Impacts of urban carbon dioxide emissions on sea-air flux and ocean acidification in nearshore waters

Devon Northcott\(^1\)*, Jeff Sevadjian\(^1,2\), Diego A. Sancho-Gallegos\(^1,3\), Chris Wahl\(^1\), Jules Friederich\(^1\), Francisco P. Chavez\(^1\)*

1 Monterey Bay Aquarium Research Institute, Moss Landing, CA, United States of America, 2 Scripps Institution of Oceanography, La Jolla, CA, United States of America, 3 Stanford University, Stanford, CA, United States of America

* chfr@mbari.org (FPC); dnorthcott@mbari.org (DN)

Abstract

Greatly enhanced atmospheric carbon dioxide (CO\(_2\)) levels relative to well-mixed marine air are observed during periods of offshore winds at coastal sensor platforms in Monterey Bay, California, USA. The highest concentrations originate from urban and agricultural areas, are driven by diurnal winds, and peak in the early morning. These enhanced atmospheric levels can be detected across a ~100km wide nearshore area and represent a significant addition to total oceanic CO\(_2\) uptake. A global estimate puts the added sea-air flux of CO\(_2\) from these greatly enhanced atmospheric CO\(_2\) levels at 25 million tonnes, roughly 1% of the ocean’s annual CO\(_2\) uptake. The increased uptake over the 100 km coastal swath is of order 20%, indicating a potentially large impact on ocean acidification in productive coastal waters.

Introduction

The increase in atmospheric carbon dioxide (CO\(_2\)) from the burning of fossil fuels has been well documented by decades of measurements from the top of Mauna Loa on the big island of Hawaii [1]. Keeling chose the iconic Mauna Loa site because it rises into the free troposphere that is less affected by local sources of carbon pollution. Oceanographers often use well-mixed atmospheric values from similar sites or global models to estimate the sea-air exchange of CO\(_2\) [2]. Global flux estimates find the current ocean uptake to be about two petagrams (two billion tonnes) of carbon per year [2,3]. This steady uptake of atmospheric CO\(_2\) by the oceans results in the so-called phenomenon of ocean acidification [4]. However, studies of atmospheric CO\(_2\) concentrations in urban environments have shown considerable enhancements of CO\(_2\) in city centers, especially in the early morning [5,6,7], an effect known as the urban CO\(_2\) dome. Agricultural practices can also impact local atmospheric CO\(_2\) on a diurnal cycle with large nighttime increases due to respiration and daytime decreases associated with photosynthesis [8,9]. Near coastlines these elevated levels of CO\(_2\) might impact marine air via atmospheric circulation and therefore increase the flux of CO\(_2\) into nearshore waters enhancing ocean acidification. Here we present novel observations from nearshore moorings and an autonomous sea surface vehicle that show the magnitude of urban CO\(_2\) pollution and allow us to calculate the contribution to sea-air fluxes from this previously unquantified source.
The diurnal cycles in CO\textsubscript{2} concentration over urban environments has been shown to peak just before sunrise at 4-5am and reach a minimum at around 4pm in the Los Angeles basin [7]. Modeling has shown similar CO\textsubscript{2} dome effects over the San Francisco and Monterey Bay areas [10]. However, little attention has been paid to the advection of these urban CO\textsubscript{2} domes over oceans and resulting impact on sea-air CO\textsubscript{2} flux. Monterey Bay is ideally situated for such observations, as large urban and agricultural areas in the Salinas and Silicon valleys are nearby, and a strong diurnally varying component in winds [11,12] can transport high levels of locally produced atmospheric CO\textsubscript{2} over the ocean. In the Monterey Bay Area the urban and agricultural CO\textsubscript{2} dome should reach its maximum concentrations at roughly the same time as the peak of the offshore phase of the diurnal wind cycle, leading to the advection of high CO\textsubscript{2} air from land sources over the coastal ocean.

As indicated above, traditional estimates of sea-air CO\textsubscript{2} flux are not able to quantify this nearshore phenomenon because they have a temporal or spatial scale that is too coarse to resolve these diurnally varying CO\textsubscript{2} anomalies. Therefore, the impact of a potentially significant source of atmospheric CO\textsubscript{2} on fluxes into the ocean has not previously been estimated. These impacts are magnified near urban or agricultural areas with strong offshore winds that can advect heavily polluted air over marine waters. Here we use high temporal resolution (1 hour over years) timeseries from multiple autonomous ocean based sensor platforms (moorings and surface vehicles) to provide a detailed assessment of the impacts of these high frequency variations in atmospheric CO\textsubscript{2} concentration on sea-air CO\textsubscript{2} fluxes in the nearshore environment.

**Material and methods**

**Data sources**

Data was collected from four different platforms operated by the Monterey Bay Aquarium Research Institute (MBARI) in the Monterey Bay region. Of these, three were moorings (OA1, OA2 and M1; Fig 1A) and one was an autonomous surface vehicle (Liquid Robotics Wave Glider [13]). All except M1 were outfitted with Airmar WX200 ultrasonic wind sensors which record wind speed and direction, Licor non-dispersive infrared gas analyzer CO\textsubscript{2} instruments which were developed to measure CO\textsubscript{2} concentrations in the air and water [13,14], and Seabird temperature and salinity sensors (herein CTD). M1 used an Aandara sonic anemometer together with the same CO\textsubscript{2} instrument and CTD. Measurements were averaged hourly in the final analysis. Wave glider measurements were taken within 5km of the Monterey Bay Time Series (MBTS) Line, a transect extending from Moss Landing out 50km along the Monterey canyon (Fig 1, red line). The moorings are located 1.5km offshore of At\’o Nuevo north of Santa Cruz, California (OA2), 300m off shore of Monterey, California (OA1), and 20km offshore of Moss Landing, CA, in the center of the Monterey Bay (M1). The OA moorings are at 20 m and M1 at 1000 m depth. Wave glider data from station M, a station 220km offshore of San Luis Obispo, California (123 W 35.14 N) were also considered. Data were taken between 2013 and 2018, and all records were used, except in cases of dropout of CO\textsubscript{2} sensors, wind measurements, or CTD sensors. M1 recorded 1417 days of data between September 2013 and April 2018, OA1 recorded 853 days between January 2014 and July 2017, OA2 sampled 461 days between May 2015 and July 2017, and the wave glider recorded 182 days of measurements on the MBTS Line between March 2014 and March 2018.

**Winds**

An analysis of the phasing and length of offshore wind events was performed using the full 29 year M1 wind record. The east-west component of the winds was isolated, and binned by hour (Fig 2A). This analysis was run on both the full record, and by month to examine seasonal variations in the phasing and strength of land-sea breezes. While the strength of offshore winds
varied by month the phasing of the cycle did not. The number of hours of offshore wind (wind direction between 0 and 165 degrees) on each fully sampled day in the record was then calculated, and these results were binned by month (Fig 2B) to give a measure of the seasonality of the persistence of offshore wind.

Atmospheric CO\textsubscript{2} anomaly calculations

In this paper “atmospheric CO\textsubscript{2} anomaly” refers to a deviation of atmospheric CO\textsubscript{2} values from a baseline CO\textsubscript{2} concentration that reflects a well-mixed atmosphere. To calculate these baseline values, a rolling median filter was applied to CO\textsubscript{2} measurements with duration of 15 days for moorings and 4 days for the wave glider. The filter window for the wave glider data was used because the short mission durations and high mobility of the platform lead to poor estimates of baseline pCO\textsubscript{2} values when the full 15 day filter window was applied. CO\textsubscript{2} measurements taken when the wind was blowing from the open ocean for more than 3.5 hours were used to calculate baseline values to ensure that baselines were representative of well-mixed marine air CO\textsubscript{2} concentrations. This baseline was then subtracted from the timeseries of atmospheric CO\textsubscript{2} values to yield a timeseries of atmospheric CO\textsubscript{2} anomalies. Comparison of the baseline pCO\textsubscript{2} values with NOAA’s Global Greenhouse Gas (GHG) Reference Network [15] Trinidad Head station (r\textsuperscript{2} = 0.71), as well as CarbonTracker [16] modeled values (r\textsuperscript{2} = 0.56), showed good agreement. We used the calculated baseline pCO\textsubscript{2} values since these sources only run until December 2016 while our timeseries extends through early 2018.

CO\textsubscript{2} Fluxes

The net sea-air CO\textsubscript{2} flux was estimated using the equation:

$$ F_{CO_2} = k \times S \times \Delta p_{CO_2} $$ (1)

Fig 1. Location of data sources and predicted terrestrial sources of atmospheric CO\textsubscript{2} from wave glider data. a) Topography of the Monterey Bay area showing the location of platforms used in this study, as well as the Hecker pass and Salinas Valley, two major terrestrial sources of enhanced atmospheric CO\textsubscript{2} to Monterey Bay waters. The Monterey Bay timeseries (MBTS) line is also shown. b) Source regions for atmospheric CO\textsubscript{2} anomalies (the difference between well-mixed marine air and enhanced atmospheric CO\textsubscript{2}; see “Anomaly Calculations” section in Methods) traced back from wave glider measurements along the MBTS Line (Fig 1A). Anomalies are particularly high in the Salinas Valley and Hecker pass areas. Anomalies are calculated for wave glider observations within 5km of the MBTS Line, and then propagated backwards for 6 hours along a path defined by wind direction and speed at the wave glider. A 5 degree uncertainty cone is drawn around the path, and all paths are averaged to yield the final figure. Coastline data republished from the Global Self-consistent, Hierarchical, High-resolution Geography Database (GSHHG) under a CC BY license, with permission from Dr. Paul Wessel, original copyright 1996.

https://doi.org/10.1371/journal.pone.0214403.g001
where $k$ is the gas transfer velocity\textsuperscript{[17]}, $S$ is the solubility of CO\textsubscript{2} in seawater \textsuperscript{[18]} and $\Delta pCO_{2}$ is the difference between $pCO_{2}$ water and $pCO_{2}$ air. Gas transfer velocity is parameterized as a function of wind speed squared, while solubility is a function of water temperature and salinity. The common convention is used whereby a negative flux indicates CO\textsubscript{2} transfer into the ocean (a sink); while a positive flux indicates release of CO\textsubscript{2} into the atmosphere (a source).

**Wave glider atmospheric CO\textsubscript{2} anomaly predictions**

In order to construct a high resolution map of enhanced atmospheric CO\textsubscript{2} sources, anomalies were calculated for each wave glider data point. At ten minute intervals the wind speed and direction were then used to calculate the source position of the wind; this process was repeated additively over the six hours prior to any given air CO\textsubscript{2} measurement to construct a probable path for the measured air parcel. This method ignores local variation in wind direction and speed, mixing, and in particular excludes the effects of topography on offshore winds. In order to reduce these effects, all pixels within a five degree cone around the calculated path were assigned the value of the calculated anomaly, and added to the developing composite image containing all previous tracks. When complete the image was then divided by the number of tracks intersecting each pixel, producing an average map of probable atmospheric CO\textsubscript{2} anomaly sources and their strengths. This method should not be seen as providing exact locations or intensities of anomaly sources but it is straightforward and provides a general picture of the origin of strong atmospheric CO\textsubscript{2} anomalies.
Fluxes driven by atmospheric CO$_2$ anomalies

To estimate the contributions of the atmospheric CO$_2$ anomalies to sea-air fluxes, two flux calculations were made. The first used the well-mixed oceanic air baseline CO$_2$ concentrations calculated as described above. This dataset represents atmospheric CO$_2$ concentrations used in studies of sea-air flux that rely on modeled or well-mixed atmospheric CO$_2$ values. This timeseries of fluxes was subtracted from a second timeseries calculated using the observed pCO$_2$ air concentrations. The difference between these two timeseries gives a measurement of flux due to our calculated atmospheric anomalies in air pCO$_2$. This method preserves the convention of negative values indicating increased transport of CO$_2$ into the ocean.

Results

Significant positive anomalies in atmospheric CO$_2$ are detected on all platforms during periods of offshore winds. A time series of atmospheric CO$_2$ from the OA1 mooring over 2014 and 2015 illustrates the extent of these anomalies (Fig 3). In Fig 3 a timeseries of atmospheric CO$_2$
from a nearby urban terrestrial station (Walnut Creek) [16] and clean marine air from Carbon Tracker [15] are plotted together with OA1. The similarity between OA1 and Walnut Creek is striking as are the mostly winter time increases in atmospheric CO$_2$ at both these sites relative to well-mixed marine air.

Using a basic advection model in conjunction with wave glider data (see methods) we found that the largest atmospheric CO$_2$ anomalies detected along the MBTS line originate from the Salinas valley and Hecker Pass (Fig 1B). These topographic features connect marine waters in the Monterey Bay region with large urban or agricultural areas inland and suggest that topography steers air with enhanced levels of CO$_2$ toward Monterey Bay. Hecker Pass represents a break in the mountains through which pollution from Silicon Valley can reach the coast, while the Salinas Valley contains urban centers and large agricultural fields. Binning atmospheric CO$_2$ anomalies measured at moorings by wind direction confirms sources at those locations. (Fig 4). The M1 mooring, which is situated at the center of the MBTS Line, shows a double peak in atmospheric CO$_2$ anomalies corresponding Hecker Pass (NE, 60 degrees) and the Salinas Valley (E, 100 degrees). OA1 is also impacted by CO$_2$ anomalies during periods of easterly winds from these topographic features. However, its largest anomalies are registered when winds blow from ~150 degrees. OA1 is situated only a few hundred meters north of the city of Monterey and the monthly averages at this buoy were much better correlated with the urban CO$_2$ measurements from Walnut Creek ($r^2 = 0.78$), than with Carbon-Tracker modeled atmospheric CO$_2$ ($r^2 = 0.47$) (Fig 3). This suggests that atmospheric CO$_2$ at OA1 is strongly influenced by air emanating from the city of Monterey. Meanwhile the largest atmospheric CO$_2$ anomalies at OA2 off Ano Nuevo came from the northeast, directly from the Silicon Valley area (Fig 4).

Average anomalies at OA1 were close to double those at M1 during periods of offshore winds. This indicates a reduction in the strength of atmospheric CO$_2$ anomalies with distance from shore. The wave glider provides the perfect platform to further explore this relationship. Frequent wave glider measurements extend 50km out to the end of the MBTS Line, and average anomalies of 6-10ppm are detected at the end of this line during periods of offshore winds. This shows that during offshore wind events the plume of high CO$_2$ air extends at least 50km from shore, although its CO$_2$ content is reduced on average from the 15 ppm anomalies seen...
nearshore (Fig 5). An outer limit is seen on anomaly propagation when data from station M is considered. This station is 220km offshore and routinely occupied by the wave glider. No
significant atmospheric CO\textsubscript{2} anomalies associated with offshore wind events are detected at this location, with all reported anomalies being within the uncertainty of the sensor. Since enhanced CO\textsubscript{2} levels are transported offshore by diurnal land-sea breezes, the area in which these high CO\textsubscript{2} levels can be observed depends on the areal extent of the land-sea breezes. The strength of offshore winds in central California varies diurnally [11,12], and the average duration of an offshore wind event at M1 is relatively short at ~6 hours. Average offshore wind speeds were 4.4 m/s, so assuming CO\textsubscript{2} is transported offshore at that wind speed for the duration of an offshore wind event, anomalies should rarely be detected more than 100km from the coast. This agrees with more thorough modeling and remote sensing studies that have concluded that the seaward influence of diurnal land breezes extends roughly 100km offshore along much of the west coast of the United States [19,20].

Total sea-air fluxes varied widely across the study area, largely owing to upwelling processes. OA2 sits directly in the upwelling plume off Año Nuevo where water high in carbon dioxide is brought to the surface during upwelling [13,21,22]. As a result, CO\textsubscript{2} flux is large and positive in recently upwelled water indicating that freshly upwelled waters are a considerable source of atmospheric CO\textsubscript{2}. M1 is downstream from the upwelling plume, in an area where phytoplankton have increased and via photosynthesis converted much of the upwelled carbon dioxide into organic carbon. As a result average fluxes are slightly negative in this area, indicating that on an annual basis M1 is a weak sink for carbon. Inside Monterey Bay and along the southern edge fluxes are increasingly negative as the slower circulation allows phytoplankton to bloom and further reduce pCO\textsubscript{2}. OA1 and the MBTS Line average, which is weighted to the inshore environment owing to uneven sampling effort, reflect this greater sink of atmospheric CO\textsubscript{2} (Fig 6).

Sea-air CO\textsubscript{2} fluxes due to enhanced atmospheric CO\textsubscript{2} were similar across all platforms. The highest atmospheric CO\textsubscript{2} anomaly driven fluxes were recorded at M1, reaching 0.046 mols C m\textsuperscript{-2} yr\textsuperscript{-1}. While atmospheric CO\textsubscript{2} anomalies were higher at OA1 owing to this mooring’s close proximity to land and urban areas, a lower average offshore wind speed and a lower incidence of offshore winds reduced the effects of the higher atmospheric CO\textsubscript{2} on fluxes, resulting in an anomaly driven flux of 0.041 mols C m\textsuperscript{-2} yr\textsuperscript{-1}. The total annual flux along the MBTS Line was 0.030 mols C m\textsuperscript{-2} yr\textsuperscript{-1}. (Fig 6) Average fluxes were reduced over the whole line relative to M1 owing to very low wind speeds close to shore and a drop-off in anomalies beyond 35km from shore (Fig 5). OA2 recorded the lowest average anomalies, as well as the lowest percentage of offshore wind hours, but still displayed wind driven CO\textsubscript{2} flux anomalies of 0.019 mols C m\textsuperscript{-2} yr\textsuperscript{-1} (Fig 6). The averaged offshore profile of anomaly driven sea-air fluxes was dependent on both wind speed and atmospheric CO\textsubscript{2} anomalies, and was depressed inshore where wind speeds are low, and offshore where wind speed are high but atmospheric CO\textsubscript{2} anomalies are lower. The combination of these factors leads to a large increase in sea-air fluxes between 20 and 30 km offshore, where relatively stronger offshore winds combine with high atmospheric CO\textsubscript{2} anomalies (Fig 5).

Sea-air CO\textsubscript{2} fluxes due to atmospheric CO\textsubscript{2} anomalies undergo a strong seasonal cycle with increased offshore wind duration (Fig 2B) in the winter months driving larger and more frequent anomalies at all platforms. OA1 exhibits a particularly strong seasonal cycle, as strong wintertime storm winds are oriented to bring polluted air directly from the city of Monterey onto the buoy (Fig 7). At M1, where southerly storm winds come off the open ocean, a strong seasonal cycle is still present due to higher incidence of easterly and southeasterly offshores during the winter season. (Fig 2B)

**Discussion**

Studies of the role of the oceans in the global CO\textsubscript{2} budget often use low-resolution information on atmospheric CO\textsubscript{2} concentration to estimate sea-air CO\textsubscript{2} fluxes. Large scale models or
relatively few stations representative of well-mixed marine air concentrations are consistently used in flux calculations [2,23]. Observations from autonomous platforms in coastal California show that these clean air open ocean carbon dioxide concentrations are often not representative of carbon dioxide concentrations in the air nearshore. Nearshore areas experience
significant enhancement from land sources and this impact can extend on the order of 100km from the coast. These atmospheric CO\textsubscript{2} anomalies, relative to well-mixed atmospheric air, contribute to sea-air fluxes regardless of the underlying oceanographic and atmospheric conditions. The association of diurnal winds to these atmospheric anomalies clearly ties them to terrestrial sources. The high frequency variability also makes it difficult to model these terrestrial sources from sparse temporal or spatial resolution datasets. High-resolution

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**Fig 7. OA1 anomaly driven flux by month and wind direction.** Atmospheric CO\textsubscript{2} anomaly driven sea-air flux averaged by month (a) and by wind direction (b) at OA1. Negative values indicate CO\textsubscript{2} flux into the ocean. Atmospheric CO\textsubscript{2} anomaly driven fluxes are strongest in the winter months when offshore winds are stronger and more frequent. Anomaly driven fluxes are restricted to periods of offshore winds.

https://doi.org/10.1371/journal.pone.0214403.g007
measurements from autonomous platforms allow us to make an initial estimate of the impacts of elevated atmospheric CO2 levels from highly polluted terrestrial sources on sea-air fluxes.

The combination of offshore winds and terrestrial carbon sources drive large gradients in sea-air CO2. However, the relatively weak offshore winds in the Monterey Bay greatly constrain CO2 flux since it is parameterized to approach zero under very low wind speeds. This is a reasonable approximation in the open ocean where average wind speeds are rarely very low; the current parameterization may not apply when wind speeds are less than 3m/s, seriously underestimating sea-air flux at low wind speeds [17]. Average offshore wind speeds recorded along the Monterey Bay time series line never exceed 7m/s, and nearshore where CO2 anomalies are largest wind speeds were around 3m/s. While open ocean fluxes may be accurately predicted using current sea-air flux models, our observations highlight the difficulty of using these same models in the nearshore realm, where wind speeds are lower.

The atmospheric CO2 anomalies described above should occur anywhere that urban or agricultural areas are found near the coast, and the winds are such that they carry terrestrial sources over the marine environment. The diurnal land-sea breezes that contribute to offshore CO2 transport observed in Monterey Bay are a widespread phenomenon worldwide [20]. While the contributions of direct CO2 pollution to sea-air fluxes is modest, areas with stronger or more frequent offshore wind events, or larger urban centers situated closer to the water’s edge should experience more significant fluxes. If the wave glider measurements in the Monterey Bay area are taken as a reasonable estimate of average anomaly driven CO2 fluxes within 100km of land we can estimate the impact of increased atmospheric CO2 globally. We calculated that coastal waters occupy 2.6*10^7 km^2 or about 7% of the global ocean so the added contribution to CO2 fluxes could be order of 25 million tonnes, or roughly 1% of the ocean’s total estimated CO2 uptake[2]. However, since this uptake is concentrated over the 100 km coastal swath the estimated sea-air flux into the ocean in these regions should be increased by around 20%.

Clearly, this additional uptake of anthropogenic CO2 will have consequences for ocean acidification [4] in nearshore regions where marine biota is concentrated. These same processes should also drive enhanced transport of other terrestrially emitted gases, aerosols, and particles over nearshore marine waters. It is therefore likely that other pollutants are also entering nearshore waters at increased rates. How these enhanced urban sources of pollution will change over time and what are their ecological impacts will need to be assessed by future studies.

Supporting information

S1 Dataset. pCO2 Dataset. Dataset containing pCO2 air and water measurements, as well as wind speed and direction, sea surface temperature, and salinity at all platforms.

(MAT)

Acknowledgments

CarbonTracker CT2016 results provided by NOAA ESRL, Boulder, Colorado, USA from the website at http://carbontracker.noaa.gov. Elevation data is from the GTOP030 dataset. This Data is available from the U.S. Geological Survey. Coastline data used is from the freely available Global Self-consistent, Hierarchical, High-resolution Geography Database (GSHHG).

This research was supported by the David and Lucile Packard Foundation and the National Science Foundation.

Author Contributions

Conceptualization: Diego A. Sancho-Gallegos, Francisco P. Chavez.
Data curation: Devon Northcott, Jeff Sevadjian, Chris Wahl, Jules Friederich.

Formal analysis: Devon Northcott, Diego A. Sancho-Gallegos.

Funding acquisition: Francisco P. Chavez.

Investigation: Diego A. Sancho-Gallegos.

Methodology: Devon Northcott, Jeff Sevadjian, Chris Wahl, Jules Friederich.

Project administration: Francisco P. Chavez.

Resources: Chris Wahl.

Software: Devon Northcott, Jeff Sevadjian.

Supervision: Francisco P. Chavez.

Visualization: Devon Northcott.

Writing – original draft: Devon Northcott.

Writing – review & editing: Francisco P. Chavez.

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