Abstract. Numerous experiments have shown that ocean acidification impedes coral calcification, but knowledge about in situ reef ecosystem response to ocean acidification is still scarce. Bahía Culebra, situated at the northern Pacific coast of Costa Rica, is a location naturally exposed to acidic conditions due to the Papagayo seasonal upwelling. We measured pH and $pCO_2$ in situ during two non-upwelling seasons (June 2012, May-June 2013), with a high temporal resolution of every 15 and 30 min, respectively, using two Submersible Autonomous Moored Instruments (SAMI-pH, SAMI-CO2). These results were compared with published data from the 2009 upwelling season. Findings revealed that the carbonate system in Bahía Culebra shows a high temporal variability. Incoming offshore waters drive inter- and intra-seasonal changes. Lowest pH (7.8) and highest $pCO_2$ (658.3 µatm) values measured during a cold-water intrusion event in the non-upwelling season were similar to those minimum values reported from upwelling season (pH = 7.8, $pCO_2 = 643.5$ µatm), unveiling that natural acidification also occurs sporadically in the non-upwelling season. This affects the interaction of photosynthesis, respiration, calcification, and carbonate dissolution and the resulting diel cycle of pH and $pCO_2$ in the reefs of Bahía Culebra. During non-upwelling season the aragonite saturation state ($\Omega_a$) rises to values of >3.3 and during upwelling season fall below 2.5. The $\Omega_a$ threshold values for coral growth were derived from the correlation between measured $\Omega_a$ and coral linear extension rates which were obtained from the literature and suggest that future ocean acidification will threaten the continued growth of reefs in Bahía Culebra. These data contribute to build a better understanding of the carbonate system dynamics and coral reefs key response (e.g. coral growth) to natural low-pH conditions, in upwelling areas in the Eastern Tropical Pacific and beyond.
1 Introduction

Ocean acidification (OA) caused by human-induced increase of atmospheric CO$_2$ (Sabine et al., 2004; Feely et al., 2009) is considered one of the major threats to marine calcifying organisms and ecosystems (Fabry et al., 2008; Hofmann et al., 2010; Doney et al., 2012; Gattuso et al., 2015). Among all marine habitats, tropical coral reefs are recognized as the most endangered (Hoegh-Guldberg et al., 2007; Kleypas and Yates, 2009; Pörtner et al., 2014), since in addition to reduced calcification (Langdon et al., 2000; Marubini et al., 2008; Doney et al., 2009; Gattuso et al., 2014), a lower pH also weakens the reef framework by favoring bioerosion and enabling carbonate dissolution (Gattuso et al., 2014; Manzello et al., 2014; Barkley et al., 2015). According to the IPCC business-as-usual scenario, about 90% of the ocean’s surface waters will become undersaturated with respect to aragonite in the next decades (Gattuso et al., 2015), emphasizing the need to study the response of natural ecosystems to OA. Nowadays, aragonite undersaturated surface waters occur naturally in some parts of the ocean, as consequence of underwater volcanic seeps (Hall-Spencer et al., 2008; Fabricius et al., 2011; Enochs et al., 2015; Fabricius et al., 2015) or upwelling that drags corrosive deep water into the surface mixed layer (Feely et al., 2008; Hauri et al., 2009; Fassbender et al., 2011; Harris et al., 2013).

Aside from some studies at volcanic seeps (Fabricius et al., 2011; Kroeker et al., 2011; Enochs et al., 2015; Fabricius et al., 2015) or at reefs in the Eastern Tropical Pacific (ETP) (Manzello, 2008, 2010a, 2010b; Manzello et al., 2008, 2014), our understanding of OA impacts on corals derives mainly from laboratory and seawater enclosure experiments (Pörtner et al., 2014; Hall-Spencer et al., 2015). These results are used to predict ecosystem responses to future OA (Kleypas et al., 2006; Kleypas and Langdon, 2006), but their reliability is challenged by the artificial conditions under which the experiments are conducted. For example, the duration of studies is often too short to allow a full adaptation or acclimatization of the organisms/systems to the changing environmental conditions, and the missing connectivity between ecosystems in seawater enclosures restricts natural interactions between organisms (Kleypas et al., 2006; Kleypas and Langdon, 2006; Hofmann et al., 2010). In situ studies in natural low-pH conditions are able to overcome some of these problems and the ETP is well known for its CO$_2$-enriched and acidic subsurface waters (Takahashi et al., 2014). Upwelling events decreases the carbonate saturation state ($\Omega$) along the Central American coast (Manzello et al., 2008; Manzello, 2010b; Rixen et al., 2012), and have the potential to produce poorly cemented coral reefs with low accretion rates that are subject to rapid bioerosion (Manzello et al., 2008; Alvarado et al., 2012).

Corals in the northern part of the Costa Rican Pacific coast are developing under the influence of the seasonal Papagayo upwelling (Jiménez et al., 2010; Rixen et al., 2012; Stuhldreier et al., 2015a, 2015b). To contribute to the general understanding of OA impacts on coral reefs, we investigated the variability of the carbonate system in the upwelling-influenced Bahía Culebra, Costa Rica. The main objectives of this study were 1) to describe the behavior of the carbonate system on diurnal and seasonal time scales, 2) to characterize the controlling processes, and 3) to determine ecological impacts of changing carbonate systems. Furthermore, our results will allow us to draw some conclusions concerning future thresholds of coral reef development within this bay.
2 Methods

2.1 Study site

Bahía Culebra, located in the Gulf of Papagayo, North Pacific coast of Costa Rica (Fig. 1), is strongly influenced by the northeasterly Papagayo winds. The strongest wind jets develop during the boreal winter (Amador et al., 2016) and are driven by large-scale variations of the trade winds (Chelton et al., 2000; Alfaro and Cortés, 2012). When Papagayo winds blow through the mountain gap between southern Nicaragua and northern Costa Rica, the resulting strong offshore winds on the Pacific side lead to upwelling of cold and nutrient-enriched subsurface waters between December and April (McCreary et al., 1989; Brenes et al., 1990; Ballesteros and Coen, 2004; Kessler, 2006;). These cyclonic eddies also influence the magnitude and location of the Costa Rica Dome (CRD), which is located ca. 300 km off the Gulf of Papagayo (Fiedler, 2002). However, the CRD changes its distance to the Costa Rican coast throughout the year, as a result of differences in wind forcing (Wyrtki, 1964; Fiedler, 2002). During the dry season, particularly between February and April, offshore moving water masses strengthen upwelling at the coast and shoal the thermocline in the Gulf of Papagayo (Wyrtki, 1965, 1966; Fiedler, 2002). In May-June, during the onset of the rainy season the CRD moves offshore (Fiedler, 2002; Fiedler and Talley, 2006) and the North Equatorial Countercurrent (NECC) can carry tropical water masses into Bahía Culebra until December, when again upwelling sets in (Wyrtki, 1965, 1966).

2.2 Measurements

We measured in situ pH, $pCO_2$ and seawater temperature (SWT) during two non-upwelling periods (15 days in June 2012 and 7 days in May-June 2013, Fig. 2). Measurements were undertaken with two Submersible Autonomous Moored Instruments (SAMI-pH and SAMI-CO$_2$) (www.sunburstssensors.com), in sampling intervals of 15 (June 2012) and 30 minutes (May-June 2013). SAMI-sensors were deployed at the pier of Marina Papagayo (85°39’21.41”W; 10°32’32.89”N), on top of a carbonate sandy bottom in the inner part of Bahía Culebra (Fig. 1). The water-depth varied approximately between 5-8 m depending on the tide, but sensors, hooked to the pier, moved up and down with the tide and were always at the same depth, 1.5 m below the surface. SAMI instruments measured pH (total hydrogen ion scale) and pCO$_2$ spectrophotometrically by using a colorimetry reagent method (DeGrandpre et al., 1995, 1999; Seidel et al., 2008). Salinity from discrete samples was measured with a WTW probe (Cond3310) and was used for correction of pH values. Calculation of aragonite saturation state ($\Omega_a$) from parameters measured in situ with SAMI sensors is accurate (Cullison Gray et al., 2011; Gray et al., 2012), but discrete water samples were collected as often as possible to validate the instruments (Fig. 3). 250 mL borosilicate bottles were filled with seawater at 30 cm below the surface and preserved with 200 µl of 50% saturated HgCl$_2$ solution to inhibit biological activity (Dickson et al., 2007). Samples were stored at 3-4 °C until analysis. Total alkalinity (TA) and Dissolved inorganic carbon (DIC) were measured using a VINDTA 3C (Versatile Instrument for the Determination of Total dissolved inorganic carbon and Alkalinity; Marianda, Kiel, Germany) coupled with a UIC CO$_2$ coulometer detector (UIC Inc., Joliet, USA). Both instruments were calibrated with Dickson Certified Reference Material
DIC concentrations as well as TA and Ωₐ were calculated with the CO2SYS program as a function of measured pH and pCO₂; with dissociation constants of Mehrbach et al. (1973) for carbonic acid as refit by Dickson and Millero (1987), and Dickson (1990) for boric acid.

Wind speeds were obtained from a station of the Instituto Metereológico Nacional (National Meteorological Institute of Costa Rica), located at the nearby Liberia airport. The Módulo de Información Oceanográfica of the University of Costa Rica (www.miocimar.ucr.ac.cr) supplied the tidal data. All coral growth values were taken from the literature; linear extension rates from Bahía Culebra were measured by Jiménez and Cortés (2003), whilst coral growth in Panamá and Galápagos was measured by Manzello (2010a). For the correlation between coral growth and Ωₐ, we used the mean Ωₐ values from Panamá and Galápagos previously reported by Manzello (2010b).

2.3 Data analysis

We compared our data with values measured during upwelling season in 2009 (Rixen et al., 2012). In 2009 xCO₂ was measured by an underway pCO₂ system (SUNDANS) equipped with an infrared gas analyzer (LI-7000), and pH was measured using an Orion ROSS electrode an Orion Star™. Correlations between tidal cycles and physicochemical parameters (pH, pCO₂, T, wind) during non-upwelling periods were tested via Pearson Correlation in Python. Differences in parameters (temperature, pH, pCO₂, TA, DIC, Ωₐ) between all periods (2009, 2012, 2013) were tested with a General Linear Model (GLM), in the statistical package R. The GLM was evaluated using graphical methods to identify violations of assumptions of homogeneity of variance and normality of residuals. All GLM assumptions were met. Additionally, we developed a simple model to improve our understanding of processes controlling the observed diel trends, as seen in the time series data of pH and pCO₂ (Fig. 2, 4). The model simulates combined effects of metabolic processes (photosynthesis, respiration, calcification and dissolution) on the carbonate chemistry. Input parameters for starting the model were the calculated DIC (in 2012: 2037 µmol kg⁻¹ at 7:00 h and 2019 µmol kg⁻¹ at 15:00 h; in 2013: 1883 µmol kg⁻¹ at 5:00 h and 1805 µmol kg⁻¹ at 15:00 h) and TA (in 2012: 2284 µmol kg⁻¹ at 7:00 h; in 2013: 2193 µmol kg⁻¹ at 5:00) values, corresponding to the highest and lowest measured pCO₂ during the day. Calculation of TA and DIC from the pair pH and pCO₂ is prone to errors (Millero, 2007; Cullison Gray et al., 2011), however the values used as input parameters in the model are in range with those reported from other studies in tropical areas (Manzello, 2010b; Cyronak et al., 2013b). The difference between the two DIC concentrations (ΔDIC) was assumed to be caused by photosynthesis/respiration and the resulting formation and decomposition of particulate organic carbon (POC) as well as calcification/dissolution and the precipitation and dissolution of particulate inorganic carbon (PIC, Eq. 1). R_OI describes the ratio between the production of organic carbon (POC) and precipitation of calcium carbonate carbon (PIC), and was used to link ΔPOC to ΔPIC (R_OI=POC/PIC) (Eq. 2, 3). The R_OI was further constrained by the determined change of TA (ΔTA). Therefore, it was considered that photosynthesis and respiration of one mole of carbon increases and reduces TA by 0.15 units, respectively (Broecker and Peng, 1982). Calcification and dissolution of one mole of carbon decreases and increases TA by two units (Eq. 4). To verify the results...
from the model, we used the output $\Delta$DIC and $\Delta$TA to calculate new $pCO_2$ and pH values, which were further compared to the measured ones (Fig. 5). The best fit between modeled and measured values was achieved with a respective $R_{OI}$ of -2.6 for 2012 and 1.0 for 2013, whereas the assumption of calcium carbonate dissolution caused the negative sign.

\[
\Delta DIC = \Delta POC + \Delta PIC 
\]

(1)

\[
\Delta PIC = \left(\frac{\Delta POC}{R_{OI}}\right)
\]

(2)

\[
\Delta POC = \frac{\Delta DIC}{\left(1 + \left(\frac{1}{R_{OI}}\right)\right)}
\]

(3)

\[
\Delta TA = (\Delta POC \times 0.15) - \left(\frac{\Delta POC}{R_{OI}}\times 2\right)
\]

(4)

This was calculated on hourly time steps, separately for 2012 and 2013, using the mean SWT (2012 = 29.61 ± 0.93 °C, 2013 = 30.08 ± 0.27 °C) and salinity (2012 = 32.5, 2013 = 32.5).

3 Results

3.1 Carbonate chemistry during non-upwelling season

In June 2012, average SWT was 29.61 ± 0.93 (average ± standard deviation) °C and ranged from 27.13 °C to 31.37 °C. In May-June 2013 SWT ranged from 29.3 °C to 30.7 °C (average 30.08 ± 0.27°C). During both periods, the salinity was 32.5 ± 0.8. During the study periods, the wind intensified during the afternoons reaching speeds of up to 8.5 m s$^{-1}$ and 6.0 m s$^{-1}$ in 2012 and 2013, respectively (Fig. 2). Average pH and $pCO_2$ in June 2012 were 7.98 ± 0.04 and 456.38 ± 69.68 µatm, respectively; the corresponding averages for May-June 2013 were 8.02 ± 0.03 and 375.67 ± 24.25 µatm. Since the tidal cycle was not significantly correlated with the variability of pH, $pCO_2$, T or wind ($p > 0.05$) during the periods of observations (Table 2), it was excluded from further discussions. Mean $\Omega_a$ values were 3.32 ± 0.46 in June 2012 and 3.50 ± 0.49 in May-June 2013 (Table 1).

3.2 Seasonal variation of the carbonate system

Measured parameters showed significant differences between study periods ($p < 0.05$). The SWT range differed among years (Table 1); 2013 was the warmest study period, followed by 2012 and 2009. Lowest measured pH was 7.81 in June 2012, 7.84 in April 2009 and 7.95 in May-June 2013. We also compared DIC and TA, in order to estimate to which extend the observed variations of $pCO_2$ were caused by changes in temperature and/or DIC concentrations. Mean DIC values were 2098.71 ± 103.81 µmol kg$^{-1}$ in April 2009, 1924.65 ± 195.07 µmol kg$^{-1}$ in June 2012 and 1800.92 ± 142.78 µmol kg$^{-1}$ in May-June 2013. Similarly, mean TA values were 2328.42 ± 118.45 µmol kg$^{-1}$ in April 2009, 2204.54 ± 212.18 µmol kg$^{-1}$ in June 2012 and 2102.66 ± 174.79 µmol kg$^{-1}$ in May-June 2013. According to average values, April 2009 was the period with
most acidic water and greater CO$_2$ enrichment, followed by June 2012 and May-June 2013 (Table 1). Mean $\Omega_a$ values were 2.71 ± 0.29 during upwelling season (April 2009) and 3.41 ± 0.13 during non-upwelling season (June 2012, May-June 2013), resulting in annual average $\Omega_a$ of 3.06 ± 0.49 at Bahía Culebra. Time series of pH and $p$CO$_2$ in June 2012 and May-June 2013 showed a pronounced daily cycle (Fig. 4), which in addition to previously described data will be discussed in the following paragraphs.

4 Discussion

4.1 Natural OA beyond the upwelling season

The observed differences in pH and $p$CO$_2$ between 2012 and 2013 suggest that the non-upwelling season exhibits a strong interannual variability (Table 1). In 2012 pH was lower and $p$CO$_2$ higher than in 2013 (Fig. 2b, c). The June 2012 time-series data showed that SWT decreased and $p$CO$_2$ increased from 300 to 650 µatm in less than a week, after several days of strong afternoon winds (Fig. 2a). Similarly, this increase in $p$CO$_2$ was accompanied by a dropped in pH form 8.04 to 7.83 (Fig. 2a). This suggests that an enhanced wind-driven vertical mixing entrained cooler and CO$_2$-enriched waters from greater water-depth into the surface layer. The associated SWT drop from 31.4 °C to 27.1 °C was similar to that observed during the onset of the 2009 upwelling event (26.2 °C to 23.7 °C; Rixen et al., 2012). Nevertheless, the higher SWT during the 2012 non-upwelling season suggests that the entrained water originated from a shallower water-depth, compared with the water upwelled in 2009. The $p$CO$_2$ values with up to 650 µatm reached the same level during both events, which is partially caused by the higher SWT in 2012. However, DIC concentrations in 2012 (1924.65 ± 195.07 µmol kg$^{-1}$) were lower than those in 2009 (2098.71 ± 103.81 µmol kg$^{-1}$), but exceeded those in 2013 (1800.92 ± 142.78 µmol kg$^{-1}$, Table 1). During the seven days that lasted the cold-water intrusion event in 2012 (June 10-17), the DIC concentrations dropped from 2355.39 µmol kg$^{-1}$ down to 1715.30 µmol kg$^{-1}$. This implies that in addition to high SWT, the entrainment of CO$_2$-enriched subsurface water increased the $p$CO$_2$ not only during the upwelling periods, but also during the 2012 non-upwelling season. Since in 2012 the $p$CO$_2$ had already increased by June 7th and the SWT decreased two days later (June 10th), the inflow of CO$_2$-enriched waters seems to have increased the $p$CO$_2$ already prior to the strengthening of local winds (Fig. 2b). Later, local wind-induced vertical mixing seems to have amplified the impact of the inflowing CO$_2$-enriched water mass on the $p$CO$_2$ in the surface water by increasing its input into surface layers. Accordingly, the CO$_2$-enriched waters were apparently supplied from a different location than they are during upwelling season. Since the NECC carries offshore waters towards the Costa Rican shore during the non-upwelling season (Wyrtki, 1965, 1966; Fiedler, 2002), it is assumed that the CO$_2$-enriched subsurface water originated somewhere south of our study area in the open ETP. The absence of such a cold-event during the non-upwelling season in 2013 suggests that the occurrence of this kind of events might be an irregular feature (Fig. 2c, d), and the driving forces are still elusive. Nevertheless, these types of events have the potential to affect the metabolic processes in the bay as will be discussed in the following section, which analyzes the daily cycles during the non-upwelling seasons in 2012 and 2013.
4.2 Processes behind the variability of the carbonate system

In 2012, the pH and the $pCO_2$ values followed a pronounced diurnal cycle with highest pH and lowest $pCO_2$ values during the late afternoon and lowest pH and highest $pCO_2$ values around sunrise in the early morning (Fig. 4a). Such daily cycles are typical for tropical regions and are assumed to be caused by photosynthesis during the day and respiration of organic matter during the night (Shaw et al., 2012; Albright et al., 2013; Cyronak et al., 2013a). The aragonite saturation state as well as the DIC/TA ratio followed this pattern, with higher $\Omega_a$ and lower DIC/TA ratio values during the day as well as lower $\Omega_a$ and higher DIC/TA values at night (Fig. 4b). Although the $pCO_2$ cycles in 2013 followed a similar pattern to 2012, pH cycles were less predictable (Fig. 4).

To characterize the relative importance of the processes responsible for the observed changes in pH and $pCO_2$ (photosynthesis, respiration, calcification and dissolution) we used the model described earlier, which is based on the determined DIC concentrations during times when pH and $pCO_2$ revealed their daily minima and maxima, respectively. For example, if photosynthesis of organic matter dominates the transition from early morning maxima of $pCO_2$ to late afternoon minima of $pCO_2$ it should be associated with a decline in DIC. Whether photosynthesis was accompanied with enhanced calcification can be detected by an associated decrease of TA. Since decreasing DIC raises the pH and a decrease in TA lowers the pH, such photosynthetic enhanced calcification hardly affects the pH and could explain the weak daily cycle observed in 2013. Alternatively, if photosynthesis is accompanied by carbonate dissolution during the day, this would amplify the daily cycle of pH and $pCO_2$ as seen during the cold-water intrusion event in 2012. Likewise, an increased photosynthesis resulting from higher nutrient concentrations (Pennington et al., 2006) could also be causing the observed large amplitude during the event in 2012. However, in our case the determined TA and DIC concentrations constrain the impact of the formation of organic matter (POC = photosynthesis – respiration) and calcification (PIC = calcification – dissolution) on the carbonate system. This sets the boundaries within which the observed diurnal cycle of pH and $pCO_2$ has to be explained (Fig. 5c, d). In order to reconstruct the diurnal cycle of pH and $pCO_2$ within these boundaries we assumed a photosynthetic-enhanced calcification during the day and vice versa, dissolution and respiration at night. Thereby the best fit between pH and $pCO_2$ measured in 2013 and the respective calculated values could be obtained by using a $R_{OI}$ of 1. This approach failed to explain the diurnal cycle of pH and $pCO_2$ as observed during the 2012 cold-water intrusion event (June 10-17). The only solution we found to explain these pronounced diurnal cycles within the given DIC and TA boundaries was to assume that photosynthesis and dissolution prevailed during the day and respiration and calcification occurred at night. The $R_{OI}$ of -2.6 resulted in the best fit between the measured and calculated pH and $pCO_2$ for the 2012 event, whereas the negative sign reflects the contrasting effects of calcification and dissolution on the DIC concentration.

Dissolution taking place during daytime is peculiar but not completely unusual, as it has been reported on tropical sandy bottoms under ambient (Yates and Halley, 2006; Cyronak et al., 2013b) and high-CO$_2$ conditions (Comeau et al., 2015). Similarly, dark-calcification is not entirely uncommon and occurs in both, sandy bottoms and coral reefs (Yates and Halley, 2006b; Albright et al., 2013). Accordingly, the entrainment of CO$_2$-enriched water from the NECC seems to shift the
carbonate chemistry of Bahía Culebra from a system where photosynthesis and calcification are the controlling processes during light hours to a system in which daytime is dominated by photosynthesis and dissolution. The net effect, as observed, is an enhanced $\rho\text{CO}_2$ and lower $\Omega_a$ during periods characterized by the inflow of CO2-enriched waters (Table 1). This has strong ecological implications for local coral reef ecosystems.

4.3 Ecological implications for coral reefs

Coral reefs in Bahía Culebra were dominated by *Pocillopora* spp. and *Pavona clavus* (Jiménez, 2001; Jiménez et al., 2010), whereas *Porites lobata* is the main reef forming coral in the southern part of the Costa Rican Pacific coast (Cortés and Jiménez, 2003; Glynn et al., 2017). Although the reefs in the north are naturally exposed to periodic high-CO2 conditions during upwelling events (Rixen et al., 2012), as well as during cold water intrusions in non-upwelling season, the linear extension rates of *Pocillopora* spp. and *P. clavus* exceeded those of the same species in other regions (Fig. 6) (Glynn, 1977; Jiménez and Cortés, 2003; Manzello, 2010a; Rixen et al., 2012). This suggests that local corals are adapted and/or acclimatized to the upwelling of cold and acidic waters.

Aragonite saturation state ($\Omega_a$) is known as one of the main variables influencing coral growth and therefore reef distribution around the world (Kleypas et al. 1999). By integrating the data from the present study and values previously reported by Rixen et al. (2012), we estimated that the annual mean $\Omega_a$ in Bahía Culebra is 3.06. Additionally, earlier studies in the ETP measured $\Omega_a$ values and coral extension rates from locations that are under the influence of upwelling events (Manzello 2010a), whilst extension rates from Bahía Culebra were measured by Jiménez and Cortés (2003). The correlation between our estimated $\Omega_a$ with the available data from Bahía Culebra, Panamá and Galápagos indicates that coral extension rates in each of those locations are predictable by their corresponding $\Omega_a$ values (Fig. 6).

The dependency of coral growth on $\Omega_a$ and the mean $\Omega_a$ (2.71) during the upwelling season (Table 1) suggests that upwelling of acidic waters should reduce corals’ relatively high annual mean growth rates in Bahía Culebra. The increased $\Omega_a$ during non-upwelling season in turn must enhance linear extension and explains corals’ high annual mean growth rates. The $\Omega_a$ values from this study suggest that most favourable conditions for coral growth occur during non-upwelling season, the period that coincides with development of the rainy season. This implies that during the main growing season the eutrophication and siltation caused by human impacts on river discharges, as well as the development of harmful algal blooms, could also strongly affect the corals’ annual mean growth rates (Cortés and Reyes-Bonilla, 2017).

Despite the corals’ high annual mean linear extension rates, studies carried out in 1973 showed that the thickness of the reef framework within our study area was with 0.6 to 3 m (mean 1.8 m) among the lowest in the ETP, where Holocene framework accumulation in *Pocillopora*-dominated reefs could reach up to 9 m (Glynn et al., 1983; Toth et al., 2017).

During the last decade it further decreased (Alvarado et al., 2012), and during the period of our observation the reef frameworks of *Pocillopora* spp. in Bahía Culebra hardly exceeded a thickness of 0.5 m. This denotes that although *Pocillopora* spp. and *P. clavus* are adapted to the entrainment of acidic waters, these reefs are growing in an environment at
the limit of reef-building corals tolerance in terms of temperature, nutrient loads and pH (Manzello et al., 2017). Gaps in coral reef accretion at the ETP are known from the geological record (Toth et al., 2012; 2015; 2017). They have been linked to increased ENSO variability (Toth et al., 2012, 2015) and stronger upwelling conditions (Glynn et al., 1983), favouring dissolution and erosion of reef frameworks while at the same time restricting coral growth.

The y-intercept of the regression equation derived from the correlation between linear extension rates and Ωa furthermore implies that linear extension of P. damicornis and P. clavus should approach zero under a carbonate saturation state of Ωa < 2.5 (P. damicornis) and < 2.2 (P. clavus). According to climate predictions, the global Ωa will reach values < 2.0 by the end of this century (IPCC, 2014), and major upwelling systems such as those off California and South America will intensify (Wang et al., 2015). Combined effects of ocean acidification and impacts of stronger upwelling on Ωa in the ETP and on Ωa in Bahía Culebra are difficult to predict. Worldwide, OA is expected to reduce coral reefs’ resilience by decreasing calcification and increasing dissolution and bioerosion (Kleypas et al., 1999; Yates and Halley, 2006a; Anthony et al., 2011). Coral reefs from the ETP are affected by chronic and acute disturbances, such as thermal stress and natural ocean acidification resulting from ENSO and upwelling events, respectively (Manzello et al., 2008; Manzello, 2010b). Historically, these reefs have shown a high resilience to both stressors by separately but their coupled interaction can cause coral reef lost within the next decades. The ETP have the lowest Ωa of the tropics, near to the threshold values for coral reef distribution, therefore the reefs from this region may be the most affected by the increasing levels of anthropogenic CO2 and also show the first negative impacts of this human induced OA (Manzello et al., 2017). This emphasizes the importance of the Paris agreement and all the global efforts to reduce the CO2 emission into the atmosphere (Figueres et al., 2017).

5 Conclusions

The present study provides data from in situ measurements from a system that is naturally exposed to low-pH conditions, and seeks to characterize the carbonate chemistry within a bay (Bahía Culebra) and its potential impact on the reefs. This study builds on previous field studies in the upwelling areas of Panamá (Manzello et al., 2008; Manzello, 2010b) and Papagayo (Rixen et al., 2012). Our results indicate that physical processes, such as the coastal upwelling and the exchange of water between the bay and the open ocean, influence the carbonate chemistry on timescales of weeks to months, where metabolic processes (photosynthesis and calcification) influence the diurnal cycle. To which extend benthic and pelagic processes control the diurnal cycle, cannot be established based on our data. However, the results from the present study also suggest that coral reefs from Bahía Culebra are exposed to a high intra- and interannual variability in the carbonate system. Challenging conditions for reef development are not restricted to the upwelling season, they occur sporadically also during non-upwelling season, when pH and CO2 concentrations reach values comparable to those during upwelling events. Previous studies reported that the linear extension rates measured in Bahía Culebra were among the highest in the ETP, thus is likely that coral growth in this bay is enhanced with increased Ωa during periods with no entrainment of low-pH waters. However, coral growth must be measured during both seasons in order to confirm this assumption. Threshold values of Ωa when coral growth likely approaches zero were derived from the correlation of Ωa and previously measured annual linear extension
rates. The $\Omega_a$ threshold values from the present study and the fact that high-CO$_2$ waters are occasionally hauled in to the bay during non-upwelling season; suggest that coral reef development in Bahía Culebra is potentially threatened by anthropogenic OA.

6 Data availability

Data are available by direct request to the corresponding author.

7 Author contribution

C. Sánchez-Noguera and T. Rixen designed the study, analyzed the data, prepared figures and/or tables and wrote the paper C. Sánchez-Noguera collected and analyzed the samples. I. Stuhldreier, J. Cortés, Á. Morales, C. Jiménez and C. Wild reviewed the paper.

8 Competing interests

The authors declare that they have no conflict of interest.

9 Disclaimer

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References


Table 1: Measured and calculated (*) parameters, during upwelling (2009) and non-upwelling seasons (2012, 2013) at Bahía Culebra, Costa Rica.

<table>
<thead>
<tr>
<th></th>
<th>pH (Total scale)</th>
<th>$p\text{CO}_2$ (µatm)</th>
<th>CO$_2$ (µmol kg$^{-1}$)</th>
<th>T (°C)</th>
<th>DIC* (µmol kg$^{-1}$)</th>
<th>TA* (µmol kg$^{-1}$)</th>
<th>Ω*</th>
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</thead>
<tbody>
<tr>
<td>2009</td>
<td>Mean ± SD</td>
<td>7.91 ± 0.32</td>
<td>578.49 ± 42.82</td>
<td>16.44 ± 1.35</td>
<td>25.09 ± 0.57</td>
<td>2098.71 ± 103.81</td>
<td>2328.42 ± 118.45</td>
</tr>
<tr>
<td>2012</td>
<td>Mean ± SD</td>
<td>7.98 ± 0.04</td>
<td>456.38 ± 69.68</td>
<td>11.77 ± 1.99</td>
<td>29.61 ± 0.93</td>
<td>1924.65 ± 195.07</td>
<td>2204.54 ± 212.18</td>
</tr>
<tr>
<td>2013</td>
<td>Mean ± SD</td>
<td>8.02 ± 0.03</td>
<td>375.67 ± 24.25</td>
<td>9.56 ± 0.64</td>
<td>30.08 ± 0.27</td>
<td>1800.92 ± 142.78</td>
<td>2102.66 ± 174.79</td>
</tr>
</tbody>
</table>
Table 2: Correlations between tide height and four parameters during non-upwelling season (2012, 2013).

<table>
<thead>
<tr>
<th>Year</th>
<th>pH</th>
<th>$\rho$CO$_2$</th>
<th>T</th>
<th>Wind</th>
</tr>
</thead>
<tbody>
<tr>
<td>2012</td>
<td>-0.004</td>
<td>0.037</td>
<td>-0.005</td>
<td>0.033</td>
</tr>
<tr>
<td>2013</td>
<td>0.111</td>
<td>0.026</td>
<td>-0.093</td>
<td>-0.126</td>
</tr>
</tbody>
</table>

All p-values > 0.05
Figure 1: Location of Bahía Culebra (square) in the Gulf of Papagayo, North Pacific coast of Costa Rica (insert). Measurements were made at Marina Papagayo (star). Main ocean currents influencing the Gulf of Papagayo (dashed arrows): NECC= North Equatorial Counter Current, CRCC= Costa Rica Coastal Current.
Figure 2: Measured parameters (wind speed, SWT, pH and $pCO_2$) during the non-upwelling seasons of June 2012 (a, b) and May-June 2013 (c, d), at Bahía Culebra. Shaded area in (a) and (b) indicates the 2012 upwelling-like event.
Figure 3: Validation of in situ measurements of pH (a) and $p$CO$_2$ (b) using discrete water samples. SAMI sensors measured pH and $p$CO$_2$ directly in the water column. The pH and $p$CO$_2$ values used for validation were calculated with the CO2SYS program as a function of measured TA and DIC; discrete samples were measured with a VINDTA 3C system.
Figure 4: Diel pattern of parameters measured in Bahía Culebra. Data points are hourly averages of 15 and 7 consecutive days in 2012 (a, b) and 2013 (c, d), respectively. The shaded area represents daylight hours.
Figure 5: Expected diel behaviour of the carbonate system in 2012 (a, b) and 2013 (c, d), based on measured parameters. Modeled parameters are shown as blue crosses and empty circles; the reference parameter used to adjust the model is shown in black triangles. Shaded area represents daylight hours.
Figure 6: Mean aragonite saturation states ($\Omega_a$) – from present and former studies - versus previously reported mean linear extension rates of (a) *Pocillopora damicornis* and (b) *Pavona clavus* from upwelling areas in Costa Rica (CR) (Jiménez and Cortés, 2003), Panamá (PAN) and Galápagos (GAL) (Manzello, 2010a). Red broken line shows the regression equation as estimated by Rixen et al. (2012). Red mark represents our estimated $\Omega_a$ threshold for Bahía Culebra, when coral growth equals zero.