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PRIMARY RESEARCH PAPER

Changes in microphytobenthos fluorescence over a tidal cycle: implications for sampling designs

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Abstract Intertidal microphytobenthos (MPB) are important primary producers and provide food for herbivores in soft sediments and on rocky shores. Methods of measuring MPB biomass that do not depend on the time of collection relative to the time of day or tidal conditions are important in any studies that need to compare temporal or spatial variation, effects of abiotic factors or activity of grazers. Pulse amplitude modulated (PAM) fluorometry is often used to estimate biomass of MPB because it is a rapid, non-destructive method, but it is not known how measures of fluorescence are altered by changing conditions during a period of low tide. We investigated this

experimentally using in situ changes in minimal fluorescence (F_o^{15}) on a rocky shore and on an estuarine mudflat around Sydney (Australia), during low tides. On rocky shores, the time when samples are taken during low tide had little direct influence on measures of fluorescence as long as the substratum is dry. Wetness from wave-splash, seepage from rock pools, run-off, rainfall, etc., had large consequences for any comparisons. On soft sediments, fluorescence was decreased if the sediment dried out, as happens during low-spring tides on particularly hot and dry days. Surface water affected the response of PAM and therefore measurements used to estimate MPB, emphasising the need for care to ensure that representative sampling is done during low tide.

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Introduction

Intertidal microphytobenthos (MPB) are important primary producers in soft sediments (Yallop et al., 1994; MacIntyre & Cullen, 1996; Underwood et al., 2005) and on rocky shores (Castenholz, 1961; Nicotri, 1977; Magalhães et al., 2003). They contribute to carbon budgets (Sullivan & Moncrieff, 1988; Underwood & Kromkamp, 1999), can stabilize sediments (Tolhurst & Chapman, 2007) and provide food for

herbivores (Castenholz, 1961; Hawkins et al., 1992). Spatially, amounts of intertidal MPB vary considerably at scales from millimetres (Blanchard, 1990; Hutchinson et al., 2006; Jackson et al., 2009) to kilometres (Jenkins et al., 2001; Thompson et al., 2005), with a lot of the variation at the smallest scales measured (e.g. Saburova et al., 1995; Chapman & Tolhurst, 2007). Measures of temporal variation on rocky shores have mostly been about changes at large temporal scales (e.g. monthly, seasonal or inter-annual variability; Underwood, 1984a; Jenkins et al., 2001; Thompson et al., 2005; Jackson et al., 2010), with no studies of short-temporal variation in MPB (e.g. diel or tidal variation). Such large-scale comparisons would, however, be compromised if measures of intertidal MBP abundance are affected by temporal variation throughout periods of low tide. When such variation is not considered, large-scale, seasonal or inter-annual comparisons would be confounded (e.g. Jenkins et al., 2001).

Many intertidal micro-algae in soft sediments have diel and/or tidal migratory behaviour and complex periodicities controlled by availability of light and timing of emersion (Round & Palmer, 1966; Serôdio et al. 1997; Perkins et al., 2003). Although Lamontagne et al. (1989) suggested that migration by MPB on rocky substrata is unlikely, some species of diatoms are endolithic and migrate in the rock (Houpt, 1994). Migratory patterns (usually upward migration during periods of emersion during daylight hours; Round & Palmer, 1966; MacIntyre et al., 1996) have been also observed primarily in muddy estuaries which are often dominated by diatoms (e.g. Blanchard et al., 2001; Tolhurst et al., 2003). Composition of MPB assemblages does, however, vary globally. For example, in south-east Australia, rocky shores are often dominated by cyanobacteria (Jackson et al., 2010) and estuaries may have patches of filamentous green algae (Chapman & Tolhurst, 2004, 2007), which are not migratory. Dominance by these algae in mangrove forests may explain Tolhurst & Chapman's (2005) results that there was little effect of time of low tide or time during a tidal cycle on concentrations of chlorophyll.

Intertidal systems have emersed and submersed conditions twice daily. Changes in environmental conditions associated with the tide (e.g. temperature, light, water-content, abrasion, etc.) can be stressful to MPB (Joint et al., 1982; Helmuth et al., 2002; Perkins et al., 2003; Easley et al., 2005). Although many

studies have shown spatial and temporal variations in chlorophyll (e.g. Blanchard, 1990; Thompson et al., 2004) and photosynthetic activity (e.g. measured by PAM fluorescence; Jesus et al., 2005; Serôdio et al., 2005; Tolhurst et al., 2006), the reasons for these variations are not always clear. They are, however, potentially confounded by variations in the timing of sampling relative to the state of the tide. For example, physiological and physical factors that vary during emersion/submersion (Serôdio et al., 1997; Perkins et al., 2001; Cohn et al., 2003; Roncarati et al., 2008; Coelho et al., 2009) affect these measurements and the timing of sampling is not consistent across sites or larger time-scales. If there is substantial variation from one time to another during the period since the tide fell, it is important to standardise the timing of sampling or to sample at several defined times throughout low tide to be able to compare data from one time or place to another. Otherwise, any difference from time to time (e.g. season to season) may simply reflect differences at different stages of the low-tidal cycle. Despite this requirement, most studies of large-scale temporal patterns in MPB biomass have simply reported that sampling was done 'during low tide', without any further information on the timing relative to the state of the tide. Thus, we do not currently know how representative any of these measures of MPB are. Knowing whether, where and when measurements of MPB biomass change during a tidal cycle is therefore essential for planning sampling that is to be comparable across studies or across large spatial or temporal scales.

Methods for sampling and quantifying MPB involve technical difficulties and are time-consuming, particularly on hard substrata (Underwood, 1984b; Hill & Hawkins, 1990; Nagarkar & Williams, 1999). Therefore, rapid, non-destructive methods, such as field spectrometry (Murphy et al., 2005a), digital infra-red photography (Murphy et al., 2004; Murphy et al., 2006) and pulse amplitude modulated (PAM) fluorometry (Serôdio et al. 1997; Honeywill et al., 2002; Consalvey et al., 2005; Jesus et al., 2005), have been developed. PAM is a rapid and widely used method, also suitable in systems (such as Mediterranean or Baltic microtidal habitats) where the presence of water (e.g. from wave-splash, seepage from rock pools, run-off, etc.) can prevent the use of alternative non-destructive methods for estimates of MPB abundance. Values of F_0^{15} (minimal fluorescence after

15 min dark adaptation) have been positively correlated with amounts of chlorophyll *a* and biomass of MPB (Serôdio et al., 1997; Honeywill et al., 2002).

We used PAM fluorometry to investigate in situ changes in minimal fluorescence (F_o^{15}) on a rocky shore and on an estuarine mudflat during periods of emersion, to test the hypothesis that fluorescence would vary throughout the period between the fall and the subsequent rise of water during a low tide. We predicted more change (if any) in the sediments, where motile micro-algae were more likely to be present and could more easily migrate vertically compared to the rocky shore. A decrease in fluorescence on the rocky shore was observed as the tide fell and an increase when it rose (see 'Results' section). We proposed the explanation that changes in F_o^{15} were caused by surficial water. This was examined experimentally. On the mudflat, fluorescence decreased during emersion, so we tested whether this was due to water in the surface of the sediment via experiments in which water was either added during low tide or was removed by draining the sediments.

Materials and methods

Studies were done on a rocky shore in the Cape Banks Scientific Marine Research Area, Botany Bay (hereafter CB; 33°59'54"S; 151°14'39"E) and on two estuarine mudflats at Tambourine Bay (hereafter TB; 33°49'41"S; 151°09"E) and Fig Tree Bridge (hereafter FTB; 33°49'44"S; 151°08'44"E) in Sydney Harbour, NSW. In these areas, tides are semidiurnal with a maximal range of ~2 m. At Cape Banks, rocky epilithic assemblages were dominated by cyanobacteria; in addition, spectral signatures were never similar to assemblages of diatoms (Jackson et al., 2010). During a study of chlorophyll distribution on the Fig Tree Bridge and Tambourine Bay mudflats in Sydney harbour (Murphy et al., 2005b), the shape of the spectra and the peak at green wavelengths indicated that MPB assemblages were dominated by green algae (mostly filamentous green algae, which were distributed on the surface of and in between grains of sediment in the upper few millimetres of the sediment surface) (R. Murphy, pers. comm.). The upper millimetres of the sediment surface has a grain size <63 µm (corresponding to mud) of 24% at Tambourine Bay and 29% at Fig Tree Bridge.

Rocky shore

At CB, two randomly selected sites (stretches of ~10 m, about 200 m apart) at each of two heights on the shore were used. The first ('Mid', about 1.6 m above Chart Datum) was approximately in the middle of the vertical distribution of an intertidal assemblage of species dominated by grazing molluscs (Underwood, 1980), the most common of which were the limpet *Cellana tramoserica* (Sowerby) and the snails *Bembicium nanum* (Lamarck) and *Nerita atramentosa* (Reeve). The second height ('Low', about 1.4 m above Chart Datum) was where there were more macro-algae, mostly the encrusting red *Hildenbrandia rubra* (Sommerfelt) Meneghini and brown *Ralfsia verrucosa* (Areschoug) Areschoug algae. At each site and height, four patches (30 × 25 cm) were scrubbed with hydrochloric acid in November 2006 to remove all organisms so that measurements of fluorescence from epilithic MPB were not confused by encrusting macro-algae (Underwood, 1984b). The cleared space was rapidly recolonised by MPB (Murphy et al., 2006) and at 6 months, the reflectance spectra indicated that patches were still colonised by cyanobacteria and possibly some diatoms (R. Murphy, pers. comm.) with no macro-algae.

Observational study

On March 28th 2007, three replicate measures of fluorescence were taken in each of three 'Mid' patches at one site. Patches were sampled every 5 min, starting when the rock was exposed by the ebbing tide and finishing when the patches were wetted by the rising tide. Care was taken to ensure that the same areas of a patch were not sampled twice.

Experimental addition of water

An experiment to test the effect of surficial water on fluorescence were done on April 13th and then repeated on May 14th, 2007. At each site and height, two patches were randomly allocated to each of Control and +Water treatments. Control treatments were allowed to dry naturally, and +Water treatments were kept wet throughout low tide by adding seawater from a tank via a perforated rubber hose. Water flow was maintained at a rate that ensured the entire patch was covered by a thin film of water. Fluorescence

Table 1 Summary of the experimental designs

| | | |
|--|--|-----------------------------|
| Rocky shore | | |
| Experimental addition of water (on April 13th and then repeated on May 14th, 2007) | | |
| Site = S ^a | Random, 2 levels (Site 1, Site 2) | Orthogonally factorial |
| Height = H | Fixed, 2 levels (Mid, Low) | |
| Time | Fixed, 6 levels (B, E, M1, M2, L, A) | Nested within S × H × Treat |
| Treatment = Treat | Fixed, 2 levels (Control, +Water) | |
| Patch | Random, 2 levels (Patch 1, Patch 2) | |
| Replicate | Random, 3 | |
| Soft sediment | | |
| Experimental addition of water (on May 15th, 2007) | | |
| Site = S | Random, 2 levels (Site 1, Site 2) | Orthogonally factorial |
| Time | Fixed, 7 levels (B, E, M1, M2, M3, L, A) | |
| Treatment = Treat | Fixed, 2 levels (Control, +Water) | |
| Replicate | Random, 3 | |
| Draining experiment (on May 29th, 2007) | | |
| Time | Fixed, 3 levels (E, M, L) | Orthogonally factorial |
| Treatment = Treat | Fixed, 2 levels (Control, Drained) | |
| Replicate | Random, 5 | |

B = just before the mud was emersed; E = early after emersion; M1, M2 and M3 = at three times in the middle in the period of emersion; L = late in the period of emersion; A = just after the water covered the substratum

^a On April 13th, the experiment was done only at Site 1

($n = 3$) was measured in each patch six times: just before emersion (B; approximately 1 h before emersion), early during emersion (E; approximately 5–10 min after emersion), twice in the middle of emersion (M1 and M2; approximately 1–1.30 and 2–2.30 h after emersion, respectively) just before the patches were wetted by the rising tide (L; approximately 3–3.30 h after emersion) and finally just after the rising tide covered the patches (A; approximately 5–10 min after water covered the plot; Table 1). The precise timing of sampling differed among patches, due to sampling constraints and differences in elevation (hence differences in the times of emersion and submersion).

Mudflat

Observational study

Changes in fluorescence during emersion were first measured on an intertidal mudflat at TB on March 15th, 2007. Replicate measurements (about 50 cm apart; $n = 168$) were taken every 2–5 min at the same

height along a 40-m stretch of shore, starting just after the tide receded and finishing before the water covered the substratum. The same areas were not sampled twice.

Experimental addition of water

The first experiment to test the effect of presence of water on fluorescence was done on May 15th, 2007, in similar habitat at FTB. In each of two sites (40 m long, 5 m apart), 42 areas of sediment (150 mm in diameter) were selected at the same height. These were randomly allocated to Control and +Water treatments. Controls were allowed to dry naturally, and +Water treatments were kept wet throughout the low tide, by retaining water in surrounding plastic rings (150 mm in diameter, 200 mm high) pushed 100 mm into the mud before the tide ebbed. These were replenished with water as required. Previous experiments had demonstrated that the rings themselves did not affect the fluorescence of algae [mean F_o^{15} in unmanipulated sediment = 429 (\pm SE 26, $n = 3$); mean F_o^{15} in sediment inside plastic rings = 416 (\pm SE 30,

$n = 3$]. Times were: just before the mud was emersed (B; approximately 5–10 min before emersion), early (E; approximately 10–20 min after emersion), at three times in the middle (M1, M2 and M3; approximately 1, 2 and 3 h after emersion, respectively), late (L; approximately 4 h after emersion) during the period of emersion and finally just after the water covered the mud again (A; approximately 10 min after being covered). The fluorescence for each area was calculated from the mean of three measurements in each plot (Table 1).

Experimental draining

An experiment to investigate the effect of desiccation of sediment on fluorescence was done on May 29th, 2007, at one of the two sites at FTB. There were two treatments: 15 cores of sediment (7 cm in diameter, 5 cm deep) were drained during a low tide (Drained) and 15 cores were left as Controls. The drained cores were removed from the mudflat at the start of low tide and placed on a rigid mesh sheet raised above the substratum. The sediment was coherent, and the cores did not disintegrate. The surfaces of the cores were not touched during this procedure. This treatment let the sediment drain faster than the Control cores which were left in situ. At each of three times [Early (E; approximately 15 min after emersion), Mid (M; approximately 1.3 h after emersion) and Late (L; approximately 3 h after emersion)], fluorescence was measured in five randomly chosen cores of each treatment (Table 1).

Measurement of fluorescence

Algae were dark-adapted for 15 min (Kromkamp et al., 1998; Honeywill et al., 2002) and minimal fluorescence (F_o^{15}) measured with a PAM fluorometer (Diving PAM, Walz, Germany; used settings: MI = 12, G = 9). The probe was kept at 4 mm above the substratum and measured an area of approximately 24 mm². The zero offset (measurement of background signal without a sample) was done by pointing the fibre-optic cable at the sky away from the sun. Where two fluorometers were used concurrently on the rocky shore, they were cross-calibrated using measurements of fluorescence standards to ensure comparability of data between devices. Correlations were linear and strong (e.g. $r = 0.99$, d.f. = 8, $P < 0.001$).

Results

Rocky shore

Observational study

Minimal fluorescence (F_o^{15}) decreased rapidly when micro-algae were first emersed, remained nearly constant during low tide and then increased rapidly again when the substratum was wetted by the rising tide (Fig. 1).

Experimental addition of water

In general, decreases in F_o^{15} were smaller where the experimental rock surfaces were kept wet through the low tide than in controls which dried (Fig. 2).

On April 13th, one PAM developed a fault, so data were only collected from one site in the first run of the experiment. There was a significant interaction between Time and Patch (Treatment) (ANOVA; $F_{20,96} = 2.31$, $P < 0.01$); but the patterns of change among patches were similar for each treatment, differing only in magnitude. SNK tests on the significant interaction between Time and Treatment ($F_{5,5} = 4.32$, $P < 0.01$) showed less fluorescence in Control samples between M1 and L than at other times (Fig. 2a), but this was not the case in patches where

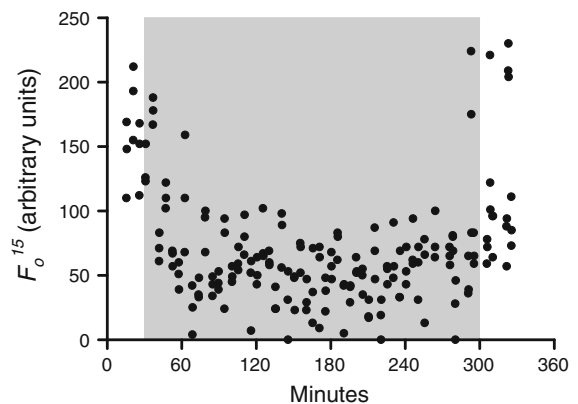
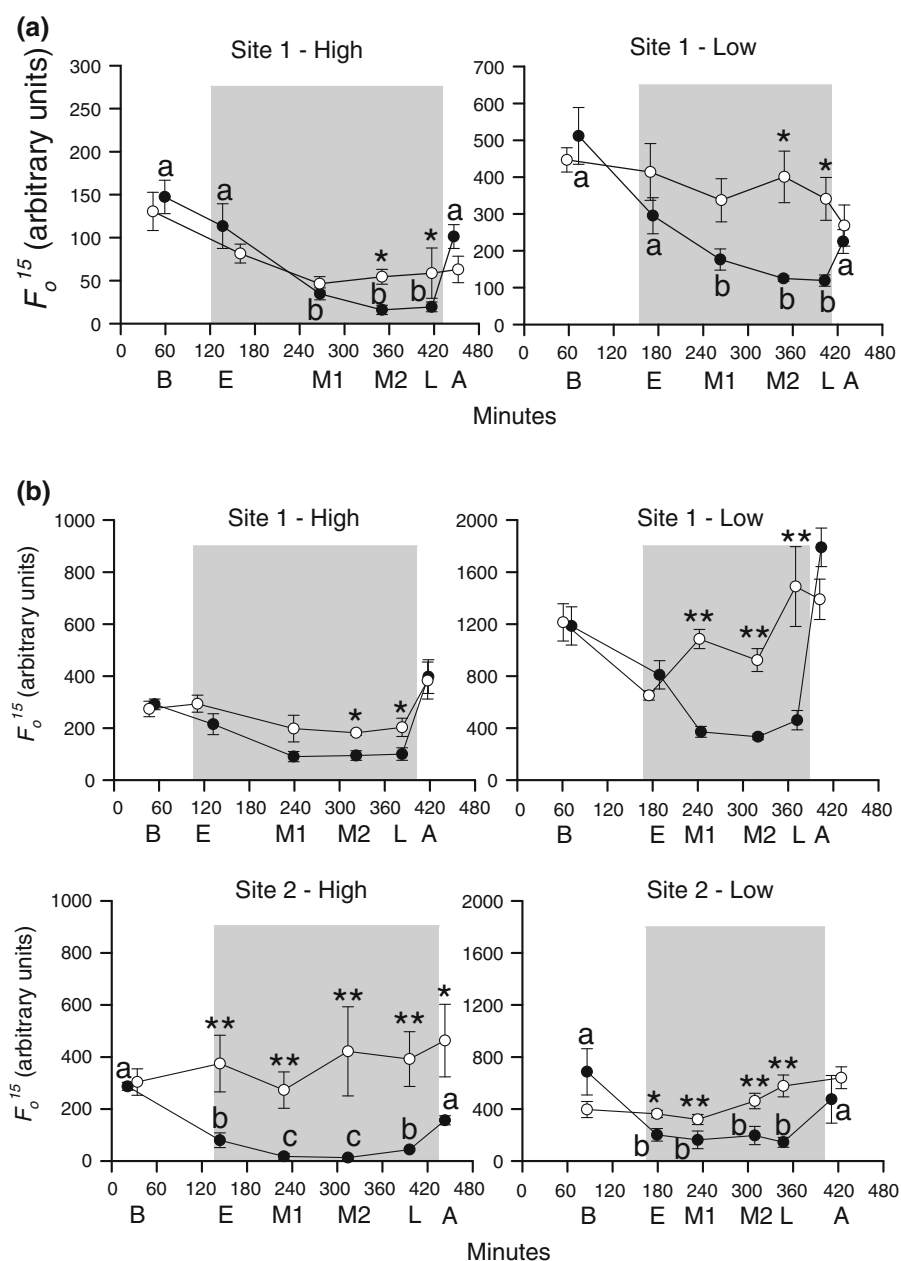


Fig. 1 Individual F_o^{15} from micro-algae on a rocky shore (CB; Site 1, high level) every 5 min during a diurnal low tide, starting when patches were still slightly submerged and finishing just after they were covered again by the rising tide. Low tide was at 11.25 a.m. on 28/03/2007. For this and the subsequent two figures, fluorescence is plotted against the time (min) elapsed since the start of the experiment and *grey shading* indicates when experimental patches were emersed

Fig. 2 Mean (\pm SE; $n = 3$) F_o^{15} values in control (filled circle) and +Water (open circle) treatments on a rocky shore (CB) at different Times (B = before emersion, E = early, M1 = middle 1, M2 = middle 2, L = late in the emersion period, A = after the rising tide re-covered the patches) for **a** April 13th 2007, low tide at 11.05, one site and **b** May 14th 2007, low tide at 11.50, two sites. Different letters indicate means that differed significantly among times ($P < 0.05$) for Control patches. Asterisks indicate means that differed significantly between Control and +Water treatments ($*P < 0.05$; $**P < 0.01$)



water was added. At Times M2 and L, fluorescence in Control patches was significantly less ($P < 0.05$) than in +Water patches (Fig. 2a). It is worth noting that, at the High height, values of F_o^{15} were very small (< 100) during the emersion period (i.e. between Times E and L), so that results have to be interpreted with caution.

In the second run of the experiment, fluorescence varied between sites and heights on the shore (a

significant interaction Height \times Site \times Treatment \times Time, $F_{5,187} = 4.59$, $P < 0.01$). In Controls at Site 2, F_o^{15} was greater when the substratum was wet (at the beginning and end of the experiment) than during emersion (Fig. 2b). This was similar at Site 1, although differences were not statistically significant (Fig. 2b). When Control patches were dry (i.e. between Times E and L), measurements of F_o^{15} were

generally larger in the +Water treatment than in the Controls. In 13 of the 16 comparisons (2 Heights, 2 Sites and 4 Times [E, M1, M2, L]), F_o^{15} was significantly greater in +Water than in Controls (at $P < 0.05$ in SNK tests). The probability of 13 of 16 cases in the same direction is small ($P < 0.05$, Binomial test).

In some analyses, variances were heterogeneous among samples, but results were considered interpretable because of the large number of degrees of freedom in the analyses (Underwood, 1997). At the last time of sampling (A) on the rocky shore, some dark adaptation chambers were washed away by large waves. To maintain a balanced design, missing data were replaced by the mean value obtained from the remaining replicates in the same treatment; the number of degrees of freedom was correspondingly reduced (Underwood, 1997).

Mudflats

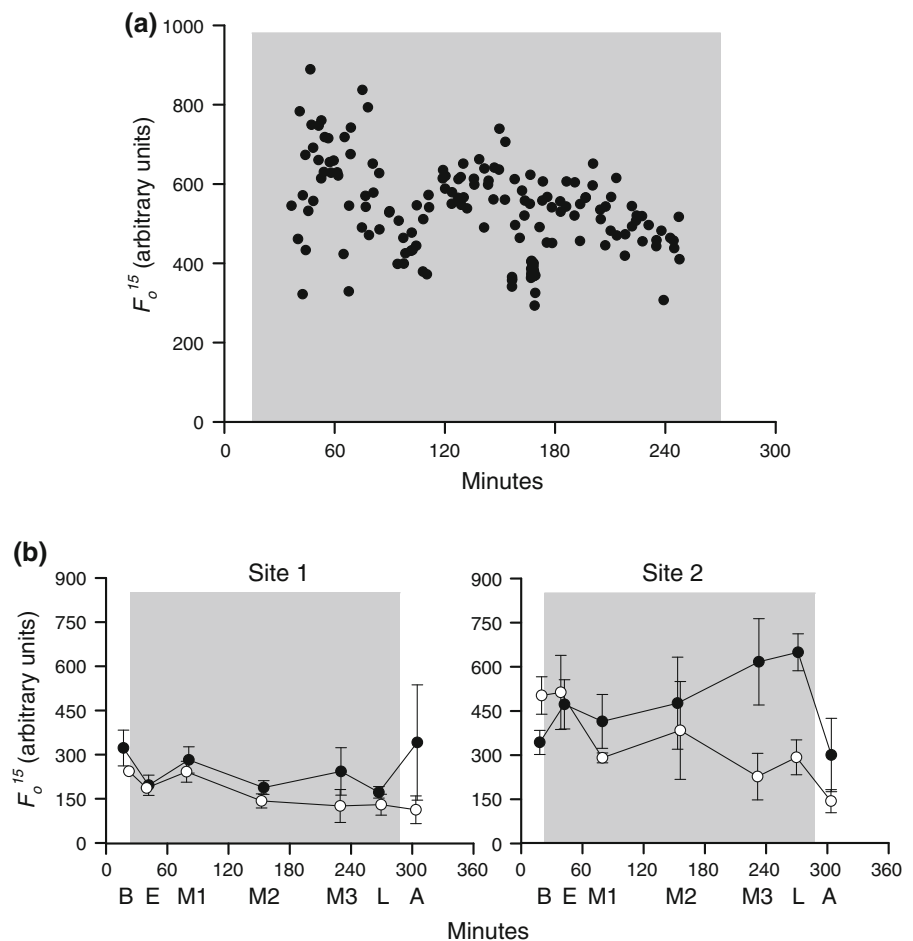
Observational study

At TB, fluorescence decreased slightly and became less variable as the period of low tide progressed (Fig. 3a), although data were only collected during the period of emersion and it was not possible to see whether fluorescence increased again on re-submersion.

Experimental addition of water

At FTB, maintaining overlying water during low tide showed no interactions among Site, Treatment and Time, so non-significant interactions ($P > 0.25$) were pooled (Underwood, 1997). F_o^{15} slightly decreased during low tide (ANOVA, $F_{6,69} = 3$, $P < 0.05$;

Fig. 3 **a** Individual F_o^{15} from micro-algae on an estuarine mudflat (TB) every 2–5 min during a diurnal low tide (at 11.45, on March 15th 2007), starting after emersion and finishing just before sediment was covered by the rising tide. **b** Mean (\pm SE; $n = 3$) F_o^{15} in Control (filled circle) and +Water (open circle) treatments on the mudflat (FTB) at different Times (B = before emersion, E = early, M1 = middle 1, M2 = middle 2, M3 = middle 3, L = late in the emersion period, A = after the rising tide re-covered the patches) on May 15th 2007, low tide at 12.30



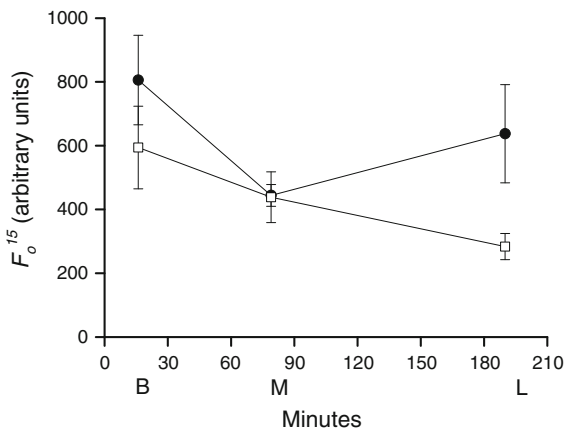


Fig. 4 Mean (\pm SE; $n = 5$) F_o^{15} in Control cores (filled circle) and experimentally drained cores (open square) on a mudflat (FTB) during low tide (at 12.00) on May 29th 2007

Fig. 3b), although SNK tests were unable to detect the differences among times. In contrast to the rocky shore, fluorescence did not increase when the Control sediment was covered again by the tide. Values of F_o^{15} were consistently and significantly greater ($F_{1,69} = 13.0$, $P < 0.01$; Fig. 3b) in Controls than in +Water treatments, which was opposite to the prediction, although both treatments decreased during low tide. Thus, there was no support for the model that the absence of water overlying the sediment during low tide decreases the measures of F_o^{15} in these MPB.

Experimental draining

When samples of sediment were drained, F_o^{15} was smaller than in Control samples ($F_{1,24} = 4.89$, $P < 0.05$) and, again, measures decreased during the period of emersion (Time: $F_{2,24} = 3.97$, $P < 0.05$; Beginning > Middle = Late, SNK tests at $P = 0.05$). Towards the end of low tide, fluorescence of Control samples increased slightly, whilst fluorescence in Drained samples continued to decrease (Fig. 4), although these trends were not significant.

Discussion

Methods of measuring fluorescence, for example, to estimate MPB biomass that do not depend on the time of collection relative to the time of day or tidal conditions are fundamentally important. In intertidal

habitats, numerous environmental variables at the surface of the substratum are likely to change with cover of water, e.g. temperature (especially of the rock, Helmuth & Hofmann, 2001), light, moisture, gas supply and pH. Each of these may affect photosynthetic reactions, such as non-photochemical quenching (NPQ), as a consequence of photo-inhibition (Falkowski & Kiefer, 1985; Hanelt, 1992; Henley et al., 1992; Hader et al., 1997). Desiccation of algae has been demonstrated to cause reductions in fluorescence (e.g. Dring & Brown, 1982; Huppertz et al., 1990; Ji & Tanaka, 2002; Coelho et al., 2009) and measures of MPB using spectrometry on muddy sediments (Coelho et al., 2009), but not using field spectrometry or digital imaging on rock platforms (Murphy et al., 2006). Resuspension of algae in the water above the sediment (Heckman, 1985) may decrease fluorescence, whilst compaction of sediment during de-watering (Perkins et al., 2003) can increase concentrations of micro-algae.

Here, we examined variation in in situ fluorescence emitted by MPB during a single tidal cycle, because many physiological and physical processes indicated that fluorescence might vary at this scale. For example, although previous studies in soft sediments have shown F_o^{15} to be strongly correlated with biomass of MPB (Serôdio et al., 1997; Honeywill et al., 2002), Jesus et al. (2006) warned against sampling at the end of the period of emersion, after showing that differing tendencies of algae to migrate can lead to biased estimates of biomass. Although vertical migration of MPB on rocky substrata may be less important (Lamontagne et al., 1989), tube-dwelling diatoms can move in a mucilaginous sheath (Houpt, 1994), thus affecting the distance from the sensor and hence intensity of fluorescence. In addition, some boring cyanobacteria (forming both epilithic and endolithic filaments) are present on intertidal rock surfaces (Golubic, 1969).

On a rocky shore, we showed that in situ fluorescence emitted by MPB changed little, on average, during most of the period of low tide (Figs. 2, 3). There was, however, a sharp decrease in F_o^{15} as the water drained, and an increase immediately after the rock was wet again by the rising tide. We demonstrated experimentally that this was due to the loss and addition of surficial water, thus excluding the possibility of an effect solely due to endogenous circadian or tidal rhythms on photosynthetic activity (Suzuky &

Johnson, 2001). In fact, although changes in abiotic factors (e.g. desiccation, pH, gas supply), vertical migration and/or NPQ processes could possibly decrease F_o^{15} values during the period of emersion in Control patches, in most of the cases, the presence of water reduced this effect. In addition, it is worth stressing that our rocky MPB assemblages were strongly dominated by cyanobacteria (Jackson et al., 2010), which are known to modulate excessive light harvesting processes mostly through state transitions (Campbell et al., 1998). Although NPQ processes in these organisms can be activated by high levels of blue light (see the review by Bailey & Grossman, 2008), this was not the case for our study, as the Diving PAM excited fluorescence by pulse modulated red light. Last, but not least, it is important to note that the observed increases in F_o^{15} values in Control patches, from just before to just after the water covered them (Fig. 2a and b, see Late and After times), are unlikely to be due to decreases in PAR (photosynthetically active radiation) values. In fact, in both runs of the experiment, the difference in time between the two measurements was about half an hour (and measurements were taken approximately between 2 and 3 p.m.), which is a very short period of time in comparison to the total length of the dry period, which encompassed about 4 h (approximately between 10 a.m. and 2.30 p.m.), during which MPB experienced greater changes in PAR values.

On rocky shores, the time of sampling during low tide may have little influence on measures of fluorescence as long as the substratum is dry. Wetness (e.g. from wave-splash, seepage from rock pools, run-off, rainfall, etc.) may have large consequences for any comparisons. For example, comparisons of MPB biomass at different heights on a shore or between sheltered and exposed sites may be confounded if low shore or wave-exposed sites are consistently wetter than high shore or sheltered sites. We are, however, not aware of studies using PAM on rocky shores that have considered surface moisture when sampling, although Murphy et al. (2006) demonstrated clearly that surface films of moisture do not affect measures of MPB biomass using field spectrometry or digital imaging. This suggests that surface water does not affect the behaviour of the algae (i.e. NPQ processes or vertical migration) on rock surfaces, but seems to affect directly the measures made by the PAM, i.e. the relationship between MPB biomass and F_o^{15} . Recent

measurements on a Mediterranean temperate rocky shore showed that the relationship between biomass of MPB (estimated as $\mu\text{g chl } a/\text{cm}^2$) and F_o^5 varied on wet versus dry sandstone rock surfaces (equations from Maggi et al., unpublished data; wet surface: MPB biomass = $0.004 \cdot F_o^5 + 0.26$; dry surface: MPB biomass = $0.013 \cdot F_o^5 + 0.32$).

On soft sediments, changes in the fluorescence during low tide differed from those on the rocky shore. Here fluorescence slightly decreased during emersion; it is worth noting, however, that this result was mainly in the '+Water' treatment. Although this is in contrast to some studies (Serôdio & Catarino, 2000; Jesus et al., 2005; Tolhurst et al., 2006), which showed increases in fluorescence during low tide, the lack of changes in Control plots during low tide agrees with past experiments at the same location (Tolhurst & Chapman, 2005). Other authors (Easley et al., 2005) showed that diatoms in a sandy beach did not always migrate vertically during low tide. Here, we cannot exclude an interactive effect between vertical migration of algae and NPQ processes (Falkowski & Kiefer, 1985). Upward migration in Controls during the period of low tide was, however, very unlikely at these sites, due to a relative scarcity of migratory micro-algae (Chapman & Tolhurst, 2004, 2007). As a consequence, we should expect only a possible decrease in F_o^{15} values, due to NPQ processes, only in the Control in comparison to the +Water plots (if the presence of some centimetres of water in +Water treatments protected MPB from NPQ processes) or in both Control and +Water plots (if the presence of water did not protect MPB). None of the two alternatives were in accordance with our results. The addition of water may have slightly decreased F_o^{15} values because of resuspension of MPB (Heckman, 1985). Easley et al. (2005) proposed that this would happen with the arrival of water as the tide rose. More interestingly, experimental draining of the sediment decreased F_o^{15} relative to sediment in situ. This pattern was observed at approximately the same time (around 1.30–2 pm) as the M2 time of the previous experiment, when no changes in F_o^{15} values were observed. Thus, although not excluding the occurrence of NPQ processes in this habitat, this result clearly shows that the absence of moisture can decrease fluorescence in soft sediments if the sediment dries out, e.g. during exceptionally low spring tides on particularly hot and dry days. This is in accordance with the negative effect

of desiccation on measures of fluorescence found by Coelho et al. (2009) in a laboratory study. This would cause a similar pattern to that shown on the rocky shore and also suggests that positive effects of compaction of sediment on fluorescence values are weak or non-existent in this estuary. Data from the same location used here showed a good correlation between MPB biomass and $F_o^{1.5}$ at two sites during the middle of low tide (with r^2 values of 0.6 and 0.7, unpublished data). This means that, as on a rocky shore, moisture content of the sediment can affect the relationship between biomass and $F_o^{1.5}$. The relationship varied not only from the middle to the end of the period of low tide (which could be due, again, both to NPQ processes and drainage), but also between sites (which probably differ in the amount of moisture of the sediment, despite their apparent similarity). Thus, on soft sediments, the time during the low tide period when one samples using PAM may provide different results depending on the amount of drainage. This varies a lot from place to place on a mudflat, depending on compaction of the sediment, the mix of grain-sizes and small micro-topographic features caused by environmental conditions or actions of biota.

Conclusions

The two studies described here emphasized the need to sample representatively when using PAM (e.g. to estimate MPB biomass) during low tide. On rocky shores and in soft sediments, it is important to consider the wetness of the substratum. If this cannot be kept consistent across larger temporal scales, it may be important to stratify sampling to incorporate wet and dry patches of habitat or to collect enough samples at each time to ensure that samples are representative of the range of conditions at each time. The latter method would then allow comparisons at different states of the tidal cycle. On rocky shores, the timing of sampling during low tide did not seem to affect the results as long as the substratum was dry. As a consequence, in conditions where the presence of water (e.g. from wave-splash, seepage from rock pools, run-off, etc.) cannot be avoided, the rock surface should be dampened manually, to guarantee that measurements are comparable. Timing of sampling during emersion appears to be more important in sediments than on

rock, although this can vary among habitats. In this case, it is probably more difficult to ensure a comparable moisture of the sediment both among sites and times during low tide, without altering other characteristics of the sediment (such as micro-topography, compaction of sediment, micro-spatial variability of MPB). In this case, stratification of sampling and increasing the number of samples at each time during low tide is probably the only way to ensure representativeness of sampling.

In conclusion, when using PAM methodology to estimate MPB biomass, it will be necessary to demonstrate that the time since emersion does not affect the results obtained, rather than simply assume that it does not. All of this indicates that as much thought needs to go into considering small temporal scales of sampling on PAM measurements as has gone into considering small-scale spatial variation. Further investigations are needed to clarify the physiological and/or physical changes occurring inside the MPB cells and/or in the first millimetres of the substratum (rock, mud or sand), which are causing a change in the relationship between fluorescence and MPB biomass in the presence of different amounts of water.

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