

1 **Title:**

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3 **Environmental calibration of coral luminescence as a proxy for terrigenous dissolved**
4 **organic carbon concentration in tropical coastal oceans**

5

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45

46 **Highlights**

- 47 • Coral luminescence green-to-blue ratio (coral G/B) is a quantitative proxy for
48 terrigenous dissolved organic carbon concentration in river-influenced tropical coastal
49 oceans.

- 50 • Coral G/B can also be used to reconstruct the absorption spectrum of terrigenous
51 chromophoric dissolved organic matter (CDOM) over the wavelength range 230-550
52 nm.
- 53 • Downcore luminescence G/B variability can therefore be used to investigate temporal
54 variability in tDOC input and its impacts on optical water quality.

55

56 **Abstract**

57 The riverine flux of terrigenous dissolved organic matter (tDOM) to the ocean is a
58 significant contributor to the global carbon cycle. In response to anthropogenic drivers such as
59 land-use change the flux is expected to increase, and this may impact both the availability of
60 sunlight in coastal ecosystems, and the seawater carbonate system and coastal CO₂ fluxes. Yet
61 despite its biogeochemical and ecological significance, there are few long-term and high-
62 resolution time series of tDOM parameters. Corals incorporate fluorescent tDOM molecules
63 from the chromophoric dissolved organic matter (CDOM) pool in their skeletons, and the
64 resulting luminescence variability in coral skeleton cores has traditionally been used to
65 reconstruct hydroclimate variation. Here, we use two replicate coral cores and concurrent in-
66 situ biogeochemical data from the Sunda Sea Shelf in Southeast Asia, where coastal peatlands
67 supply high tDOM inputs, to show that variability in coral luminescence green-to-blue ratios
68 (coral G/B) can be used to quantitatively reconstruct the concentration of terrigenous dissolved
69 organic carbon (tDOC). Moreover, coral G/B can be used to reconstruct the full absorption
70 spectrum of CDOM from 230–550 nm, as well as the specific ultraviolet absorbance at 254 nm
71 (SUVA₂₅₄) of the DOM pool. Comparison to a core from Borneo shows that there may be site-
72 specific offsets in the G/B–CDOM absorption relationship, but that the slope of the relationship
73 is very similar, validating the robustness of the proxy. By demonstrating that coral cores can
74 be used to estimate past changes in coastal tDOC and CDOM, we establish a method to study

75 natural and anthropogenic drivers of land–ocean tDOM fluxes and their ecological
76 consequences in tropical coastal seas over decadal to centennial time scales.

77

78 **Keywords**

79 corals, luminescence G/B, tDOC, CDOM, dissolved organic matter, Southeast Asia,
80 peatlands

81

82 **1. Introduction**

83

84 The transfer of terrigenous Dissolved Organic Carbon (tDOC) from land to the coastal
85 ocean is a significant flux in the global carbon cycle (Le Quéré et al., 2013). Tropical rivers
86 contribute nearly two-thirds of the global land-to-ocean tDOC flux (Dai et al., 2012).
87 Moreover, this flux appears to have been considerably perturbed by human activity in recent
88 times (Butman et al., 2015; Monteith et al., 2007; Noacco et al., 2017). Long-term increases in
89 surface water DOC across North America and Europe are largely attributed to the recovery of
90 ecosystems from historical atmospheric pollution (Monteith et al., 2007), although
91 anthropogenic landscape alterations (Butman et al., 2015; Noacco et al., 2017) and climatic
92 changes and permafrost thaw (de Wit et al., 2016; Larsen et al., 2011; Wauthy et al., 2018) are
93 also implicated. In tropical regions, deforestation destabilizes soil organic carbon pools (Evans
94 et al., 2014) and can lead to either net increases (Moore et al., 2013; Sanwlani et al., 2022) or
95 net decreases (Drake et al., 2019) in riverine tDOC flux, depending on the lability of the newly
96 mobilized soil carbon.

97

98 The environmental consequences of changing tDOC fluxes to the ocean depend on the
99 biogeochemical fate of tDOC at sea, which remains poorly understood (Ciais et al., 2014). In

100 parts of the Arctic and Southeast Asia, extensive remineralization of tDOC results in coastal
101 seawater acidification and eventually degassing of CO₂ to the atmosphere (Wit et al., 2018;
102 Zhou et al., 2021). tDOC is also rich in chromophoric dissolved organic matter (CDOM), which
103 is that fraction of DOM that absorbs light. Increased fluxes of terrigenous CDOM can thus
104 reduce underwater light availability and spectral quality in coastal waters (Urtizberea et al.,
105 2013).

106

107 Despite the significance of tDOC and CDOM, long time series of their fluxes and
108 concentrations in coastal waters are only available in parts of Europe and North America.
109 Although CDOM and DOC can be estimated by satellite remote sensing (Sanwlani et al., 2022)
110 accurate remote sensing in optically complex coastal waters requires extensive optical and
111 biogeochemical field data for algorithm development, which limits the widespread use of this
112 technique. Moreover, satellite remote sensing can only provide data over the most recent few
113 decades. Longer records are required to understand the impact of anthropogenic changes that
114 pre-date the satellite era, as well as long-term and cyclical drivers of terrigenous CDOM and
115 tDOC variability such as temperature and hydrology. It is therefore necessary to find a
116 paleoproxy to reconstruct terrigenous dissolved organic matter (tDOM) parameters.

117

118 In the absence of measured instrumental records, we look to natural archives such as corals
119 for geochemical paleoproxies. Corals offer exceptional chronological constraints and allow for
120 approximately monthly-resolution climate and environmental proxy reconstruction over
121 hundreds of years (Thompson, 2022 and references therein). It has long been known that
122 skeleton cores from corals such as *Porites* spp., composed of the calcium carbonate polymorph
123 aragonite, show luminescent layers under UV light (Isdale, 1984). The luminescence intensity
124 of these bands correlates with freshwater run-off, as shown in multiple locations including

125 Australia (Rodriguez-Ramirez et al., 2014), Florida Bay (Smith et al., 1989) and Madagascar
126 (Grove et al., 2013).

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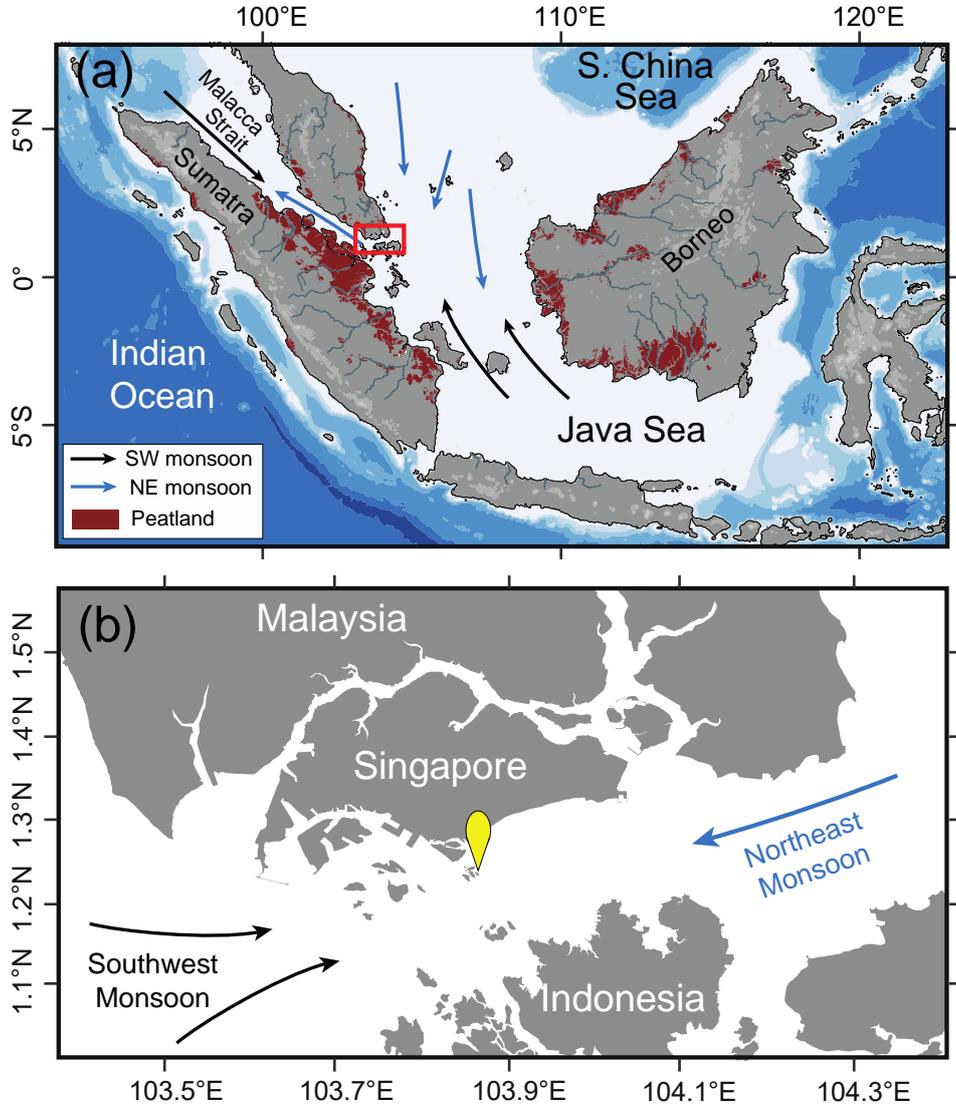
128 The luminescence is caused by the incorporation of humic-like substances into the coral
129 skeleton during growth (Kaushal et al., 2020; Susic et al., 1991). Humic-like substances on
130 land are formed from the breakdown of plant tissue, and after leaching into aquatic ecosystems
131 they form an integral component of terrigenous CDOM and tDOC. Humic-like substances are
132 rich in fluorescent aromatic moieties, and this fluorescence extends to longer wavelengths than
133 that of coral skeletal aragonite. Therefore, when a coral core is illuminated with ultraviolet
134 light, the ratios of emitted luminescence in the green wavelength band to that in the blue
135 wavelength band (G/B ratio) measures downcore variability in the concentration of terrigenous
136 humic-like substances, while normalizing for luminescence variation caused by changing coral
137 skeletal structure (Grove et al., 2010; Kaushal et al., 2020). Moreover, we recently showed that
138 luminescence G/B in a coral from Borneo tracks sub-annual terrigenous CDOM variability, as
139 estimated from satellite-based measurements of the CDOM absorption coefficient at 440 nm
140 (Kaushal et al., 2021). While luminescence G/B thus clearly has high potential as a paleoproxy
141 for tDOM, reconstructing just CDOM absorption at a single wavelength can only give limited
142 insights into land–ocean carbon flux dynamics. Here, we analyse two replicate coral cores
143 collected from Singapore, where we collected ~monthly measurements over 2 years of CDOM
144 absorption spectra, DOC concentration, and estimated tDOC concentration based on carbon
145 stable isotopes (Zhou et al., 2021). We show that coral luminescence G/B can be used as a
146 quantitative proxy to estimate tDOC concentration as well as the full CDOM absorption
147 spectrum, demonstrating that this proxy can provide considerable insights into coastal carbon
148 cycling in the tropics.

149

150 **2. Material and methods**

151

152 **2.1 Study site and coral core collection**



153

154 Figure 1: (a) Map of our study region showing the location of tropical peatlands and seasonal

155 monsoon wind directions. During the northeast (NE) monsoon months, the currents transport

156 waters from the South China Sea to our study region. During the southwest (SW) monsoon

157 months, the currents reverse, transporting waters with inputs from tropical peatlands located

158 on the east coast of Sumatra to our study site. Red rectangle indicates the region shown in panel

159 (b). (b) Site for time series sampling and coral core collection in the Singapore Strait (yellow

160 marker).

161 The Singapore Strait is located in the central Sunda Shelf Sea (Fig. 1) where water
162 depths are mostly <50 m, and strong tidal currents fully mix the water column (Mayer and
163 Pohlmann, 2014). The annual net circulation runs from the South China Sea into the Indian
164 Ocean via the Malacca Strait owing to the pressure gradient between the South China Sea and
165 the Andaman Sea that results from the basin-wide response to the monsoon winds. The
166 monsoon system also causes seasonal reversal of currents, and during the peak southwest (SW)
167 monsoon months from June to August, the current flows eastward from the coast of Sumatra
168 through the Singapore Strait (van Maren and Gerritsen, 2012). Sumatra harbours the largest
169 deposits of peat in Southeast Asia (Page et al., 2011) and the rivers draining these peatlands
170 (which carry up to 5000 $\mu\text{mol l}^{-1}$ DOC) are one of the most significant sources of tDOC to
171 Southeast Asian coastal waters (Alkhatib et al., 2007; Baum et al., 2007; Rixen et al., 2008;
172 Wit et al., 2018). The seasonally reversing currents thus result in seasonally high concentrations
173 of tDOC and terrigenous CDOM in the Singapore Strait (Martin et al., 2021; Zhou et al., 2021).

174
175 One plug core was collected from the main growth axis of each two massive *Porites* spp.
176 corals (KU-K and KU-L) off Kusu Island in the Singapore Strait (1.226°N 103.860°E, Fig. 1)
177 at 2-3 m depth in Nov-2020, using an underwater electric drill with a 3 cm diameter, 30 cm
178 length diamond bit core barrel. The cores were cleaned with water and sliced (~0.7 cm thick)
179 using a rotary saw, and then further cleaned for 48 hours in a 1:4 dilution of household bleach
180 (NaOCl, 3–7% reactive chlorine) and sonicated in deionized water for a total of 30 min (water
181 changed every 10 min); this removes contaminants and increases the luminescence intensity
182 (Nagtegaal et al., 2012). The KU-K core (hereafter KU-K) measures ~13 cm in length and has
183 11 visible couplets of dark and light bands. The KU-L plug core (hereafter KU-L) measures ~7
184 cm in length and has 7 visible couplets of dark and light bands (Figure 3). The seasonal timing
185 of luminescent band formation in Singapore was previously ascertained through alizarin

186 staining and repeated subsampling of tagged colonies (Tanzil et al., 2016). The dark bands in
187 true colour images (bright bands in luminescence images) represent coral skeletal growth
188 during the southwest monsoon period of high tDOC influx (Tanzil et al., 2016; Zhou et al.,
189 2021). Here, we subsampled 4.8 cm of the KU-K and 3.0 cm of the KU-L cores, covering the
190 growth period from the end of the SW monsoon in 2017 to core collection in Nov-2020.

191

192 **2.2 Kusu seawater biogeochemical sampling**

193

194 Biogeochemical sampling was conducted at Kusu once or twice per month at 5 m depth
195 from Oct-2017 to Nov-2020. Details of collection, preservation, measurements and analysis of
196 water samples are described in detail in Zhou et al., 2021 (Zhou et al., 2021), and are only
197 briefly mentioned here and below (Section 2.3.1). Profiles of salinity and temperature were
198 measured using a FastCTD Profiler (Valeport Ltd). Water samples for CDOM, DOC and
199 $\delta^{13}\text{C}_{\text{DOC}}$ measurements were collected using a Niskin bottle and immediately filtered on the
200 boat through a pre-rinsed 47 mm diameter, 0.22 μm pore-size polyethersulfone membrane filter
201 (Supor, Merck Millipore) in an in-line filter housing connected to a peristaltic pump.

202

203 **2.3 Experimental methods**

204

205 **2.3.1 Water sample analysis**

206

207 CDOM absorption spectra were measured on a Thermo Evolution 300 dual-beam
208 spectrophotometer in a 10-cm quartz cuvette from 230 to 900 nm. Ultrapure deionized water
209 ($18.2 \text{ M}\Omega \text{ cm}^{-1}$) was used as the reference. The CDOM absorbance (A_λ) was converted to
210 absorption coefficient (a_λ) using:

211 $a_{\lambda} = 2.303 \times (A_{\lambda} \div l)$ (1)

212 where, a_{λ} and A_{λ} are, respectively, the absorption coefficient and the absorbance at wavelength
213 λ , and l is the cuvette length in m. We refer to CDOM absorption at wavelength λ as $a_{\text{CDOM}}(\lambda)$.
214 SUVA_{254} is the specific UV absorbance at 254 nm and was calculated from the absorbance at
215 254 nm and the DOC concentration (Weishaar et al., 2003). The CDOM spectral slope $S_{275-295}$
216 and spectral slope ratio S_R (the ratio of $S_{275-295}$ to the spectral slope between 350-400 nm) were
217 calculated as in Helms et al. (2008) (Helms et al., 2008). DOC samples (30 ml) were acidified
218 with 100 μl 50% v/v H_2SO_4 and analyzed on a Shimadzu TOC-L system. Potassium hydrogen
219 phthalate was used for calibration and accuracy was monitored using deep-sea certified
220 reference material from the University of Miami, USA with a long-term reproducibility of 48.0
221 $\pm 3.9 \mu\text{mol L}^{-1}$. The $\delta^{13}\text{C}_{\text{DOC}}$ samples were measured at the Jan Veizer Stable Isotope
222 Laboratory, University of Ottawa Canada using an OI Analytical Aurora Model 1030W TOC
223 Analyzer interfaced to a Finnigan Mat DeltaPlusXP isotope ratio mass spectrometer with a
224 long-term reproducibility of $\pm 0.4 \%$. The spectral slope ratios were published in Martin et al.,
225 2021 (Martin et al., 2021), while the remaining parameters (water temperature, salinity, CDOM
226 absorption, $S_{275-295}$, SUVA_{254} , DOC concentration, $\delta^{13}\text{C}_{\text{DOC}}$) were published in Zhou et al, 2021
227 (Zhou et al., 2021). Here, we used monthly averaged values of the water measurements for
228 comparison against coral data. We used the higher-resolution $a_{\text{CDOM}}(350)$ data *only* for age-
229 depth model development.

230 **2.3.2 Coral core luminescence, $\delta^{18}\text{O}$ and Sr/Ca measurements**

231

232 Luminescence was measured parallel to the sampling track for Sr/Ca and $\delta^{18}\text{O}$
233 measurements. The cleaned KU-K and KU-L sections were scanned under ultraviolet light
234 (365 nm) using spectral line scanning on an Avaatech XRF core scanner at the Asian School

235 of the Environment following Grove et al. (2010) and Tanzil et al. (2016). The line scan camera
236 records luminescence emission with the light source and recording camera progressing down
237 the coral section as a single unit scanning multiple lines that are stitched together to produce a
238 continuous core section image. The incoming light from the sample is split into three
239 wavelength bands (red, green and blue) by a dichroic RGB beam splitter prism and recorded
240 by separate sensors. Luminescence green-to-blue (G/B) measurements (or coral G/B) were
241 obtained along the main growth axis for a 2 mm wide track at a resolution of 143 pixels cm⁻¹
242 (~0.07 mm), with wavelength bands of green: 525-575 nm and blue: 425-475 nm. Ratioing to
243 the blue band normalizes for changes to the luminescence signal resulting from downcore
244 changes in coral density and architecture (Grove et al., 2010). Coral samples for Sr/Ca and $\delta^{18}\text{O}$
245 were drilled continuously at 1 mm resolution for Sr/Ca and $\delta^{18}\text{O}$ measurements. Further details
246 of these measurements have been provided in Supplementary Information.

247

248 **2.4 Theory and calculation**

249

250 **2.4.1 Estimating terrigenous Dissolved Organic Carbon (tDOC)** 251 **concentration**

252

253 Zhou et al. (2021) used a two-endmember mixing model to estimate the concentration of
254 tDOC from our measured time series of DOC concentration and $\delta^{13}\text{C}_{\text{DOC}}$ in the Singapore
255 Strait, and the estimated endmember $\delta^{13}\text{C}_{\text{DOC}}$ values for peatland-derived DOC (i.e., pure
256 tDOC) and for marine DOC. The $\delta^{13}\text{C}_{\text{DOC}}$ endmember values (mean $\pm 1\sigma$ uncertainty) were
257 taken as $-29 \pm 1\text{‰}$ for peatland DOC, based on data from Sumatra, Borneo, and peninsular
258 Malaysia, and as $-21.88 \pm 0.79\text{‰}$ for autochthonous marine DOC, based on the Singapore
259 Strait data during the late NE monsoon and inter-monsoon (late February-March, when the

260 Singapore Strait receives waters from the open South China Sea). The tDOC concentration for
261 each sampling date was calculated by Zhou et al. (2021) as follows:

262 In our coastal sites, the measured DOC concentration ($[DOC]_{meas}$) is the sum of the
263 terrigenous fraction ($[tDOC]$) and the marine fraction ($[mDOC]$):

$$264 \quad [DOC]_{meas} = [tDOC] + [mDOC] \quad (2)$$

265 Also, the measured concentration of $DO^{13}C$ ($[DO^{13}C]_{meas}$) is the sum of the terrigenous fraction
266 ($[DO^{13}C]_{tDOC}$) and the marine fraction ($[DO^{13}C]_{mDOC}$):

$$267 \quad [DO^{13}C]_{meas} = [DO^{13}C]_{tDOC} + [DO^{13}C]_{mDOC} \quad (3)$$

268 The $[DO^{13}C]_{meas}$ is calculated from the measured DOC and $\delta^{13}C_{DOC}$ as:

$$269 \quad [DO^{13}C]_{meas} \approx [DOC]_{meas} \times R_{meas} \quad (4)$$

270 where R_{meas} is the measured carbon isotope ratio, calculated as:

$$271 \quad R_{meas} = (\delta^{13}C_{DOCmeas}(\text{‰}) \div 1000 + 1) \times R_{VPDB} \quad (5)$$

272 The $[DO^{13}C]_{tDOC}$ is approximated as:

$$273 \quad [DO^{13}C]_{tDOC} \approx [tDOC] \times R_{tDOC} \quad (6)$$

274 where R_{tDOC} is the carbon isotope ratio of tDOC, and calculated using the riverine endmember
275 of carbon isotopic composition of DOC:

$$276 \quad R_{tDOC} = (\delta^{13}C_{DOCriver}(\text{‰}) \div 1000 + 1) \times R_{VPDB} \quad (7)$$

277 Similarly, R_{mDOC} is the carbon isotope ratio of marine DOC and is calculated in the same
278 fashion from the marine DOC (mDOC) $\delta^{13}C$ endmember value. Equation 2 can then be
279 rewritten as:

$$280 \quad [DOC]_{meas} \times R_{meas} = [tDOC] \times R_{tDOC} + [mDOC] \times R_{mDOC} \quad (8)$$

281 where R_{mDOC} is the carbon isotope ratio of marine DOC, calculated using the marine
282 endmember of carbon isotopic composition of DOC.

283 Solving Eqn. 2 and Eqn. 8 will return the concentration of terrigenous DOC.

284

2.4.2 Coral age-depth model

The age-depth models were created using coral G/B and our measured $a_{\text{CDOM}(350)}$, which has a pronounced SW monsoon peak (Zhou et al., 2021). Seasonal coral luminescence G/B at this site are distinct and cyclically reproducible between colonies from the same location, while skeletal density variations show a lack of discernable annual patterns (Tanzil et al., 2016). Sixty-three $a_{\text{CDOM}(350)}$ measurements were made at Kusu from Oct-2017 to Oct-2020, which provides a less ambiguous seasonal marker than water temperature or salinity (Supp. Fig. 1). QAnalySeries software was used to peak-align the 2018 and 2019 SW monsoon peaks in $a_{\text{CDOM}(350)}$ with corresponding peaks in coral G/B, providing an age-depth model for each core, yielding a growth rate of 16 mm year⁻¹ for KU-K and 10 mm year⁻¹ for KU-L. To further validate our age-depth model, we compared our temperature and salinity time series data to the coral Sr/Ca and $\delta^{18}\text{O}$ measurements.

2.4.3 Reconstructing full CDOM absorption spectra from coral G/B

We used two different methods to test whether coral G/B can be used to reconstruct full CDOM absorption spectra. In Method 1, we used coral G/B to reconstruct CDOM absorption at 300 nm and 400 nm, set the absorption at 700 nm to 0 m⁻¹ (Green and Blough, 1994) and fit a first-order exponential curve through the three points. We also test a second approach (Method 2), in which we used coral G/B to reconstruct absorption at wavelength 350 and took the average the *measured* water CDOM spectral slope over the range 300–550 nm, and calculated the rest of the absorption spectrum following (Twardowski et al., 2004):

$$a_{\lambda} = a_{\lambda_{\text{ref}}} e^{-S(\lambda - \lambda_{\text{ref}})} \quad (9)$$

310

311 where a_λ and $a_{\lambda_{ref}}$ are the absorption coefficients (m^{-1}) at wavelengths λ and λ_{ref} , λ_{ref} is the
312 reference wavelength (350 nm in our case), and S is the spectral slope between 300–550 nm.

313

314 **3. Results**

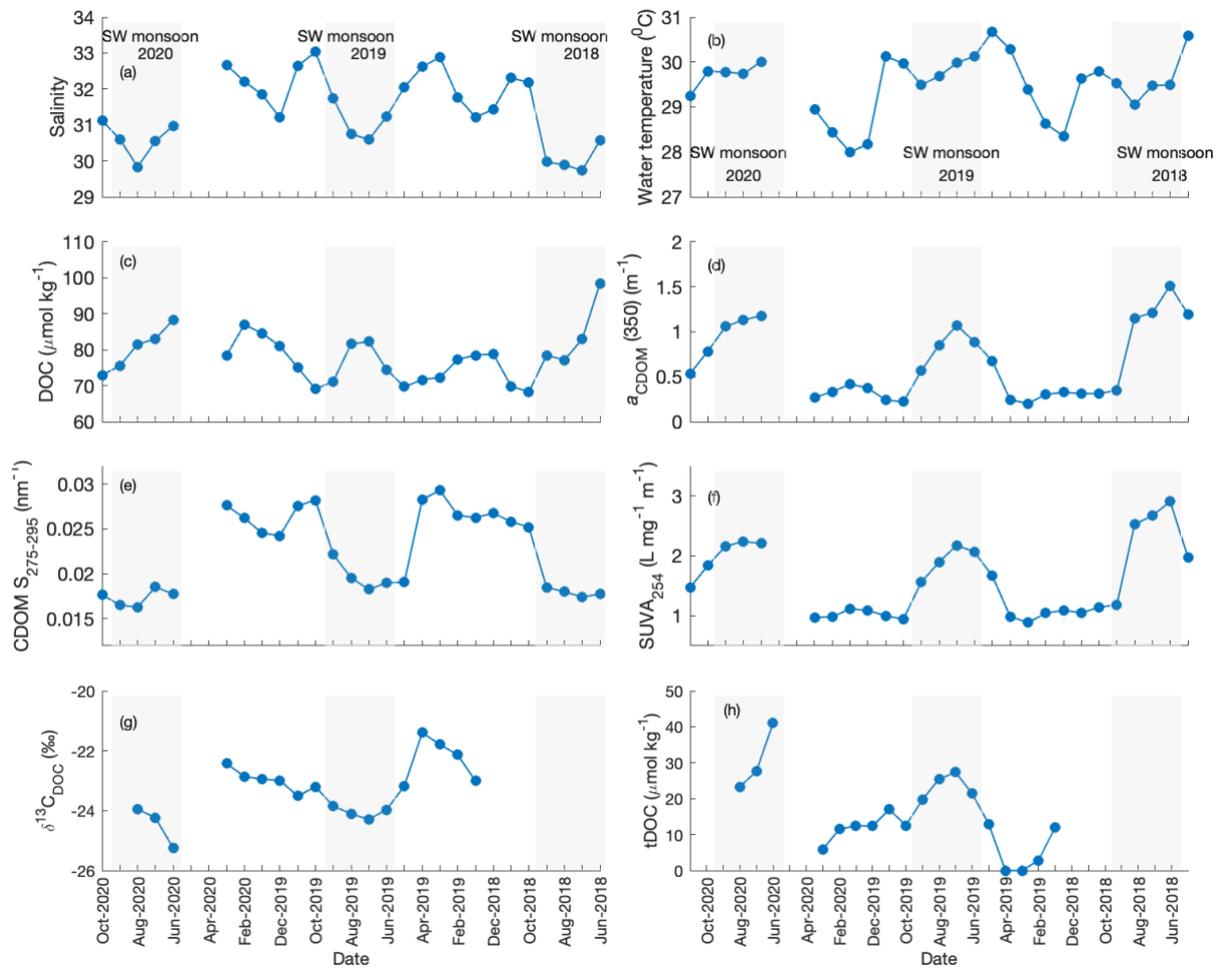
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316 **3.1 Monsoon-driven seasonality of tDOC in the Singapore Strait**

317

318 As previously described (Martin et al., 2021; Zhou et al., 2021), the Singapore Strait
319 experiences a large input of tDOC and terrigenous CDOM during the SW monsoon (mid-May
320 to mid-September, Fig. 3), when currents transport peatland-influenced waters from the east
321 coast of Sumatra through the Singapore Strait. Consequently, there is a drop in salinity from
322 ~33 to ~30, spikes in DOC concentration ($>80 \mu\text{mol kg}^{-1}$) and in CDOM absorption
323 ($a_{\text{CDOM}(350)} > 0.5 \text{ m}^{-1}$) and $\text{SUVA}_{254} (> 1.5 \text{ L mg}^{-1} \text{ m}^{-1})$, large decreases in CDOM spectral slope
324 $S_{275-295} (< 0.020 \text{ nm}^{-1})$ and in $\delta^{13}\text{C}_{\text{DOC}} (< -24 \text{ ‰})$, and high estimated tDOC concentrations ($>$
325 $20 \mu\text{mol kg}^{-1}$).

326 The early NE monsoon (December-January), when currents begin to transport open South
327 China Sea water to the Singapore Strait, is also the season with most rainfall in Singapore and
328 southern Malaysia. At this time, there is a smaller drop in salinity from 33 to 31, and lowest
329 annual water temperatures of 28°C . DOC concentration increases to $\sim 80 \mu\text{mol kg}^{-1}$, comparable
330 to the SW monsoon, but indicators of tDOM input show much smaller changes than during the
331 SW monsoon: $a_{\text{CDOM}(350)}$ is $< 0.5 \text{ m}^{-1}$, $\text{SUVA}_{254} \sim 1 \text{ L mg}^{-1} \text{ m}^{-1}$, $\delta^{13}\text{C}_{\text{DOC}} \sim -22.5\text{‰}$, and $S_{275-295}$
332 $\sim 0.025 \text{ nm}^{-1}$. The estimated tDOC concentration is therefore consistently $< 20 \mu\text{mol kg}^{-1}$.



333

334 **Figure 2:** Monthly average biogeochemical data at Kusu Island (for full time series with all

335 individual data, see (Martin et al., 2021; Zhou et al., 2021)). Note: The X-axis goes from

336 younger-to-older from left-to-right to facilitate comparison with the coral data. (a) Salinity

337 indicates a large input of freshwater during the SW monsoon (grey shading), while (b) water

338 temperature shows limited seasonal variation with lowest temperature during the NE monsoon.

339 (c) The bulk DOC pool shows clear peaks during the SW and NE monsoon, but (d-f) the CDOM

340 parameters and (g) $\delta^{13}\text{C}_{\text{DOC}}$ show that a large input of tDOM only takes place during the SW

341 monsoon. (h) Concentration of tDOC as calculated from the DOC concentration and the

342 $\delta^{13}\text{C}_{\text{DOC}}$ data. The tDOC for March 2019 and April 2019 are less than zero, which is a result of

343 using a single, fixed marine endmember value for $\delta^{13}\text{C}_{\text{DOC}}$ (from late February-March).

344

345

3.2 Coral age-depth model

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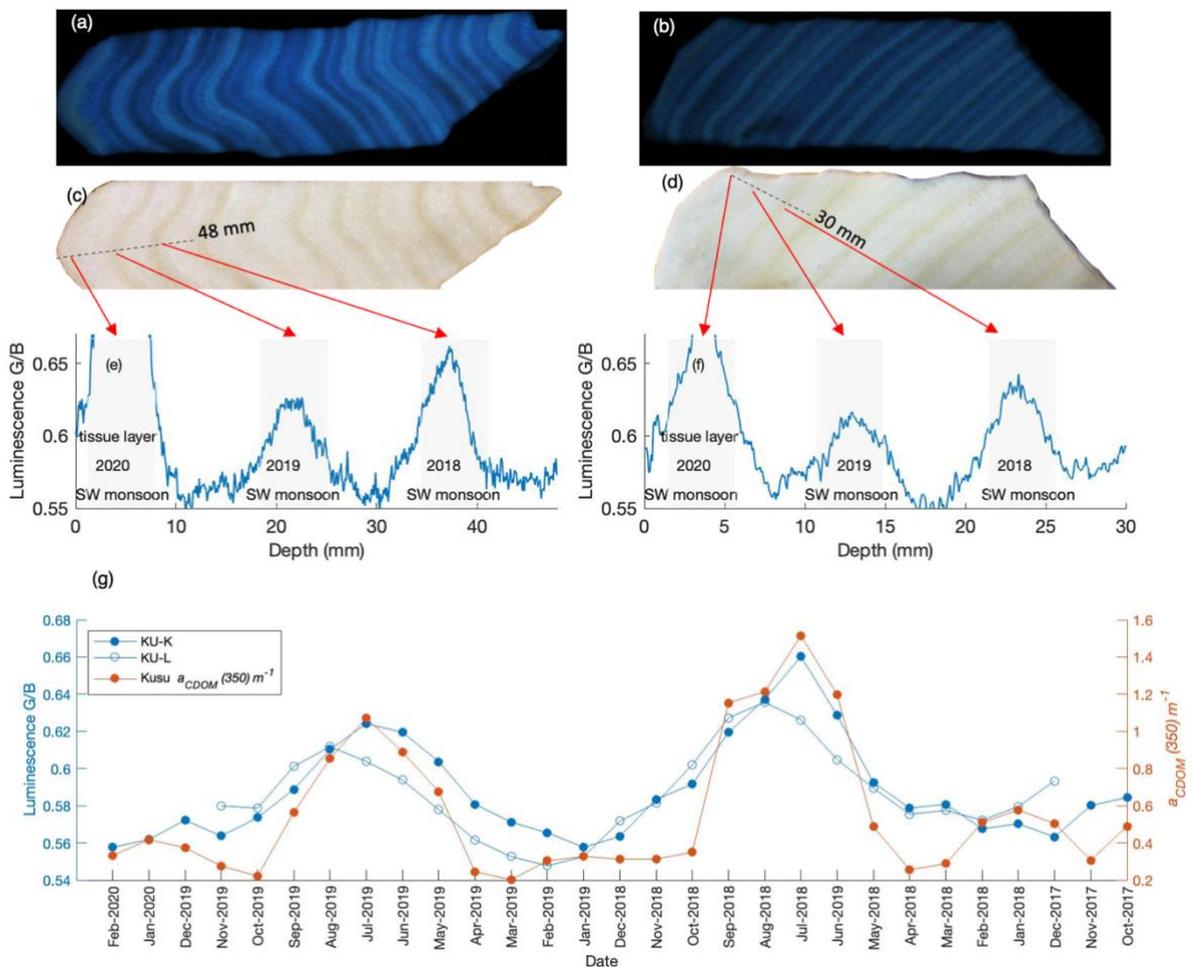
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352

Following the results from peak-aligning we use the KU-K section that grew from Jan-2018 to Feb-2020, and the KU-L section that grew from Oct-2017 to Nov-2019 for analysis in this study. These sections cover both the 2018 and 2019 SW monsoons, represented by bright layers in the coral luminescence images (Fig. 3a and 3b), dark brown colored bands in the true-colour images (Fig. 3c and 3d) and peaks in coral G/B ratio (Fig. 3e and 3f). The $a_{CDOM}(350)$ peak-aligned coral G/B values are shown in Fig. 3g.



353

354

355

356

Figure 3: Coral luminescence G/B measurements. Luminescence (under ultraviolet excitation) and true-color images of coral plug cores (a,c) KU-K and (b,d) KU-L from Kusu Island. Dashed black lines in (c,d) indicate the growth axis along which luminescence G/B was quantified.

357 (e,f) The raw coral G/B measurements have been plotted against core depth. The core top
358 ‘tissue layers’ show very high G/B and are omitted from our analysis. The approximate timing
359 of the 2019 and 2018 SW monsoon periods are marked. (g) A monthly resolution age model
360 was developed for the coral G/B by peak-aligning with measured $a_{CDOM}(350)$.

361

362 We then compared the measured temperature and salinity data to the corresponding coral
363 Sr/Ca and $\delta^{18}O$ data according to our age-depth model (Supp. Fig. 2). In both cores, high water
364 temperature coincides with low Sr/Ca, and high salinity coincides with high $\delta^{18}O$, which
365 provides additional, independent validation of our age model.

366

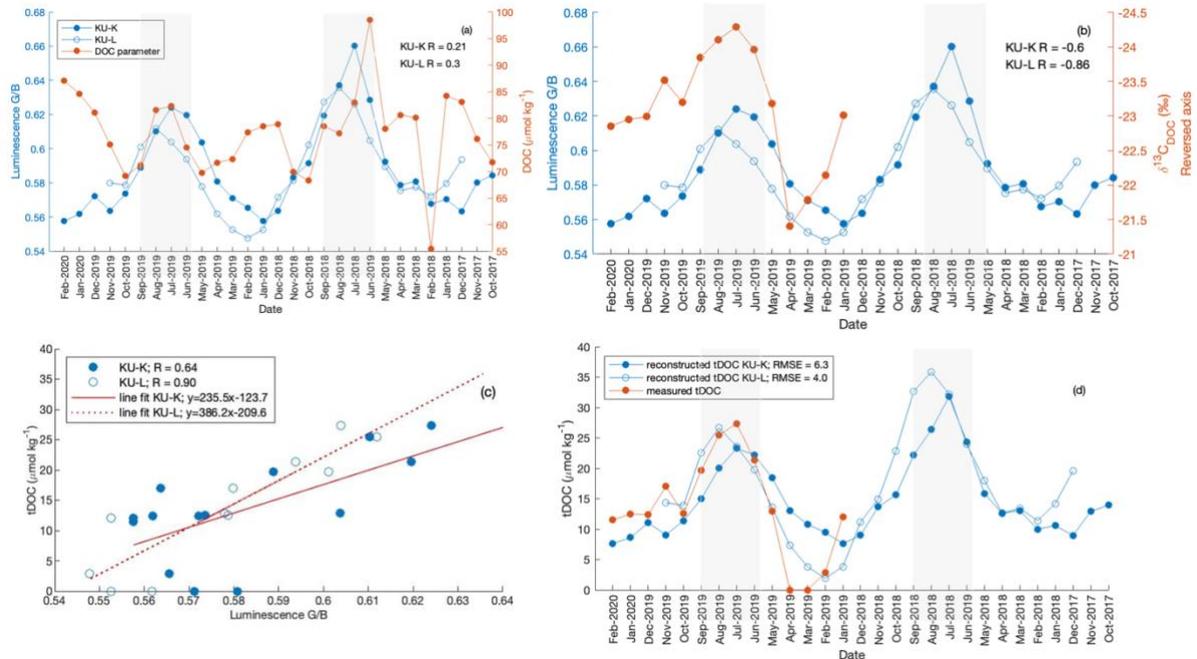
367 **3.3 Reconstructing tDOC and CDOM using coral G/B as a proxy**

368

369 To test whether coral G/B ratios can track tDOC in coastal waters we compared coral G/B
370 ratios with bulk DOC (which includes both the marine and terrigenous components of DOC;
371 Fig. 4a), with $\delta^{13}C_{DOC}$ (where more negative values indicate a greater contribution from
372 terrigenous DOC to the bulk DOC pool; Fig. 4b), and with our calculated tDOC concentration
373 (Fig. 4c and 4d). The coral G/B ratio varied between 0.58 to 0.68 with distinct peaks in the
374 SW monsoon, which coincided with the SW monsoon peak in bulk DOC (Fig. 4a). However,
375 the coral G/B did not reflect the NE monsoon peak in bulk DOC, and coral G/B was therefore
376 only poorly correlated with bulk DOC (KU-K $R=0.21$; KU-L $R = 0.3$; Fig. 4a). Coral G/B
377 instead correlated well with $\delta^{13}C_{DOC}$ (KU-K $R = -0.60$; KU-L $R = -0.86$; Fig. 4b) and with
378 calculated tDOC concentration (KU-K $R = 0.64$; KU-L $R = 0.90$; Fig. 4c,d). We then used the
379 two regression models from Fig. 4c to reconstruct tDOC from the coral G/B records; this
380 reconstruction has an RMSE of $\pm 6.3 \mu\text{mol kg}^{-1}$. Our reconstructed tDOC time series varies

381 from 3 to 37 $\mu\text{mol kg}^{-1}$ between Oct-2017 and Feb-2020, with both cores indicating higher
 382 tDOC concentrations during the 2018 SW monsoon than in 2019 (Fig. 4d).

383

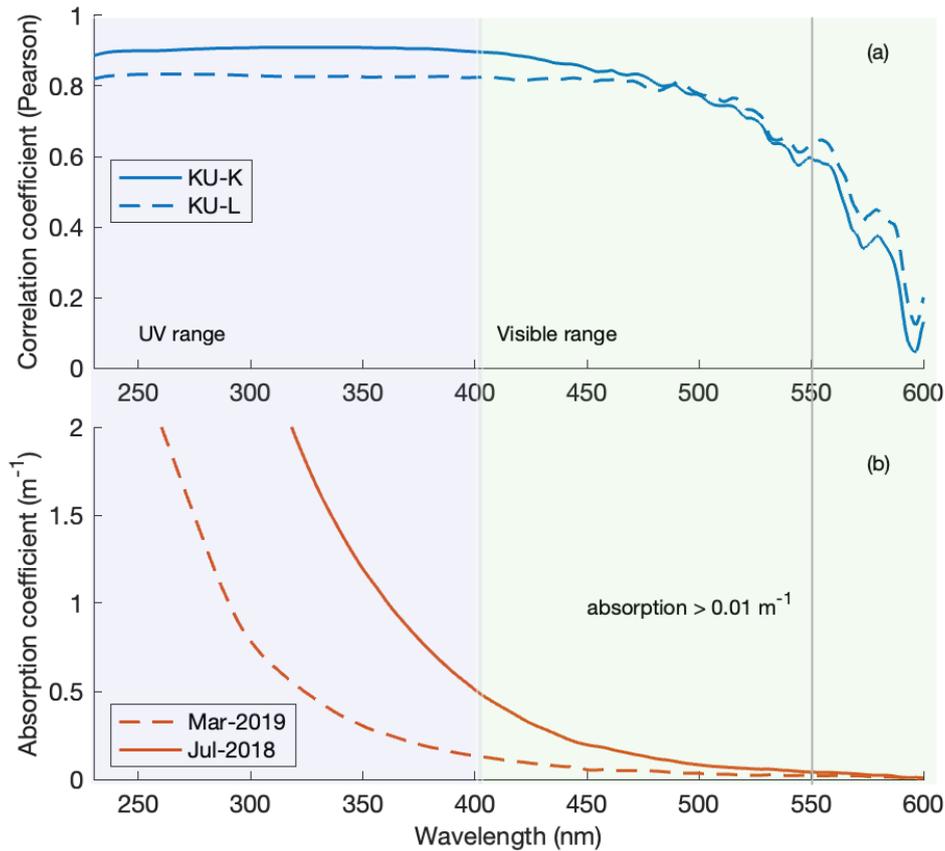


384

385 **Figure 4:** tDOC reconstruction using coral G/B measurements. (a) bulk DOC concentration,
 386 (b) $\delta^{13}\text{C}_{\text{DOC}}$, (c) coral G/B versus tDOC (d) and reconstructed tDOC have been plotted. tDOC
 387 has been calculated using the bulk DOC concentration and $\delta^{13}\text{C}_{\text{DOC}}$ measurements and an
 388 isotope mass balance approach. The relationship between measured coral G/B and tDOC has
 389 been used to reconstruct tDOC going back to Oct-2017. The measured tDOC and reconstructed
 390 tDOC values show RMSE values of 6.3 $\mu\text{mol kg}^{-1}$ during the overlapping period from Jan-
 391 2019 to Feb-2020.

392

393 Moreover, coral G/B was strongly correlated ($R > 0.6$) with monthly average CDOM
 394 absorption at every wavelength between 230 to 550 nm (Fig. 5a), which essentially covers the
 395 full wavelength spectrum of CDOM absorption. At wavelengths greater than 550 nm, monthly
 396 mean CDOM absorption was always $< 0.01 \text{ m}^{-1}$ (Fig. 5b).



397

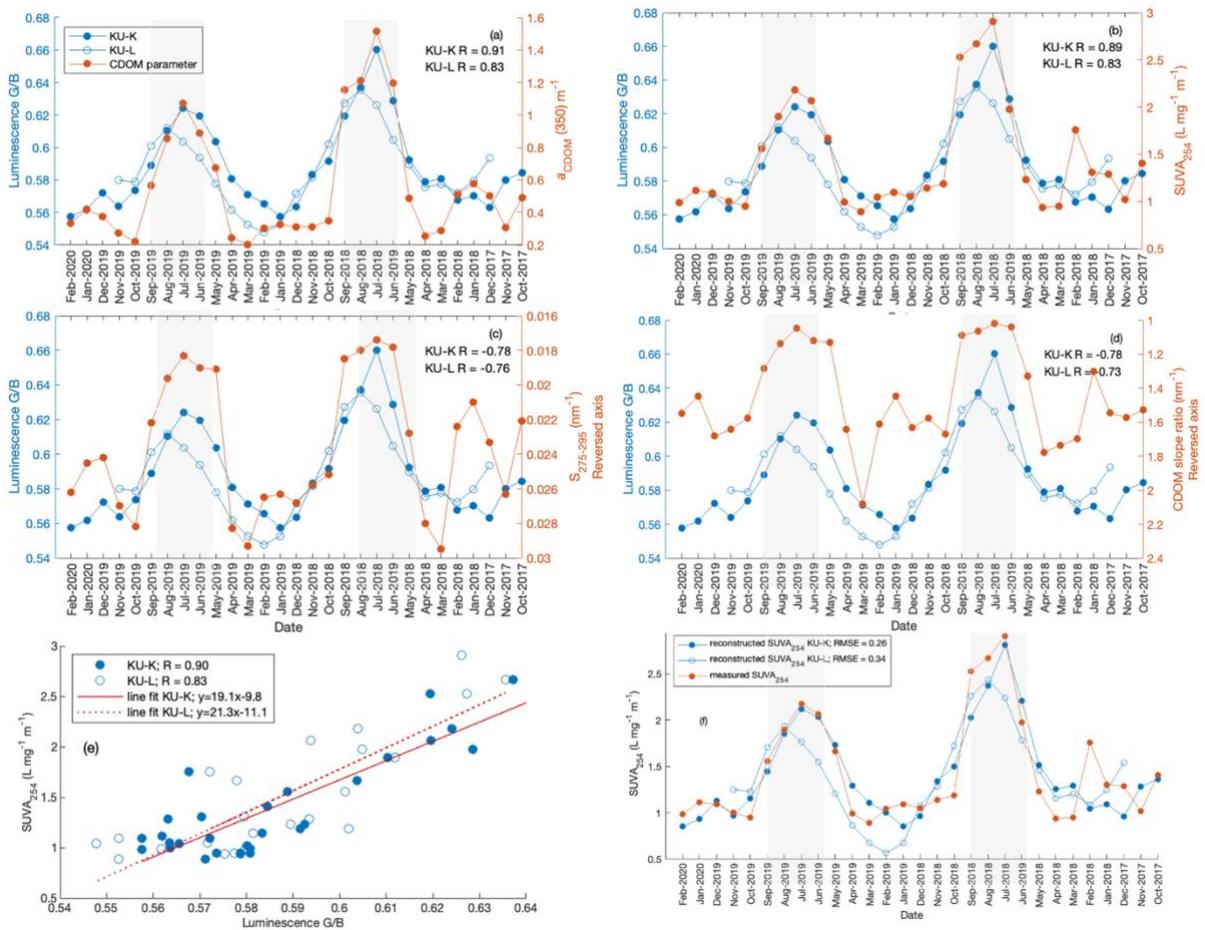
398 **Figure 5:** (a) Pearson's correlation coefficient between coral G/B and CDOM absorption
 399 coefficient at every wavelength between 230-600 nm. Strong correlations ($R \geq 0.6$) are seen at
 400 all wavelengths up to 550 nm. (b) CDOM absorption spectra measured at Kusu in Mar-2019
 401 and Jul-2018, which mark the lowest and highest CDOM absorption measured in our time
 402 series, respectively. At wavelengths greater than 550 nm, CDOM absorption was always < 0.01
 403 m^{-1} , which indicates that coral G/B correlates with CDOM absorption throughout the full
 404 CDOM absorption spectrum.

405

406 CDOM absorption and spectral slope parameters are a valuable way of comparing CDOM
 407 variability in different environments (Vantrepotte et al., 2015) in addition to providing
 408 independent information on the composition and biogeochemical processing of CDOM (e.g.
 409 Helms et al., 2008; Weishaar et al., 2003). Of the CDOM parameters, coral G/B was most
 410 strongly correlated with $a_{\text{CDOM}}(350)$ (KU-K $R = 0.91$; KU-L $R = 0.83$; Fig 6a) and with

411 SUVA₂₅₄ (KU-K R = 0.89; KU-L R = 0.83; Fig. 6b), which are good tracers of terrigenous
 412 DOM in this region (Martin et al. 2018; 2021; Zhou et al. 2021). Coral G/B was also quite
 413 strongly correlated with the CDOM spectral slope parameters $S_{275-295}$ (KU-K R = -0.78; KU-L
 414 R = -0.76; Fig 6c) and slope ratio (KU-K R = -0.78; KU-L R = -0.73; Fig. 6d), although these
 415 correlations are weaker than for $a_{CDOM}(350)$ and SUVA₂₅₄ because $S_{275-295}$ and the slope ratio
 416 showed little difference between the 2018 and 2019 SW monsoons. SUVA₂₅₄ is emerging as a
 417 key measure to understand and model tDOC in the ocean (Anderson et al., 2019) and we found
 418 that coral G/B can also be used as a proxy to reconstruct SUVA₂₅₄, with low RMSE values of
 419 ± 0.34 L mg⁻¹ m⁻¹ (Fig. 6e,f).

420



421

422 **Figure 6:** Coral G/B has been correlated (Pearson's correlation coefficient) with CDOM
 423 parameters relevant for tracing tDOM in the coastal ocean. The highest correlations are seen

424 with (a) $a_{\text{CDOM}}(350)$, and (b) SUVA_{254} , both of which trace terrigenous CDOM concentration
425 in this region. The correlation is slightly lower for the spectral slope parameters (c) $S_{275-295}$ and
426 (d) the slope ratio. Panels (e) and (f) demonstrate that coral G/B can be used to reconstruct
427 SUVA_{254} with low RMSE values of $\pm 0.34 \text{ L mg}^{-1} \text{ m}^{-1}$.

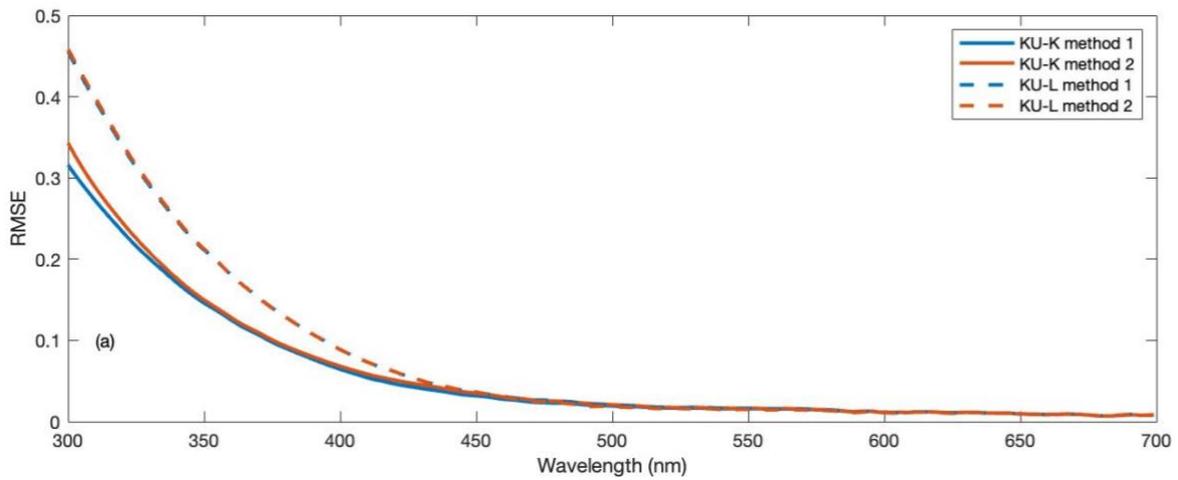
428

429 **3.4 Reconstructing full CDOM spectra using coral G/B as a proxy**

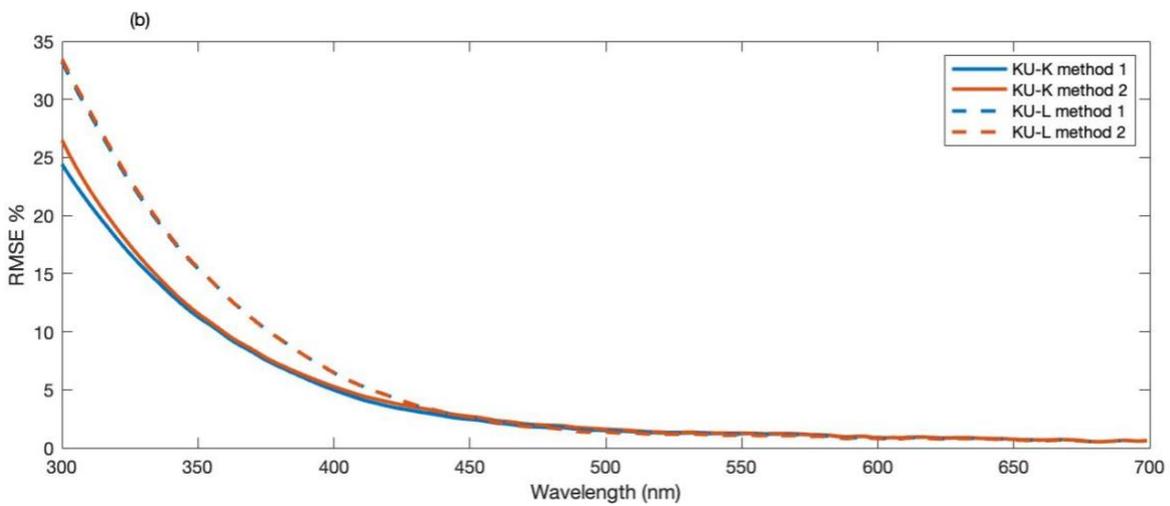
430

431 We tested two methods for reconstructing the full CDOM absorption spectrum: Method 1
432 simply reconstructs absorption at a limited number of points and estimates the full spectrum
433 by fitting an exponential curve to the reconstructed points, while Method 2 requires an estimate
434 of the CDOM spectral slope from field measurements. We find that both methods provide
435 similarly accurate reconstructions of the CDOM absorption spectrum, as measured by the
436 absolute and percentage RMSE between reconstructed and measured absorption at each
437 wavelength, although Method 1 may perform very slightly better (Fig. 7). The linear models
438 for reconstruction of absorption spectra, spectral slope measurements (from 300 to 550 nm)
439 and plots of measured and reconstructed spectra are shown in supplementary figures 3, 4 and
440 5.

441



442



443

444

445 **Figure 7:** RMSE (300-700 nm) (a) and RMSE percentage (b) against wavelength between the
446 measured CDOM spectra and spectra reconstructed from coral G/B using both reconstruction
447 methods. Method 1 uses coral G/B to reconstruct absorption at wavelengths 300 and 400 nm,
448 assumes 0 absorption at 700 nm, and fits a first-order exponential curve through these three
449 points. Method 2 uses coral G/B to reconstruct absorption at wavelength 350 and uses the
450 average CDOM spectral slope between 300-550 nm from in-situ CDOM measurements.

451

452 4. Discussion

453

454 **4.1 tDOC source and seasonal variability**

455

456 In this study, we use tDOC estimates calculated from carbon isotope mass balance
457 calculations and CDOM parameters as measured by Zhou et al, 2021 to quantify tDOM at our
458 study site. The terrigenous organic carbon in peatland-draining rivers into the Sunda Sea Shelf
459 is overwhelmingly in the form of DOC (typically >95%) rather than POC (Alkhatib et al., 2007;
460 Moore et al., 2011; Müller et al., 2015), so that our proposed proxy for tDOC captures the bulk
461 of the terrigenous carbon input to the Sunda Shelf Sea. Coastal waters in the Sunda Shelf Sea
462 receive relatively high tDOC inputs year-round (Sanwlanı et al., 2022; Wit et al., 2018; Zhou
463 et al., 2021). Although year-to-year variation in rainfall may cause some variation in the tDOC
464 flux from peatlands, the seasonal variability in tDOC seen in Singapore is caused by the
465 monsoonal reversals in ocean currents that bring peatland-influenced waters from the Sumatran
466 coast in the SW monsoon, and open marine waters from the South China Sea during the NE
467 monsoon (Zhou et al., 2021). The large and regular seasonal variation in tDOC at our site
468 therefore make this a particularly suitable location for coral paleoproxy development.

469

470 **4.2 Coral G/B is a sensitive proxy for tDOC concentration**

471

472 Based on a simple linear relationship, we have found that tDOC can be reconstructed from
473 coral G/B measurements with RMSE values of $\pm 6.3 \mu\text{mol kg}^{-1}$. The measured tDOC ranged
474 from 0 to $30 \mu\text{mol kg}^{-1}$ with SW monsoon values of $>20 \mu\text{mol kg}^{-1}$. From our coral G/B-based
475 reconstruction, we inferred a $10 \mu\text{mol kg}^{-1}$ difference in tDOC between the 2018 and 2019 SW
476 monsoon peaks with a higher concentration in 2018 than 2019. This is also reflected in the
477 measured CDOM parameters, bulk DOC concentration, and salinity, which suggested more

478 freshwater input and higher concentration of tDOM in the Singapore Strait in 2018 compared
479 to 2019 (Martin et al., 2021; Zhou et al., 2021).

480

481 Although the $\delta^{13}\text{C}_{\text{DOC}}$ suggests that there might be up to $10 \mu\text{mol kg}^{-1}$ tDOC during the NE
482 monsoon period, the CDOM parameters such as $a_{\text{CDOM}}(350)$ and SUVA_{254} show only small
483 changes at this time (Fig. 6). This may indicate that the tDOC pool at this time is inherently
484 less CDOM-rich and less aromatic, or that it has undergone more extensive prior
485 biogeochemical processing that preferentially removed CDOM (e.g. photobleaching; see
486 Helms et al., 2008; Martin et al., 2018). It is possible that less CDOM-rich and less aromatic-
487 rich tDOM may be less well incorporated by corals. Abiotic precipitation experiments have
488 shown that aragonite does not incorporate DOM indiscriminately, but preferentially
489 incorporates peatland-derived DOM over marine DOM (Kaushal et al., 2020), which may be
490 due to the variable structures of humic-like aromatic ring compounds and available sites for
491 binding with calcium carbonate. However, coral cores also show luminescence due to
492 terrestrial input in regions without peatlands (e.g. Grove et al., 2013), which shows that the
493 G/B proxy is not just specific to peatland-derived tDOC.

494

495 Instead, it is possible that tDOC during the NE monsoon was overestimated, since it was
496 calculated using a fixed value for the marine end-member $\delta^{13}\text{C}_{\text{DOC}}$, even though marine
497 autochthonous carbon can span a relatively wide range of $\delta^{13}\text{C}$ values (Verwega et al., 2021).
498 It is also not clear whether the peatland tDOC end-member $\delta^{13}\text{C}$ value used by (Zhou et al.,
499 2021) is appropriate during the NE monsoon, because the most likely sources of tDOC during
500 the NE monsoon would be local or regional input from the south and east coasts of the Malay
501 Peninsula, where few peatlands are found (Fig. 1).

502

503 Overall, our data show that coral G/B can be a sensitive proxy for tDOC concentration, and
504 that G/B is not affected by variation in the bulk concentration of marine autochthonous DOC.
505 However, it is possible that coral G/B might not be less sensitive to some types of tDOC that
506 are less aromatic and CDOM-rich, or that have undergone very extensive prior photobleaching.
507 Knowledge of the regional tDOC sources their CDOM:DOC ratios, and of the possible extent
508 of biogeochemical processing of the tDOC, is therefore advisable before applying this proxy
509 in different regions.

510

511 **4.3 Coral G/B can be used to reconstruct additional CDOM parameters that reflect** 512 **DOM composition**

513

514 CDOM parameters measure the absorption of light by DOM at different wavelengths, and
515 DOM that originates from partial degradation of terrigenous vegetation is particularly light-
516 absorbent (Massicotte et al., 2017; Vantrepotte et al., 2015a). This is reflected in the high
517 CDOM absorption measured during the SW monsoon (Fig. 5). The high correlation of > 0.6
518 between the measured CDOM absorption and coral G/B ratio is consistent with the corals
519 incorporating humic-like material of terrestrial origin. Of the different measured CDOM
520 parameters, the coral G/B measurements showed highest correlation with $a_{\text{CDOM}(350)}$ and with
521 SUVA_{254} ($R > 0.83$). SUVA_{254} is strongly correlated with aromaticity (Weishaar et al., 2003),
522 which fits well with our understanding of coral luminescence. SUVA_{254} was also recently
523 proposed as a measure to distinguish tDOC pools according to their source and reactivity
524 (especially as photolabile *versus* biolabile tDOC) in marine carbon cycle models (Anderson et
525 al., 2019). However, SUVA_{254} data from tropical coastal oceans are scarce, so the fact that this
526 parameter can be reconstructed with relatively good accuracy (RMSE of $\pm 0.34 \text{ L mg}^{-1} \text{ m}^{-1}$)
527 from long coral cores may help to inform better marine carbon cycle models.

528

529 Compared to CDOM absorption coefficients and to $SUVA_{254}$, we found that coral G/B was
530 slightly less strongly correlated with the CDOM spectral slope parameters $S_{275-295}$ and slope
531 ratio. In particular, the spectral slope parameters did not reflect the difference between the 2018
532 and 2019 SW monsoons in tDOM input (as inferred from our in-situ CDOM, DOC, and salinity
533 measurements). While both $S_{275-295}$ and slope ratio are very sensitive measures of the presence
534 of tDOM (Fichot and Benner, 2011; Vantrepotte et al., 2015; Zhou et al., 2021), CDOM
535 spectral slope properties change in a strongly non-linear manner when tDOM mixes with low-
536 CDOM marine DOM, such that even small quantities of terrigenous CDOM initially lead to
537 large changes in spectral slopes, but spectral slopes are then relatively insensitive to further
538 increases in terrigenous CDOM (Stedmon and Markager, 2003). Hence, slope ratios might not
539 be ideal measures of absolute concentration of tDOM and tDOC unless the terrigenous
540 contribution is relatively low. The fact that coral G/B reflects variation in CDOM absorption
541 coefficients and in $SUVA_{254}$ more closely than variation in CDOM spectral slope properties
542 therefore suggests that coral G/B primarily traces the variation in absolute tDOM
543 concentration.

544

545 The two methods of CDOM reconstruction from coral G/B work comparably well in our
546 case. Method 1 fits an exponential curve through reconstructed points and has a slightly lower
547 RMSE compared to Method 2 which uses spectral slope of CDOM from in-situ measurements.
548 Method 2 requires in-situ measurements, which are also likely to have varied in the past, thus
549 adding uncertainty to the paleo-reconstruction. Therefore, Method 1 provides a stronger
550 approach for the reconstruction of CDOM.

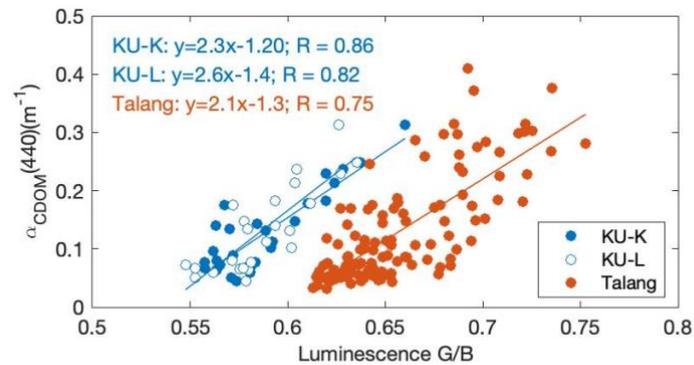
551

552 **4.4 Comparison to previous coral core calibration**

553

554 In our previous work, we calibrated G/B ratios of a coral core collected off the coast of
555 Borneo (Talang) with $a_{\text{CDOM}}(440)$ from satellite remote sensing, and also found a strong and
556 linear relationship (Kaushal et al., 2021). Comparing the relationships for coral G/B to
557 $a_{\text{CDOM}}(440)$ between our new, in-situ calibration from Kusu with the previous calibration from
558 the Talang core (Fig. 8) reveals that the slopes of liner regressions are statistically
559 indistinguishable since the standard errors (s.e.) of the slopes overlap (KU-K slope = 2.3 ± 0.3 ;
560 KU-L slope = 2.6 ± 0.4 ; Talang slope = 2.1 ± 0.2). Moreover, a regression analysis of the
561 combined dataset for all cores showed no significant interaction term between luminescence
562 and coral core. However, there is a clear offset, with the Talang core having higher G/B for a
563 given seawater $a_{\text{CDOM}}(440)$ value than either of the Kusu cores. This might simply reflect
564 individual differences between coral colonies, although the fact that the two Kusu colonies
565 have nearly identical calibrations might also indicate that there is a consistent difference in the
566 tDOM pool between the sites. The Talang coral is located close to a peatland-draining river,
567 and thus receives tDOM that is likely comparatively fresh and undegraded, given the
568 predominantly conservative mixing pattern of tDOM in Talang coastal waters (Martin et al.,
569 2018). In contrast, the Kusu cores receive tDOM from more distant rivers on Sumatra, and this
570 tDOM has already undergone extensive remineralization and likely significant photobleaching
571 in the coastal ocean prior to reaching the Singapore Strait (Zhou et al., 2021). Coral G/B is
572 caused by the incorporation of a humic- and aromatic-rich fraction of the fluorescent tDOM
573 pool, which itself is a subset of the light-absorbing CDOM pool. It is therefore possible that
574 corals preferentially incorporate a relatively labile fraction of the tDOM pool (perhaps
575 especially to photobleaching), in which case the observed offset between corals might be linked
576 to how rich the terrigenous CDOM pool at each site is in this putatively labile tDOM fraction
577 incorporated by the corals. Further research is clearly warranted to identify which chemical

578 fractions of the tDOM pool contribute to coral G/B. Importantly, while our results suggest that
579 local calibrations of coral G/B to measured tDOM parameters may be desirable, the similarity
580 in calibration slopes between both sites suggests that at least relative variation in terrigenous
581 CDOM can be inferred from coral G/B without a local calibration.



582
583 **Figure 8:** The relationship between coral G/B measurements of the KU-K and KU-L cores and
584 measured $a_{CDOM}(440)$ is compared to the relationship between the Talang coral core G/B
585 measurements and $a_{CDOM}(440)$ as estimated from satellite remote sensing (Kaushal et al.,
586 2021). The offset on the luminescence axis may suggest that corals preferentially incorporate
587 a relatively labile fraction of the tDOM pool; however, the similarity in slopes suggest that
588 variability within a core nevertheless accurately reflects relative changes in CDOM absorption.

589
590 We find that we can reliably reconstruct full CDOM absorption spectra during high CDOM
591 absorption periods (SW monsoon) at our study site. This provides information about changes
592 in optical water quality that are important to fully understand the ecological effects of tDOM
593 fluxes to coastal waters. In Singapore, the large seasonal variation in CDOM absorption leads
594 to significant shoaling of the euphotic zone and causes a shift in underwater irradiance
595 spectrum to longer wavelengths, which likely contributes to the seasonal change in
596 phytoplankton pigment composition and to limiting the depth distribution of photosynthetic
597 corals (Martin et al., 2021). Anthropogenic changes to tDOM input have been inferred to have

598 similar impacts at several temperate sites as well (Aksnes et al., 2009). While coral G/B only
599 provides information about variation in the terrigenous CDOM absorption spectrum, and thus
600 cannot provide a complete description of the underwater light environment, the proxy may help
601 to constrain relative variation in light penetration, especially in regions where information
602 about particulate optical properties is available.

603

604 **5. Conclusion**

605

606 Our study demonstrates that monthly-resolution tDOC and CDOM variability in tropical
607 coastal oceans can be reconstructed using coral cores. Given the long growth periods of coral
608 cores extending back to several decades, our method provides a way to understand long-term
609 drivers of tDOM to the coastal ocean. This is vital for better quantification of variability in
610 land-to-ocean transfer of carbon, a significant component of the global carbon cycle, as well as
611 to examine the ecological effects of coastal terrigenous CDOM variability.

612

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614

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626

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628

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