Drivers of CO$_2$ along a mangrove-seagrass transect in a tropical bay: delayed groundwater seepage and seagrass uptake.

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Abstract

Water-to-air carbon dioxide fluxes from tropical coastal waters are an important but understudied component of the marine carbon budget. Here, we investigate drivers of carbon dioxide partial pressure (\(p\text{CO}_2\)) in a relatively pristine mangrove-seagrass embayment on a tropical island (Bali, Indonesia). Observations were performed over eight underway seasonal surveys and a fixed location time series for 55 hours. There was a large spatial variability of \(p\text{CO}_2\) across the continuum of mangrove forests, seagrass meadows and the coastal ocean. Overall, the embayment waters surrounded by mangroves released \(\text{CO}_2\) to the atmosphere with a net flux rate of 18.1 ± 5.8 mmol m\(^{-2}\) d\(^{-1}\). Seagrass beds produced an overall \(\text{CO}_2\) net flux rate of 2.5 ± 3.4 mmol m\(^{-2}\) d\(^{-1}\), although 2 out of 8 surveys revealed a sink of \(\text{CO}_2\) in the seagrass area. The mouth of the bay where coral calcification occurs was a minor source of \(\text{CO}_2\) (0.3 ± 0.4 mmol m\(^{-2}\) d\(^{-1}\)). The overall average \(\text{CO}_2\) flux to the atmosphere along the transect was 9.8 ± 6.0 mmol m\(^{-2}\) d\(^{-1}\), or 3.6 x 10\(^3\) mol d\(^{-1}\) \(\text{CO}_2\) when upscaled to the entire embayment area. There were no clear seasonal patterns in contrast to better studied temperate systems. \(p\text{CO}_2\) significantly correlated with antecedent rainfall and the natural groundwater tracer radon (\(^{222}\text{Rn}\)) during each survey. We suggest that the \(\text{CO}_2\) source in the mangrove dominated upper bay was associated with delayed groundwater inputs, and a shifting \(\text{CO}_2\) source-sink in the lower bay was driven by the uptake of \(\text{CO}_2\) by seagrass and mixing with oceanic waters. This differs from modified landscapes where potential uptake of \(\text{CO}_2\) is weakened due to the degradation of seagrass beds, or emissions are increased due to drainage of coastal wetlands.

Keywords: blue carbon; submarine groundwater discharge; greenhouse gas emissions; mangrove forests; seagrass beds; ecosystem connectivity.
1. Introduction

The large net primary production of the world’s coastal embayments are exported to coastal waters (Robertson et al., 1992) primarily through the interplay of tidal dynamics and seasonal river discharge. With a global area of \( \sim 45000 \text{ km}^2 \), intertidal areas of temperate embayments predominantly comprise of salt marsh habitats (Greenberg et al., 2006) occupying low-lying topographic zones (Scott et al., 2014) and high distributions of seagrass beds in subtidal zones (Short et al., 2007). In contrast, tropical coastal embayments typically consist of a continuum of fringing coral reefs, seagrass beds and mangrove forests (Torres-Pulliza et al., 2013). Near-shore tropical mangrove forests are the most carbon-rich forests on earth, storing and sequestering globally significant amounts of carbon in their soils (Donato et al., 2011). Occupying only 0.02% of global surface area, mangrove forests are responsible for approximately 11% of the total terrestrial organic carbon delivery to oceans (Jennerjahn & Ittekkot, 2002; Sippo et al., 2017). Mangroves are tightly connected with their adjacent habitats (Signa et al., 2017) and support marine biodiversity, regulate water quality and protect tropical coastlines against storms (Ganguly et al., 2017).

Tropical seagrass beds are located shoreward of coral reefs and seaward of mangrove forests in areas with high light availability and favourable water quality (Guannel et al., 2016) and have been reported to be largely net autotrophic (i.e. a net atmospheric CO\(_2\) sink) (Duarte & Cebrian, 1996). Coastal geomorphology is recognised as being important in seagrass abundance, distribution and diversity, as these habitats usually exist near fringing reefs in protected, shallow coastal lagoons (Torres-Pulliza et al., 2013). Combined, mangrove forests and seagrass beds play a major role in biological connectivity of coastal embayments, acting as coastal buffers by filtering sediment and nutrient loads to adjacent coral reefs (Hemminga & Duarte, 2000).
Indonesia, lying between latitudes 6 °N and 11 °S, has a coastline of more than 95,180 km, the second longest coastline in the world (Spalding et al., 1997) and 2.9 Mha of mangrove cover, larger than any continent on earth (Atwood et al., 2017). With such an extent and high carbon stocks, Indonesia’s mangrove forests store on average 3.14 PgC (Murdiyarso et al., 2015). However, in three decades (1985-2005), Indonesia has lost 40% of its mangroves, mainly as a result of aquaculture development (Giri et al., 2011). This has resulted in potential global annual emissions of 0.07 to 0.21 Pg CO2 (Murdiyarso et al., 2015). Seagrass beds cover an estimated 30,000 km² of Indonesian coastline (Green & Short, 2003), and combined with mangrove forests, account for approximately 3.4 Pg C (~17%) of the global blue carbon reservoir (Alongi et al., 2016).

Mangrove-seagrass connectivity research has usually centred on the exchange of dissolved organic carbon (DOC) and particulate organic carbon (POC) (Dittmar et al., 2009; Hemminga et al., 1994; Maher et al., 2013; Müller et al., 2015). Little is known about CO2 interactions between near-mangrove forest surrounding waters (described as mangrove forest water from here on) and adjacent seagrass beds. Stable isotope studies show that seagrasses close to mangroves have a more depleted δ¹³C value than those further away (Bouillon, Connolly, et al., 2008; Hemminga et al., 1994), suggesting that seagrasses are fixing DIC sourced from mangrove respiration.

Mangrove groundwater and porewater exchange can be an important source of carbon to coastal waters (Bouillon et al., 2007; Maher et al., 2013; Maher et al., 2017; Sadat-Noori et al., 2016). A recent literature review demonstrates that groundwater fluxes in mangroves can be a major component of tropical coastal carbon budgets with fluxes on the same order of magnitude as rivers (Chen et al., 2018). Since mangrove forests usually coexist with seagrass beds and coral reefs (Fourqurean et al., 1992), and carbon exchange along this continuum supports cross-productivity (Unsworth et al., 2008), understanding the relationship between
groundwater seepage, carbon dynamics and ecosystem connectivity in transition zones is important.

Here, we investigate the drivers of $pCO_2$ dynamics along a mangrove-seagrass transect in Bali, Indonesia. We performed coupled, automated seasonal $pCO_2$ and radon ($^{222}$Rn; a natural groundwater tracer) investigations to assess whether CO$_2$ is derived from groundwater or porewater pathways. We investigate temporal and spatial scales of $pCO_2$ dynamics, hydrological drivers such as groundwater seepage, delayed antecedent rainfall, and interplay along the mangrove-seagrass continuum in a non-impacted embayment.

2. Material and methods

2.1 Area description

Gilimanuk Bay, a 3.7 km$^2$ coastal embayment, is located in Jembrana Regency on the northwest coast of Bali, Indonesia. Including two small islands, Kalong Island and Burung Island, the area contains some of Bali’s most pristine mangrove forests (Thoha, 2007) (Figure 1). Oceanic upwelling and tidal exchange from the deep Java Strait supply nutrients to Gilimanuk Bay (Ningsih et al., 2013; Siswanto, 2008). The average depth of the embayment is ~2m, with intertidal zones in the upper embayment unnavigable at low tides (<0.5m; Figure 1). Patchy coral reefs are present in the channel at the ocean mouth of the embayment, while the lower and upper embayment consists of seagrass beds and sand flats. Due to the low-lying topography and geometry, predominant southeast winds (June-September) and northeast winds (December-March) produce small wind-generated waves (<0.5m), which influence the mangrove forest water in the upper embayment. Microtides with an average range of ~1 m occur in the bay, preventing significant current-driven turbulence. The bordering West Bali National Park on the northern side is comprised of mangrove forests (310 ha), lowland rain forest, savanna and sea grasses (40 ha), coral reefs (810 ha) and both
shallow and deep sea waters (3,520 ha) (Utama, 2015). The regional geology includes alluvial deposits and Prapat Agung Formation which consists of limestone, calcareous sandstone and marts (Purbo-Hadiwidjojo, 1971). Soils are hydromorphic alluvial in the near-shore zone and grey-brown alluvial surrounding the embayment. Dominant mangrove species in Gilimanuk Bay include *Rhizophora apiculata*, *Excoecaria agallocha*, and *Ceriops tagal* (Marbawa et al., 2015). Twelve of the world’s sixty known species of seagrass are present in West Bali National Park (Purnomo et al., 2017). Dominant seagrass species include *Cymodocea rotundata*, *Halophila ovalis* and *Enhalus acoroides* (Purnomo et al., 2017; Zulkarnaen et al., 2014).

Figure 1. Gilimanuk Bay study site in western Bali. The A to B line and 0 to 3 along the line indicates the seasonal underway sampling route and distance travelled, respectively. The square near the mouth shows the location of the time series deployment. The location of seagrass beds and mangroves is indicated for the line A-B on the bottom right.

2.2 Approach and methods

To investigate hydrologic variations, groundwater seepage and CO$_2$ fluxes, we combined
eight underway spatial surveys (S1-S8) with a detailed 55 hour fixed location time series (TS) in Gilimanuk Bay between 20th November 2015 and 15th November 2017 (see Table 1). The average tidal range throughout the surveys was 1.3m. As S1 was conducted at the end of a subsequent drought period and followed by a period of prolonged rainfall, following surveys (S2-S8) were intended to replicate rainfall events and possible groundwater relationships.

During the underway surveys, we measured high resolution spatial variations at 10 minute intervals while location was tracked and logged by a Garmin GPS72 continuously. Each survey commenced at high tide beginning at the ocean mouth and ending upstream at the mangrove forest, in a small research vessel travelling between 4 and 6 km/hr, while the time series was conducted at the ocean mouth (Figure 1).

A Li-820 CO₂ detector and a radon-in-air monitor (RAD7, Durridge) were deployed to measure \( p\text{CO}_2 \) and \( ^{222}\text{Rn} \) concentrations at approximately 1m depth. The Li-820 and the RAD7 were connected with a closed-air-loop to a shower head gas exchange (Dulaiova et al., 2005; Santos et al., 2012) using the methodology of Santos et al. (2012) and references therein. A Li-820 (calibrated before underway surveys with 0, 400 and 10 000 ppm spans) was checked for accuracy alongside a calibrated Li-820 which was deployed in the 55 hr time series at the end of the study campaign. The RAD 7 was, pre-calibrated by the manufacturer (Durridge) and is expected to hold calibration for at least one year. Mole fraction measurements provided by the Li-820 CO₂ detector were later calculated to \( p\text{CO}_2 \) according to the recommendations of Pierrot et al. (2009). Atmospheric \( p\text{CO}_2 \) was assumed to be constant at 400 µatm. A Hydrolab DS-5 water quality sonde was calibrated before each survey and deployed to measure temperature, salinity and dissolved oxygen at 10 min intervals. We used a calibrated handheld YSI EcoSence EC300 meter and a Hatch 40D LDO Sensor for field measurements every 10 to 15 minutes, which were comparable to the Hydrolab DS-5 measurements.
CO₂ fluxes at the water-air interface were then calculated as function of air-water CO₂ gradient (ΔpCO₂), temperature and salinity dependant solubility (k₀ (Weiss, 1974)) and gas transfer velocity (k) according to:

\[ F_{CO₂} = k \cdot k₀ \cdot ΔpCO₂, \]

where \( k \) is the CO₂ gas transfer velocity, \( K₀ \) is the solubility of CO₂ (Weiss, 1974) and \( ΔpCO₂ \) is the difference between sea and air (\( pCO₂_{sea} - pCO₂_{air} \)).

To calculate \( k \), we used four wind-speed based parameterizations which provide a reasonable range in evasion rate estimates.

\[ k = 0.0283u^3 (\text{Sc/660})^{-1/2} \]

Wanninkhof and McGillis (1999) (2)
\[ k = 1.91 \text{e}^{0.35u} \left( \frac{\text{Sc}}{600} \right)^{1/2} \]

Raymond and Cole (2001) (3)

\[ k = 5.141u^{0.758} \left( \frac{\text{Sc}}{600} \right)^{1/2} \]

Borges et al. (2004) (4)

\[ k = 0.251u^2 \left( \frac{\text{Sc}}{660} \right)^{1/2} \]

Wanninkhof (2014) (5)

where \( k \) is the transfer velocity (cm h\(^{-1}\)), \( u \) is the wind speed (ms\(^{-1}\)) at a height of 10m and \( \text{Sc} \) is the Schmidt number of CO\(_2\) at in situ temperature and salinity (Wanninkhof, 2014).

Positive CO\(_2\) flux values signify a CO\(_2\) exchange from water-to-air (CO\(_2\) source) while negative CO\(_2\) flux values signify an exchange from air-to-water (CO\(_2\) sink). Estimations of CO\(_2\) fluxes were calculated by using 10 minute sampling times for both underway and time series measured \( p\text{CO}_2 \) data and average wind speeds from 28 days prior to each survey and the time series (Table 1). Rainfall and wind data were sourced from Banyuwangi Weather Station (8.21700°S, 114.38300°E; average atmospheric pressure=1021.5 hPa (1.0 atm); 10 m above sea level) located 7 km from the study site (www.dataonline.bmkg.go.id).

Table 1. Date, start and finish times (24hrs), tidal ranges (m), windspeeds (m sec\(^{-1}\)) 24hrs, 48hrs, 7 days, 14 days and 28 days prior, for each survey (S1-S8) and the time series (TS) at the ocean mouth of the embayment.

| Survey | Date    | Start (24 hrs) | Finish (24 hrs) | Tidal range (m) | 24 hrs | 48 hrs | 7 days | 14 days | 28 days |
|--------|---------|----------------|-----------------|-----------------|--------|--------|--------|---------|---------|
| S1     | 20/11/2105 | 15:10          | 18:00           | 1.1             | 2.2    | 2.2    | 2      | 1.8     | 2       |
| S2     | 9/01/2016  | 9:58           | 13:50           | 0.9             | 1.4    | 1.1    | 1.3    | 1.3     | 1.6     |
| S3     | 30/06/2016 | 6:45           | 10:25           | 1.4             | 1.3    | 1.2    | 1.3    | 1.3     | 1.3     |
| S4     | 14/07/2016 | 8:25           | 12:40           | 1.3             | 1.1    | 0.9    | 0.9    | 0.8     | 0.9     |
| S5     | 30/10/2016 | 9:50           | 13:40           | 1.6             | 3.7    | 2.1    | 1.7    | 1.6     | 1.4     |
| S6     | 15/02/2017 | 12:25          | 16:20           | 2.0             | 1.4    | 1.7    | 1.3    | 1.4     | 1.4     |
| S7     | 5/05/2017  | 6:40           | 10:25           | 0.6             | 1.3    | 1      | 1.2    | 1.2     | 1.1     |
| S8     | 15/11/2017 | 8:00           | 12:00           | 1.1             | 1.2    | 1.1    | 1.4    | 1.2     | 1.4     |
| TS     | 31/08/2017 | 16:25          | 03/09/2017 at 1:20 | 1.5          | 2.1    | 1.7    | 1.5    | 1.6     | 1.5     |
3. Results.

3.1 Seasonal spatial surveys

Water temperature was lowest at the ocean mouth increasing towards the shallow mangrove forest water endmember (max=34.7 °C; Table 2) throughout the eight underway surveys. The lowest salinity was observed in the mangrove forest water endmember (26.2; Survey 1) however overall average salinity ranges were relatively close to seawater (31.5 - 33.0) reflecting the characteristics of an ocean dominated embayment. Dissolved oxygen (DO) was slightly undersaturated at the ocean entrance and increased significantly in the seagrass beds in 5 out of 8 surveys (reaching 166.3 %; Survey 5, Figure 3), while the lowest DO was observed in the mangrove forest water area (≤ 60% in 5 surveys). There was minimal tidal variation on survey days (mean= ~1.3m although Survey 6 tidal range was ~2m.

Table 2. Mean, standard deviation, minimum and maximum for pCO₂, radon, DO, salinity and temperature for eight underway spatial surveys and time series in Gilimanuk Bay.

| Survey     | Date     | pCO₂ (µatm) | ²²²Rn (dpm/L) | DO (%) | Salinity | Temp. (°C) |
|------------|----------|-------------|---------------|--------|----------|------------|
| Survey 1   | 20-Nov-15| 516         | 2.3           | 101    | 32.2     | 30.8       |
| Survey 2   | 9-Jan-16  | 1018        | 3.4           | 87     | 32.3     | 31.2       |
| Survey 3   | 30-Jun-16 | 796         | 2.6           | 87     | 31.5     | 30.7       |
| Survey   |       |       |       |       |       |
|----------|-------|-------|-------|-------|-------|
|          | Min   | Max   | Min   | Max   |       |
|          | 360   | 1965  | 399   | 1388  |       |
| 14-Sep-16| Mean  | 908   | 385   | 1388  |       |
|          | 2.0   | 1.5   | 0.1   | 4.7   |       |
|          | 77    | 9     | 60    | 90    |       |
|          | 32.7  | 1.1   | 31.1  | 34.8  |       |
|          | 26.5  | 1.7   | 23.8  | 29.9  |       |
| 30-Oct-16| Mean  | 573   | 288   | 1167  |       |
|          | 2.3   | 1.4   | 0.2   | 4.4   |       |
|          | 96    | 29    | 70    | 166   |       |
|          | 32.8  | 1.2   | 30.4  | 33.9  |       |
|          | 28.2  | 2.6   | 25.8  | 33.9  |       |
| 5-Jan-17 | Mean  | 1164  | 597   | 2101  |       |
|          | 2.7   | 2.6   | 0.0   | 7.4   |       |
|          | 77    | 13    | 56    | 96    |       |
|          | 32.5  | 1.3   | 29.4  | 34.7  |       |
|          | 27.1  | 1.1   | 25.4  | 28.8  |       |
| 5-May-17 | Mean  | 560   | 214   | 1001  |       |
|          | 0.9   | 0.6   | 0.1   | 1.9   |       |
|          | 85    | 11    | 69    | 100   |       |
|          | 33.1  | 1.1   | 31.6  | 34.8  |       |
|          | 29.8  | 1.9   | 27.4  | 34.7  |       |
| 17-Nov-17| Mean  | 849   | 519   | 1600  |       |
|          | 2.2   | 1.6   | 0.1   | 4.9   |       |
|          | 86    | 21    | 59    | 141   |       |
|          | 33.0  | 0.7   | 32.1  | 34.5  |       |
|          | 27.3  | 1.6   | 25    | 30.6  |       |
| Time Series | Mean | 458   | 519   |       |       |
|          | 0.8   | 3.1   |       |       |       |
|          | 94    | 104   |       |       |       |
|          | 35.0  | 35.2  |       |       |       |
|          | 26.6  | 27.7  |       |       |       |
Figure 3. Distance plots for 8 underway seasonal surveys for CO$_2$ fluxes from the parameterization of Raymond and Cole (2001) (red line = source/sink boundary), $p$CO$_2$ measurements (blue line represents atmospheric $p$CO$_2$ concentrations ~400 µatm), $^{222}$Rn, DO (grey line = 100% supersaturation), salinity and temperature from the ocean mouth of Gilimanuk Bay to the mangrove forest water.

$^{222}$Rn (groundwater tracer) was lowest (0.1 dpm/L) at the ocean entrance and increased markedly towards the mangrove forest water where concentrations reached 7.4 dpm/L.
Stronger correlations between $^{222}$Rn and $pCO_2$ than between dissolved DO and $pCO_2$ (Figure 4) in 7 out of 8 surveys suggest that the seepage of groundwater, not pelagic respiration and photosynthesis, drove $pCO_2$ supersaturation within the embayment, particularly in the mangrove forest water (Figure 4). $pCO_2$ was mostly above atmospheric equilibrium (>400 µatm) with up to a five-fold increase in the mangrove forest water (max = 2101 µatm). On average, $pCO_2$ in the mangrove forest water was ~four-fold that of the overall embayment (Figure 3). In contrast, the seagrass beds were undersaturated in 30% of surveys suggesting that, seagrass was fixing mangrove water-derived CO2 from the upper embayment.

Figure 4. Correlations between $pCO_2$ and proxies of potential drivers ($^{222}$Rn, DO, salinity and temperature) during the 8 spatial surveys in Gilimanuk Bay.

Annual rainfall average over the three year sampling span was 695 mm in 2015 (drought year), 1357 mm in 2016 and 1247 mm in 2017, with 2609 mm of rainfall recorded over the 24 months of observations. Historical rainfall for the region is 1316 mm/year. The highest
lagged correlation coefficients (Figure 5) between rainfall and radon, CO₂, and salinity were used to interpret the lag time between rainfall, groundwater discharge, and subsequent response in groundwater-derived pCO₂ or radon in Gilimanuk Bay. In spite of the relatively dry conditions during sampling, cumulative antecedent rainfall 17 to 63 days prior to surveys had significant correlations to pCO₂ (p≤0.05) with peak correlations (R² = 0.81; p<0.002) when 29 days of cumulative rainfall was used. Significant correlations between pCO₂ and cumulative antecedent rainfall was observed 15 and 84 days prior to our survey in the mangrove waters (p<0.05) with the strongest correlation at 48 days cumulative antecedent rainfall (R² = 0.90; Figure 5). Correlations in the seagrass area were significant between 29 and 104 days cumulative antecedent rainfall (p<0.05) with the strongest correlation for 73 days of cumulative antecedent rainfall (R² = 0.77). This wide range of significant lagged correlations shown may imply that these correlations may not necessarily represent a causation in the seagrass area. Seagrasses and the embayment mouth only showed significant salinity correlations up to 11 days and 4 days, suggesting direct rainfall is the major influence on local salinity. Interestingly, groundwater seepage (as traced by ²²²Rn) mimicked pCO₂ in the mangrove forest water, but was decoupled in the highly productive seagrass (Figure 5). This suggests that in-situ productivity was driving pCO₂ rather than groundwater seepage in the seagrasses. In the ocean mouth, ²²²Rn had significant correlations with the widest range of delayed rainfall (29-268 days; p<0.05) also implying that these correlations may not necessarily represent a causation in this case. Overall, the lagged correlations imply that delayed groundwater seepage following seasonal rainfall plays a significant role in CO₂ concentration and distribution in this embayment.
Figure 5. Lagged correlation coefficients (R²) of the relationship between average pCO₂ (circles), ²²²Rn (diamonds) and salinity (squares) and accumulated antecedent rainfall for eight underway surveys in mangrove forest water (brown), seagrass areas (green) and ocean mouth (blue) in Gilimanuk Bay.

CO₂ fluxes revealed shifts from a strong source to the atmosphere in the mangrove forest water, an intermittent sink in the seagrass beds to equilibrium or a weak source at the ocean mouth (Table 3). The total mangrove forest area covered ~30 % of the embayment transect.
and seagrass beds covered ~40 %. Measurements suggested that the release of CO₂ to the atmosphere occurred in mangrove forest water (total average CO₂ flux= 18.1 ± 5.8 mmol m⁻² d⁻¹) was much greater than in the seagrass beds where the CO₂ flux was 2.5 ± 3.4 mmol m⁻² d⁻¹. Seagrass beds were a sink of CO₂ in 2 out of the 8 surveys (Figure 3). Average fluxes along the transect were 9.8 ± 6.0 mmol m⁻² d⁻¹, a source of CO₂ (Table 3). No clear seasonal patterns were observed.

Table 3. Average CO₂ fluxes (mmols m⁻² day⁻¹) for eight underway surveys categorised into three classes: ocean, seagrass beds and mangrove forest water.

| Survey | Ocean   | Seagrass | Mangrove |
|--------|---------|----------|----------|
| S1     | 0.0 ± 0.0 | -1.3 ± 1.8 | 10.5 ± 3.2 |
| S2     | 0.0 ± 0.6 | 1.8 ± 2.5  | 22.6 ± 7.6 |
| S3     | 0.0 ± 0.2 | 2.1 ± 1.7  | 16.0 ± 12.4 |
| S4     | 0.0 ± 1.8 | 0.3 ± 4.2  | 18.1 ± 2.2 |
| S5     | 0.5 ± 0.5 | 6.3 ± 2.2  | 11.2 ± 5.7 |
| S6     | 0.9 ± 0.1 | -0.9 ± 3.3 | 30.1 ± 7.6 |
| S7     | 0.1 ± 0.0 | 5.8 ± 1.6  | 10.7 ± 2.8 |
| S8     | 0.5 ± 0.2 | 6.1 ± 10.0 | 25.8 ± 4.8 |
| Total mean | 0.3 ± 0.4 | 2.5 ± 3.4 | 18.1 ± 5.8 |

We selected Raymond and Cole (2001) as the most reasonable k₆₀₀ model for the local conditions, and use these values in all figures. Raymond and Cole (2001) estimated k₆₀₀ as a function of windspeed using a range of deliberate gas tracer and floating dome experiments in estuaries. Currents were assumed to be a minor driver of k₆₀₀ since there was no river inflow and microtides prevented significant current-driven turbulence. Borges et al. (2004) k₆₀₀ model resulted in unrealistic high emissions values since their gas transfer velocities estimates were developed in the microtidal Scheldt estuary which experiences strong currents.
(Table 3). Wanninkhof (2014) had the lowest values, likely due to this parameterization being
developed for the open ocean, where wind is the only driver of near surface turbulence.

Raymond and Cole (2001) presented a mid-range flux value, which is likely more
representative of the study site, which does have some tidal flow, but much lower current
velocities than the macrotidal Scheldt. Table 4 also includes the oceanic wind speed
parameterisations of Wanninkhof and McGillis (1999) and Wanninkhof (2014) for reference.

Table 4. Survey (1-8) and time series, average windspeeds 28 days prior (AWS) and CO₂
flux calculations using four author's transfer velocity parameterizations for equations 2-5:
Wanninkhof and McGillis (1999), Raymond and Cole (2001), (Borges et al., 2004) and
Wanninkhof (2014), including minimum, maximum, mean, standard deviations.

| Survey | CO₂ fluxes (mmol m⁻² d⁻¹) | AWS | W&M99 | R&C01 | BO4 | W2014 | Total Mean |
|--------|---------------------------|-----|-------|-------|-----|-------|------------|
|        |                           | Mean| 0.1   | 2.8   | 6.7 | 0.8   | 2.6        |
| Survey 1 |                           | St. Dev ± | 0.2 | 6.4 | 15.1 | 1.7 | 5.9        |
|         |                           | Min  | -0.1  | -3.2 | -7.5 | -0.9 | -2.9       |
|         |                           | Max  | 0.5   | 16.5 | 38.9 | 4.5  | 15.1       |
|         |                           |        |       |       |       |       |            |
| Survey 2 |                           | Mean | 0.3   | 16.9 | 35.8 | 3.1  | 14.0       |
|         |                           | St. Dev ± | 0.2 | 11.3 | 26.0 | 2.3  | 9.9        |
|         |                           | Min  | 0.0   | -0.2 | -0.3 | 0.0  | -0.1       |
|         |                           | Max  | 0.7   | 35.4 | 81.9 | 7.1  | 31.3       |
|         |                           |        |       |       |       |       |            |
| Survey 3 |                           | Mean | 0.1   | 9.8  | 21.6 | 1.5  | 8.2        |
|         |                           | St. Dev ± | 0.1 | 12.4 | 27.2 | 1.8  | 10.4       |
|         |                           | Min  | 0.0   | -0.9 | -2.0 | -0.1 | -0.8       |
|         |                           | Max  | 0.4   | 37.8 | 83.0 | 5.6  | 31.7       |
|         |                           | 1.3 m/s |       |       |       |       |            |
| Survey 4 |                           | Mean | 0.1   | 11.7 | 21.0 | 0.9  | 8.4        |
|         |                           | St. Dev ± | 0.0 | 7.4  | 14.7 | 0.6  | 5.7        |
|         |                           | Min  | 0.0   | 0.0  | 0.0  | 0.0  | 0.0        |
### Survey 5

| Mean | 0.1 | 4.3 | 9.6 | 0.7 | 3.7 |
|------|-----|-----|-----|-----|-----|
| St. Dev ± | 0.1 | 6.5 | 14.5 | 1.1 | 5.6 |
| Min | -0.1 | -5.1 | -11.5 | -0.9 | -4.4 |
| Max | 0.2 | 19.6 | 44.0 | 3.3 | 16.8 |

**1.4 m/s**

### Survey 6

| Mean | 0.4 | 19.9 | 44.9 | 3.4 | 17.1 |
|------|-----|-----|-----|-----|-----|
| St. Dev ± | 0.3 | 13.9 | 31.2 | 2.4 | 11.9 |
| Min | 0.0 | 1.0 | 2.2 | 0.2 | 0.8 |
| Max | 0.7 | 41.2 | 92.5 | 7.0 | 35.4 |

### Survey 7

| Mean | 0.0 | 3.8 | 7.6 | 0.4 | 3.0 |
|------|-----|-----|-----|-----|-----|
| St. Dev ± | 0.0 | 4.8 | 9.8 | 0.5 | 3.8 |
| Min | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| Max | 0.1 | 13.3 | 27.3 | 1.4 | 10.5 |

### Survey 8

| Mean | 0.1 | 9.8 | 22.1 | 1.7 | 8.4 |
|------|-----|-----|-----|-----|-----|
| St. Dev ± | 0.3 | 13.1 | 29.5 | 2.2 | 11.3 |
| Min | -1.0 | -13.1 | -28.1 | -2.2 | -11.1 |
| Max | 0.5 | 29.0 | 65.4 | 5.0 | 25.0 |

### Time Series

| Mean | 0.0 | 1.4 | 3.1 | 0.3 | 1.2 |
|------|-----|-----|-----|-----|-----|
| St. Dev ± | 0.0 | 0.4 | 0.9 | 0.1 | 0.3 |
| Min | 0.0 | 0.6 | 1.4 | 0.1 | 0.5 |
| Max | 0.1 | 2.7 | 6.3 | 0.5 | 2.4 |

### 3.2 Time Series at the ocean entrance

A 55 hr time series was conducted at the embayment entrance from 16:25, 31 August 2017 to 00:30, 3 September 2017 to investigate aquatic CO₂ exchanges between the embayment and the ocean (Figure 6). There was no precipitation during observations and 11.0 mm of rain was recorded in the month preceding the time series observations. Water temperatures ranged
from 25.1 to 27.7 °C and were lowest between midnight and early morning. Salinity ranged from 34.9 to 35.2. The higher salinity measurements on the outgoing tide may be a result of evaporation within this shallow embayment (data not shown). DO ranged from 82.5 % to 104.3 %. In spite of some data gaps, DO appeared to have diel trends reflecting photosynthesis during the day and respiration at night (Figure 7). The $p$CO$_2$ trends did not follow DO. Indeed, $p$CO$_2$ had stronger correlations to $^{222}$Rn than DO during the day (Figure 7) implying the groundwater CO$_2$ source was stronger than the photosynthesis sink. $^{222}$Rn followed a tidal cycle and correlated with $p$CO$_2$ during both incoming and outgoing tides ($R^2=0.53 & R^2=0.22$, respectively; data not shown). The highest $^{222}$Rn, salinity and CO$_2$ values were observed during the two lowest tides (overall tidal range = 1.5m) which is consistent with groundwater-derived inputs that are well known to occur at low tide (Atkins et al., 2013; Call et al., 2015; McMahon & Santos, 2017; Santos et al., 2009). It is difficult to explain the increase in $p$CO$_2$ from ~440 to 500 µatm in the last 4 hours of the time series since both radon (groundwater proxy) and DO (respiration proxy) had no similar changes. $p$CO$_2$ was above atmospheric equilibrium throughout with the highest $p$CO$_2$ observations at low tides and lowest at high tides. Low $p$CO$_2$ at the embayment mouth are consistent with the survey observations of CO$_2$ outgassing and/or uptake by the seagrass beds (Table 4; R&C01: 1.4 ± 0.4 mmol m$^{-2}$ d$^{-1}$).
Figure 6. Observations during the 55 hour time series conducted at the ocean mouth of Gilimanuk Bay.
4. Discussion

Aquatic systems in Southeast Asia are recognised as significant sources of CO$_2$ to the atmosphere but remain poorly represented in global databases (Müller et al., 2015). The few studies available focus on tropical river-dominated estuaries which produce significant CO$_2$ fluxes (Borges & Abril, 2011; Müller et al., 2015). Global summaries of water-to-air CO$_2$ fluxes are generally confined to human impacted river-dominated estuarine systems (Cai, 2011). A recent study in a temperate autotrophic marine dominated system in Australia reported ten-fold lower and reversed CO$_2$ fluxes than the more studied river-dominated counterparts (Maher & Eyre, 2012). Since there is a paucity of data from marine-dominated tropical coastal embayments, our investigation contributes to filling gaps in global estuarine CO$_2$ fluxes.

By categorizing CO$_2$ fluxes into three classes (mangrove forest water, seagrass beds and the ocean-dominated mouth), comparisons with temperate coastal water CO$_2$ fluxes could be made. For instance, mangrove forest water had CO$_2$ fluxes ($18.1 \pm 5.8$ mmol m$^{-2}$ d$^{-1}$) similar to estuarine systems such as the York River estuary in the U.S.A. ($17$ mmol m$^{-2}$ d$^{-1}$) (Raymond et al., 2000) and the Pearl River in China ($24$ mmol m$^{-2}$ d$^{-1}$)(Yuan et al., 2011). Overall, the mangrove forest water dominated surrounding water CO$_2$ fluxes and were within estimated flux ranges for many global coastal waters (Chen et al., 2013). A study by Ho et al. (2017) reported average CO$_2$ fluxes of $105 \pm 9$ and $99 \pm 6$ mmol m$^{-2}$ d$^{-1}$ at the end of the wet
season, where $p$CO$_2$ values ranged from 1000 to 6200 µatm (present study ranges = 516 (S1; Nov.2015) to 1164 µatm (S6; Jan. 2017; Table 2). We report overall embayment CO$_2$ fluxes of 9.8 ± 6.0 mmol m$^{-2}$ d$^{-1}$ comparable to the lower range of the Everglades study, estimated global mangrove fluxes of 4.6 to 113.5 mmol m$^{-2}$ d$^{-1}$ (Borges et al., 2003) and revised global estimates of 56.8 ± 8.9 mmol m$^{-2}$ d$^{-1}$ (Rosentreter et al., 2018). This is most likely due to lack of river inputs and large uptake by seagrass beds in the mid-embayment.

Seasonality seems to play a greater role in temperate waters (Guo et al., 2009). In the Changjiang Estuary (China) seasonal ranges were 52.9 mmol m$^{-2}$ d$^{-1}$ (December) to 92.9 mmol m$^{-2}$ d$^{-1}$ (August) based on the parameterization of Raymond and Cole (2001) (Zhai et al., 2007). Seasonality was not observed throughout our study. Gilimanuk Bay CO$_2$ emissions ranged from 2.8 mmol m$^{-2}$ d$^{-1}$ (S1; October 2015) to 16.9 mmol m$^{-2}$ d$^{-1}$ (S2; January 2016; Table 4) with no clear changes over an annual temperature cycle. Autotrophic coastal systems have been reported as sinks of atmospheric derived CO$_2$ (Borges and Abril, 2011; Maher and Eyre, 2012). The seagrass-dominated area of Gilimanuk Bay alternated between a CO$_2$ source and sink. The more consistent CO$_2$ source in the near shore ocean was possibly due to coral calcification in Gilimanuk Bay’s fringing coral reefs (Figure 8).
Groundwater seepage, porewater exchange and rainfall events have been linked to delayed groundwater discharge as a source of CO$_2$ in subtropical systems (Ruiz-Halpern et al., 2015). Rainfall, particularly flood events, rapidly transport carbon stored in upper soil profiles across the land-to-water interface (Atkins et al., 2013; Gatland et al., 2014; Jeffrey et al., 2016; Webb et al., 2016). Paquay et al. (2007) reported that water residence times and periodic floods drove the distribution of $p$CO$_2$ in Hilo Bay (Hawaii) with groundwater seepage suggested to explain a 3 day delay of elevated $p$CO$_2$ after a heavy rainfall event. In contrast to Hilo Bay, Gilimanuk Bay has no riverine input, making groundwater seepage a dominant source of CO$_2$ year-round (average $R^2$=0.74) followed by temperature (average $R^2$=0.65) and salinity (average $R^2$=0.63; Figure 4).

Tidally-driven porewater exchange (tidal pumping) has been suggested to release carbon dioxide and nutrients to estuaries (Sadat-Noori et al., 2016) and intertidal flats (Bouillon, Connolly, et al., 2008; Santos et al., 2014). Large tidal amplitudes (spring tides) have been related to enhanced $^{222}$Rn in coastal waters as observed off a sandy beach in Korea (Kim & Hwang, 2002) and in Florida (Santos et al., 2009) and mangrove forest water in Australia (Call et al., 2015). Radon traces any water in contact with sediments regardless of salinity. However, our dataset cannot resolve whether tidally-driven porewater exchange or fresh groundwater discharge are the source of radon enrichments in the mangrove forest water. Considering the small tidal amplitude, lack of well-defined tidal creeks in the mangroves, steep topography surrounding the upper embayment, and the significant correlations between radon and antecedent rainfall, we suggest that delayed fresh groundwater discharge (rather
than tidal pumping) is more likely to be the radon and CO₂ source to the upper embayment. Groundwater discharge at the shoreline is well known to lag rainfall for several months. Seasonal oscillation in groundwater level and inland recharge can explain large saline groundwater discharge several months after rainfall as observed in a Massachusetts aquifer (Michael et al., 2005). Unfortunately, no groundwater level data are available for the Gilimanuk Bay area to build on this hypothesis.

CO₂ derived from the mangrove forest in the upper embayment appears to be fixed by seagrass beds in the lower embayment, intermittently transforming the lower embayment into a net sink of CO₂ (Figure 1; Figure 3; Figure 8). However, there was higher pCO₂ and water-to-air pCO₂ fluxes in wetter periods in the seagrass beds (Figure 9; Surveys 3, 6 & 8) which coincided with the highest antecedent rainfall (Table 3). Seagrasses, although known to be net autotrophic, are largely under-represented in global carbon budgets (Duarte et al., 2010). Additionally, there are large uncertainties in the extent of seagrass beds globally, resulting from a) many regions in Indonesia being known to support extensive seagrass beds but have not been surveyed and b) the continuing loss of seagrass bed areas as a result of well-reported anthropogenic degradation (Alongi et al., 2016; Duarte et al., 2010; Unsworth & Cullen, 2010).
Figure 9. Average water-to-air CO₂ fluxes using piston velocities from the parameterization of Raymond and Cole (2001) for eight underway surveys and time series conducted in Gilimanuk Bay.

The ocean’s blue carbon sinks (mangroves, saltmarshes and seagrasses) capture and store approximately 70% of the carbon perpetually stored in aquatic systems (Nellemann & Corcoran, 2009). In contrast to mangrove habitats, which occur only in warmer tropical and subtropical climates, seagrasses are also present in colder northern and southern latitudes (Orth et al., 2006). The importance of their combined presence with mangrove forests in tropical systems is noteworthy as both mangrove forests and seagrass beds stabilise sediment (Hogarth, 2015). About 50% of net primary production produced by seagrasses is buried within seagrass sediments (Kennedy et al., 2010), and ~ 8% of mangrove net primary production is retained in mangrove forest soils (Bouillon, Borges, et al., 2008).

Degradation or loss of mangrove forests and seagrass beds may cause the release of large stores of sedimentary carbon to both the atmosphere and the coastal ocean (Mcleod et al.,
2011). Therefore, continued degradation may increase atmospheric CO₂. Due to the high carbon stores in blue carbon systems, and in particular with the tropical regions such as Indonesia, prevention of ecosystem degradation is crucial for limiting the potential release of this carbon to the atmosphere as CO₂ (Tollefson, 2018). Strategies to counter the drainage of blue carbon ecosystems would be an effective measure in maintaining carbon in soils (Crooks et al., 2011). For example, construction or artificial canals draining coastal wetlands more than doubled CO₂ emissions from waterways on the highly urbanized Gold Coast, Australia (Macklin et al., 2014). Our investigation further demonstrates the role seagrass beds play in taking up dissolved inorganic carbon and preventing emissions from groundwater-derived CO₂ in nearby ecosystems.

5. Conclusion

Our investigations across a coral reef-seagrass-mangrove continuum revealed a CO₂ source in the mangrove dominated upper bay apparently associated with delayed groundwater inputs, and differing CO₂ dynamics in the lower bay driven by the uptake of CO₂ by seagrass. The bay mouth was a source of CO₂ possibly due to production of CO₂ during fringing coral reef calcification. The average CO₂ water-to-air flux along the transect was 9.8 ± 6.0 mmol m⁻² d⁻¹. Antecedent rainfall and radon were the best predictors of CO₂ dynamics, with no clear seasonality observed, in contrast to better studied seasonal temperate systems. Potential changes in rainfall events due to climate change in Indonesia (Overpeck & Cole, 2007) as well as ecosystem degradation may alter aquatic CO₂ emissions. This study may assist when determining anthropogenic modification buffer zones in mangrove forest embayments. As pCO₂ data in tropical coastal embayments are scarce, more studies are needed to assess the role of these systems in the global carbon cycle.

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Highlights

- An aquatic CO2 source in the mangrove dominated upper embayment waters was associated with delayed groundwater inputs.
- A shifting CO2 source-sink in the lower embayment bay was driven by the uptake of CO2 by seagrass and mixing with oceanic waters.
- This more pristine system differs to modified landscapes, where potential uptake of CO2 is weakened due to the degradation of seagrass beds, or emissions are increased due to drainage of coastal wetlands.
- Antecedent rainfall and radon were found to be the best predictors of CO2 dynamics, with no clear seasonality observed, in contrast to better studied seasonal temperate systems.
- Extended delays of groundwater-derived CO2 in the coastal waters compared to mangrove forest waters highlight the importance of comparative studies with more modified systems which are known to have more rapid delivery of carbon to coastal waters.
- This study may assist when determining anthropogenic modification buffer zones in mangrove forest embayments.