# Detiding time series of dissolved oxygen for improved estimates of ecosystem metabolism

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*Running head: Noise in Estuary Metabolism*

## *Acknowledgments*

# *Abstract*

In aquatic ecosystems, time series of dissolved oxygen (DO) can be used to infer integrated ecosystem processes such as primary production, respiration, and net metabolism. However, continuous monitoring data at estuaries may reflect variation from both biological and physical processes, such that observed data may produce inaccurate or misleading process estimates. Statistical techniques that dynamically quantify variation in DO and tidal changes over time have the potential to isolate biological signals in DO variation. A weighted regression method was developed to normalize, or detide, the predicted DO signal to remove the influence of physical advection on ecosystem metabolism estimates. The method was tested using a simulation approach to create multiple DO time series with known additive components of biological and physical variation on different periods. The method was further validated using one year of continuous monitoring data at four water quality stations that are part of the National Estuarine Research Reserve System. We provide a detailed discussion on use of the method for improving certainty in evaluation of DO measurements from sites with strong tidal influences. Moreover, we propose that the method could expand use of the open-water method for estimating ecosystem metabolism in estuaries given that the approach can produce robust estimates of DO that are independent of tidal advection. In particular, this could facilitate the use of shorter deployment periods for water quality monitors or incomplete time series given that known biases related to water movement could be removed.

# *Introduction*

Time series of dissolved oxygen are increasingly used to estimate ecosystem metabolism (Kemp and Testa 2012, Needoba et al. 2012). Integrated measures of metabolism describe the balance between production and respiration processes that create and consume organic matter, respectively. Although metabolic rates vary naturally between systems (Ziegler and Benner 1998, Caffrey 2004, Russell and Montagna 2007), anthropogenic nutrient sources are often contributing factors that increase rates of production (Diaz and Rosenberg 2008). Inputs of limiting nutrients beyond background concentrations may decrease the resilience of an ecosystem such that higher rates of production are coupled with higher biological oxygen demand (Yin et al. 2004, Kemp et al. 2009). Cultural eutrophication is frequently linked to declines in water quality through lower levels of dissolved oxygen, degradation in aquatic vegetation habitat, and increased frequency of harmful algal blooms (Cloern 1996, Short and Wyllie-Echeverria 1996, Rabalais et al. 2002). Reliables estimates of ecosystem metabolism are critical for measuring both background rates of production and potential impacts of human activities on ecosytem condition.

Ecosystem metabolism can be estimated using two basic techniques, each of which is appropriate under different conditions or assumptions (Kemp and Testa 2012). Bottle-based techniques rely on rate measurements from discrete water quality samples, whereas open-water techniques infer metabolic rates using *in situ* measurements from continuous monitoring data. Bottle-based techniques partition metabolic contributions into discrete habitats, such as pelagic production rates during specific time periods (Kemp and Testa 2012). However, such measurements cannot estimate whole ecosystem metabolism if significant metabolism occurs in habitats that are not sampled, such as the benthos. As such, open-water metabolism provides an

integrative measure of process rates from *in situ*, continuous monitoring data. Originally

conceived for streams (Odum 1956), the open-water method has been used with varying success

in lakes (Staehr et al. 2010, Coloso et al. 2011, Batt and Carpenter 2012) and estuaries (Caffrey

2004, Hopkinson and Smith 2005, Caffrey et al. 2013). As with any method, the ability to

accurately estimate whole system metabolism depends on the degree to which assumptions of the

theory are met. Such assumptions are often only implicity verified in practice, leading to potential

biases.

The open-water method uses the diel fluctuation of dissolved oxygen to infer rates of

ecosytem metabolism, after correcting for air-water gas exchange (Kemp and Testa 2012). Daily

integrated measurements of metabolism represent the balance between daytime production and

nighttime respiration. The fundamental assumption is that the time series of dissolved

oxygen (DO) describes the same water mass over time (Needoba et al. 2012). Estimates of

metabolism from a single location may be inaccurate if substantial variation in water column

mixing occurs throughout the period of observation (Russell and Montagna 2007). Application to

lakes or estuaries have often assumed that a single sampling station provides sufficient data for

estimating metabolism Staehr et al. (2010). While single stations may be valid under specific

conditions, numerous studies have shown that the open-water method may be inappropriate given

the effects of physical mixing in lakes or estuaries (Ziegler and Benner 1998, Caffrey 2003,

Coloso et al. 2011, Batt and Carpenter 2012, Nidzieko et al. 2014). An extensive analysis by

Caffrey (2003) applied the open-water method to estimate metabolim at 28 continuous

monitoring stations at 14 US estuaries. A significant portion of the estimates were negative for

production and positive for respiration, i.e., opposite in sign given the assumptions of the method.

These 'anamolous' values were attributed to the effects of tidal advection such that water masses

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with different metabolic histories were sampled at multiple sites. Moreover, anomalous estimates were included in annual averages of ecosystem metabolism such that inferred rates could have been biased.

The effects of tidal advection on estimates of ecosystem metabolism have been emphasized in numerous studies (Ziegler and Benner 1998, Collins et al. 2013, Howarth et al. 2014, Nidzieko et al. 2014), although systematic estimates of its effects have been minimal. Analytical techniques to evaluate and correct for tidal advection could improve certainty in metabolism estimates and increase the use of data from shorter deployment periods if sources of bias are removed. This article describes the application of a method for filtering an observed DO time series for estimated tidal effects to more accurately quantify estimates of ecosystem metabolim for estuaries. Specifically, the effects of tidal advection on DO observations are removed to improve estimates of open-water metabolism with continuous water quality data. We use a weighted regression approach developed to resolve trends in pollutant concentrations in streams and rivers (Hirsch et al. 2010). The weighted regression approach creates dynamic predictions of DO as a function of time and tidal height change, which are then used to filter, or detide, the DO signal. First, we use a simulation approach to create time series of DO observations with known characteristics to evaluate ability of the weighted regression to remove the effects of tidal advection. Second, four case studies were chosen from the National Estuarine Research Reserve System (NERRS, Wenner et al. 2004) to illustrate the method for adjusting estimates of ecosystem metabolism. Overall, the analysis is meant to better characterize the relative roles of biological and physical processes in estuarine systems.

## Materials and Procedures

### Weighted regression for modelling and detiding DO time series

The weighted regression model for filtering DO time series for tidal effects was adapted

from the weighted regression on time, discharge, and season (WRTDS) method (Hirsch et al.

2010). The functional form of the model is a simple linear regression:

$$DO_{obs} = \beta_0 + \beta_1 t + \beta_2 H \tag{1}$$

where observed DO is a linear function of time $t$ and tidal height $H$. Time is a continuous

variable for the day and time of each obervation with time as a proportion of the number of total

observations added to each day. The beginning of each day was considered the nearest thirty

minute observation to sunrise for the location. Our model also differed from the original WRTDS

method that included parameters to estimate variation of the response variable on a sinuisoidal

period. DO variation was not modeled using this approach to avoid constraining parameter

estimates by periodic components.

Weighted regression was implemented as a moving window that allowed for estimation of

DO throughout the time series by adapting to variation through time as a function of tide.

Regression models were estimated sequentially for each observation in the time series using

dynamic weight vectors that change with the center of the window. Weight vectors quantified the

relevance of observations to the center of the window in respect to time, hour of the day, and tidal

height. Specifically, weights were assigned to each variable using a tri-cube weighting function

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(Tukey 1977, Hirsch et al. 2010):

$$
w = \begin{cases} \left(1 - (d/h)^3\right)^3 & \text{if } |d| \leq h \\ 0 & \text{if } |d| > h \end{cases} \tag{2}
$$

where the weight $w$ of each observation is inversely proportional to the distance $d$ from the center

of the window such that observations more similar to the point of reference are given higher

importance in the regression. Weights exceeding the maximum width of the window $h$ are equal

to zero. The tri-cube weighting function is similar to a Gaussian distribution such that weights

decrease gradually from the center until the maximum window width is reached. Regressions that

use simpler windows (e.g., boxcar approach) are more sensitive to influential observations as they

enter or leave the window, whereas the tri-cube function minimizes their effect through gradual

weighting of observations from the center (Hirsch et al. 2010). The final weight vector for each

observation is the product of three separate weight vectors for decimal time (day), hour, and tidal

height. Windows for time and hour weight observations based on distance (time) from the center

of the window. The window for tidal height weights observations based on the difference from the

center as a proportion of the total tidal height range. For example, a half-window width of 0.5

means that observations are weighted proportionately within +/- 50% the total range referenced to

the tidal height in the center of the window. A low weight is given to an observation if any of the

three weighting values were not similar to the center of the window since the final weight vector

is the product of three weight vectors for each variable (see the link in the multimedia section for

graphical display of different weights).

      The choice of window widths for weight vectors strongly affects the model results.

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Excessively large or small window widths may respectively under- or over-fit the observed data.

Accordingly, appropriate window widths depend on the objective for using the model. The

weighted regression approach can be used for both predicting observed DO and filtering the

observed time series to remove the variance that coincided with the tidal cycle. Window widths

that minimize prediction error or fit to the observed data are typically smaller than the widths for

filtering the time series. Similarly, window widths that more effectively detide the DO signal may

produce imprecise predictions for the observed data. Evaluations of the weighted regression

method with simulated DO time series, described below, used multiple window widths to evaluate

the ability of the model to filter the DO signal. The ability to predict observed DO was not a

primary objective such that the window widths were evaluated only in the context of removing

tidal variation from the DO time series.

The normalization approach to filter tidal variation from the observed DO time series

differs slightly from methods in Hirsch et al. (2010). The previous approach used a

two-dimensional grid predicted for stream pollutant concentrations across the time series and the

range of discharge values observed in the study system (Hirsch et al. 2010). Normalized or

discharge-independent values for pollutant concentration were obtained by averaging grid

predictions across the discharge values that were likely to occur on a given day. Rather than

creating a two-dimensional grid of DO related to time and tidal height change, the normalized

time series herein were the model predictions conditional on time and constant tidal height set to

the mean:

$$DO_{nrm} = f(DO_{obs}|\bar{H}, t) \tag{3}$$

such that the normalized time series represents DO variation related to biological processes

154 (Fig. 1).

## *Assessment*

### Simulation of DO time series

157       To test the ability of the weighted regression to filter the DO signal for apparent tide

158 effects, multiple time series with known characteristics were simulated and filtered. Observed DO

159 time series were simulated as the sum of variation from biological processes and physical effects

160 related to tidal advection:

$$DO_{obs} = DO_{bio} + DO_{adv} \tag{4}$$

161 Biological DO signals are inherently noisy (Batt and Carpenter 2012) and variance can be further

162 described as:

$$DO_{bio} = DO_{die} + DO_{unc} \tag{5}$$

163

$$DO_{unc} = \epsilon_{obs} + \epsilon_{proc} \tag{6}$$

164 where the biological DO signal is the sum of diel variation plus uncertainty or noise. Total

165 uncertainty in the biological DO signal is described as variation from observation and process

166 uncertainty (Hilborn and Mangel 1997). Multiple time series at 30 minute time steps over 30 days

167 were created by varying the relative magnitudes of each of the four components of observed DO

168 in eqs. (4) to (6) to test the effectiveness of weighted regression under different scenarios.

169 Accordingly, observed DO was generalized as the additive combination of four time series

170 (Fig. 2):

$$DO_{obs} = DO_{adv} + DO_{die} + \epsilon_{obs} + \epsilon_{pro} \tag{7}$$

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171    Each component of the simulated time series was created as follows. First, the diel

172  component, $DO_{die}$, was estimated using a sine/cosine function (Cryer and Chan 2008):

$$DO_{die} = \alpha + \beta \cos\left(2\pi ft + \Phi\right) \tag{8}$$

173  such that the mean DO $\alpha$ was 8, amplitude $\beta$ was 1, $f$ was 1/48 to repeat on a 24 hour period

174  every 30 minutes, $t$ was the time series vector and $\Phi$ was the x-axis origin set for an arbitrary

175  sunrise at 630am. The diel signal was increasing during the day and decreasing during the night

176  for each 24 hour period and ranged from 7 to 9 mg $L^{-1}$. Noise or uncertainty was added to the

177  diel DO signal to simulate natural variation in DO throughout the time series (Fig. 2). Total

178  uncertainty was the sum of observation and process uncertainty for $n = 1440$ (30 minutes by 30

179  days) observations (Hilborn and Mangel 1997), such that:

$$DO_{unc,n} = \epsilon_{obs,n} + \int_{t=1}^{n} \epsilon_{pro,t} \tag{9}$$

180  where observation and process uncertainty ($\epsilon_{obs}$, $\epsilon_{pro}$) were simulated as normally distributed

181  random variables with mean zero and standard deviation varying from zero to an upper limit,

182  described below. To induce auto-correlation, process uncertainty was estimated as the cumulative

183  sum of $n$ observations where the noise at time $t + 1$ was equal to the noise at time $t$ plus

184  additional variation drawn from the normal distribution. The noise vector for process uncertainty

185  was rescaled to constrain the variation within the bounds for standard deviation defined by the

186  random variable. The total uncertainty, $DO_{unc}$, was added to the diel DO time series to create the

187  biological DO time series (eq. (5) and Fig. 2).

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A semidiurnal tidal series was simulated as a sine wave with a period of 12.5 hours to

represent the principal lunar component (Foreman and Henry 1989). The amplitude was set to 1

meter and centered at 4 meters. Initial assessments indicated that tide type (i.e., diurnal,

semidiurnal, mixed) did not signifcantly affect the outcome of the results and a semidiurnal time

series was used to reduce the total number of simulations. The tidal time series was added to the

biological DO series to simulate DO changes with advection, $DO_{adv}$ (eq. (7) and Fig. 2).

Conceptually, this vector represents the rate of change in DO as a function of horizontal water

movement from tidal advection such that:

$$\frac{\delta DO_{adv}}{\delta t} = \frac{\delta DO}{\delta x} \cdot \frac{\delta x}{\delta t} \tag{10}$$

$$\frac{\delta x}{\delta t} = k \cdot \frac{\delta H}{\delta t} \tag{11}$$

where the first derivative of the tidal time series, as change in height over time $\delta H/\delta t$, is

multiplied by a constant $k$, to estimate horizontal tidal excursion over time, $\delta x/\delta t$. The horizontal

excursion is assumed to be associated with a horizontal DO change, $\delta DO/\delta x$, such that the

product of the two estimates the DO change at each time step from advection, $DO_{adv}$. In practice,

the simulated tidal signal was used to estimate $DO_{adv}$:

$$DO_{adv} \propto H \tag{12}$$

$$DO_{adv} = 2 \cdot a + a \cdot \frac{H - \min H}{\max H - \min H} \tag{13}$$

where $a$ is analogous to $k$ in eq. (11) and is chosen as the transformation parameter to standardize

change in DO from tidal height change to desired units. For example, $a = 1$ will convert $H$ to a

scale that simulates changes in DO from tidal advection that range from +/- 1 mg L$^{-1}$. The final

time series for observed DO was the sum of biological DO and advection DO (eq. (4) and Fig. 2).

## Evaluation of weighted regression with simulated DO time series

Multiple time series were simulated by varying the conditions in eqs. (4) to (13).

Specifically, the simulated data varied in the relative amount of noise in the measurement ($e_{pro}$,

$e_{obs}$), relative amplitude of the diel DO component ($DO_{die}$), and degree of association of the tide

with the DO signal ($DO_{adv}$). Three levels were evaluated for each variable: relative noise as 0, 1,

and 2 standard deviations for both process and observation uncertainty, amplitude of diel

biological DO as 0, 1, and 2 mg L$^{-1}$, and DO change from tidal advection as 0, 1, and 2 mg L$^{-1}$.

A total of 81 time series were created based on the unique combinations of parameters (Fig. 3).

Three half window widths for each variable (day, hour of day, and tide height) were evaluated for

each simulation: decimal time as 1, 3, and 6 days, time of day as 1, 3, and 6 hours, and tidal

height as 0.25, 0.5, and 1 as a proportion of the total range given the height at the center of the

window. The window widths were chosen based on preliminary assessments that suggested

variation in model performance was adequately captured by these values. In total, 27 window

width combinations were evaluated for each of 81 simulated time series, producing results for

2187 weighted regressions.

The detided or normalized values for each regression were compared to the simulated data

to evaluate the ability of weighted regression to reproduce the biological DO signals. Results

were summarized using Pearson correlation coefficients and the root mean square error (RMSE)

between the detided time series and the biological DO time series as a known component of the

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observed. Overall, the weighted regressions provided accurate results for the detided time series compared to the 'true' biological time series regardless of the simulation parameters (Table 1) or window widths (Table 2, results for each simulation can be viewed using the link in the multimedia section). Mean correlation for all time series and window widths between the detided and biological values was 0.60, with values ranging from -0.78 to 1.00. Mean error was 1.21, with values ranging from 0 to 2.40. Simulations with very poor performance (e.g., negative correlations) were those that had minimum widths for day windows and maximum widths for hour windows, or were those with the DO signal composed entirely of noise from observation uncertainty. Conversely, simulations with detided time series that were identical to the true time series (e.g., correlation of one, RMSE of zero) were those for which there was no biological or tidal influence. Although these examples do not represent real-world scenarios, they were included in the simulations to provide verification that the weighted regression provided reasonable results given extremes.

Characteristics of DO time series that contributed to improved model performance were increasing amplitude of the diel DO component ($DO_{die}$) and increasing process uncertainty ($e_{pro}$), whereas increasing observation uncertainty contributed to decreased performance (Table 1 and Fig. 4). Model performance was minimally influenced by magnitude of the tidal advection component ($DO_{adv}$), although performance decreased slightly with increasing tidal effects. Increasing widths for day and tidal proportion windows contributed to increasing model performance, whereas the opposite was true for increasing hour windows (Table 2 and Fig. 5). Graphical summaries of model performance by simulation parameters (Fig. 4) and half window widths (Fig. 5) support the general trends described by Tables 1 and 2.

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## Validation of weighted regression with case studies

NERRS is a federally-funded network of 28 protected estuaries established for long-term research, water-quality monitoring, education, and coastal stewardship (Wenner et al. 2004). Continuous water quality data have been collected at NERRS sites since 1994 through the System Wide Monitoring Program (SWMP). In addition to providing a basis for trend evaluation, data from SWMP provides an ideal opportunity to evaluate long-term variation in water quality parameters from biological and physical processes. Continuous SWMP data can be used to describe DO variation at sites with different characteristics, including variation from ranges in tidal regime (Sanger et al. 2002) and rates of ecosystem production (Caffrey 2003, 2004).

Continuous DO time series and tidal height measurements at four sites from the SWMP database (CDMO 2014) were used to validate the detiding model with real data. Monitoring data from January 1$^{st}$ to December 31$^{st}$ 2012 were obtained from a range of geographic locations (Fig. 6 and Table 3). Astronomical tidal heights were predicted for each site using harmonic regression applied to the sonde depth data (`oce` package in R, Foreman and Henry 1989, RDCT 2014). Although, the depth data represent tidal height variation from both astronomical (i.e., gravitational effects) and meteorological (e.g., wind, precipitation inflows) sources, we isolated the former given that daily metabolism estimates were more likely to be affected by repeated diel cycling from normal tidal changes. Each station was also chosen based on highly significant correlations between DO and tidal changes. The four sites included the Vierra Mouth station at Elkhorn Slough (California, 36.81°N, 121.78°W), Bayview Channel at Padilla Bay (Washington, 48.50°N 122.50°W), Middle Blackwater River station at Rookery Bay (Florida, 25.93°N 81.60°W), and Dean Creek station at Sapelo Island (Georgia, 31.39°N 81.28°W). The stations are

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generally macrotidal semidiurnal or mixed semidiurnal and net heterotrophic on an annual basis

(Table 3). Net heterotrophy (i.e., respiration exceeding production) is typical for shallow water

systems at temperate latitudes (Caffrey 2003), although values in Table 3 are from observed DO

time series that are strongly influenced by tidal advection.

**Estimates of ecosystem metabolism before and after detiding**

The weighted regression method was applied to the time series for each station to obtain a

detided DO time series for estimating metabolism. Half window widths of six days, one hour, and

a tidal proportion of one half were chosen based on a balance between large and small window

widths, although the chosen window widths are arbitrary and a more exhaustive evaluation should

be conducted prior to using the results to inform management. Unlike the simulated data, the true

biological DO signal was unknown for the case studies. Accordingly, results were evaluated using

correlations of DO and metabolism estimates with tidal height before and after application of the

model. Daily metabolism estimates before and after detiding were compared to the mean rate of

tidal height change (i.e., first derivative of the predicted tidal height) for each day during separate

solar periods. Production rates were compared to mean rates of tidal height change during the

day, respiration rates were compared to mean rates of change during the night, and net

metabolism rates were compared to mean rates of change for the total 24 hour period each day.

Results were also evaluated based on the occurrence of 'anomalous' daily production or

respiration estimates, where anomalous was defined as negative production during the day and

positive respiration estimates during the night. Anomalous values have been previously attributed

to the effects of physical processes on DO time series (Caffrey 2003). We hypothesized that

metabolism esimates using the detided signal would contain less 'anomalous' values than those

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from the observed DO time series. Although anomalies could be caused by processes other than tidal advection, e.g., abiotic dark oxygen production (Pamatmat 1997), we assume that physical processes are the dominant sources of these values given the tidal characteristics at each site. Finally, means and standard errors of metabolism estimates were evaluated before and after detiding to determine if annual aggregations were significantly different.

Ecosystem metabolism was estimated from the DO time series using the open-water technique (Odum 1956) as described in Caffrey et al. (2013). The method is used to infer net ecosystem metabolism using the mass balance equation:

$$\frac{\delta DO}{\delta t} = P - R + D \tag{14}$$

where the change in DO concentration ($\delta DO$, g $O_2$ m$^{-3}$) over time ($\delta t$, hours) is equal to photosynthetic rate ($P$, g $O_2$ m$^{-3}$ hr$^{-1}$) minus respiration rate ($R$, g $O_2$ m$^{-3}$ hr$^{-1}$), corrected for air-sea gas exchange at the surface ($D$, g $O_2$ m$^{-3}$ hr$^{-1}$) (Caffrey et al. 2013). $D$ is estimated as the difference between the DO saturation concentration and observed DO, multiplied by a volumetric reaeration coefficient, $k_a$ (Thébault et al. 2008). The diffusion-corrected DO flux estimates were averaged during day and night for each 24 hour period in the time series, where flux is an hourly rate of DO change as the difference between observations at time $t$ and $t + 1$ on an hourly scale. Areal respiration rates were assumed constant during the night and substracted from daily gross production estimates to yield net ecosystem metabolism (Table 3).

Detiding had significant effects on the correlations between tidal height changes, DO time series, and metabolism estimates (Table 4, see the link in the multimedia section for graphical results of each case study). Correlations of observed DO time series with predicted tidal height

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were highly significant, with all sites indicating positive relationships, except Padilla Bay where tidal increases were associated with declines in DO concentration. This suggests that seaward water masses were less anoxic than landward masses, with the opposite being true for Padilla Bay. The detided DO time series had greatly reduced correlations with tidal height, although relationships were still significant after detiding likely because of the large sample size for each site (n $\approx$ 17500). Comparison of metabolic rates to tidal changes before and after detiding produced inconsistent results (Table 4). Correlations for Elkhorn Slough and Sapelo Island showed consistent reductions in all three metabolims estimates after detiding. Correlations for Padilla Bay and Rookery Bay were of opposite sign and greater magnitude after detiding for production and respiration, although net metabolism estimates had reduced correlations.

The proportion of daily integrated metabolism estimates that were anomalous (negative production, positive respiration) were significantly reduced for most sites after detiding (Table 5). Before detiding, anomalous values ranged from 0.09 (Rookery Bay) to 0.22 (Padilla Bay) for production and 0.08 (Rookery Bay) to 0.21 (Elkhorn Slough) for respiration. Anomalous values were reduced to near zero for Rookery Bay and Sapelo Island, by approximately half for Padilla Bay (0.13 for production, 0.13 for respiration), and only slightly reduced for Elkhorn Slough (0.17 for production, 0.17 for respiration). Metabolism estimates using detided DO time series had decreased mean production and respiration for Elkhorn Slough, increased mean production and respiration for Padilla Bay, and generally unchanged mean production and respiration for Rookery Bay and Sapelo Island (Table 5). Mean net ecosystem metabolism was unchanged for all sites. Decreases in the standard erorr for all metabolism estimates (production, respiration, and net) were observed for all cases after detiding.

An example from Sapelo Island illustrates the effects of weighted regression on DO and

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metabolism estimates (Figs. 7 to 9). A two-week period in February shows when the tidal

changes were both in and out of phase with the diel cycling. Maximum tide heights were

generally out of phase with the diel cycle during the first week such that low tides were observed

during the middle of the night and the middle of the day (Fig. 8). The opposite scenario occurred

during the second week when the maximum tide height occured during the day and night (Fig. 8).

The effects of tidal height change on the observed DO time series are visually apparent in the

plots. The first week illustrates a strong negative bias (less respiration, less production) in the

observed DO signal from low tides at mid-day and mid-night, whereas the second example

illustrates a strong positive bias (more respiration, more production) in the observed DO from

high tides. These biases are apparent in the metabolism estimates using the observed data (Fig. 9).

Anomalous estimates occur when low tides are in phase with the solar cycle, whereas metabolism

estimates are likely over-estimated when high tides are in phase with the solar cycle. The detided

time series shows noticeable changes given the direction of bias from the phasing between tidal

height and diel period. DO values were higher after detiding when low tides occurred during

night and day periods, whereas DO values were lower after detiding when high tides occurred

during day and night periods (Figs. 7 and 8). Changes in metabolism estimates after detiding

were also apparent, such that the anomalous values were removed during the first week and the

positive bias in the second week is decreased. Detiding had similar effects for the remaining sites,

particularly when tidal changes were strongly in or out of phase with diel periods.

## Effects of aggregation and importance of detiding

A final point of concern is the period of observation within which observed DO is affected

by tidal height changes and the extent to which this affects the interpretation of ecosystem

metabolism. From a management or ecological perspective, the effects of tidal variation on daily estimates may not be a primary concern given that seasonal or annual rates may be more relevant for evaluating ecosystem dynamics with continuous monitoring data. The example from Sapelo Island in the previous section highlights this point given that mean production and respiration estimates before and after detiding were generally unchanged for the two-week period. Table 5 also indicated that mean annual estimates of production and respiration were unchanged for Rookery Bay and Sapelo Island, whereas production and respiration estimates were significantly different for Elkhorn Slough and Padilla Bay. Although we acknowledge that the specific results may be related to the window widths, this suggests that tidal effects contribute to changes in metabolic estimates from observed time series that are aggregated on longer time periods. Therefore, an evaluation of the effects of tidal variation on ecosystem metabolism for different periods of observation is critical for understanding practical implications of weighted regression. Specifically, when should detiding be applied if aggregation of observed data on longer time periods removes potential bias? A comparison of observed and detided estimates that are aggregated over different periods of observation (e.g., annual, seasonal, monthly) could help address this question.

The observed and detided daily estimates were averaged by month and season (Fall, Spring, Summer, and Winter) for each case study to evaluate effects of aggregation on mean production and respiration estimates. Mean annual estimates in Table 5 also provided a basis of comparison with monthly and seasonal aggregation. Significant variation in aggregated production and respiration estimates for month and season was observed for each case study (Figs. 10 and 11). Detided production and respiration estimates for Padilla Bay and Rookery Bay exhibited seasonal and monthly variation that was more characteristic of expected trends with

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increases in metabolism during warmer months. Specifically, production estimates based on observed DO were substantially muted for both Padilla Bay (Fig. 10) and Rookery Bay (Fig. 11) during summer months, whereas values were significantly higher based on the detided data. Results for Sapelo Island suggested that winter and summer months were under- and over-estimated, respectively, based on the observed data. Results for Elkhorn Slough varied significantly such that production and respiration were significantly reduced after detiding regardless of the aggregation period. Overall, these trends emphasize the importance of considering different aggregation periods for interpreting metabolism estimates. Each case study showed differences in observed and detided values at monthly and seasonal aggregations, whereas only two of the four case studies had mean aggregated estimates that were substantially different (Elkhorn Slough and Padilla Bay, Table 5). Periods of observation as long as one year may include significant sources of bias from tidal advection, suggesting the need for applying weighted regression given careful consideration of appropriate window widths.

## *Discussion*

The primary objective for development and application of the weighted regression technique was to provide a method for more accurately estimating ecosystem metabolism by removing variation associated with tidal change in observed DO time series. The application of weighted regression to simulated DO time series with known characteristics and extension to continuous monitoring data from selected NERRS sites provided a proof-of-concept that the method can isolate and remove variation in observed DO from tidal change. Further, aggregation of metabolism estimates using the detided DO time series were significantly different than those using the observed data, particularly for relatively long periods of observation. These results

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suggest that previous estimates of annual means may not accurately reflect true metabolic signals if the effects of tidal variation confound biological signals in observed DO time series. Additionally, variation of aggregated metabolism estimates were substantially reduced after detiding, suggesting greater confidence in interpreting detided estimates even if the mean values are similar. Monitoring data for periods of observation up to one year may also produce significantly different metabolism estimates if observed data are not detided.

Comparisons between detided and biological DO time series from the simulations indicated that adequate results can be obtained from weighted regressions for a range of characteristics of DO time series. An examination of scenarios that produced abnormal results can provide additional insight into factors that affect the performance of weighted regression. For example, poor performance was observed when the observation uncertainty ($\epsilon_{obs}$) was high and both process uncertainty ($\epsilon_{pro}$) and tidal advection ($DO_{adv}$) were low. These examples represent time series with excessive random variation, no auto-correlation, and no tidal influence. Poor performance is expected because the weighted regression models a non-existent tidal signal in a very noisy DO time series. These results were observed even for time series with a large diel component of the biological DO signal, suggesting that the model will produce random results in microtidal systems with high noise and no serial correlation. From a practical perspective, weighted regression should not be applied to noisy time series if there is not sufficient evidence to suggest the variation is related to tidal changes. Similarly, results with perfect or near-perfect correlations between detided and biological DO time series were observed when observation uncertainy and tidal effects were not components of the simulated time series. Although there is no logical basis for applying weighted regression to time series with no apparent tidal influences, the results will be as expected, as was true for cases with low tidal advection, high observation

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uncertainty, and low process uncertainty. We emphasize that the weighted regression should only be applied to time series for which specific conditions apply, as described below.

The performance metrics used to evaluate weighted regression with the case studies suggested that detiding provided more accuate estimates of ecosystem metabolism. Correlations of metabolism estimates with tidal height changes after detiding were generally reduced, although trends were not always consistent as correlations were reduced in some cases (Sapelo Island) or reversed in others (Padilla Bay). However, correlations of net metabolism estimates were reduced in all cases. Tidal height change provided a proxy for horizontal advection that directly affects the measured rate of change of oxygen. Further, changes in DO concentration represent integrated measures of both production and respiration (eq. (14)). The inconsistent results in Table 4 are potentially related to the effects of horizontal advection on the integrated DO signal, given that production and respiration each represent a unique component of the diel DO variation that is directly affected by tidal variation. Regardless, the proportion of anomalous metabolism estimates was reduced by detiding for all case studies, although this measure may also be an incomplete indication of the combined effects of tidal variation. Negative production and positive respiration estimates suggest assumptions of the open-water method are violated (Needoba et al. 2012), whereas 'normal' estimates (positive production and negative respiration) may still include a significant source of bias from physical advection by providing over-estimates of true values. For example, Nidzieko et al. (2014) observed that net metabolism at Elkhorn Slough was more often heterotrophic during maximum spring tides that occurred at nighttime, as a substantially larger area of salt marsh was inundated leading to higher respiration estimates. Although this result supports our general conclusions, a broader discussion regarding whether or not this represents an actual bias in metabolism from physical advection may be needed.

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A strength of the weighted regression approach is a lack of assumptions describing the relationships between DO and tidal variation over time. Although the functional form of the model is a simple linear regression with only two explanatory variables (eq. (1)), the moving window approach combined with the adaptive weighting scheme allows for quantification of complex tidal effects that may not be possible using alternative approaches. For example, Batt and Carpenter (2012) described the use of a moving window or Kalman filter (Harvey 1989) to improve estimates of ecosystem metabolism in lakes. The approach minimizes the influences of process and observation uncertainty on observed DO time series, which is attributed to the effects of water movement on metabolic signals. Although a similar approach may be useful for estuaries if process and observation uncertainty are the only sources of variation in the DO series, the weighted regression approach is more appropriate if periodic tidal advection is the primary confounding factor. Additionally, results from the case studies illustrated the ability of the weighted regression approach to model changes over time in the relationships between tidal change and DO. Results for Padilla Bay and Rookery Bay suggested that detiding had the largest effect during the summer, whereas the results for cooler months were not significantly different from the observed. The weighted regression method produced detided time series that accommodated seasonal variation in DO conditional on tidal height change, whereas moving window filters or standard regression techniques would likely not have characterized these dynamic relationships.

## *Comments and recommendations*

Results from the simulations and case studies suggested that weighted regression can be a practical approach for detiding DO time series to remove the effects of physical advection on

estimates of ecosystem metabolism. However, application of the method may only be appropriate under specific situations. The case studies were chosen based on the relatively high proportion of metabolism estimates that were anomalous and the strength of correlation between the observed DO time series and tidal height. Despite these similarites among the case studies, detiding had variable effects on metabolism estimates. The results for Elkhorn Slough and Padilla Bay are of particular concern given that mean annual estimates were substantially different compared to those from the observed DO time series. Although the correlation of DO and tidal height was reduced for both cases, in addition to a reduction of anomalous estimates, the relative change in mean metabolism before and after detiding suggests a more careful evaluation of the method is needed. In particular, alternative window widths should evaluated for the ability to remove tidal effects while preserving the biological signal. The window widths in the above analysis may have removed variation in the DO signal from both of these sources.

Although the above analyses suggest the approach has merit, the case studies emphasize a critical challenge in applying weighted regression to monitoring data. Specifically, the true biological signal is not known and the relative contribution of horizontal advection to bias is not accurately quantified with the available data. Comparative analyses between systems with varying tidal influence or within-system evaluations of multiple sites at fixed distances are necessary to further validate the performance of weighted regression. In the absence of additional validation, we propose a precautionary approach for application of the weighted regression to monitoring data. Weighted regression may be most effective at macrotidal sites with strong evidence of the effects of tidal advection on biological signals. A weight-of-evidence approach should be used such that the occurrence of anomalous metabolism estimates, strong correlations between observed DO and tide height, and clear visual patterns of tide change on DO would suggest

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detiding is appropriate. The choice of window widths may also produce varying results. Window

widths that produce large changes in mean annual estimates should be interpreted with caution. In

general, a pragmatic approach is emphasized such that results should be evaluated based on the

preservation of diel variation from production while exhibiting minimal changes with the tide.

Such an approach, combined with further validation, will support informed management

decisions through more accurate estimates of ecosystem metabolism.

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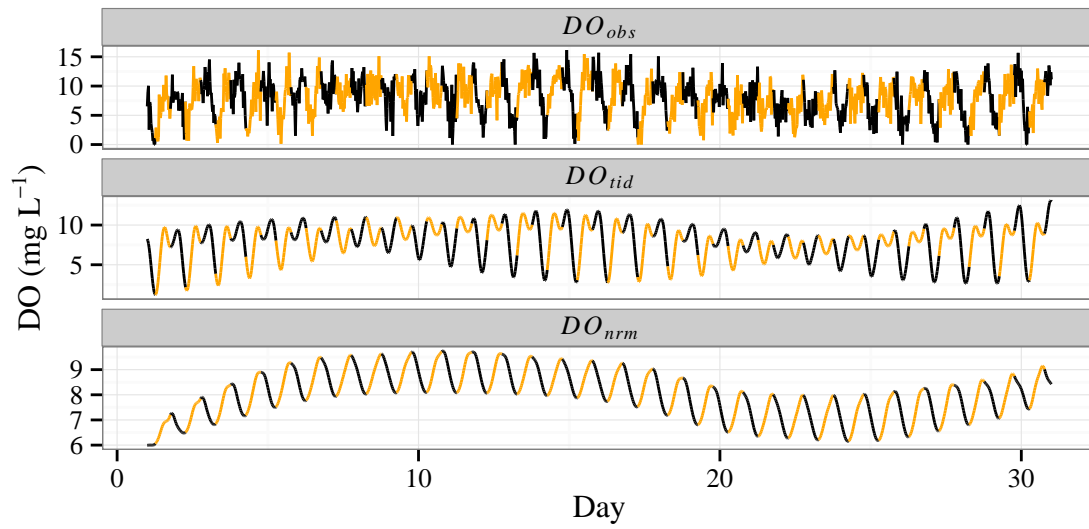581      Marine Ecology Progress Series, 173:1–12.

*Figures*



Fig. 1: Example of detiding a simulated DO time series. $DO_{obs}$ represents an additive time series as in eq. (7), $DO_{tid}$ represents the predicted values of DO conditional on tidal height and time (**??**), and $DO_{nrm}$ represents the detided values of DO conditional on constant tidal height and time (eq. (3)). Yellow indicates daylight periods.
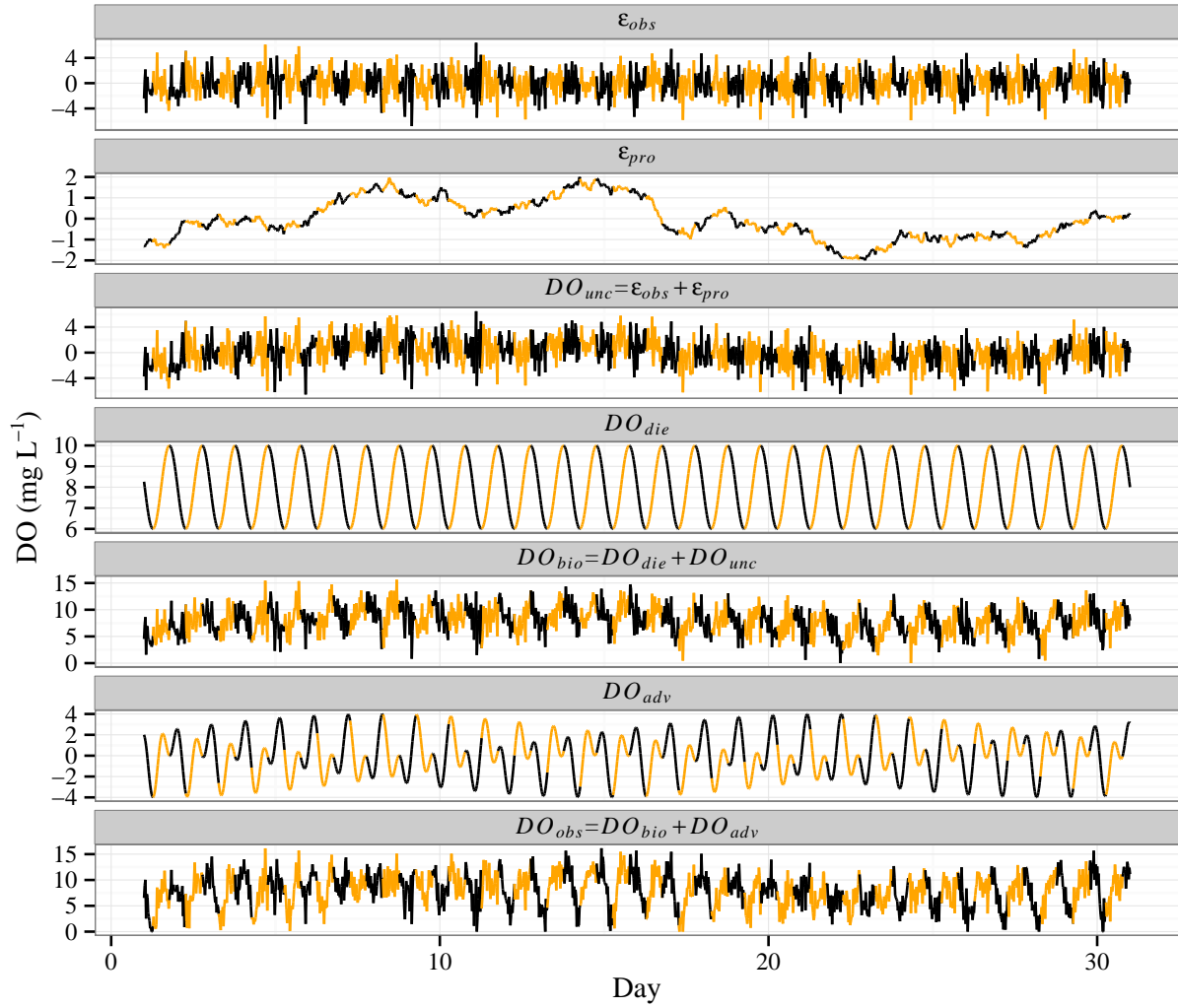
Fig. 2: Example of each component of a simulated DO time series for testing weighted regression. The time series were created using eqs. (4) to (13). Yellow indicates daylight periods.
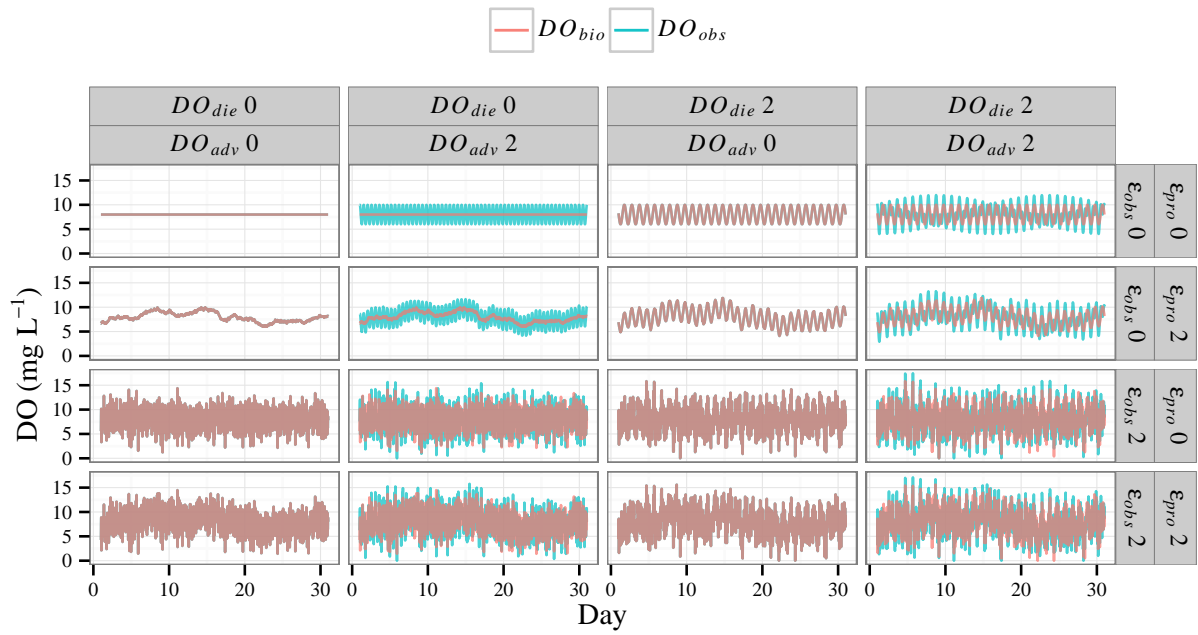
Fig. 3: Representative examples of simulated time series of observed DO ($DO_{obs}$) and biological DO ($DO_{bio}$, as a component of observed) created by varying each of four parameters: strength of tidal association with DO signal ($DO_{adv}$), amount of process uncertainty ($\epsilon_{pro}$), amount of observation observation uncertainty ($\epsilon_{obs}$), and strength of diel DO component ($DO_{die}$). Parameter values represent the minimum and maximum used in the simulations as mg L$^{-1}$ of DO.
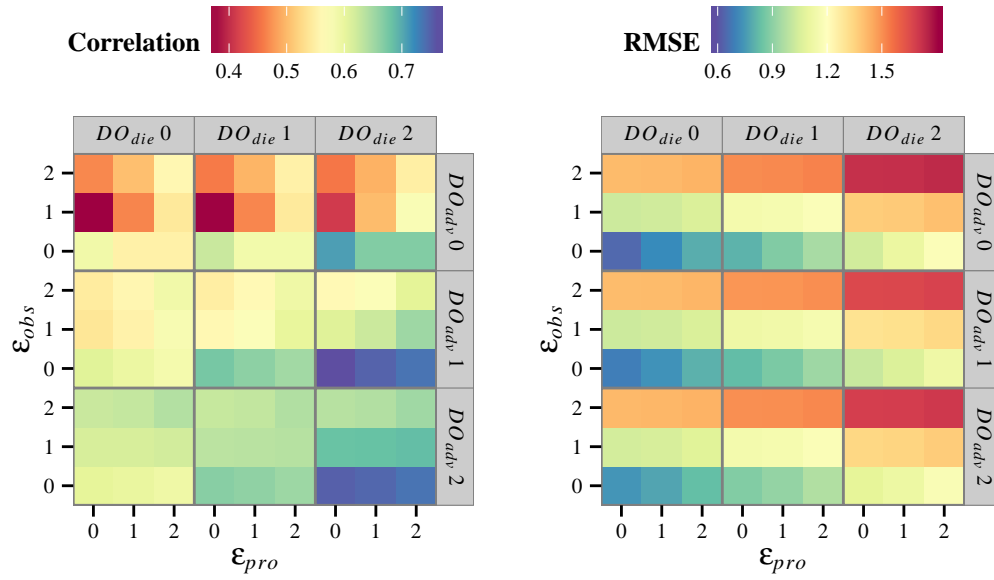
Fig. 4: Correlations and errors (RMSE) for detided DO time series ($DO_{dtd}$) from weighted regression with 'true' biological DO ($DO_{bio}$) for varying simulation parameters: strength of tidal association with DO signal ($DO_{adv}$), amount of process uncertainty ($\epsilon_{pro}$), amount of observation observation uncertainty ($\epsilon_{obs}$), and strength of diel DO component ($DO_{die}$). Each tile represents the correlation or error from results for a given combination of simulation parameters averaged for all window widths (Fig. 5).
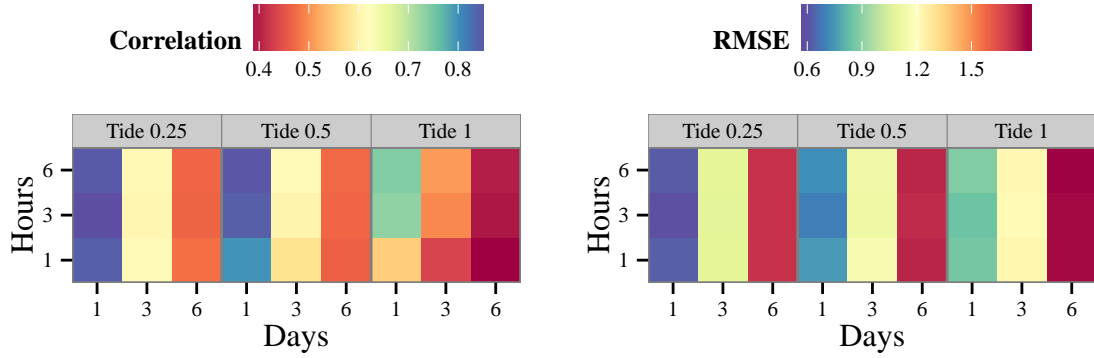
Fig. 5: Correlations and errors (RMSE) for detided DO time series ($DO_{dtd}$) from weighted regression with 'true' biological DO ($DO_{bio}$) for varying half window widths: days, hour of day, and proportion of tidal range. Each tile represents the correlation or error from results for a given combination of window widths averaged for all simulation parameters (Fig. 5).
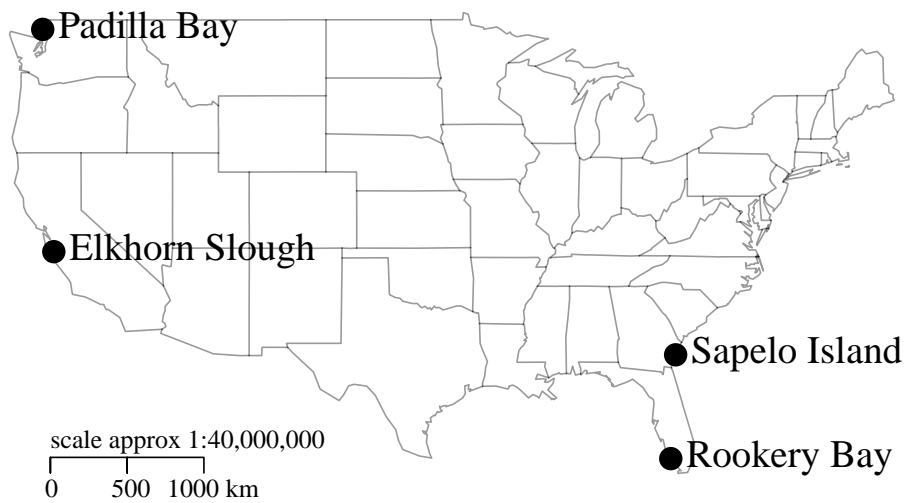
Fig. 6: Locations of NERRS sites used as case studies to validate weighted regression. Stations at each reserve are ELKVM (Vierra Mouth at Elkhorn Slough), PDBBY (Bayview Channel at Padilla Bay), RKBMB (Middle Blackwater River at Rookery Bay), and SAPDC (Dean Creek at Sapelo Island).

Fig. 7: Continuous DO time series before (observed) and after (detided) detiding with weighted regression (top) and tidal height colored by total photosynthetically active radiation (bottom, mmol $m^{-2}$). Results are for the Sapelo Island station for a seven day period when high tide events were out of phase with diel periods, creating lower than expected observed DO during night and day periods. Detided values are based on a weighted regression with half window widths of six days, one hour within each day, and tidal height proportion of one half.
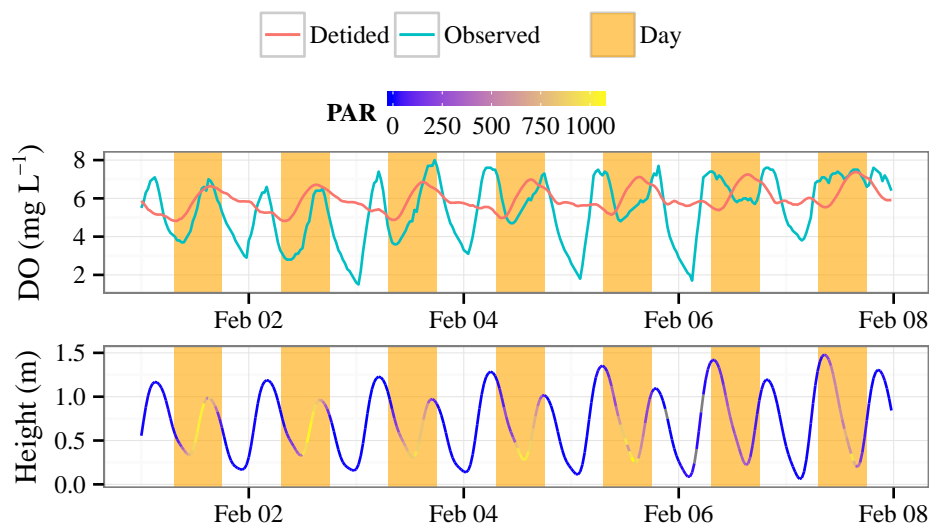
Fig. 8: Continuous DO time series before (observed) and after (detided) detiding with weighted regression (top) and tidal height colored by total photosynthetically active radiation (bottom, mmol $m^{-2}$). Results are for the Sapelo Island station for a seven day period when high tide events were in phase with diel periods, creating higher than expected observed DO during night and day periods. Detided values are based on a weighted regression with half window widths of six days, one hour within each day, and tidal height proportion of one half.
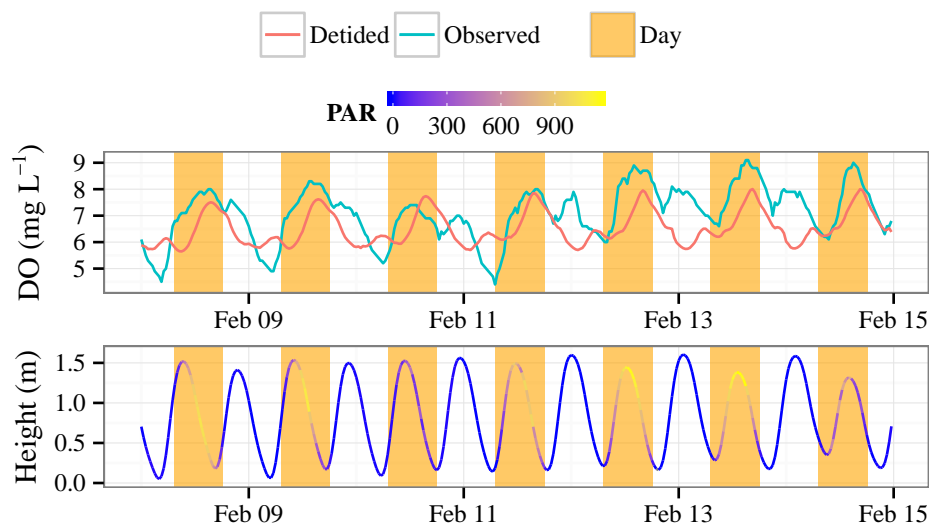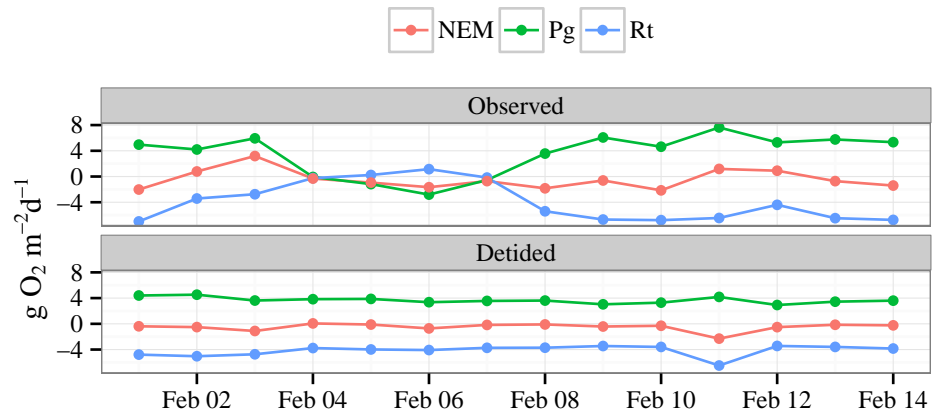
Fig. 9: Example of daily mean metabolism (net ecosystem metabolism, gross production, and total respiration) before (observed) and after (detided) detiding with weighted regression. Results are for the Sapelo Island station for a two week period in February, 2012 when high tide was out of phase with the diel cycle during the first week (Fig. 7) and in phase during the second week (Fig. 8).

Fig. 10: Means and standard errors of daily metabolism estimates (gross production, total respiration) aggregated by month and season. Aggregated estimates are for Elkhorn Slough and Padilla Bay from observed and detided DO time series.

Fig. 11: Means and standard errors of daily metabolism estimates (gross production, total respiration) aggregated by month and season. Aggregated estimates are for Rookery Bay and Sapelo Island from observed and detided DO time series. May was removed from Rookery Bay because of incomplete data.

*Tables*

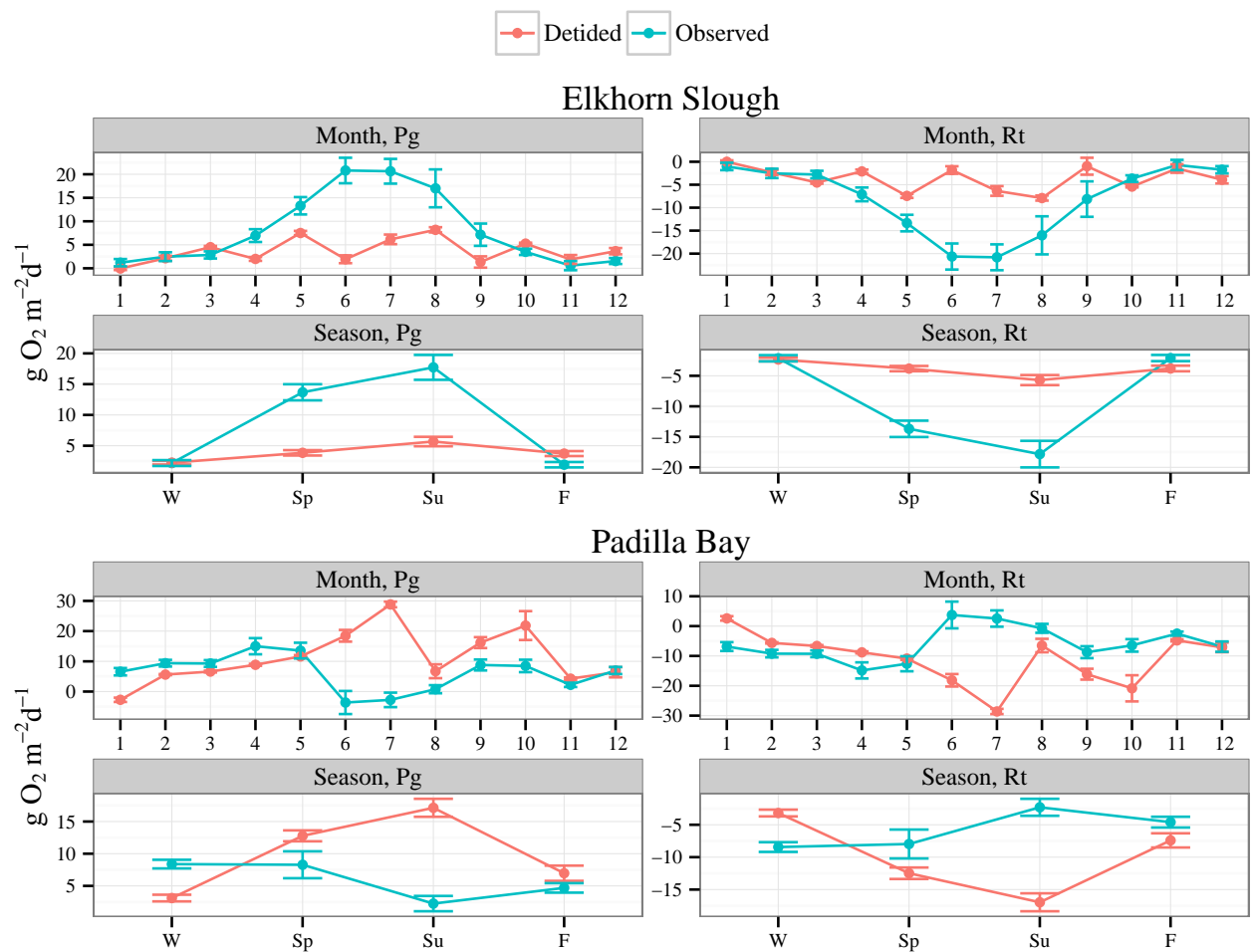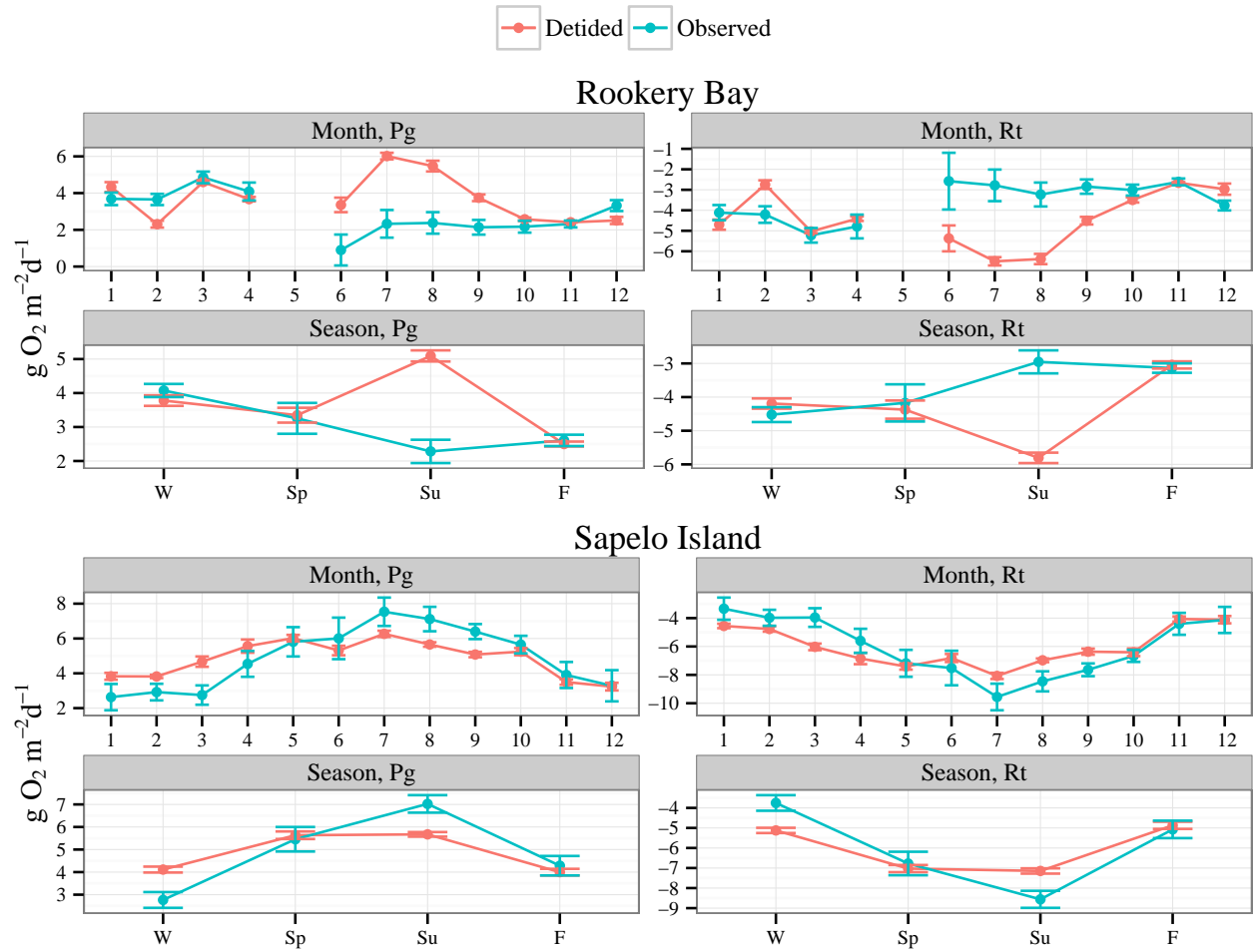Table 1: Summary (range, mean, percentiles) of correlations and error estimates comparing detided and biological DO time series for different simulation parameters ($DO_{die}$, $DO_{adv}$, $\epsilon_{pro}$, $\epsilon_{obs}$). Values represent averages from multiple simulations with common parameters (e.g., row one is a summary of all simulations for which diel DO component was zero).

| Parameter | Correlation | | | | | RMSE | | | | |
|---|---|---|---|---|---|---|---|---|---|---|
| | Min | 25th | Mean | 75th | Max | Min | 25th | Mean | 75th | Max |
| $DO_{die}$ | | | | | | | | | | |
| 0 | -0.78 | 0.30 | 0.53 | 0.82 | 1.00 | 0.00 | 0.68 | 1.22 | 1.97 | 2.39 |
| 1 | -0.28 | 0.38 | 0.61 | 0.88 | 1.00 | 0.00 | 0.59 | 1.20 | 1.96 | 2.40 |
| 2 | -0.39 | 0.46 | 0.65 | 0.90 | 1.00 | 0.00 | 0.62 | 1.22 | 1.97 | 2.40 |
| $DO_{adv}$ | | | | | | | | | | |
| 0 | 0.00 | 0.27 | 0.57 | 0.93 | 1.00 | 0.00 | 0.34 | 1.07 | 1.96 | 2.12 |
| 1 | -0.78 | 0.37 | 0.59 | 0.83 | 1.00 | 0.00 | 0.63 | 1.18 | 1.98 | 2.12 |
| 2 | -0.78 | 0.47 | 0.63 | 0.82 | 1.00 | 0.00 | 0.98 | 1.38 | 1.99 | 2.40 |
| $\epsilon_{pro}$ | | | | | | | | | | |
| 0 | -0.78 | 0.34 | 0.58 | 0.86 | 1.00 | 0.00 | 0.63 | 1.19 | 1.96 | 2.40 |
| 1 | -0.78 | 0.37 | 0.59 | 0.85 | 1.00 | 0.00 | 0.63 | 1.21 | 1.97 | 2.40 |
| 2 | -0.78 | 0.41 | 0.61 | 0.85 | 1.00 | 0.00 | 0.63 | 1.24 | 1.98 | 2.40 |
| $\epsilon_{obs}$ | | | | | | | | | | |
| 0 | -0.78 | 0.31 | 0.65 | 0.98 | 1.00 | 0.00 | 0.29 | 0.92 | 1.50 | 2.40 |
| 1 | 0.05 | 0.37 | 0.57 | 0.81 | 0.99 | 0.07 | 0.98 | 1.18 | 1.49 | 2.39 |
| 2 | 0.05 | 0.40 | 0.57 | 0.70 | 0.99 | 0.15 | 1.06 | 1.54 | 2.01 | 2.40 |

Table 2: Summary (range, mean, percentiles) of correlations and error estimates comparing detided and biological DO time series for simulations using different half window widths in the weighted regressions (days, hours, and proportion of tidal range). Values represent averages from multiple simulations with common window values (e.g., row one is a summary of all simulations for which the half window width was one day).

| Window | Correlation | | | | | RMSE | | | | |
|---|---|---|---|---|---|---|---|---|---|---|
| | Min | 25th | Mean | 75th | Max | Min | 25th | Mean | 75th | Max |
| **Days** | | | | | | | | | | |
| 1 | -0.78 | 0.63 | 0.78 | 0.97 | 1.00 | 0.00 | 0.28 | 0.74 | 1.04 | 2.12 |
| 3 | -0.07 | 0.40 | 0.56 | 0.75 | 1.00 | 0.00 | 0.99 | 1.15 | 1.28 | 2.08 |
| 6 | 0.00 | 0.26 | 0.45 | 0.58 | 1.00 | 0.00 | 1.95 | 1.75 | 2.05 | 2.40 |
| **Hours** | | | | | | | | | | |
| 1 | -0.78 | 0.36 | 0.57 | 0.82 | 1.00 | 0.00 | 0.63 | 1.22 | 1.96 | 2.40 |
| 3 | 0.00 | 0.40 | 0.61 | 0.87 | 1.00 | 0.00 | 0.58 | 1.20 | 1.97 | 2.36 |
| 6 | 0.03 | 0.37 | 0.61 | 0.85 | 1.00 | 0.00 | 0.64 | 1.22 | 1.98 | 2.40 |
| **Tide** | | | | | | | | | | |
| 0.25 | 0.00 | 0.42 | 0.64 | 0.91 | 1.00 | 0.00 | 0.51 | 1.14 | 1.97 | 2.21 |
| 0.5 | 0.06 | 0.43 | 0.63 | 0.88 | 1.00 | 0.00 | 0.61 | 1.20 | 1.97 | 2.27 |
| 1 | -0.78 | 0.30 | 0.52 | 0.79 | 1.00 | 0.00 | 0.73 | 1.30 | 1.97 | 2.40 |

Table 3: Summary statistics of tidal component amplitudes (m), selected water quality parameters (DO mg L$^{-1}$, chlorophyll-a $\mu$g L$^{-1}$, salinity psu, water temperature °C) and metabolism estimates (gross production, respiration, and net ecosystem metabolism as g m$^{-2}$ d$^{-1}$) for each case study. Tidal components are principal lunar semidiurnal (O1, frequency 25.82 hours), solar diurnal (P1, 24.07 hours), lunar semidiurnal (M2, 12.42 hours), and solar semidiurnal (S2, 12 hours) estimated from harmonic regressions of tidal height (`oce` package in R, Foreman and Henry 1989, RDCT 2014). Water quality data are averages for the entire period of record (30 minute observations) for each site. Metabolism estimates are means of daily integrated values.

| Site | Tidal amplitude | | | | Water quality | | | | Metabolism[a] | | |
|---|---|---|---|---|---|---|---|---|---|---|---|
| | O1 | P1 | M2 | S2 | DO | Chl | Sal | Temp | Pg | Rt | NEM |
| ELKVM | 0.24 | 0.12 | 0.48 | 0.13 | 7.87 | 3.87 | 32.43 | 13.78 | 8.14 | -8.19 | -0.05 |
| PDBBY | 0.46 | 0.23 | 0.63 | 0.15 | 8.97 | 2.24 | 29.17 | 10.44 | 5.95 | -5.90 | 0.05 |
| RKBMB | 0.13 | 0.04 | 0.36 | 0.10 | 4.48 | 4.50 | 30.53 | 25.85 | 3.02 | -3.62 | -0.60 |
| SAPDC | 0.10 | 0.02 | 0.54 | 0.07 | 4.96 | 5.98 | 27.30 | 21.77 | 4.89 | -6.04 | -1.16 |

[a]Pg: gross production, Rt: respiration, NEM: net ecosystem metabolism

Table 4: Correlations of tidal changes at each site with continuous DO observations and metabolism estimates (gross production, respiration, and net metabolism) before (observed) and after (detided) detiding with weighted regression. DO values are correlated with predicted tidal height at each observation, whereas metabolism estimates are correlated with mean tidal height change between observations during day, night, or total day periods for production, respiration, and net metabolism, respectively.

| Site | DO | Pg[a] | Rt | NEM |
|------|------|------|------|------|
| **ELKVM** | | | | |
| Observed | 0.47*** | 0.60*** | 0.73*** | 0.35*** |
| Detided | 0.02* | 0.19*** | 0.13* | 0.06 |
| **PDBBY** | | | | |
| Observed | -0.45*** | -0.33*** | -0.46*** | -0.25*** |
| Detided | 0.07*** | 0.48*** | 0.47*** | -0.21*** |
| **RKBMB** | | | | |
| Observed | 0.28*** | 0.34*** | 0.39*** | 0.24*** |
| Detided | -0.02** | -0.31*** | -0.36*** | 0.12* |
| **SAPDC** | | | | |
| Observed | 0.48*** | 0.54*** | 0.71*** | 0.41*** |
| Detided | -0.03*** | 0.16** | 0.18*** | -0.05 |

*$p < 0.05$; **$p < 0.01$; ***$p < 0.001$
[a]Pg: gross production, Rt: respiration, NEM: net ecosystem metabolism

Table 5: Summary of metabolism estimates (gross production, respiration, and net metabolism) for case studies using DO time series before (observed) and after (detided) detiding with weighted regression. Means and standard errors are based on daily integrated metabolism estimates. Anomalous values are the proportion of metabolism estimates that were negative for gross production and positive for respiration. Results are for weighted regressions with half window widths of six days, one hour within each day, and a tidal height proportion of one half.

| Site | Pg[a] | | | Rt | | | NEM | |
|---|---|---|---|---|---|---|---|---|
| | Mean | Std. Err. | Anom | Mean | Std. Err. | Anom | Mean | Std. Err. |
| **ELKVM** | | | | | | | | |
| Observed | 8.14 | 0.67 | 0.19 | -8.19 | 0.69 | 0.21 | -0.05 | 0.16 |
| Detided | 3.63 | 0.23 | 0.17 | -3.67 | 0.24 | 0.17 | -0.04 | 0.05 |
| **PDBBY** | | | | | | | | |
| Observed | 5.95 | 0.69 | 0.22 | -5.90 | 0.74 | 0.19 | 0.05 | 0.22 |
| Detided | 10.36 | 0.63 | 0.13 | -10.32 | 0.63 | 0.13 | 0.04 | 0.08 |
| **RKBMB** | | | | | | | | |
| Observed | 3.02 | 0.14 | 0.09 | -3.62 | 0.15 | 0.08 | -0.60 | 0.06 |
| Detided | 3.73 | 0.09 | 0.01 | -4.35 | 0.10 | 0.00 | -0.62 | 0.04 |
| **SAPDC** | | | | | | | | |
| Observed | 4.89 | 0.23 | 0.13 | -6.04 | 0.25 | 0.11 | -1.16 | 0.09 |
| Detided | 4.85 | 0.08 | 0.00 | -6.04 | 0.10 | 0.00 | -1.19 | 0.05 |

[a]Pg: gross production, Rt: respiration, NEM: net ecosystem metabolism

## *Multimedia*

The supporting information for this manuscript includes a graphical illustration of the

weighting scheme described in the material and procedures section

(http://spark.rstudio.com/beckmw/weights_widget), results for each simulation

(http://spark.rstudio.com/beckmw/detiding_sims), and results for each case study

(http://spark.rstudio.com/beckmw/detiding_cases). Each link is a graphical summary of data

based on interactive inputs to support the results in the manuscript.