaborative Carbon Column Observing  
1719 Network (COCCON): long-term stability and ensemble performance of the EM27/SUN  
1720 Fourier transform spectrometer. *Atmospheric Measurement Technologies*, **12**, 1513–  
1721 1530. doi:10.5194/amt-12-1513-2019
- 1722 Friedlingstein, P., Jones, M. W., O'Sullivan, M., Andrew, R. M., Hauck, J., Peters, G. P., Peters,  
1723 W., Pongratz, J., Sitch, S., Le Quéré, C., Bakker, D. C. E., Canadell, J. G., Ciais, P.,  
1724 Jackson, R. B., Anthoni, P., Barbero, L., Bastos, A., Bastrikov, V., Becker, M., Bopp, L.,  
1725 Buitenhuis, E., Chandra, N., Chevallier, F., Chini, L. P., Currie, K. I., Feely, R. A.,  
1726 Gehlen, M., Gilfillan, D., Gkritzalis, T., Goll, D. S., Gruber, N., Gutekunst, S., Harris, I.,  
1727 Haverd, V., Houghton, R. A., Hurtt, G., Ilyina, T., Jain, A. K., Joetzjer, E., Kaplan, J. O.,  
1728 Kato, E., Klein Goldewijk, K., Korsbakken, J. I., Landschützer, P., Lauvset, S. K.,  
1729 Lefèvre, N., Lenton, A., Lienert, S., Lombardozzi, D., Marland, G., McGuire, P. C.,  
1730 Melton, J. R., Metzl, N., Munro, D. R., Nabel, J. E. M. S., Nakaoka, S.-I., Neill, C.,  
1731 Omar, A. M., Ono, T., Peregón, A., Pierrot, D., Poulter, B., Rehder, G., Resplandy, L.,  
1732 Robertson, E., Rödenbeck, C., Séférian, R., Schwinger, J., Smith, N., Tans, P. P., Tian,  
1733 H., Tilbrook, B., Tubiello, F. N., van der Werf, G. R., Wiltshire, A. J. and Zaehle, S.  
1734 (2019). Global Carbon Budget 2019. *Earth Syst. Sci. Data*, **11**, 1783–1838.  
1735 doi:10.5194/essd-11-1783-2019
- 1736 Friedlingstein, P., O'Sullivan, M., Jones, M. W., Andrew, R. M., Hauck, J., Olsen, A., Peters, G.  
1737 P., Peters, W., Pongratz, J., Sitch, S., Le Quéré, C., Canadell, J. G., Ciais, P., Jackson, R.  
1738 B., Alin, S., Aragão, L. E. O. C., Arneeth, A., Arora, V., Bates, N. R., Becker, M., Benoit-  
1739 Cattin, A., Bittig, H. C., Bopp, L., Bultan, S., Chandra, N., Chevallier, F., Chini, L. P.,  
1740 Evans, W., Florentie, L., Forster, P. M., Gasser, T., Gehlen, M., Gilfillan, D., Gkritzalis,  
1741 T., Gregor, L., Gruber, N., Harris, I., Hartung, K., Haverd, V., Houghton, R. A., Ilyina,  
1742 T., Jain, A. K., Joetzjer, E., Kadono, K., Kato, E., Kitidis, V., Korsbakken, J. I.,  
1743 Landschützer, P., Lefèvre, N., Lenton, A., Lienert, S., Liu, Z., Lombardozzi, D., Marland,  
1744 G., Metzl, N., Munro, D. R., Nabel, J. E. M. S., Nakaoka, S.-I., Niwa, Y., O'Brien, K.,  
1745 Ono, T., Palmer, P. I., Pierrot, D., Poulter, B., Resplandy, L., Robertson, E., Rödenbeck,  
1746 C., Schwinger, J., Séférian, R., Skjelvan, I., Smith, A. J. P., Sutton, A. J., Tanhua, T.,  
1747 Tans, P. P., Tian, H., Tilbrook, B., van der Werf, G., Vuichard, N., Walker, A. P.,  
1748 Wanninkhof, R., Watson, A. J., Willis, D., Wiltshire, A. J., Yuan, W., Yue, X. and  
1749 Zaehle, S. (2020). Global Carbon Budget 2020. *Earth Syst. Sci. Data*, **12**, 3269–3340.  
1750 doi:10.5194/essd-12-3269-2020



- 1751 Friis, K., Körtzinger, A., Pätsch, J. and Wallace, D. W. R. (2005). On the temporal increase of  
1752 anthropogenic CO<sub>2</sub> in the subpolar North Atlantic. *Deep-Sea Research.* **52**, 681-698.  
1753 doi:10.1016/j.dsr.2004.11.017
- 1754 Frölicher, T. L., Joos, F., Raible, C. C. and Sarmiento, J. L. (2013). Atmospheric CO<sub>2</sub> response  
1755 to volcanic eruptions: The role of ENSO, season, and variability. *Global Biogeochemical*  
1756 *Cycles*, **27**, 239-251. doi:10.1002/gbc.20028
- 1757 Frölicher, T.L., Rodgers, K. B., Stock, C. A., Cheung, W. W. L. (2016). Sources of uncertainties  
1758 in 21st century projections of potential ocean ecosystem stressors: Uncertainties in  
1759 stressor projections. *Global Biogeochemical Cycles*, **30**, 1224–1243.  
1760 doi:10.1002/2015GB005338
- 1761 Galbraith, E. D., and Skinner, L. C. (2020). The biological pump during the last glacial  
1762 maximum. *Annual Reviews of Marine Science* **12**, 559–586. doi:10.1146/annurev-marine-  
1763 010419-010906
- 1764 Gatti L.V., Gloor M., Miller J. B., Doughty C. E., Malhi Y., Domingues L. G., Basso L. S.,  
1765 Martinewski A., Correia C. S., Borges V. F., Freitas S., Braz R., Anderson L. O., Rocha  
1766 H., Grace J., Phillips O. L., and Lloyd J. (2014). Drought sensitivity of Amazonian  
1767 carbon balance revealed by atmospheric measurements. *Nature*, **506**, 76-80.  
1768 doi:10.1038/nature12957
- 1769 Gloege, L., McKinley, G. A., Landschützer, P., Fay, A., Frölicher, T., Fyfe, J. C., Illyina ,T.,  
1770 Jones, S.D., Lovenduski, N. S., Rödenbeck ,C., Rodgers K. B., Schlunegger, S. and  
1771 Takano, Y. (2020). Quantifying errors in observationally-based estimates of ocean carbon  
1772 sink variability. *Global Biogeochemical Cycles*, in review.  
1773 doi:10.1002/essoar.10502036.1
- 1774 Gonsamo, A., Chen, J. M., He, L., Sun, Y., Rogers, C. and Liu, J. (2019). Exploring SMAP and  
1775 OCO-2 observations to monitor soil moisture control on photosynthetic activity of global  
1776 drylands and croplands, *Remote Sensing of Environment*, **232**, 111314.  
1777 doi:10.1016/j.rse.2019.111314
- 1778 Goris, N., Tjiputra, J. F., Olsen, A., Schwinger, J., Lauvset S. K. and Jeansson E. (2018).  
1779 Constraining projection-based estimates of the future North Atlantic carbon uptake.  
1780 *Journal of Climate*, **31**(10), 3959–3978, doi:10.1175/JCLI-D-17-0564.1
- 1781 Grassi, G., House, J., Kurz, W. A., Cescatti, A., Houghton, R. A., Peters, G. P., Sanz, M. J.,  
1782 Viñas, R. A., Alkama, R., Arneeth, A., Bondeau, A., Dentener, F., Fader, M., Federici, S.,  
1783 Friedlingstein, P., Jain, A. K., Kato, E., Koven, C. D., Lee, D., Nabel, J. E. M. S.,  
1784 Nassikas, A. A., Perugini, L., Rossi, S., Sitch, S., Viovy, N., Wiltshire, A. and Zaehle, S.  
1785 (2018). Reconciling global-model estimates and country reporting of anthropogenic  
1786 forest CO<sub>2</sub> sinks. *Nature Climate Change*, **8**, 914–920. doi.org/10.1038/s41558-018-  
1787 0283-x
- 1788 Graven, H. D., Keeling, R. F., Piper, S. C., Patra, P. K., Stephens, B. B., Wofsy, S. C., Welp, L.  
1789 R., Sweeney, C. and Tans, P. P. (2013). Enhanced seasonal exchange of CO<sub>2</sub> by northern  
1790 ecosystems since 1960, *Science*, **341**(6150): 1085–1089. doi:10.1126/science.1239207
- 1791 Gregor, L., Lebehot, A. D. Kok, S. and Scheel Monteiro, P. M. (2019). A comparative  
1792 assessment of the uncertainties of global surface-ocean CO<sub>2</sub> estimates using a machine

- 1793 learning ensemble (CSIR-ML6 version 2019a); have we hit the wall? *Geosci. Model*  
1794 *Dev.*, **12**, 5113–5136. doi:10.5194/gmd-12-5113-2019
- 1795 Gruber, N., Clement, D., Carter, B.R., Feely, R.A., Van Heuven, S., Hoppema, M., Ishii, M.,  
1796 Key, R.M., Kozyr, A., Lauvset, S.K., Lo Monaco, C., Mathis, J.T., Murata, A., Olsen, A.,  
1797 Perez, F.F., Sabine, C.L., Tanhua, T., and Wanninkhof, R. (2019a). The oceanic sink for  
1798 anthropogenic CO<sub>2</sub> from 1994 to 2007. *Science* **363**, 1193–1199.  
1799 doi:10.1126/science.aau5153
- 1800 Gruber, N., Landschützer, P., and Lovendus, N. S. (2019b). The variable Southern Ocean  
1801 carbon sink. *Annual Reviews of Marine Science*, **11**, 159–186. doi:10.1146/annurev-  
1802 marine-121916-063407
- 1803 Guan, K., Berry, J. A., Zhang, Y., Joiner, J., Guanter, L., Badgley, G. and Lobell, D. B. (2016).  
1804 Improving the monitoring of crop productivity using spaceborne solar-induced  
1805 fluorescence. *Global Change Biology*, **22**(2), 716–726. doi:10.1111/gcb.13136
- 1806 Guan, K., Wu, J., Kimball, J. S., Anderson, M. C., Frohling, S., Li, B., Hain, C. R. and Lobell, D.  
1807 B. (2017). The shared and unique values of optical, fluorescence, thermal and microwave  
1808 satellite data for estimating large-scale crop yields. *Remote Sensing of Environment*, **199**,  
1809 333–349. doi:10.1016/j.rse.2017.06.043
- 1810 Gurney, K. R., Liang, J., O’Keefe, D., Patarasuk, R., Hutchins, M., Huang, J., Rao, P. and Song,  
1811 Y. (2019). Comparison of global downscaled versus bottom-up fossil fuel CO<sub>2</sub> emissions  
1812 at the urban scale in four U.S. urban areas. *Journal of Geophysical Research:*  
1813 *Atmospheres*, **124**(5), 2823–2840. doi:10.1029/2018JD028859
- 1814 Hakkarainen, J., Ialongo, I. and Tamminen, J. (2016). Direct space-based observations of  
1815 anthropogenic CO<sub>2</sub> emission areas from OCO-2. *Geophysical Research Letters*, **43**,  
1816 11,400–11,406. doi:10.1002/2016GL070885
- 1817 Hakkarainen, J., Ialongo, I., Maksyutov, S., and Crisp, D. (2019). Analysis of Four Years of  
1818 Global XCO<sub>2</sub> Anomalies as Seen by Orbiting Carbon Observatory-2, *Remote Sensing*, **11**,  
1819 850. doi:10.3390/rs11070850
- 1820 Hansis, E., Davis, S. J. and Pongratz, J. (2015). Relevance of methodological choices for  
1821 accounting of land use change carbon fluxes. *Global Biogeochemical Cycles*, **29**, 1230–  
1822 1246. doi:10.1002/2014GB004997
- 1823 Hauck, J., Völker, C., Wang, T., Hoppema, M., Losch, M. and Wolf-Gladrow, D.A. (2013).  
1824 Seasonally different carbon flux changes in the Southern Ocean in response to the  
1825 southern annular mode. *Global Biogeochemical Cycles*, **27**, 1236–1245.  
1826 doi:10.1002/2013GB004600
- 1827 Hauck, J., Völker, C., Wolf-Gladrow, D. A., Laufkötter, C., Vogt, M., Aumont, O., Bopp, L.,  
1828 Buitenhuis, E. T., Doney, S. C., Dunne, J., Gruber, N., Hashioka, T., John, J., Le Quéré,  
1829 C., Lima, I. D., Nakano, H., Séférian, R. and Totterdell, I. (2015). On the Southern Ocean  
1830 CO<sub>2</sub> uptake and the role of the biological carbon pump in the 21st century. *Global*  
1831 *Biogeochemical Cycles*, **29**, 1451–1470. doi:10.1002/2015GB005140
- 1832 Hauck, J., Zeising, M., Le Quéré, C., Gruber, N., Bakker, D. C. E., Bopp, L., Chau, T. T. T.,  
1833 Gürses, Ö., Ilyina, T., Landschützer, P., Lenton, A., Resplandy, L., Rödenbeck, C.,  
1834 Schwinger, J. and Séférian, R. (2020). Consistency and challenges in the ocean carbon

- 1835 sink estimate for the Global Carbon Budget. *Frontiers in Marine Science*, **7**, 571720.  
1836 doi:10.3389/fmars.2020.571720
- 1837 Hausfather, Z. and Peters, G. P. (2020). Emissions - the “business as usual” story is misleading.  
1838 *Nature*, **577**(7792), 618–620, doi:10.1038/d41586-020-00177-3
- 1839 He, L., Magney, T., Dutta, D., Yin, Y., Köhler, P., Grossmann, K., Stutz, J., Dold, C., Hatfield,  
1840 J., Guan, K., Peng, B. and Frankenberg, C. (2020). From the ground to space: Using  
1841 solar-induced chlorophyll fluorescence to estimate crop productivity. *Geophysical*  
1842 *Research Letters*, **47**, e2020GL087474. doi:10.1029/2020GL087474
- 1843 Hedelius, J. K., Liu, J., Oda, T., Maksyutov, S., Roehl, C. M., Iraci, L., Podolske, J., Hillyard, P.,  
1844 Liang, J., Gurney, K., Wunch, D. and Wennberg P. (2018). Southern California megacity  
1845 CO<sub>2</sub>, CH<sub>4</sub>, and CO flux estimates using ground- and space-based remote sensing and a  
1846 Lagrangian model, *Atmospheric Chemistry and Physics*, **18**, 16271–16291.  
1847 doi:10.5194/acp-18-16271-2018
- 1848 Henson, S. A., Beaulieu, C. and Lampitt, R. (2016). Observing climate change trends in ocean  
1849 biogeochemistry: when and where. *Global Change Biology*, **22**, 1561–1571.  
1850 doi:10.1111/gcb.13152
- 1851 Heymann, J., Reuter, M., Buchwitz, M., Schneising, O., Bovensmann, H., Burrows, J. P.,  
1852 Massart, S., Kaiser, J. W., and Crisp, D. (2017). CO<sub>2</sub> emission of Indonesian fires in 2015  
1853 estimated from satellite-derived atmospheric CO<sub>2</sub> concentrations. *Geophysical Research*  
1854 *Letters*, **44**, 1537–1544. doi: 10.1002/2016GL072042
- 1855 Hoppema, M., Bakker, K., van Heuven, S. M. A. C., van Ooijen, J. C. and de Baar, H. J. W.  
1856 (2015). Distributions, trends and inter-annual variability of nutrients along a repeat  
1857 section through the Weddell Sea (1996–2011). *Marine Chemistry*, **177**, 545–553.  
1858 doi:10.1016/j.marchem.2015.08.007
- 1859 Houghton, R. A. (2003). Revised estimates of the annual net flux of carbon to the atmosphere  
1860 from changes in land use and land management 1850–2000. *Tellus*, **55B**, 378–390.  
1861 doi:10.1034/j.1600-0889.2003.01450.x
- 1862 Houghton, R. A. and Nassikas, A. A. (2017). Global and regional fluxes of carbon from land use  
1863 and land cover change 1850–2015. *Global Biogeochemical Cycles*, **31**, 456–472.  
1864 doi:10.1002/2016GB005546
- 1865 Hubau, W., Lewis, S. L., Phillips O. L., Affum-Baffoe, Beeckman, K. H., Cuni-Sanchez, A.,  
1866 Ewango, C. E. N., Fauset, S., Sheil, D., Sonké, B., Sullivan, M. J. P., Sunderland, T.,  
1867 Thomas, S. C., Abernethy, K. A., Adu-Bredu, S., Amani C. A., Baker, T. R., Banin, L. F.,  
1868 Baya, F., Begne, S. K., Bennett, A. C., Benedet, F., Bitariho, R., Bocko, Y. E., Boeckx,  
1869 P., Boundja, P., Brienen, R. J. W., Brncic, T., Chezeaux, E., Chuyong, G. B., Clark, C. J.,  
1870 Collins, M., Comiskey, J. A., Coomes, D. A., Dargie, G. C., de Haulleville, T., Kamdem,  
1871 M. N. D., Doucet, J. L., Esquivel-Muelbert, A., Feldpausch, T. R., Fofanah, A., Foli, E.  
1872 G., Gilpin, M., Gloor, E., Gonmadje, C., Gourlet-Fleury, S., Hall, J. S., Hamilton, A. C.,  
1873 Harris, D. J., Hart, T. B., Hockemba, M. B. N., Hladik, A., Ifo, S. F., Jeffery, K. J.,  
1874 Jucker, T., Yakusu, E. K., Kearsley, E., Kenfack, D., Koch, A., Leal, M. E., Levesley, A.,  
1875 Lindsell, J. A., Lisingo, J., Lopez-Gonzalez, G., Lovett, J. C., Makana, J. R., Malhi, Y.,  
1876 Marshall, A. R., Martin, J., Martin, E. H., Mbayu, F. M., Medjibe, V. P., Mitchard, E. T.  
1877 A., Moore, S., Munishi, P. K. T., Bengone, N. N., Ojo, L., Ondo, F. E., Peh, K. S.,

- 1878 Pickavance, G. C., Poulsen, A. D., Poulsen, J. R., Qie, L., Reitsma, J., Rovero, F.,  
1879 Swaine, M. D., Talbot, J., Taplin J., Taylor, D. D., Thomas, D. W., Toirambe, B.,  
1880 Mukendi, J. T., Tuagben, D., Umunay, P. M., van der Heijden, G. M. F., Verbeeck, H.,  
1881 Vleminckx, J., Willcock, S., Wöll, H., Woods, J. T. and Zemagho, L. (2020).  
1882 Asynchronous carbon sink saturation in African and Amazonian tropical forests. *Nature*,  
1883 **579**, 80–87. doi:10.1038/s41586-020-2035-0
- 1884 Huber, M. B. and Zanna, L. (2017). Drivers of uncertainty in simulated ocean circulation and  
1885 heat uptake. *Geophysical Research Letters*, **44**(3), 1402-1413.  
1886 doi:10.1002/2016GL071587
- 1887 Hurtt, G. C., Chini, L., Sahajpal, R., Frolking, S., Boudirsky, B. L., Calvin, K., Doelman, J. C.,  
1888 Fisk, J., Fujimori, S., Klein Goldewijk, K., Hasegawa, T., Havlik, P., Heinemann, A.,  
1889 Humenöder, F., Jungclaus, J., Kaplan, J. O., Kennedy, J., Krisztin, T., Lawrence, D.,  
1890 Lawrence, P., Ma, L., Mertz, O., Pongratz, J., Popp, A., Poulter, B., Riahi, K.,  
1891 Shevliakova, E., Stehfest, E., Thornton, P., Tubiello, F. N., van Vuuren, D. P. and Zhang,  
1892 X. (2020). Harmonization of global land use change and management for the period 850–  
1893 2100 (LUH2) for CMIP6. *Geosci. Model Dev.*, **13**, 5425–5464. doi:10.5194/gmd-  
1894 135425-2020
- 1895 Iida, T., Odate, T., Fukuchi, M. (2013). Long-Term Trends of Nutrients and Apparent Oxygen  
1896 Utilization South of the Polar Front in Southern Ocean Intermediate Water from 1965 to  
1897 2008. *PLoS ONE*, **8**, e71766. doi:10.1371/journal.pone.0071766
- 1898 IEA World Energy Balances, 2020 Edition. Available from [https://www.iea.org/subscribe-to-](https://www.iea.org/subscribe-to-data-services/world-energy-balances-and-statistics)  
1899 [data-services/world-energy-balances-and-statistics](https://www.iea.org/subscribe-to-data-services/world-energy-balances-and-statistics), last viewed on 2 February 2021.
- 1900 IPCC 2006, 2006 IPCC Guidelines for National Greenhouse Gas Inventories, Prepared by the  
1901 National Greenhouse Gas Inventories Programme, Eggleston H.S., Buendia L., Miwa K.,  
1902 Ngara T. and Tanabe K. (eds). Published: IGES, Japan.
- 1903 IPCC 2019, 2019 Refinement to the 2006 IPCC Guidelines for National Greenhouse Gas  
1904 Inventories, Calvo Buendia, E., Tanabe, K., Kranjc, A., Baasansuren, J., Fukuda, M.,  
1905 Ngarize S., Osako, A., Pyrozhenko, Y., Shermanau, P. and Federici, S. (eds). Published:  
1906 IPCC, Switzerland.
- 1907 Ito, A. (2020). Constraining size-dependence of vegetation respiration rates, *Scientific Reports*,  
1908 **10**, 4304. doi:10.1038/s41598-020-61239-0
- 1909 Iudicone, D., Rodgers, K. B., Plancherel, Y., Aumont, O., Ito, T., Key, R. M., Madec, G. and  
1910 Ishii, M. (2016). The formation of the ocean’s anthropogenic carbon reservoir. *Scientific*  
1911 *Reports*, **6**, 35473. doi:10.1038/srep35473
- 1912 Jacobson, A. R., Mikaloff Fletcher, S. E., Gruber, N., Sarmiento, J. L. and Gloor, M. (2007). A  
1913 joint atmosphere-ocean inversion for surface fluxes of carbon dioxide: 1. Methods and  
1914 global-scale fluxes. *Global Biogeochemical Cycles*, **21**(1), 273.  
1915 doi:10.1029/1999GL900363
- 1916 Janssens-Maenhout, G., Crippa, M., Guizzardi, D., Muntean, M., Schaaf, E., Dentener, F.,  
1917 Bergamaschi, P., Pagliari, V., Olivier, J. G. J., Peters, J. A. H. W., van Aardenne, J. A.,  
1918 Monni, S., Doering, U., Petrescu, A. M. R., Solazzo, E. and Oreggioni, G. D. (2019).

- 1919 EDGAR v4.3.2 Global Atlas of the three major greenhouse gas emissions for the period  
1920 1970–2012. *Earth Syst. Sci. Data*, **11**, 959–1002. doi:10.5194/essd-11-959-2019
- 1921 Janssens-Maenhout, G., Pinty, B., Dowell, M., Zunker, H., Andersson, E., Balsamo, G., Bézy, J.  
1922 L., Brunhes, T., Bösch, H., Bojkov, B., Brunner, D., Buchwitz, M., Crisp, D., Ciais, P.,  
1923 Counet, P., Dee, D., Denier van der Gon, H., Dolman, H., Drinkwater, M., Dubovnik O.,  
1924 Engelen, R., Fehr, T., Fernandez, V., Heimann, M., Holmlund, K., Houweling, S.,  
1925 Husband, R., Juvyns, O., Kentarchos, A., Landgraf, J., Lang, R., Löschner, A., Marshall,  
1926 J., Meijer, Y., Nakajima, M., Palmer, P. I., Peylin, P., Rayner, P., Scholze, M., Sierk, B.,  
1927 Tamminen, J., Veeffkind, P. (2020). Toward an operational anthropogenic CO<sub>2</sub> emissions  
1928 monitoring and verification support capacity. *Bulletin of the American Meteorological*  
1929 *Society*, **101**(8), E1439–E1451. doi:10.1175/BAMS-D-19-0017.1
- 1930 Jian, J., Vargas, R., Anderson-Teixeira, K., Stell, E., Herrmann, V., Horn, M., Kholod, N.,  
1931 Manzon, J., Marchesi, R., Paredes, D. and Bond-Lamberty, B. (2020). A restructured and  
1932 updated global soil respiration database (SRDB-V5). *Earth Syst. Sci. Data Discuss.*  
1933 [preprint], doi:10.5194/essd-2020-136, in review.
- 1934 Johnson, K. S., Jannasch, H. W., Coletti, L. J., Elrod, V. A., Martz, T. R., Takeshita, Y., Carlson,  
1935 R. J. and Connery, J. G. (2016). Deep-Sea DuraFET: A pressure tolerant pH sensor  
1936 designed for global sensor networks. *Analytical Chemistry*, **88**, 3249–3256.  
1937 doi:10.1021/acs.analchem.5b04653
- 1938 Jones, C. D. and P. M. Cox (2005). On the significance of atmospheric CO<sub>2</sub> growth rate  
1939 anomalies in 2002– 2003. *Geophysical Research Letters*, **32**, L14816.  
1940 doi:10.1029/2005GL023027
- 1941 Joos, F., Bruno, M., Fink, R., Siegenthaler, U., Stocker, T. F., Le Quéré, C. and Sarmiento, J. L.  
1942 (1996). An efficient and accurate representation of complex oceanic and biospheric  
1943 models of anthropogenic carbon uptake. *Tellus B*, **48**, 397. doi:10.1034/j.1600-  
1944 0889.1996.t01-2-00006.x
- 1945 Jung, M., Reichstein, M. and Bondeau, A. (2009). Towards global empirical upscaling of  
1946 FLUXNET eddy covariance observations: validation of a model tree ensemble approach  
1947 using a biosphere model. *Biogeosciences*, **6**, 2001–2013. doi:10.5194/bg-6-2001-2009
- 1948 Jung, M., Schwalm, C. Migliavacca, M., Walther, S., Camps-Valls, G., Koirala, S., Anthoni, P.,  
1949 Besnard, S., Bodesheim, P., Carvalhais, N., Chevallier, F., Gans, F., Goll, D. S., Haverd,  
1950 V., Koehler, P., Ichii, K., Jain, A. K., Liu, J., Lombardozzi, D., Nabel, J. E. M. S.,  
1951 Nelson, J. A., O’Sullivan, M., Pallandt, M., Papale, D., Peters, W., Pongratz, J.,  
1952 Roedenbeck, C., Sitch, S., Tramontana, G., Walker, A., Weber, U. and Reichstein, M.  
1953 (2020). Scaling carbon fluxes from eddy covariance sites to globe: synthesis and  
1954 evaluation of the FLUXCOM approach. *Biogeosciences*, **17**, 1343–1365. doi:  
1955 10.5194/bg-17-1343-2020
- 1956 Karion, A., Sweeney, C., Tans, P. P. and Newberger, T. (2010). AirCore: An innovative  
1957 atmospheric sampling system. *Journal of Atmospheric and Oceanic Technology*, **27**,  
1958 1839–1853. doi:10.1175/2010JTECHA1448.1
- 1959 Keenan, T. F., and Williams, C. A. (2018). The terrestrial carbon sink. *Annual Review of*  
1960 *Environment and Resources*. **43**, 219–243. doi:10.1146/annurev-environ- 102017-030204

- 1961    Keppler, L., Landschützer, P. (2019). Regional wind variability modulates the Southern Ocean  
1962       carbon sink. *Scientific Reports*, **9**, 7384. doi:10.1038/s41598-019-43826-y
- 1963    Key, R. M., Kozyr, A., Sabine, C. L., Lee, K., Wanninkhof, R., Bullister, J. L., Feely, R. A.,  
1964       Millero, F. J., Mordy, C. and Peng, T. H. (2004). A global ocean carbon climatology:  
1965       Results from Global Data Analysis Project (GLODAP). *Global Biogeochemical Cycles*,  
1966       **18**(4), GB4031. doi:10.1029/2004GB002247
- 1967    Khatiwala, S., Primeau, F. and Hall, T. (2009). Reconstruction of the history of anthropogenic  
1968       CO<sub>2</sub> concentrations in the ocean. *Nature*, **462**, 346–349. doi:10.1038/nature08526
- 1969    Khatiwala, S., Tanhua, T., Mikaloff Fletcher, S., Gerber, M., Doney, C. S., Graven, H. D.,  
1970       Gruber, N., Mckinley, G. A., Murata, A. and Sabine, C. (2013). Global storage of  
1971       anthropogenic carbon. *Biogeosciences*, **10**, 2169-2191, 2013. doi: 10.519/bg-10-2169-  
1972       2013
- 1973    Kiel, M., O'Dell, C. W., Fisher, B., Eldering, A., Nassar, R., MacDonald, C. G. and Wennberg,  
1974       P. O. (2019). How bias correction goes wrong: measurement of XCO<sub>2</sub> affected by  
1975       erroneous surface pressure estimates. *Atmospheric Measurement Techniques*, **12**, 2241-  
1976       2259. doi: 10.5194/amt-12-2241-2019
- 1977    Knorr, W. (2009). Is the airborne fraction of anthropogenic CO<sub>2</sub> emissions increasing?  
1978       *Geophysical Research Letters*, **36**, L21710. doi:10.1029/2009GL040613
- 1979    Koren, G., van Schaik, E., Araujo, A. C., Boersma, K. F., Gartner, A., Killaars, L., Kooreman,  
1980       M. L., Kruijt, B., van der Laan-Luijkx, I. T., von Randow, C., Smith, N. E., and Peters,  
1981       W. (2018). Widespread reduction in sun-induced fluorescence from the Amazon during  
1982       the 2015/2016 El Nino. *Philosophical Transactions of the Royal Society of London.*  
1983       *Series B: Biological Sciences*, **373**(1760), 20170408. doi:10.1098/rstb.2017.0408
- 1984    Lacroix, F., T. Ilyina, and J. Hartmann (2020). Oceanic CO<sub>2</sub> outgassing and biological  
1985       production hotspots induced by pre-industrial river loads of nutrients and carbon in a  
1986       global modeling approach. *Biogeosciences*, **17**(1), 55–88. doi:10.5194/bg-17-55-2020
- 1987    Landschützer, P., Gruber, N., Bakker, D. C. E., Schuster, U., Nakaoka, S., Payne, M. R., Sasse,  
1988       T. P. and Zeng, J. (2013). A neural network-based estimate of the seasonal to inter-annual  
1989       variability of the Atlantic Ocean carbon sink. *Biogeosciences*, **10**, 7793–7815.  
1990       doi:10.5194/bg-10-7793-2013
- 1991    Landschützer, P., Gruber, N., Bakker, D. C. E. and Schuster, U. (2014). Recent variability of the  
1992       global ocean carbon sink. *Global Biogeochemical Cycles*, **28**, 927–949.  
1993       doi:10.1002/2014GB004853
- 1994    Landschützer, P., Gruber, N., Haumann, F. A., Rödenbeck, C., Bakker, D. C. E., van Heuven, S.,  
1995       Hoppema, M., Metzl, N., Sweeney, C., Takahashi, T., Tilbrook, B. and Wanninkhof, R.  
1996       (2015). The reinvigoration of the Southern Ocean carbon sink. *Science* **349**, 1221–1224.  
1997       doi:10.1126/science.aab2620
- 1998    Landschützer, P., Laruelle, G. G. , Roobaert, A. and Regnier, P. (2020). A uniform pCO<sub>2</sub>  
1999       climatology combining open and coastal oceans. *Earth Syst. Sci. Data*, **12**, 2537–2553.  
2000       doi:10.5194/essd-2020-90

- 2001 Langlais, C. E., Lenton, A., Matear, R., Monselesan, D., Legresy, B., Cougnon, E. and Rintoul,  
2002 S. (2017). Stationary Rossby waves dominate subduction of anthropogenic carbon in the  
2003 Southern Ocean. *Scientific Reports*, **7**, 17076. doi:10.1038/s41598-017-17292-3
- 2004 Laufkötter, C., Vogt, M., Gruber, N., Aita-Noguchi, M., Aumont, O., Bopp, L., Buitenhuis, E.,  
2005 Doney, S. C., Dunne, J., Hashioka, T., Hauck, J., Hirata, T., John, J., Le Quéré, C., Lima,  
2006 I.D., Nakano, H., Seferian, R., Totterdell, I., Vichi, M. and Völker, C. (2015). Drivers  
2007 and uncertainties of future global marine primary production in marine ecosystem  
2008 models. *Biogeosciences*, **12**, 6955–6984. doi:10.5194/bg-12-6955-2015
- 2009 Laufkötter, C., Vogt, M., Gruber, N., Aumont, O., Bopp, L., Doney, S.C., Dunne, J.P., Hauck, J.,  
2010 John, J.G., Lima, I.D., Seferian, R. and Völker, C. (2016). Projected decreases in future  
2011 marine export production: the role of the carbon flux through the upper ocean ecosystem.  
2012 *Biogeosciences*, **13**, 4023–4047. doi:10.5194/bg-13-4023-2016
- 2013 Lenton, A. and Matear, R. J. (2007). Role of the Southern Annular Mode (SAM) in Southern  
2014 Ocean CO<sub>2</sub> uptake. *Global Biogeochemical Cycles*, **21**, GB2016.  
2015 doi:10.1029/2006GB002714
- 2016 Le Quéré, C., Rödenbeck, C., Buitenhuis, E. T., Conway, T. J., Langenfelds, R., Gomez, A.,  
2017 Labuschagne, C., Ramonet, M., Nakazawa, T., Metzl, N., Gillett, N. and Heimann, M.  
2018 (2007). Saturation of the Southern Ocean CO<sub>2</sub> Sink Due to Recent Climate Change.  
2019 *Science*, **316**, 1735–1738. doi:10.1126/science.1136188
- 2020 Le Quéré, C., Raupach, M. R., Canadell, J. G., Marland, G., Bopp, L., Ciais, P., Conway, T. J.,  
2021 Doney, S. C., Feely, R. A., Foster, P., Friedlingstein, P., Gurney, K., Houghton, R. A.,  
2022 House, J. I., Huntingford, C., Levy, P. E., Lomas, M. R., Majkut, J., Metzl, N., Ometto, J.  
2023 P., Peters, G. P., Prentice, I. C., Randerson, J. T., Running, S. W., Sarmiento, J. L.,  
2024 Schuster, U., Sitch, S., Takahashi, T., Viovy, N., van der Werf, G. R. and Woodward, F.  
2025 I. (2009). Trends in the sources and sinks of carbon dioxide. *Nature Geosciences*, **2**, 831–  
2026 836. doi:10.1038/ngeo689
- 2027 Le Quéré, C., Takahashi, T., Buitenhuis, E. T., Rödenbeck, C. and Sutherland, S. C. (2010).  
2028 Impact of climate change and variability on the global oceanic sink of CO<sub>2</sub>. *Global*  
2029 *Biogeochemical Cycles*, **24**(4), GB4007. doi:10.1029/2009GB003599
- 2030 Le Quéré, C., Andres, R. J., Boden, T., Conway, T., Houghton, R. A., House, J. I., Marland, G.,  
2031 Peters, G. P., van der Werf, G. R., Ahlström, A., Andrew, R. M., Bopp, L., Canadell, J.  
2032 G., Ciais, P., Doney, S. C., Enright, C., Friedlingstein, P., Huntingford, C., Jain, A. K.,  
2033 Jourdain, C., Kato, E., Keeling, R. F., Klein Goldewijk, K., Levis, S., Levy, P., Lomas,  
2034 M., Poulter, B., Raupach, M. R., Schwinger, J., Sitch, S., Stocker, B. D., Viovy, N.,  
2035 Zaehle and S. and Zeng, N. (2013). The global carbon budget 1959–2011. *Earth Syst. Sci.*  
2036 *Data*, **5**, 165–185. doi:10.5194/essd-5-165-2013
- 2037 Le Quéré, C., Peters, G. P., Andres, R. J., Andrew, R. M., Boden, T. A., Ciais, P., Friedlingstein,  
2038 P., Houghton, R. A., Marland, G., Moriarty, R., Sitch, S., Tans, P., Arneeth, A., Arvanitis,  
2039 A., Bakker, D. C. E., Bopp, L., Canadell, J. G., Chini, L. P., Doney, S. C., Harper, A.,  
2040 Harris, I., House, J. I., Jain, A. K., Jones, S. D., Kato, E., Keeling, R. F., Klein  
2041 Goldewijk, K., Körtzinger, A., Koven, C., Lefèvre, N., Maignan, F., Omar, A., Ono, T.,  
2042 Park, G.-H., Pfeil, B., Poulter, B., Raupach, M.R., Regnier, P., Rödenbeck, C., Saito, S.,  
2043 Schwinger, J., Segschneider, J., Stocker, B.D., Takahashi, T., Tilbrook, B., van Heuven,



- 2044 S., Viovy, N., Wanninkhof, R., Wiltshire, A. and Zaehle, S. (2014). Global carbon budget  
2045 2013. *Earth Syst. Sci. Data*, **6**, 235–263. doi:10.5194/essd-6-235-2014
- 2046 Le Quéré, C., Moriarty, R., Andrew, R. M., Peters, G. P., Ciais, P., Friedlingstein, P., Jones, S.  
2047 D., Sitch, S., Tans, P., Arneth, A., Boden, T. A., Bopp, L., Bozec, Y., Canadell, J. G.,  
2048 Chini, L. P., Chevallier, F., Cosca, C. E., Harris, I., Hoppema, M., Houghton, R. A.,  
2049 House, J. I., Jain, A. K., Johannessen, T., Kato, E., Keeling, R. F., Kitidis, V., Klein  
2050 Goldewijk, K., Koven, C., Landa, C. S., Landschützer, P., Lenton, A., Lima, I. D.,  
2051 Marland, G., Mathis, J. T., Metzl, N., Nojiri, Y., Olsen, A., Ono, T., Peng, S., Peters, W.,  
2052 Pfeil, B., Poulter, B., Raupach, M. R., Regnier, P., Rödenbeck, C., Saito, S., Salisbury, J.  
2053 E., Schuster, U., Schwinger, J., Séférian, R., Segsneider, J., Steinhoff, T., Stocker, B.  
2054 D., Sutton, A. J., Takahashi, T., Tilbrook, B., van der Werf, G. R., Viovy, N., Wang, Y.-  
2055 P., Wanninkhof, R., Wiltshire, A. and Zeng, N. (2015). Global carbon budget 2014. *Earth*  
2056 *Syst. Sci. Data*, **7**, 47–85. doi:10.5194/essd-7-47-2015
- 2057 Le Quéré, C., Moriarty, R., Andrew, R. M., Canadell, J. G., Sitch, S., Korsbakken, J. I.,  
2058 Friedlingstein, P., Peters, G. P., Andres, R.J., Boden, T. A., Houghton, R. A., House, J. I.,  
2059 Keeling, R. F., Tans, P., Arneth, A., Bakker, D. C .E., Barbero, L., Bopp, L., Chang, J.,  
2060 Chevallier, F., Chini, L. P., Ciais, P., Fader, M., Feely, R. A., Gkritzalis, T., Harris, I.,  
2061 Hauck, J., Ilyina, T., Jain, A. K., Kato, E., Kitidis, V., Klein Goldewijk, K., Koven, C.,  
2062 Landschützer, P., Lauvset, S. K., Lefèvre, N., Lenton, A., Lima, I. D., Metzl, N., Millero,  
2063 F., Munro, D. R., Murata, A., Nabel, J. E. M. S., Nakaoka, S., Nojiri, Y., O’Brien, K.,  
2064 Olsen, A., Ono, T., Pérez, F. F., Pfeil, B., Pierrot, D., Poulter, B., Rehder, G., Rödenbeck,  
2065 C., Saito, S., Schuster, U., Schwinger, J., Séférian, R., Steinhoff, T., Stocker, B. D.,  
2066 Sutton, A. J., Takahashi, T., Tilbrook, B., van der Laan-Luijkx, I. T., van der Werf, G. R.,  
2067 van Heuven, S., Vandemark, D., Viovy, N., Wiltshire, A., Zaehle, S. and Zeng, N.  
2068 (2015). Global Carbon Budget 2015. *Earth Syst. Sci. Data*, **7**, 349–396. doi:10.5194/essd-  
2069 7-349-2015
- 2070 Le Quéré, C., Andrew, R. M., Canadell, J. G., Sitch, S., Korsbakken, J. I., Peters, G. P.,  
2071 Manning, A. C., Boden, T. A., Tans, P. P., Houghton, R. A., Keeling, R. F., Alin, S.,  
2072 Andrews, O. D., Anthoni, P., Barbero, L., Bopp, L., Chevallier, F., Chini, L. P., Ciais, P.,  
2073 Currie, K., Delire, C., Doney, S. C., Friedlingstein, P., Gkritzalis, T., Harris, I., Hauck, J.,  
2074 Haverd, V., Hoppema, M., Klein Goldewijk, K., Jain, A. K., Kato, E., Körtzinger, A.,  
2075 Landschützer, P., Lefèvre, N., Lenton, A., Lienert, S., Lombardozzi, D., Melton, J. R.,  
2076 Metzl, N., Millero, F., Monteiro, P. M. S., Munro, D. R., Nabel, J. E. M. S., Nakaoka, S.,  
2077 O’Brien, K., Olsen, A., Omar, A. M., Ono, T., Pierrot, D., Poulter, B., Rödenbeck, C.,  
2078 Salisbury, J., Schuster, U., Schwinger, J., Séférian, R., Skjelvan, I., Stocker, B. D.,  
2079 Sutton, A. J., Takahashi, T., Tian, H., Tilbrook, B., van der Laan-Luijkx, I. T., van der  
2080 Werf, G. R., Viovy, N., Walker, A. P., Wiltshire, A. J. and Zaehle, S. (2016). Global  
2081 Carbon Budget 2016. *Earth Syst. Sci. Data*, **8**, 605–649. doi:10.5194/essd-8-605-2016
- 2082 Le Quéré, C., Andrew, R. M., Friedlingstein, P., Sitch, S., Hauck, J., Pongratz, J., Pickers, P. A.,  
2083 Korsbakken, J. I., Peters, G. P., Canadell, J. G., Arneth, A., Arora, V. K., Barbero, L.,  
2084 Bastos, A., Bopp, L., Chevallier, F., Chini, L.P., Ciais, P., Doney, S. C., Gkritzalis, T.,  
2085 Goll, D. S., Harris, I., Haverd, V., Hoffman, F. M., Hoppema, M., Houghton, R. A.,  
2086 Hurtt, G., Ilyina, T., Jain, A.K., Johannessen, T., Jones, C. D., Kato, E., Keeling, R. F.,  
2087 Goldewijk, K. K., Landschützer, P., Lefèvre, N., Lienert, S., Liu, Z., Lombardozzi, D.,  
2088 Metzl, N., Munro, D. R., Nabel, J. E. M. S., Nakaoka, S., Neill, C., Olsen, A., Ono, T.,

- 2089 Patra, P., Peregon, A., Peters, W., Peylin, P., Pfeil, B., Pierrot, D., Poulter, B., Rehder,  
2090 G., Resplandy, L., Robertson, E., Rocher, M., Rödenbeck, C., Schuster, U., Schwinger,  
2091 J., Séférian, R., Skjelvan, I., Steinhoff, T., Sutton, A., Tans, P. P., Tian, H., Tilbrook, B.,  
2092 Tubiello, F. N., van der Laan-Luijkx, I. T., van der Werf, G. R., Viovy, N., Walker, A. P.,  
2093 Wiltshire, A. J., Wright, R., Zaehle, S. and Zheng, B. (2018a). Global Carbon Budget  
2094 2018. *Earth Syst. Sci. Data*, **10**, 2141–2194. doi:10.5194/essd-10-2141-2018
- 2095 Le Quéré, C., Andrew, R. M., Friedlingstein, P., Sitch, S., Pongratz, J., Manning, A. C.,  
2096 Korsbakken, J. I., Peters, G. P., Canadell, J. G., Jackson, R. B., Boden, T. A., Tans, P. P.,  
2097 Andrews, O. D., Arora, V. K., Bakker, D. C. E., Barbero, L., Becker, M., Betts, R. A.,  
2098 Bopp, L., Chevallier, F., Chini, L. P., Ciais, P., Cosca, C. E., Cross, J., Currie, K., Gasser,  
2099 T., Harris, I., Hauck, J., Haverd, V., Houghton, R. A., Hunt, C. W., Hurtt, G., Ilyina, T.,  
2100 Jain, A. K., Kato, E., Kautz, M., Keeling, R. F., Klein Goldewijk, K., Körtzinger, A.,  
2101 Landschützer, P., Lefèvre, N., Lenton, A., Lienert, S., Lima, I., Lombardozzi, D., Metzl,  
2102 N., Millero, F., Monteiro, P. M. S., Munro, D. R., Nabel, J. E. M. S., Nakaoka, S., Nojiri,  
2103 Y., Padin, X. A., Peregon, A., Pfeil, B., Pierrot, D., Poulter, B., Rehder, G., Reimer, J.,  
2104 Rödenbeck, C., Schwinger, J., Séférian, R., Skjelvan, I., Stocker, B. D., Tian, H.,  
2105 Tilbrook, B., Tubiello, F. N., van der Laan-Luijkx, I. T., van der Werf, G. R., van  
2106 Heuven, S., Viovy, N., Vuichard, N., Walker, A. P., Watson, A. J., Wiltshire, A. J.,  
2107 Zaehle, S. and Zhu, D. (2018b). Global Carbon Budget 2017. *Earth Syst. Sci. Data*, **10**,  
2108 405–448. doi:10.5194/essd-10-405-2018
- 2109 Liu, J., Bowman, K., Schimel, D., Parazoo, N., Jiang, Z., Lee, M., Bloom, A., Wunch, D.,  
2110 Gurney, K., Menemenlis, D., Girerach, M., Crisp, D. and Eldering A. (2017). Contrasting  
2111 carbon cycle responses of the tropical continents to the 2015–2016 El Niño. *Science*, **358**,  
2112 eaam5690. doi: 10.1126/science.aam5690
- 2113 Liu, J., Wennberg, P. O., Parazoo, N. C., Yin, Y. and Frankenberg, C. (2020). Observational  
2114 constraints on the response of high-latitude northern forests to warming. *AGU Advances*,  
2115 **2**, e2020AV000228. doi:10.1029/2020AV000228
- 2116 Liu, Z., Dreybrodt, W. and Wang, H. (2010). A new direction in effective accounting for the  
2117 atmospheric CO<sub>2</sub> budget: Considering the combined action of carbonate dissolution, the  
2118 global water cycle and photosynthetic uptake of DIC by aquatic organisms. *Earth Science*  
2119 *Reviews*, **99**, 169–172. doi:10.1016/j.earscirev.2010.03.001
- 2120 Long, M. C., Lindsay, K., Peacock, S., Moore, J. K., and Doney, S. C. (2013). Twentieth-century  
2121 oceanic carbon uptake and storage in CESM1(BGC). *Journal of Climate*, **26**, 6775–6800.  
2122 doi:10.1175/JCLI-D-12-00184.1
- 2123 Lovenduski, N. S., Gruber, N. and Doney, S.C. (2008). Toward a mechanistic understanding of  
2124 the decadal trends in the Southern Ocean carbon sink: Southern Ocean CO<sub>2</sub> flux trends.  
2125 *Global Biogeochemical Cycles*, **22**(3), GB3016. doi:10.1029/2007GB003139
- 2126 Lovenduski, N. S., Gruber, N., Doney, S. C. and Lima, I. D. (2007). Enhanced CO<sub>2</sub> outgassing in  
2127 the Southern Ocean from a positive phase of the Southern Annular Mode. *Global*  
2128 *Biogeochemical Cycles*, **21**(2), GB2026. doi:10.1029/2006GB002900
- 2129 Lucht, W., Prentice, C., Myneni, R. B., Sitch, S., Friedlingstein, P., Cramer, W., Bousquet, P.,  
2130 Buermann, W. and Smith, B. (2002). Climatic control of the high-latitude vegetation

- 2131 greening trend and Pinatubo effect. *Science*, **296**(5573), 1687–1689.  
2132 doi:10.1126/science.1071828
- 2133 MacDougall, A. H., Frölicher, T. L., Jones, C. D., Rogelj, J., Matthews, H. D., Zickfeld, K.,  
2134 Arora, V. K., Barrett, N. J., Brovkin, V., Burger, F. A., Eby, M., Eliseev, A. V., Hajima,  
2135 T., Holden, P. B., Jeltsch-Thömmes, A., Koven, C., Mengis, N., Menviel, L., Michou,  
2136 M., Mokhov, I. I., Oka, A., Schwinger, J., Séférian, R., Shaffer, G., Sokolov, A., Tachiiri,  
2137 K., Tjiputra, J., Wiltshire, A. and Ziehn, T. (2020). Is there warming in the pipeline? A  
2138 multi-model analysis of the zero emissions commitment from CO<sub>2</sub>. *Biogeosciences*, **17**,  
2139 2987–3016. doi:10.5194/bg-17-2987-2020
- 2140 Maier-Reimer, E. (1993). Geochemical cycles in an ocean general circulation model.  
2141 Preindustrial tracer distributions. *Global Biogeochemical Cycles*, **7**, 645–677, 1993.  
2142 doi:10.1029/93GB01355
- 2143 Marsay, C.M., Sanders, R. J., Henson, S. A., Pabortsava, K., Achterberg, E. P. and Lampitt, R. S.  
2144 (2015). Attenuation of sinking particulate organic carbon flux through the mesopelagic  
2145 ocean. *Proceedings of the National Academy of Sciences*, **112**, 1089.  
2146 doi:10.1073/pnas.1415311112
- 2147 Mau, A. C., Reed, S. C., Wood, T. E. and Cavaleri, M. A. (2018). Temperate and tropical forest  
2148 canopies are already functioning beyond their thermal thresholds for photosynthesis.  
2149 *Forests*, **9**, 47. doi:10.3390/f9010047
- 2150 McKinley, G., Follows, M., and Marshall, J. (2004). Mechanisms of air-sea CO<sub>2</sub> flux variability  
2151 in the equatorial Pacific and the North Atlantic. *Global Biogeochemical Cycles*, **18**(2),  
2152 GB2011. doi:10.1029/2003GB002179
- 2153 McKinley, G. A., Fay, A. R., Takahashi, T. and Metzl, N. (2011). Convergence of atmospheric  
2154 and North Atlantic carbon dioxide trends on multidecadal timescales. *Nature Geosci*, **4**,  
2155 606–610. doi:10.1038/ngeo1193
- 2156 McKinley, G. A., Pilcher, D. J., Fay, A. R., Lindsay, K., Long, M. C. and Lovenduski, N. S.  
2157 (2016). Timescales for detection of trends in the ocean carbon sink. *Nature*, **530**(7591),  
2158 469–472. doi:10.1038/nature16958
- 2159 McKinley, G. A., Fay, A. R., Lovenduski, N. S. and Pilcher, D. J. (2017). Natural variability and  
2160 anthropogenic trends in the ocean carbon sink. *Annual Review of Marine Science*, **9**(1),  
2161 125–150, doi:10.1146/annurev-marine-010816-060529
- 2162 McKinley, G. A., Fay, A. R., Eddebbar, Y. A., Gloege L. and Lovenduski, N. S. (2020). External  
2163 forcing explains recent decadal variability of the ocean carbon sink. *AGU Advances*, **1**(2),  
2164 1, e2019AV000149. doi:10.1029/2019AV000149
- 2165 Mikaloff Fletcher, S. E., Gruber, N., Jacobson, A. R., Doney, S., C., Sutkiewicz, S., Gerber, M.,  
2166 Follows, M., Joos, F., Lindsay, K., Menemenlis, D., Mouchet, A., Müller, S., A. and  
2167 Sarmiento, J. L. (2006). Inverse estimates of anthropogenic CO<sub>2</sub> uptake, transport, and  
2168 storage by the ocean, *Global Biogeochemical Cycles*, **20**, GB2002.  
2169 doi:10.1029/2005GB002530
- 2170 Miller, J. B., Lehman, S. J., Montzka, S. A., Sweeney, C., Miller, B. R., Karion, A., Wolak, C.,  
2171 Dlugokencky, E. J., Southon, J., Turnbull, J. C. and Tans, P. P. (2012). Linking emissions

- of fossil fuel CO<sub>2</sub> and other anthropogenic trace gases using atmospheric <sup>14</sup>CO<sub>2</sub>. *Journal of Geophysical Research: Atmospheres*, **117**, D08302. doi:10.1029/2011JD017048
- Miller, J. B., Lehman, S. J., Verhulst, K. R., Miler, C., E., Duren, R. M., Yadav, V., Newman, S. and Sloop, C. D. (2020). Large and seasonally varying biospheric CO<sub>2</sub> fluxes in the Los Angeles megacity revealed by atmospheric radiocarbon. *Proceedings of the National Academy of Sciences*, **117**, 26681–26687. doi:10.1073/pnas.2005253117
- Mongwe, N. P., Vichi, M. and Monteiro, P. M. S. (2018). The seasonal cycle of pCO<sub>2</sub> and CO<sub>2</sub> fluxes in the Southern Ocean: diagnosing anomalies in CMIP5 Earth system models. *Biogeosciences*, **15**, 2851–2872. doi:10.5194/bg-15-2851-2018
- Moore, J. K., Fu, W., Primeau, F., Britten, G. L., Lindsay, K., Long, M., Doney, S. C., Mahowald, N., Hoffman, F. and Randerson, J. T. (2018). Sustained climate warming drives declining marine biological productivity. *Science*, 359, 1139–1143. doi:10.1126/science.aao6379
- Nassar, R., Hill, T. G., McLinden, C. A., Wunch, D., Jones, D. B.A. and Crisp D. (2017). Quantifying CO<sub>2</sub> emissions from individual power plants from space. *Geophysical Research Letters*, **44**(19), 10045–10053. doi:10.1002/2017GL074702
- Oda, T., Maksyutov, S. and Andres, S. J. (2018). The Open-source Data Inventory for Anthropogenic Carbon dioxide (CO<sub>2</sub>), version 2016 (ODIAC2016): A global, monthly fossil-fuel CO<sub>2</sub> gridded emission data product for tracer transport simulations and surface flux inversions. *Earth Syst. Sci. Data*, **10**, 87–107. doi:10.5194/essd-10-87-2018
- O'Dell, C. W., Eldering, A., Wennberg, P. O., Crisp, D., Gunson, M. R., Fisher, B., Frankenberg, C., Kiel, M., Lindqvist, H., Mandrake, L., Merrelli, A., Natraj, V., Nelson, R. R., Osterman, G. B., Payne, V. H., Taylor, T. E., Wunch, D., Drouin, B. J., Oyafuso, F., Chang, A., McDuffie, J., Smyth, M., Baker, D. F., Basu, S., Chevallier, F., Crowell, S. M. R., Feng, L., Palmer, P. I., Dubey, M., García, O. E., Griffith, D. W. T., Hase, F., Iraci, L. T., Kivi, R., Morino, I., Notholt, J., Ohyama, H., Petri, C., Roehl, C. M., Sha, M. K., Strong, K., Sussmann, R., Te, Y., Uchino, O. and Velasco, V. A. (2018). Improved retrievals of carbon dioxide from Orbiting Carbon Observatory-2 with the version 8 ACOS algorithm, *Atmospheric Measurement Techniques*, **11**(12): 6539–6576. doi:10.5194/amt-11-6539-2018
- Olsen, A., Lange, N., Key, R. M., Tanhua, T., Bittig, H. C., Kozyr, A., Álvarez, M., Azetsu-Scott, K., Becker, S., Brown, P. J., Carter, B. R., Cotrim da Cunha, L., Feely, R. A., van Heuven, S., Hoppema, M., Ishii, M., Jeansson, E., Jutterström, S., Landa, C. S., Lauvset, S. K., Michaelis, P., Murata, A., Pérez, F. F., Pfeil, B., Schirnick, C., Steinfeldt, R., Suzuki, T., Tilbrook, B., Velo, A., Wanninkhof, R., and Woosley, R. J. (2021). GLODAPv2.2020 – the second update of GLODAPv2. *Earth Syst. Sci. Data Discuss.* doi:10.5194/essd-2020-165, in review, 2020.
- Palmer, P. I., Feng, L., Baker, D., Chevallier, F., Bösch, H. and Somkuti, P. (2019). Net carbon emissions from African biosphere dominate pan-tropical atmospheric CO<sub>2</sub> signal, *Nature Communications*, **10**, 3344. doi:10.1038/s41467-019-11097-w
- Pan, Y., Birdsey, R. A., Fang, J., Houghton, R., Kauppi, P. E., Kurz, W. A., Phillips, O. L., Shvidenko, A., Lewis, S. L., Canadell, J. G., Ciais, P., Jackson, R. B., Pacala, S. W., McGuire, A. S., Piao, S., Rautiainen, A., Sitch, S. and Hayes, D. (2011). A large and

- 2214 persistent carbon sink in the world's forests. *Science*, 333, 988-993.  
2215 doi:10.1126/science.1201609
- 2216 Panassa, E., Santana-Casiano, J. M., González-Dávila, M., Hoppema, M., van Heuven, S. M. A.  
2217 C., Völker, C., Wolf-Gladrow, D. and Hauck, J. (2018). Variability of nutrients and  
2218 carbon dioxide in the Antarctic Intermediate Water between 1990 and 2014. *Ocean*  
2219 *Dynamics*, **68**, 295–308. doi:10.1007/s10236-018-1131-2
- 2220 Parazoo, N. C., Magney, T., Norton, A., Raczka, B., Bacour, C., Maignan, F., Baker, I., Zhang,  
2221 Y., Qiu, B., Shi, M., MacBean, N., Bowling, D. R., Burns, S., Blanken, P. D., Stutz, J.,  
2222 Grossmann, K. and Frankenberg, C. (2020). Wide discrepancies in the magnitude and  
2223 direction of modeled solar-induced chlorophyll fluorescence in response to light  
2224 conditions. *Biogeosciences*, **17**, 3733–3755. doi:10.5194/bg-17-3733-2020
- 2225 Pardo, P. C., Tilbrook, B., Langlais, C., Trull, T.W. and Rintoul, S. R. (2017). Carbon uptake  
2226 and biogeochemical change in the Southern Ocean, south of Tasmania. *Biogeosciences*,  
2227 **14**, 5217–5237. doi:10.5194/bg-14-5217-2017
- 2228 Pearson, T. R. H., Brown, S., Murray, L. and Sidman, G. (2017). Greenhouse gas emissions from  
2229 tropical forest degradation: an underestimated source. *Carbon Balance and Management*,  
2230 **12**, 3. doi:10.1186/s13021-017-0072-2
- 2231 Peng, B., Guan, K. Y., Zhou, W. , Jiang, C. Y., Frankenberg, C., Sun, Y., He, L. Y. and Kohler,  
2232 P. (2020). Assessing the benefit of satellite-based Solar-Induced Chlorophyll  
2233 Fluorescence in crop yield prediction. *International Journal of Applied Earth*  
2234 *Observation and Geoinformation*, **90**, 102126. doi:10.1016/j.jag.2020.102126
- 2235 Petrescu, A. M. R., Peters, G. P., Janssens-Maenhout, G., Ciais, P., Tubiello, F. N., Grassi, G.,  
2236 Nabuurs, G.-J., Leip, A., Carmona-Garcia, G., Winiwarter, W., Höglund-Isaksson, L.,  
2237 Günther, D., Solazzo, E., Kiesow, A., Bastos, A., Pongratz, J., Nabel, J. E. M. S.,  
2238 Conchedda, G., Pilli, R., Andrew, R. M., Schelhaas, M.-J. and Dolman, A. J. (2020).  
2239 European anthropogenic AFOLU greenhouse gas emissions: a review and benchmark  
2240 data. *Earth Syst. Sci. Data*, **12**, 961–1001, doi:10.5194/essd-12-961-2020
- 2241 Pfeil, B., Olsen, A., Bakker, D. C. E., Hankin, S., Koyuk, H., Kozyr, A., Malczyk, J., Manke, A.,  
2242 Metzl, N., Sabine, C. L., Akl, J., Alin, S. R., Bates, N., Bellerby, R. G. J., Borges, A.,  
2243 Boutin, J., Brown, P. J., Cai, W.-J., Chavez, F. P., Chen, A., Cosca, C., Fassbender, A. J.,  
2244 Feely, R. A., González-Dávila, M., Goyet, C., Hales, B., Hardman-Mountford, N.,  
2245 Heinze, C., Hood, M., Hoppema, M., Hunt, C. W., Hydes, D., Ishii, M., Johannessen, T.,  
2246 Jones, S. D., Key, R. M., Körtzinger, A., Landschützer, P., Lauvset, S. K., Lefèvre, N.,  
2247 Lenton, A., Laurantou, A., Merlivat, L., Midorikawa, T., Mintrop, L., Miyazaki, C.,  
2248 Murata, A., Nakadate, A., Nakano, Y., Nakaoka, S., Nojiri, Y., Omar, A. M., Padin, X.  
2249 A., Park, G.-H., Paterson, K., Perez, F. F., Pierrot, D., Poisson, A., Ríos, A. F., Santana-  
2250 Casiano, J. M., Salisbury, J., Sarma, V. V. S. S., Schlitzer, R., Schneider, B., Schuster,  
2251 U., Sieger, R., Skjelvan, I., Steinhoff, T., Suzuki, T., Takahashi, T., Tedesco, K.,  
2252 Telszewski, M., Thomas, H., Tilbrook, B., Tjiputra, J., Vandemark, D., Veness, T.,  
2253 Wanninkhof, R., Watson, A. J., Weiss, R., Wong, C. S. and Yoshikawa-Inoue, H. (2013).  
2254 A uniform, quality controlled Surface Ocean CO<sub>2</sub> Atlas (SOCAT). *Earth Syst. Sci. Data*,  
2255 **5**, 125–143. doi:10.5194/essd-5-125-2013

- 2256 Piao, S., Wang, X., Park, T., Chen, C., Lian, X., He, Y. Bjerke, J. W., Chen, A., Ciais, P.,  
2257 Tommervik, H., Nemani, R. R. and Myneni, R. B. (2020). Characteristics, drivers and  
2258 feedbacks of global greening. *Nat. Rev. Earth Environ.*, **1**, 14–27. doi:10.1038/s43017-  
2259 019-0001-x
- 2260 Quegan, S., Toan, L. T., Chave, J., Dall, J., Exbrayat, J. F., Minh, D. H. T., Lomas, M.,  
2261 D'Alessandro, M. M., Paillou, P., Papathanassiou, K., Rocca, F., Saatchi, S., Scipal, K.,  
2262 Shugart, H., Smallman, T. L., Soja, M. J., Tebaldini, S., Ulander, L., Villard, L. and  
2263 Williams, M. (2019). The European Space Agency BIOMASS mission: measuring forest  
2264 above-ground biomass from space. *Remote Sens. Environ.*, **227**, 44–60.  
2265 doi:10.1016/j.rse.2019.03.032
- 2266 Qiu, B., Ge, J., Guo, W. D., Pittman, A. J. and Mu, M. Y. (2020). Responses of Australian  
2267 dryland vegetation to the 2019 heat wave at a sub daily scale. *Geophysical Research*  
2268 *Letters*, **47**, e2019GL086569. doi:10.1029/2019GL086569
- 2269 Ramankutty N., Gibbs, H. K., Achard, F., Defries, R., Foley, J. A. and Houghton, R. A. (2007).  
2270 Challenges to estimating carbon emissions from tropical deforestation. *Global Change*  
2271 *Biology*, **13**, 51–66. doi: 10.1111/j.1365-2486.2006.01272.x
- 2272 Randerson, J. T., Hoffman, F. M., Thornton, P. E., Mahowald, N. M., Lindsay, K., Lee, Y. H.,  
2273 Nevison, C. D. Doney, S. C., Bonan, G., Stöckli, R., Covey, C., Running, S. W. and  
2274 Fung, I. Y. (2009). Systematic assessment of terrestrial biogeochemistry in coupled  
2275 climate–carbon models. *Global Change Biology*, **15**(10), 2462–2484. doi: 10.1111/j.1365-  
2276 2486.2009.01912.x
- 2277 Randerson, J. T., Lindsay, K., Munoz, E., Fu, W., Moore, J. K., Hoffman, F. M., Mahowald, N.  
2278 M. and Doney, S. C. (2015). Multi-century changes in ocean and land contributions to the  
2279 climate-carbon feedback. *Global Biogeochemical Cycles*, **29**, 744–759.  
2280 doi:10.1002/2014GB005079
- 2281 Raupach, M. R., Canadell, J. G. and Le Quéré, C. (2008). Anthropogenic and biophysical  
2282 contributions to increasing atmospheric CO<sub>2</sub> growth rate and airborne fraction.  
2283 *Biogeosciences*, **5**, 1601–1613. doi:10.5194/bg-5-1601-2008
- 2284 Raupach, M. R., Gloor, M., Sarmiento, J. L., Canadell, J. G., Frölicher, T. L., Gasser, T.,  
2285 Houghton, R. A., Le Quéré, C. and Trudinger, C. M. (2014). The declining uptake rate of  
2286 atmospheric CO<sub>2</sub> by land and ocean sinks. *Biogeosciences*, **11**(13), 3453–3475.  
2287 doi:10.1007/s10584-009-9596-0
- 2288 Regnier, P., Friedlingstein, P., Ciais, P., Mackenzie, F. T., Gruber, N., Janssens, I. A., Laruelle,  
2289 G. G., Lauerwald, R., Luyssaert, S., Andersson, A. J., Arndt, S., Arnosti, C., Borges, A.  
2290 V., Dale, A. W., Gallego-Sala, A., Goddérís, Y., Goossens, N., Hartmann, J., Heinze, C.,  
2291 Ilyina, T., Joos, F., LaRowe, D. E., Leifeld, J., Meysman, F. J. R., Munhoven, G.,  
2292 Raymond, P. A., Spahni, R., Suntharalingam, P. and Thullner, M. (2013). Anthropogenic  
2293 perturbation of the carbon fluxes from land to ocean. *Nature Geosciences*, **6**, 597–607.  
2294 doi:10.1038/ngeo1830
- 2295 Resplandy, L., Keeling, R. F., Rodenbeck, C., Stephens, B. B, Khatiwala, S., Rodgers, K. B.,  
2296 Long, M. C., Bopp, L. and Tans, P. P. (2018). Revision of global carbon fluxes based on  
2297 a reassessment of oceanic and riverine carbon transport. *Nature Geoscience*, **11**, 504–  
2298 509. doi:10.1038/s41561-018-0151-3

- 2299 Reuter, M., Buchwitz, M., Schneising, O., Noël, S., Bovensmann, H. and Burrows, J. P. (2017).  
2300 A fast atmospheric trace gas retrieval for hyperspectral instruments approximating  
2301 multiple scattering—Part 2: Application to XCO<sub>2</sub> retrievals from OCO-2. *Remote*  
2302 *Sensing*, **9**, 1102. doi: 10.3390/rs9111102
- 2303 Reuter, M., Buchwitz, M., Schneising, O., Krautwurst, S., O'Dell, C. W., Richter, A.,  
2304 Bovensmann, H. and Burrows, J. P. (2019). Towards monitoring localized CO<sub>2</sub> emissions  
2305 from space: Co-located regional CO<sub>2</sub> and NO<sub>2</sub> enhancements observed by the OCO-2 and  
2306 S5P satellites. *Atmospheric Chemistry and Physics*, **19**, 9371–9383. doi:10.5194/acp-19-  
2307 9371-2019
- 2308 Ridge, S. M. and McKinley, G. A. (2020a). Advective controls on the North Atlantic  
2309 anthropogenic carbon sink. *Global Biogeochemical Cycles*, **34**(7), 1138.  
2310 doi:10.1029/2019GB006457
- 2311 Ridge, S. M. and G. A. McKinley (2020b). Ocean carbon uptake under aggressive emission  
2312 mitigation. *Biogeosciences Discussions* doi:10.5194/bg-2020-254, in review, 2020.
- 2313 Rödenbeck, C., Bakker, D. C. E., Metzl, N., Olsen, A., Sabine, C., Cassar, N., Reum, F., Keeling,  
2314 R. F and Heimann, M. (2014). Interannual sea-air CO<sub>2</sub> flux variability from an  
2315 observation-driven ocean mixed-layer scheme. *Biogeosciences*, **11**(1), 4599–4613.  
2316 doi:10.5194/bg-11-4599-2014
- 2317 Rödenbeck, C., Bakker, D. C. E., Gruber, N., Iida, Y., Jacobson, A. R., Jones, S., Landschützer,  
2318 P., Metzl, N., Nakaoka, S., Olsen, A., Park, G.-H., Peylin, P., Rodgers, K. B., Sasse, T.  
2319 P., Schuster, U., Shutler, J. D., Valsala, V., Wanninkhof, R. and Zeng, J. (2015). Data-  
2320 based estimates of the ocean carbon sink variability – first results of the Surface Ocean  
2321 pCO<sub>2</sub> Mapping intercomparison (SOCOM). *Biogeosciences*, **12**, 7251–7278.  
2322 doi:10.5194/bg-12-7251-2015
- 2323 Rodgers, K. B., Schlunegger, S., Slater, R. D., Ishii, M., Frolicher, T. L., Toyama, K.,  
2324 Plancherel, Y., Aumont, O. and Fassbender, A. J. (2020). Reemergence of anthropogenic  
2325 carbon into the ocean's mixed layer strongly amplifies transient climate sensitivity.  
2326 *Geophysics Research Letters*, **47**(18), 130. doi:10.1002/2017GL073758
- 2327 Rosen, P., Hensley, S., Shaffer, S., Edelstein, W., Kim, Y., Kumar, R., Misra, T., Bhan, R.,  
2328 Satish, R. and Sagi, R. (2016). An update on the NASA-ISRO dual-frequency dbf SAR  
2329 (NISAR) mission. *2016 IEEE International Geoscience and Remote Sensing Symposium*.  
2330 IEEE, New York, pp. 2106–2108. doi:10.1109/IGARSS.2016.7729543
- 2331 Rudnick, D. L. (2016). Ocean Research Enabled by Underwater Gliders. *Annual Review of*  
2332 *Marine Science*, **8**, 519-541. doi:10.1146/annurev-marine-122414-033913
- 2333 Saatchi, S. S., Harris, N. L., Brown, S., Lefsky, M., Mitchard, E. T. A., Salas, W., Zutta, B. R.,  
2334 Buermann, W., Lewis, S. L., Hagen, S., Petrova, S., White, L., Silman, M. and Morel, A.  
2335 (2011). Benchmark map of forest carbon stocks in tropical regions across three  
2336 continents. *Proceedings of the National Academy of Sciences*, **108**, 9899-9904. doi:  
2337 10.1073/pnas.1019576108
- 2338 Sabine, C. L., Feely, R. A., Gruber, N., Key, R. M., Lee, K., Bullister, K. L., Wanninkhof, R.  
2339 Wong, C. S., Wallace, D. W. R. Wallace, Tilbrook, B., Millero, F. J., Peng, T.-H., Kozyr,



- 2340 A., Ono, T. and Rois, A. F. (2004). The oceanic sink for anthropogenic CO<sub>2</sub>. *Science*,  
2341 **305**, 367–371. doi:10.1126/science.1097403
- 2342 Sabine, C. L. and Tanhua, T. (2010). Estimation of anthropogenic CO<sub>2</sub> inventories in the ocean.  
2343 *Annual Reviews of Marine Sciences*, **2**, 175-198. doi:10.1146/annurev-marine-120308-  
2344 080947
- 2345 Sabine, C., Sutton, A., McCabe, K., Lawrence-Slavas, N., Alin, S., Feely, R., Jenkins, R.,  
2346 Maenner, S., Meinig, C., Thomas, J., van Ooijen, E., Passmore, A. and Tilbrook, B.  
2347 (2020). Evaluation of a new carbon dioxide system for autonomous surface vehicles. *J.*  
2348 *Atmos. Ocean. Technol.*, **37**, 1305-1317. doi:10.1175/JTECH-D-20-0010.1
- 2349 Sarmiento, J. L., and Sundquist E. T. (1992). Revised budget for the oceanic uptake of  
2350 anthropogenic carbon dioxide, *Nature*, **356**, 589–593. doi:10.1038/356589a0
- 2351 Sarmiento, J. L. and Gruber, N. (2006). Ocean Biogeochemical Dynamics. Princeton University  
2352 Press. ISBN: 0-691-01707-7. doi:10.1017/S0016756807003755
- 2353 Scharlemann, J. P. W., Tanner, E. V. J., Hiederer, R. and Kapos, V. (2014). Global soil carbon:  
2354 understanding and managing the largest terrestrial carbon pool. *Carbon Management*, **5**,  
2355 81-91. doi:10.4155/cmt.13.77
- 2356 Schepaschenko, D., Moltchanova, E., Shvidenko, A., Blyshchyk, V., Dmitriev, E., Martynenko,  
2357 O., See, L. and Kraxner F. (2018). Improved estimates of biomass expansion factors for  
2358 russian forests. *Forests*, **9**, 312. doi:10.3390/f9060312
- 2359 Schourup-Kristensen, V., Sidorenko, D., Wolf-Gladrow, D. A. and Völker, C. (2014). A skill  
2360 assessment of the biogeochemical model REcoM2 coupled to the Finite Element Sea Ice–  
2361 Ocean Model (FESOM 1.3). *Geosci. Model Dev.*, **7**, 2769–2802. doi:10.5194/gmd-7-  
2362 2769-2014
- 2363 Schuur, E. A. G., McGuire, A. D., Schädel, C., Grosse, G., Harden, J. W., Hayes, D. J.,  
2364 Hugelius, G., Koven, C. D., Kuhry, P., Lawrence, D. M., Natali, S. M., Olefeldt, D.,  
2365 Romonovsky, V. E. Schaefer, K., Truetsky, M. R., Treat, C. C. and Vonk, J. E. (2015).  
2366 Climate change and the permafrost carbon feedback. *Nature*, **520**, 171-179.  
2367 doi:10.1038/nature14338
- 2368 Schwinger, J., Tjiputra, J. F., Heinze, C., Bopp, L., Christian, J. R., Gehlen, M., Ilyina, T., Jones,  
2369 C. D., Salas-Méla, D., Segschneider, J., Séférian, R. and Totterdell, I. (2014).  
2370 Nonlinearity of ocean carbon cycle feedbacks in CMIP5 Earth system models. *Journal of*  
2371 *Climate*, **27**, 3869–3888. doi:10.1175/JCLI-D-13-00452.1
- 2372 Schwinger, J., Goris, N., Tjiputra, J. F., Kriest, I., Bentsen, M., Bethke, I., Ilicak, M., Assmann,  
2373 K. M. and Heinze, C. (2016). Evaluation of NorESM-OC (versions 1 and 1.2), the ocean  
2374 carbon-cycle stand-alone configuration of the Norwegian Earth System Model  
2375 (NorESM1). *Geosci. Model Dev.*, **9**, 2589–2622. doi:10.5194/gmd-9-2589-2016
- 2376 Schwinger, J. and Tjiputra, J. (2018). Ocean carbon cycle feedbacks under negative emissions.  
2377 *Geophysical Research Letters*, **26**(3), 5289. doi:10.1088/1748-9326/11/5/055006
- 2378 Seelmann, K., Aßmann, S. and Körtzinger, A. (2019). Characterization of a novel autonomous  
2379 analyzer for seawater total alkalinity: Results from laboratory and field tests. *Limnology*  
2380 *and Oceanography: Methods*, **17**, 515-532. doi:10.1002/lom3.10329

- 2381 Sellers, P. J., Schimel, D. S., Moore III, B., Liu, J. and Eldering, A. (2018). Observing Carbon  
2382 Cycle-Climate Feedbacks from Space, *Proceedings of the National Academy of Sciences*,  
2383 **115**(31), 7860-7868. doi:10.1073/pnas.1716613115
- 2384 Sigman, D. M., Hain, M. P. and Haug, G. H. (2010). The polar ocean and glacial cycles in  
2385 atmospheric CO<sub>2</sub> concentration. *Nature*, **466**, 47–55. doi:10.1038/nature09149
- 2386 Sitch, S., Friedlingstein, P., Gruber, N., Jones, S. D., Murray-Tortarolo, G., Ahlström, A.,  
2387 Doney, S. C., Graven, H., Heinze, C., Huntingford, C., Levis, S., Levy, P. E., Lomas, M.,  
2388 Poulter, B., Viovy, N., Zaehle, S., Zeng, N., Arneth, A., Bonan, G., Bopp, L., Canadell, J.  
2389 G., Chevallier, F., Ciais, P., Ellis, R., Gloor, M., Peylin, P., Piao, S. L., Le Quéré, C.,  
2390 Smith, B., Zhu, Z. and Myneni, R. (2015). Recent trends and drivers of regional sources  
2391 and sinks of carbon dioxide. *Biogeosciences*, **12**, 653–679. doi:10.5194/bg-12-653-2015
- 2392 Smith P., Bustamante, M., Ahammad, H., Clark, H., Dong, H., Elsiddig, E. A., Haberl, H.,  
2393 Harper, R., House, J., Jafari M., Masera, O., Mbow, C., Ravindranath N. H., Rice C. W.,  
2394 Robledo Abad, C., Romanovskaya, A., Sperling, F. and Tubiello F. (2014) Agriculture,  
2395 Forestry and Other Land Use (AFOLU). In: Climate Change 2014: Mitigation of Climate  
2396 Change. Contribution of Working Group III to the Fifth Assessment Report of the  
2397 Intergovernmental Panel on Climate Change [Edenhofer, O., R. Pichs-Madruga, Y.  
2398 Sokona, E. Farahani, S. Kadner, K. Seyboth, A. Adler, I. Baum, S. Brunner, P.  
2399 Eickemeier, B. Kriemann, J. Savolainen, S. Schlömer, C. von Stechow, T. Zwickel and  
2400 J.C. Minx (eds.)]. Cambridge University Press, Cambridge, United Kingdom and New  
2401 York, NY, USA.
- 2402 Song, X.-P., Hansen, M. C., Stehman, S. V., Potapov, P. V., Tyukavina, A., Vermote, E. F. and  
2403 Townshend, J. R. (2018). Global land change from 1982 to 2016. *Nature*, **560**., 639-343.  
2404 doi:10.1038/s41586-018-0411-9
- 2405 Soong, J. L., Fuchslueger, L., Marañon-Jimenez, S., Torn, M. S., Janssens, I. A., Penuelas, J.,  
2406 and Richter, A. (2019). Microbial carbon limitation, 2021. The need for integrating  
2407 microorganisms into our understanding of ecosystem carbon cycling. *Global Change*  
2408 *Biology*, **26**, 1953–1961. doi:10.1111/gcb.14962
- 2409 Spawn, S. A., Sullivan, C. C., Lark, T. J. and Gibbs, H. K. (2020). Harmonized global maps of  
2410 above and belowground biomass carbon density in the year 2010. *Scientific Data*, **7**, 112.  
2411 doi:10.1038/s41597-020-0444-4
- 2412 Stamell, J., Rustagi, R. R., Gloege, L. and McKinley, G. A. (2021). Strengths and weaknesses of  
2413 three Machine Learning methods for pCO<sub>2</sub> interpolation. *Geosci. Model Dev. Discuss.*  
2414 [preprint], <https://doi.org/10.5194/gmd-2020-311>, in review, 2020.
- 2415 Stock, C. A., Dunne, J. P., Fan, S., Ginoux, P., John, J., Krasting, J. P., Laufkötter, C., Paulot, F.  
2416 and Zadeh, N. (2020). Ocean biogeochemistry in GFDL's Earth System Model 4.1 and its  
2417 response to increasing atmospheric CO<sub>2</sub>. *J. Adv. Model. Earth Syst.*, **12**,  
2418 e2019MS002043. doi:10.1029/2019MS002043
- 2419 Sun, Y., Frankenberg, C., Jung, J., Joiner, J., Guanter, L., Köhler, P. and Magney, T. (2018).  
2420 Overview of solar-Induced chlorophyll fluorescence (SIF) from the Orbiting Carbon  
2421 Observatory-2: Retrieval, cross-mission comparison, and global monitoring for GPP.  
2422 *Remote Sensing of Environment*, **209**, 808-823. doi:10.1016/j.rse.2018.02.016

- 2423 Sundquist, E.T., (1993). The global carbon dioxide budget. *Science*, **259**, 934–941.  
2424 doi:10.1126/science.259.5097.934
- 2425 Sutton, A. J., Sabine, C. L., Maenner-Jones, S., Lawrence-Slavas, N., Meinig, C., Feely, R. A.,  
2426 Mathis, J. T., Musielewicz, S., Bott, R., McLain, P. D., Fought, H. J. and Kozyr, A.  
2427 (2014). A high-frequency atmospheric and seawater pCO<sub>2</sub> data set from 14 open-ocean  
2428 sites using a moored autonomous system. *Earth Syst. Sci. Data*, **6**, 353–366.  
2429 doi:10.5194/essd-6-353-2014
- 2430 Takahashi, T., Sutherland, S. C., Sweeney, C., Poisson, A., Metzl, N., Tilbrook, B., Bates, N.,  
2431 Wanninkhof, R., Feely, R. A., Sabine, C., Olafsson, J., Nojiri, Y. (2002). Global sea–air  
2432 CO<sub>2</sub> flux based on climatological surface ocean pCO<sub>2</sub>, and seasonal biological and  
2433 temperature effects. *Deep Sea Research Part II: Topical Studies in Oceanography*, **49**, 9–  
2434 10, 1601–1622. doi:10.1016/S0967-0645(02)00003-6
- 2435 Takahashi, T., Sutherland, S. C., Wanninkhof, R., Sweeney, C., Feely, R. A., Chipman, D. W.,  
2436 Hales, B., Friederich, G., Chavez, F., Sabine, C., Watson, A., Bakker, D. C. E., Schuster,  
2437 U., Metzl, N., Yoshikawa-Inoue, H., Ishii, M., Midorikawa, T., Nojiri, Y., Körtzinger, A.,  
2438 Steinhoff, T., Hoppema, M., Olafsson, J., Arnarson, T.S., Tilbrook, B., Johannessen, T.,  
2439 Olsen, A., Bellerby, R., Wong, C. S., Delille, B., Bates, N. R. and de Baar, H. J. W.  
2440 (2009). Climatological mean and decadal change in surface ocean pCO<sub>2</sub>, and net sea–air  
2441 CO<sub>2</sub> flux over the global oceans. *Deep Sea Research Part II: Topical Studies in*  
2442 *Oceanography*, **56**, 554–577. doi:10.1016/j.dsr2.2008.12.009
- 2443 Takeshita, Y., Johnson, K. S., Coletti, L. J., Jannasch, H. W., Walz, P. M. and Warren, J. K.  
2444 (2020). Assessment of pH dependent errors in spectrophotometric pH measurements of  
2445 seawater. *Marine Chemistry*, **223**, 103801. doi:10.1016/j.marchem.2020.103801
- 2446 Takeshita, Y., Johnson, K. S., Martz, T. R., Plant, J. N. and Sarmiento, J. L. (2018). Assessment  
2447 of autonomous pH measurements for determining surface seawater partial pressure of  
2448 CO<sub>2</sub>. *Journal of Geophysical Research: Oceans*, **123**, 4003–4013.  
2449 doi:10.1029/2017jc013387
- 2450 Tanhua, T., van Heuven, S., Key, R. M., Velo, A., Olsen, A. and Schirnick, C. (2010). Quality  
2451 control procedures and methods of the CARINA database. *Earth Syst. Sci. Data* **2**, 35–49.  
2452 doi:10.5194/essd-2-35-2010
- 2453 Tanhua, T. and Keeling, R. F. (2012). Changes in column inventories of carbon and oxygen in  
2454 the Atlantic Ocean. *Biogeosciences*, **9**, 4819–4833. doi:10.5194/bg-9-4819-2012
- 2455 van der Werf, G. R., Randerson, J. T., Giglio, L., van Leeuwen, T. T., Chen, Y., Rogers, B. M.,  
2456 Mu, M., van Marle, M. J. E., Morton, D. C., Collatz, G. J., Yokelson, R. J. and  
2457 Kasibhatla, P. S. (2017). Global fire emissions estimates during 1997–2016. *Earth Syst.*  
2458 *Sci. Data*, **9**, 697–720. doi:10.5194/essd-9-697-2017
- 2459 Verdy, A. Mazloff, M. R. (2017). A data assimilating model for estimating Southern Ocean  
2460 biogeochemistry. *Journal of Geophysical Research: Oceans*, **122**, 6968–6988.  
2461 doi:10.1002/2016JC012650
- 2462 Wang, S., Zhang, Y., Hakkarainen, J., Ju, W., Liu, Y., Jiang, F. and He, W. (2018).  
2463 Distinguishing anthropogenic CO<sub>2</sub> emissions from different energy intensive industrial

- 2464 sources using OCO-2 observations: A case study in northern China. *Journal of*  
2465 *Geophysical Research: Atmospheres*, **123**, 9462–9473. doi:10.1029/2018JD029005
- 2466 Wang, S., Zhang, Y., Ju, W., Chen, J. M., Ciais, P., Cescatti, A., Sardans, J., Janssens, I. A., Wu,  
2467 M., Berry, J. A., Campbell, E., Fernandez-Martinez, M., Alkama, R., Sitch, S.,  
2468 Friedlingstein, P., Smith, W. K., Yuan, W., He, W., Lombardozzi, D., Kautz, M., Zhu,  
2469 D., Lienert, S., Kato, E., Poulter, B., Sanders, T. G. M., Kruger, I., Wang, R., Zeng, N.,  
2470 Tian, H., Vulchard, N., Jian, A., K., Wiltshire, A., Haverd, V., Goll, D. S. and Penuelas,  
2471 J. (2020). Recent global decline of CO<sub>2</sub> fertilization effects on vegetation photosynthesis.  
2472 *Science*, **370**, 1295–1300. doi: 10.1126/science.abb7772
- 2473 Watson, A. J., Schuster, U., Shutler, J. D., Holding, T., Ashton, I. G. C., Landschützer, P. ,  
2474 Woolf, D. K. and Goddijn-Murphy, L. (2020). Revised estimates of ocean-atmosphere  
2475 CO<sub>2</sub> flux are consistent with ocean carbon inventory. *Nature Communications*, **11**, 4422.  
2476 doi:10.1038/s41467-020-18203-3
- 2477 Waugh, D. W., Hall, T. M., McNeil, B. I., Key, R. and Matear, R. J. (2006). Anthropogenic CO<sub>2</sub>  
2478 in the oceans estimated using transit-time distributions. *Tellus*, **58B**, 376–389.  
2479 doi:10.1111/j.1600-0889.2006.00222.x
- 2480 Welp, L., Keeling, R. and Meijer, H. A. J. (2011). Interannual variability in the oxygen isotopes  
2481 of atmospheric CO<sub>2</sub> driven by El Niño. *Nature*, **477**, 579–582. doi:10.1038/nature10421
- 2482 Wigneron, J.-P., Fan, L., Ciais, P., Bastos, A., Brandt, M., Chave, J., Saatchi, S., Baccini, A. and  
2483 Fensholt, R. (2020). Tropical forests did not recover from the strong 2015–2016 El Niño  
2484 event. *Science Advances*, **6**, eaay4603. doi:10.1126/sciadv.aay4603
- 2485 Williams, N. L., Juranek, L. W., Feely, R. A., Johnson, K. S., Sarmiento, J. L., Talley, L. D.,  
2486 Dickson, A. G., Gray, A. R., Wanninkhof, R., Russell, J. L., Riser, S. C. and Takeshita,  
2487 Y. (2017). Calculating surface ocean pCO<sub>2</sub> from biogeochemical Argo floats equipped  
2488 with pH: An uncertainty analysis. *Global Biogeochemical Cycles*, **31**, 591–604.  
2489 doi:10.1002/2016GB005541
- 2490 Willey, J. D., Kieber, R. J., Eyman, M. S. and Avery Jr., G. B. (2000). Rainwater dissolved  
2491 organic carbon: Concentrations and global flux. *Global Biogeochemical Cycles*, **14**, 139-  
2492 148. doi: 10.1007/s11270-008-9774-0
- 2493 Woolf, D. K., Shutler, J. D., Goddijn-Murphy, L., Watson, A. J., Chapron, B., Nightingale, P. D.,  
2494 Donlon, C. J., Piskozub, J., Yelland, M. J., Ashton, I., Holding, T., Schuster, U., Girard-  
2495 Ardhuin, F., Grouazel, A., Piolle, J.-F., Warren, M., Wrobel-Niedzwiecka, I., Land, P. E.,  
2496 Torres, R., Prytherch, J., Moat, B., Hanafin, J., Ardhuin, F. and Paul, F. (2019). Key  
2497 uncertainties in the recent air-sea flux of CO<sub>2</sub>. *Global Biogeochemical Cycles*, **33**, 1548-  
2498 1563. doi:10.1029/2018GB006041
- 2499 Wu, D., Lin, J. C., Fasoli, B., Oda, T., Ye, X., Lauvaux, T., Yang, E. G. and Kort, E. A. (2018). A  
2500 Lagrangian approach towards extracting signals of urban CO<sub>2</sub> emissions from satellite  
2501 observations of atmospheric column CO<sub>2</sub> (XCO<sub>2</sub>): X-Stochastic Time-Inverted Lagrangian  
2502 Transport model (“X-STILT v1”). *Geoscientific Model Development*, **11**(12): 4843–4871.  
2503 doi:10.5194/gmd-11-4843-2018
- 2504 Wu, D. E., Lin, J. C., Oda, T. and Kort, E. A. (2020). Space-based quantification of per capita  
2505 CO<sub>2</sub> emissions from cities, *Environmental Research Letters*, **15**, 035004. doi:10.1088/1748-  
2506 9326/ab68eb

- 2507 Wunch, D., Wennberg, P. O., Osterman, G., Fisher, B., Naylor, B., Roehl, C. M., O'Dell, C.,  
2508 Mandrake, L., Viatte, C., Griffith, D. W., Deutscher, N. M., Velasco, V. A., Notholt, J.,  
2509 Warneke, T., Petri, C., De Maziere, M., Sha, M. K., Sussmann, R., Rettinger, M., Pollard,  
2510 D., Robinson, J., Morino, I., Uchino, O., Hase, F., Blumenstock, T., Kiel, M., Feist, D.  
2511 G., Arnold, S. G., Strong, K., Mendonca, J., Kivi, R., Heikkinen, P., Iraci, L., Podolske,  
2512 J., Hillyard, P. W., Kawakami, S., Dubey, M. K., Parker, H. A., Sepulveda, E.,  
2513 Rodriguez, O. E. G., Te, Y., Jeseck, P., Gunson, M. R., Crisp, D. and Eldering, A. (2017).  
2514 Comparisons of the Orbiting Carbon Observatory-2 (OCO-2) XCO<sub>2</sub> measurements with  
2515 TCCON, *Atmospheric Measurement Techniques*, **10**, 2209–2238. doi:10.5194/amt-10-  
2516 2209-2017
- 2517 Xiao, J., J. Chen, Davis, K. J. and Reichstein, M. (2012). Advances in upscaling of eddy  
2518 covariance measurements of carbon and water fluxes. *Journal Geophysical Research:*  
2519 *Biogeosciences*, **117**, G00J01. doi:10.1029/2011JG001889
- 2520 Xiao, J., Chevallier, F., Gomez, C., Guanter, L., Hicke, J. A., Huete, A. R., Ichii, K., Ni, W.,  
2521 Pang, Y., Rahman, A. F. Sun, G., Yuan, W., Zhang, L. and Zhang, X. (2019). Remote  
2522 sensing of the terrestrial carbon cycle: A review of advances over 50 years. *Remote*  
2523 *Sensing of Environment*, 233, 111383. doi:10.1016/j.rse.2019.111383
- 2524 Yin, Y., Byrne, B., Liu, J., Wennberg, P., Davis, K. J., Magney, T., Koehler, P., He, L., Jeyaram,  
2525 R., Humphrey, V., Gerken, T., Feng, S., Digangi, J. P. and Frankenberg, C. (2020).  
2526 Cropland carbon uptake delayed and reduced by 2019 Midwest floods. *AGU Advances*, **1**,  
2527 e2019AV000140. doi:10.1029/2019AV000140
- 2528 Zavarisky, A. and Marandino, C. A. (2019). The influence of transformed Reynolds number  
2529 suppression on gas transfer parameterizations and global DMS and CO<sub>2</sub> fluxes.  
2530 *Atmospheric Chemistry and Physics*, **19**, 1819-1834. doi:10.5194/acp-19-1819-2019
- 2531 Zeebe, R. and Wolf-Gladrow, D. (2001). CO<sub>2</sub> in seawater: Equilibrium, kinetics, isotopes.  
2532 *Elsevier Oceanogr. Ser.*, **65**, Elsevier, Amsterdam.
- 2533 Zeng, N., Zhao, F., Collatz, G. J., Kalnay, E., Salawitch, R., West, T. O. and Guanter, L. (2014).  
2534 Agricultural Green Revolution as a driver of increasing atmospheric CO<sub>2</sub> seasonal  
2535 amplitude. *Nature*, **515**, 394–397. doi: 10.1038/nature13893
- 2536 Zickfeld, K., MacDougall, A. H., and Matthews, H. D. (2016). On the proportionality between  
2537 global temperature change and cumulative CO<sub>2</sub> emissions during periods of net negative  
2538 CO<sub>2</sub> emissions. *Environmental Research Letters*, **11**(5), 055006, doi:10.1088/1748-  
2539 9326/11/5/055006