



1 Coastal Atmosphere & Sea Time Series (CoASTS) and Bio-Optical mapping of
2 Marine optical Properties (BiOMaP): the hyperspectral absorption coefficients by
3 optically significant constituents.

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ABSTRACT

The *Coastal Atmosphere & Sea Time Series* (CoASTS) and the *Bio-Optical mapping of Marine optical Properties* (BiOMaP) programs were conceived and implemented to assist ocean color applications with field measurements of apparent and inherent optical properties, and concentration of optically significant water constituents. The CoASTS program led to the creation of time-series of bio-optical measurements at the Acqua Alta Oceanographic Tower (AAOT) site the northern Adriatic Sea continued from 1995 up to 2016. The BiOMaP program supported the collection of equivalent bio-optical measurements across European Seas from 2000 up to 2022. This work focusses on CoASTS and BiOMaP hyperspectral absorption coefficients of optically significant constituents as determined applying standardized instruments, community measurement methods, extended quality control schemes and consolidated processing codes. The work, which complements a previous one by Zibordi and Berthon (2024) centred on multi-spectral data, presents the CoASTS and BiOMaP hyperspectral absorption coefficients of pigmented and non-pigmented particles in the 400-750 nm, and of colored dissolved organic matter in the 350-680 nm interval, both with a 3 nm spectral resolution at 2 nm increment.



32 1. Introduction

33 Over a period exceeding two decades, the *Coastal Atmosphere & Sea Time Series*
34 (CoASTS) and the *Bio-Optical mapping of Marine optical Properties* (BiOMaP) field programs
35 led to the collection of *in situ* bio-optical measurements to support ocean color applications. The
36 CoASTS and BiOMaP programs, established by the Marine Optical Laboratory of the Joint
37 Research Centre (JRC) in collaboration with several European research institutions, ensured: *i.*
38 time-series measurements at the Acqua Alta Oceanographic Tower (AAOT) site in the northern
39 Adriatic Sea characterized by moderately optically complex waters (Berthon et al. 2002; Zibordi
40 et al. 2002); and *ii.* geographically distributed ship-based measurements across major European
41 Seas exhibiting a variety of water types (Berthon et al. 2008, Zibordi et al. 2011).

42 This work presents the CoAST-BiOMaP data set of hyperspectral absorption coefficients
43 determined applying spectrophotometric methods to field marine water samples. Opposite to the
44 multi-spectral absorption coefficients accompanying a previous dataset of radiometric data
45 products (Zibordi and Berthon 2024), this complementary dataset comprises absorption
46 coefficients exhibiting 3 nm spectral resolutions at 2 nm increment for pigmented and non-
47 pigmented particles in the 400-750 nm, and for colored dissolved organic matter in the 350-680
48 nm interval. This unique dataset is expected to fulfil application needs for bio-optical
49 investigations supporting the new generation of hyperspectral satellite ocean color sensors (e.g.,
50 Werdell et al. 2026).

51 2. The CoASTS and BiOMaP paired measurement programs

52 CoASTS time-series started in 1995 with monthly sampling frequency, later reduced to
53 bi-monthly (or less) up to the end of the program in 2016. CoASTS measurements comprised in-
54 water optical and hydrographic profiles, water samples from near surface, 8 m and 14 m depth,
55 meteorological data, and cloud cover and sea state observations. Bio-optical characteristics
56 (Berthon et al. 2002) are representative of marine regions with optical properties generally
57 determined by moderate concentrations of sediments and colored dissolved organic matter
58 (CDOM) and or occasionally by phytoplankton and its degradation components, *i.e.*, Case-1
59 waters (IOCCG 2000).

60 BiOMaP ship-based campaigns were performed from 2000 up to 2022 in the major
61 European seas (see Berthon et al. 2008): the Baltic Sea exhibiting waters characterized by a high
62 concentration of CDOM; the Adriatic Sea, Black Sea, Iberian Shelf, Greenland Sea, Ligurian
63 Sea, and North Sea, exhibiting diverse concentrations of CDOM and suspended particulate
64 matter (*SPM*); and finally the Eastern and Western Mediterranean Seas with oligotrophic and
65 mesotrophic Case-1 waters. BiOMaP stations included the same measurements and observations
66 as CoASTS, but water sampling was limited to the near-surface layer.

67 Data consistency across the two programs was ensured through the application of
68 standardized instruments, community measurement methods, extended quality control schemes
69 and consolidated processing codes.

70 Out of the 176 CoASTS field campaigns, only the 125 performed from December 1998
71 and leading to a total of 637 stations are included in the current dataset. This choice has been
72 suggested by the objective to populate the dataset with those data fulfilling standardized methods
73 and data reduction processes. Conversely, all the 36 BiOMaP campaigns for a total of 1915
74 stations, are included in the dataset.

75 Details on the CoASTS and BiOMaP measurement stations are summarized in Tables 1 and
76 2, respectively, reproduced from Zibordi and Berthon (2024). The reader is also addressed to the
77



78 same publication for figures presenting the temporal and geographic distribution of the CoASTS
 79 and BiOMaP stations, respectively.

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 81

82 Table 1. CoASTS campaign identifiers, marine region, year, number of stations, research
 83 platform, collaborating institution (after Zibordi and Berthon 2024).

84

Campaign ID	Location	Year	Stations #	Research platform	Collaborating Institution
V03-V99	Northern Adriatic Sea (AAOT)	1998-2011	481	<i>Acqua Alta Oceanog. Tower (AAOT)</i>	Italian National Research Council (IT)
W01-W28	Northern Adriatic Sea (AAOT)	2011-2016	136	<i>Acqua Alta Oceanog. Tower (AAOT)</i>	Italian National Research Council (IT)

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 86

87 Table 2. BiOMaP campaign identifiers, marine regions, year, number of stations, research
 88 vessels, collaborating institutions (after Zibordi and Berthon 2024).

89

Campaign ID	Region	Year	Stations #	Research vessel	Collaborating Institution
A01	Adriatic Sea (ADRS)	2000	55	<i>R/V Friuli-Venezia Giulia (FVG)</i>	University of Trieste (IT)
A02	Adriatic Sea (ADRS)	2014	66	<i>R/V Minerva-1</i>	Italian National Research Council (IT)
B01	Baltic Sea (BLTS)	2004	52	<i>R/V Oceania</i>	Institute of Oceanology (PL)
B02	Baltic Sea (BLTS)	2004	52	<i>R/V Oceania</i>	Institute of Oceanology (PL)
B03	Baltic Sea (BLTS)	2005	63	<i>R/V Oceania</i>	Institute of Oceanology (PL)
B04	Baltic Sea (BLTS)	2006	23	<i>R/V Aranda</i>	Institute of Marine Research (FI)
B05	Baltic Sea (BLTS)	2007	38	<i>R/V Aranda</i>	Institute of Marine Research (FI)
B06	Baltic Sea (BLTS)	2008	47	<i>R/V Aranda</i>	Institute of Marine Research (FI)
E01	Eastern Med. Sea (EMED)	2006	62	<i>R/V Urania</i>	Italian National Research Council (IT)
E02	Eastern Med. Sea (EMED)	2007	69	<i>R/V Urania</i>	Italian National Research Council (IT)
E03	Eastern Med. Sea (EMED)	2017	51	<i>R/V Minerva-1</i>	Italian National Research Council (IT)
E04	Eastern Med. Sea (EMED)	2022	31	<i>R/V Philia</i>	Hellenic Centre for Marine Research (GR)
I01	Iberian Shelf (IBSH)	2011	68	<i>NRP Almirante Gago Coutinho</i>	Portuguese Hydrographic Institute (PT)
I02	Iberian Shelf (IBSH)	2017	62	<i>NRP Almirante Gago Coutinho</i>	Portuguese Hydrographic Institute (PT)
K01	Black Sea (BLKS)	2006	93	<i>R/V Akademik</i>	Institute of Oceanology (BG)
K02	Black Sea (BLKS)	2009	73	<i>R/V Akademik</i>	Institute of Oceanology (BG)
K03	Black Sea (BLKS)	2009	40	<i>R/V Akademik</i>	Institute of Oceanology (BG)



K04	Black Sea (BLKS)	2011	38	<i>R/V Mare Nigrum</i>	National Institute of Marine Geology and Geoecology (RO)
K05	Black Sea (BLKS)	2011	24	<i>R/V Akademik</i>	Institute of Oceanology (BG)
K06	Black Sea (BLKS)	2011	59	<i>R/V Akademik</i>	Institute of Oceanology (BG)
K07	Black Sea (BLKS)	2012	93	<i>R/V Akademik</i>	Institute of Oceanology (BG)
K08	Black Sea (BLKS)	2012	14	<i>R/V Akademik</i>	Institute of Oceanology (BG)
K09	Black Sea (BLKS)	2016	54	<i>R/V Akademik</i>	Institute of Oceanology (BG)
K10	Black Sea (BLKS)	2016	83	<i>R/V Akademik</i>	Institute of Oceanology (BG)
K11	Black Sea (BLKS)	2019	80	<i>R/V Akademik</i>	Institute of Oceanology (BG)
K12	Black Sea (BLKS)	2019	44	<i>R/V Akademik</i>	Institute of Oceanology (BG)
L01	Ligurian Sea (LIGS)	2008	41	<i>R/V Alliance</i>	Undersea Research Center (NATO)
L02	Ligurian Sea (LIGS)	2009	63	<i>R/V Alliance</i>	Undersea Research Center (NATO)
L04	Ligurian Sea (LIGS)	2013	25	<i>R/V Alliance</i>	Undersea Research Center (NATO)
N01	English Channel & North Sea (NORS)	2004	55	<i>R/V Côtes de la Manche</i>	Université du Littoral Côte d'Opale (FR)
N02	North Sea (NORS)	2015	52	<i>R/V Belgica</i>	Royal Belgian Institute of Natural Sciences (BE)
O01	Western Med. Sea (WMED)	2012	73	<i>R/V Urania</i>	Italian National Research Council (IT)
O02	Western Med. Sea (WMED)	2014	64	<i>R/V Urania</i>	Italian National Research Council (IT)
O03	Western Med. Sea (WMED)	2021	53	<i>R/V Garcia del Cid</i>	Institute of Marine Science (ES)
P01	Greenland Sea (GRLS)	2018	15	<i>R/V Alliance</i>	Undersea Research Center (NATO)
P03	Greenland Sea ¹ (GRLS)	2021	40	<i>R/V Alliance</i>	Italian Hydrographic Institute (IT)

90 ¹ It includes stations from the Norwegian Sea.



91 3. Details on the absorption measurement methods

92 CoASTS and BiOMaP laboratory measurements ensured the determination of the following
93 spectral absorption coefficients from discrete water samples collected at approximately 1 m
94 depth:

- 95 - a_{ph} and a_{dt} for pigmented and for non-pigmented particles, respectively;
- 96 - a_{ys} for CDOM.

97 Details on the applied measurement methods are given in the following subsections.

98 3.1 Absorption coefficient of particulate matter

99 The Transmission-Reflection (*T-R*) method by Tassan and Ferrari (1995, 2002), which was
100 shown appropriate for any particle type including highly absorbing sediments or highly back-
101 scattering mineral particles, was applied to determine the absorption coefficients by water
102 particles collected on glass fiber filter. The *T-R* method combines light transmittance (*T*) and light
103 reflectance (*R*) measurements carried out using an integrating sphere combined with a dual-beam
104 spectrophotometer. The method enables the measurement of both forward-scattered (*T*-mode)
105 and backward-scattered (*R*-mode) light resulting from the interaction with the particles on the
106 filter. Data reduction is supported by a theoretical model that minimizes the impact of
107 differences in the light backscattered by the particles. The method was initially applied using a
108 Perkin Elmer Lambda-19 spectrometer, replaced by a Lambda-950 from 2004 onward.

109 Whatman GF/F glass fibre filters with 0.7 μm pore size and 25 mm diameter were used to
110 collect water particles from known water volumes. Following Tassan and Ferrari (2002), the
111 total absorption coefficient $a_p(\lambda)$ of the equivalent particle suspension was determined in the
112 400-750 nm spectral interval with 1 nm resolution from
113

$$114 \quad a_p(\lambda) = 2.3 A_s(\lambda) (F_a / V_w)^{-1} \quad (1)$$

115 where V_w , F_a and $A_s(\lambda)$ are the filtered water volume, the filter clearance area and the equivalent
116 particle suspension absorbance obtained with the *T-R* method, respectively.

117 A solution of sodium hypochlorite (NaClO) was applied to the sample to discriminate
118 the non-pigmented fraction $a_{dp}(\lambda)$ from the total $a_p(\lambda)$, which allowed to determine the pigmented
119 fractions as $a_{ph}(\lambda) = a_p(\lambda) - a_{dp}(\lambda)$.

120 The bleaching approach, which rapidly acts on pigment molecules and only slowly on
121 detritus, is comprehensively discussed in Tassan and Ferrari (1995) and in Ferrari and Tassan
122 (1999). Building on that approach, since 2003 a more controlled depigmentation methodology
123 was implemented. This combines the chemical bleaching based on sodium hypochlorite solution
124 with photo-bleaching, and additionally applies a real-time control of the depigmentation process
125 through the monitoring of the sample reflectance in the red spectral region (Zibordi and Sciuto
126 2022). Such a methodology accelerates the bleaching process and prevents successive
127 measurement trials to achieve an optimal depigmentation of the particles on the filter. In fact, the
128 original approach requires successive additions of oxidizing agent and rinsing of the filter to
129 remove the oxidation products. These imply an incremental determination of the absorbance by
130 bleached particles, but also an unavoidable perturbation of particles distribution on the filter.
131 Definitively, the implementation of the improved methodology allowed for the standardization of
132 the bleaching process and warranted a better accuracy and reproducibility of the derived
133 absorption coefficients.
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136 Final $a_{dt}(\lambda)$ and $a_{ph}(\lambda)$ data products were determined with 3 nm spectral resolution at 2
137 nm increment. The spectral averaging applied to the actual laboratory measurements performed
138 with 1 nm resolution, makes the $a_{dt}(\lambda)$ and $a_{ph}(\lambda)$ final data products less dependent on
139 measurement noise, still at the expense of a reduced spectral resolution.

140 Well appreciating any previous effort addressing the accuracy of the T - R method (*e.g.*,
141 Tassan and Ferrari (1995) and Tassan et al. (2000)), uncertainties for $a_{ph}(\lambda)$ and $a_{dp}(\lambda)$ are not
142 available. Because of this, results from the analysis of $a_p(\lambda)$ measurement precision by Zibordi et
143 al. (2002) are further recalled. That specific investigation addressed the precision of $a_p(\lambda)$
144 determined with the T - R method through: *i.* $a_p(\lambda)$ from repeated measurements of the same
145 sample and *ii.* $a_p(\lambda)$ from replicated measurements of independent samples, still obtained from
146 the same water volume. Results are summarized in Table 3 at the 443, 555 and 665 nm center-
147 wavelengths through mean absolute percent differences (MAPD). Results show increasing
148 MAPD values with wavelength. MADP for the absorption coefficients from repeated
149 measurements of the same sample are mostly explained by: *i.* the sensitivity of the method, and
150 *ii.* the lack of precision in re-positioning the filter in front of the aperture of the integrating
151 sphere. The increased MADP of replicated measurements with respect to the repeated ones, are
152 explained by: *i.* differences resulting from uneven distributions of particles on filters across
153 replicates; *ii.* inhomogeneity of particles spreading in the water volumes collected to produce the
154 samples; *iii.* uncertainty in the determination of the water volume.

155
156 Table 3. Absolute mean percent differences (MAPD) \pm their standard deviations for repeated and
157 replicated measurements of a_p at 443, 555 and 665 nm. The values in parenthesis indicate the
158 average of the a_p spectral values \pm their standard deviation, in units of m^{-1} . N is the number of
159 samples.

160

λ [nm]	443	555	665
MAPD for repeated a_p [%] N=21	2.9 \pm 2.3 (0.082 \pm 0.042)	7.4 \pm 6.0 (0.023 \pm 0.011)	7.2 \pm 5.8 (0.027 \pm 0.015)
MAPD for replicated a_p [%] N=21	8.9 \pm 5.9 (0.090 \pm 0.049)	9.8 \pm 7.0 (0.024 \pm 0.012)	10.7 \pm 7.6 (0.028 \pm 0.017)

161

162 3.2 Absorption coefficient by CDOM

163 A Perkin Elmer Lambda-12 dual-beam spectrometer, and a Lambda-35 from 2010
164 onward, were used to determine $a_{ys}(\lambda)$ applying the method by Ferrari et al. (1996). Field
165 samples were obtained by filtering water on Millipore 0.22 μ m pore size cellulose filters. To
166 prevent bacteria growth, 1mL of a solution of 10 g L⁻¹ of NaN₃ was added to a 100 mL sample.

167 Spectrometric measurements were performed in the 350-750 nm spectral interval with 1
168 nm resolution by placing a 10 cm quartz cuvette containing the sample in the optical path of the
169 spectrometer beam and an identical cuvette containing milli-Q water in the optical path of the
170 reference beam. Definitely, 10 cm path-lengths may decrease the accuracy of measurements
171 performed on samples exhibiting low CDOM absorption. Still, the need for standardizing
172 measurements across the CoASTS and BiOMaP measurement programs embracing diverse
173 geographic regions was a priority.



174 With reference to Ferrari et al. (1996) the absorption coefficient $a_{ys}(\lambda)$ was computed
175 from the absorbance $A_{ys}(\lambda)$ resulting from the difference between the sample and reference ones,
176 as

$$177 \quad \quad \quad 178 \quad \quad \quad a_{ys}(\lambda) = 2.3 A_{ys}(\lambda) L_c^{-1} \quad (2)$$

179 where L_c is the pathlength of the cuvette.

180 Assuming that CDOM absorption is nil in the red, the final absorption coefficients were
181 obtained by subtracting the mean of the $a_{ys}(\lambda_i)$ spectral values for any $\lambda_i \in 670-680$ nm to the
182 measured $a_{ys}(\lambda)$. Consistent with $a_{ph}(\lambda)$ and $a_{dt}(\lambda)$, also the final $a_{ys}(\lambda)$ values are made available
183 with 3 nm spectral resolution at 2 nm increment, but restricted to the 350-680nm interval.

184 Comprehensive uncertainty analysis are also not available for $a_{ys}(\lambda)$. Because of this,
185 results from $a_{ys}(\lambda)$ precision analysis from Zibordi et al. (2002) are summarized too in Table 4 at
186 the center-wavelengths 412, 443 and 555 nm. This specific investigation looked at measurement
187 precision through: $a_{ys}(\lambda)$ from repeated measurements of the same sample and *ii.* $a_{ys}(\lambda)$ from
188 replicated measurements of independent samples, still obtained from the same water volume.
189 Similar to results obtained for $a_p(\lambda)$, MAPD determined for $a_{ys}(\lambda)$ exhibit values largely
190 increasing with wavelength consistently with the exponential decrease of $a_{ys}(\lambda)$ toward the red.
191 MADP from repeated measurements are largely due to the precision of the method. Unavoidable
192 differences affecting sample replicates are responsible for the increase of MAPD of replicated
193 measurements with respect to those of the repeated ones.

194
195
196 Table 4. Absolute MAPD \pm their standard deviations, determined for repeated and replicated
197 measurements of a_{ys} at 412, 443 and 555 nm. The values in parenthesis indicate the average of the
198 a_{ys} spectral values \pm their standard deviation, in units of m^{-1} . N is the number of samples.
199

λ [nm]	412	443	555
APD for repeated a_{ys} [%] N=21	10.1 \pm 7.3 (0.168 \pm 0.037)	12.4 \pm 8.4 (0.097 \pm 0.022)	24.2 \pm 19.8 (0.015 \pm 0.005)
APD for replicated a_{ys} [%] N=21	12.1 \pm 6.3 (0.175 \pm 0.038)	15.8 \pm 8.8 (0.103 \pm 0.026)	30.3 \pm 23.8 (0.018 \pm 0.005)

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202 4. The CoASTS and BiOMaP dataset of hyperspectral absorption coefficients

203 CoASTS-BiOMaP hyperspectral data of absorption coefficients are accessible in tabular
204 form at <https://doi.org/10.1594/PANGAEA.XXXXXX>.

205 Details on the quantities constituting the dataset are provided in Table 5. Nevertheless, the
206 reader is addressed to Zibordi and Berthon (2024) for a comprehensive description of all the field
207 and laboratory measurements complementing the hyperspectral absorption coefficients matter of
208 this work. It is anticipated that the number of measurement stations included in the present
209 dataset is slightly higher than the one constituting the multi-spectral dataset. This is explained by
210 the quality control applied to hyperspectral data, fully independent of other measured quantities
211 (see Zibordi and Berthon, 2024). Still some a_{ys} spectra were not included in the hyperspectral
212 dataset because of unrealistically low values in the ultraviolet and blue spectral region.



213 Figures 1-3 display BioMaP and CoASTS $a_{ph}(\lambda)$, $a_{dt}(\lambda)$ and $a_{ys}(\lambda)$ spectra for the different
 214 marine regions. Some of these spectra refer to bio-optical cases typical of Eastern Mediterranean
 215 Sea (EMED) oligotrophic waters and Baltic Sea (BLTS) CDOM dominated waters. Between
 216 these two extremes, there are spectra from the North Sea (NORS), characterized by relatively
 217 high concentrations of non-organic particles, and from the Black Sea (BLKS) and northern
 218 Adriatic Sea (AAOT) exhibiting optical characteristics determined by variable concentrations of
 219 SPM and CDOM.

220
 221

222 Table 5. Details on the quantities included in the CoASTS-BiOMaP hyperspectral dataset
 223 defined in agreement with the notations already used in Zibordi and Berthon (2024).

224

Symbol	Description	Units	Details
<i>Station ID</i>	Station identifier	Code	<i>Gccssii</i> ⁽¹⁾
<i>Date&Time</i>	Date and time	GMT	<i>yyyy-mm-ddT hh:mm:ss</i> ⁽²⁾
<i>Lon</i>	Longitude	Degrees	
<i>Lat</i>	Latitude	Degrees	
$a_{ph}(\lambda)$	Absorption coefficient by pigmented particles at 1 m depth	m^{-1}	3 nm spectral resolution at 2 nm increment in 400-750 nm interval
$a_{dt}(\lambda)$	Absorption coefficient by non-pigmented part. at 1 m depth	m^{-1}	3 nm spectral resolution at 2 nm increment in 400-750 nm interval
$a_{ys}(\lambda)$	Absorption coefficient by CDOM at 1 m depth	m^{-1}	3 nm spectral resolution at 2 nm increment in 350-680 nm interval

225

226 ¹ *G* indicates the site or geographic region (V and W for AAOT, A for Adriatic Sea, B for Baltic Sea, E for Eastern
 227 Mediterranean Sea, K for Black Sea, L for Ligurian Sea, N for North Sea, O for Western Mediterranean Sea, I for
 228 Iberian Shelf, P for Greenland Sea), while *cc* indicates the campaign number for the specific region, *ss* the station
 229 number and *ii* the cast number.

230 ² The letters *yyyy* indicate the year, *mm* the month, *dd* the day *hh*, the hours, *mm* the minutes and *ss* the seconds.

231

232 Figure 1 displays the a_{ph} spectra for the various marine regions. The increase in the mean
 233 values of $a_{ph}(\lambda)$ around 440 nm from approximately $0.01 m^{-1}$ for the Eastern Mediterranean Sea
 234 (EMED) waters to values approaching $0.2 m^{-1}$ for the Baltic Sea (BLTS) and North Sea (NORS)
 235 ones, is noteworthy. Some North Sea (NORS) spectra from samples collected off the Belgian
 236 coast exhibit values of a_{ph} higher at 400 nm than at 440 nm. This is explained by high
 237 concentrations of pheophytin leading to an increase of the absorption coefficient toward 400 nm
 238 (Zibordi and Berthon 2024). Finally, a few a_{ph} spectra were noted to exhibit unrealistic negative
 239 values beyond approximately 500 nm. This is explained by limits of the measurement
 240 methodology in the presence of a low concentration of particles largely associated with
 241 oligotrophic regions. These spectra were not excluded from the data set expecting their
 242 application can be still relevant if the absorption coefficients are corrected for their background
 243 in the near-infrared.

244 The $a_{dt}(\lambda)$ spectra displayed in Fig. 2 exhibit the expected exponential decay from 400 nm to
 245 750 nm, even though sometimes affected by minor irregularities such as the shoulders around
 246 480-490 nm, which indicate a potential uncomplete bleaching of accessory pigments. Mean
 247 values of a_{dt} at 400 nm consistently vary from approximately $0.01 m^{-1}$ for the Mediterranean Sea
 248 (either EMED and WMED) waters, exceed $0.06 m^{-1}$ for the Adriatic Sea (AAOT) ones, and



249 reach 0.11 m^{-1} for the Baltic Sea (BLTS). The residual values of $a_{dt}(\lambda)$ in the near-infrared reflect
250 actual absorption by non-living particles.

251 Figure 3 displays the $a_{ys}(\lambda)$ spectra in the 350-680 nm spectral interval. The mean values of
252 a_{ys} at 350 nm largely vary across marine regions. In agreement with the oligotrophic nature of
253 the Eastern Mediterranean Sea (EMED) waters and the highly absorbing Baltic Sea (BLTS)
254 ones, a minimum value of 0.13 m^{-1} is observed for the first while a maximum of 2.2 m^{-1}
255 characterizes the second.

256 Of major interest are the slopes S_{dt} and S_{ys} defining the exponential decay of $a_{dt}(\lambda)$ and $a_{ys}(\lambda)$,
257 for which the readers are addressed to Zibordi and Berthon (2024). Those authors determined the
258 parameters defining the best exponential fit versus wavelength of $a_{dt}(\lambda)$ and $a_{ys}(\lambda)$ within the
259 412-665 nm spectral interval using

260

$$261 \quad a_{dt}(\lambda) = A_{dt} \exp(-S_{dt}(\lambda - 412)) + B_{dt} \quad (3)$$

262

263 and

$$264 \quad a_{ys}(\lambda) = A_{ys} \exp(-S_{ys}(\lambda - 412)) + B_{ys}, \quad (4)$$

265

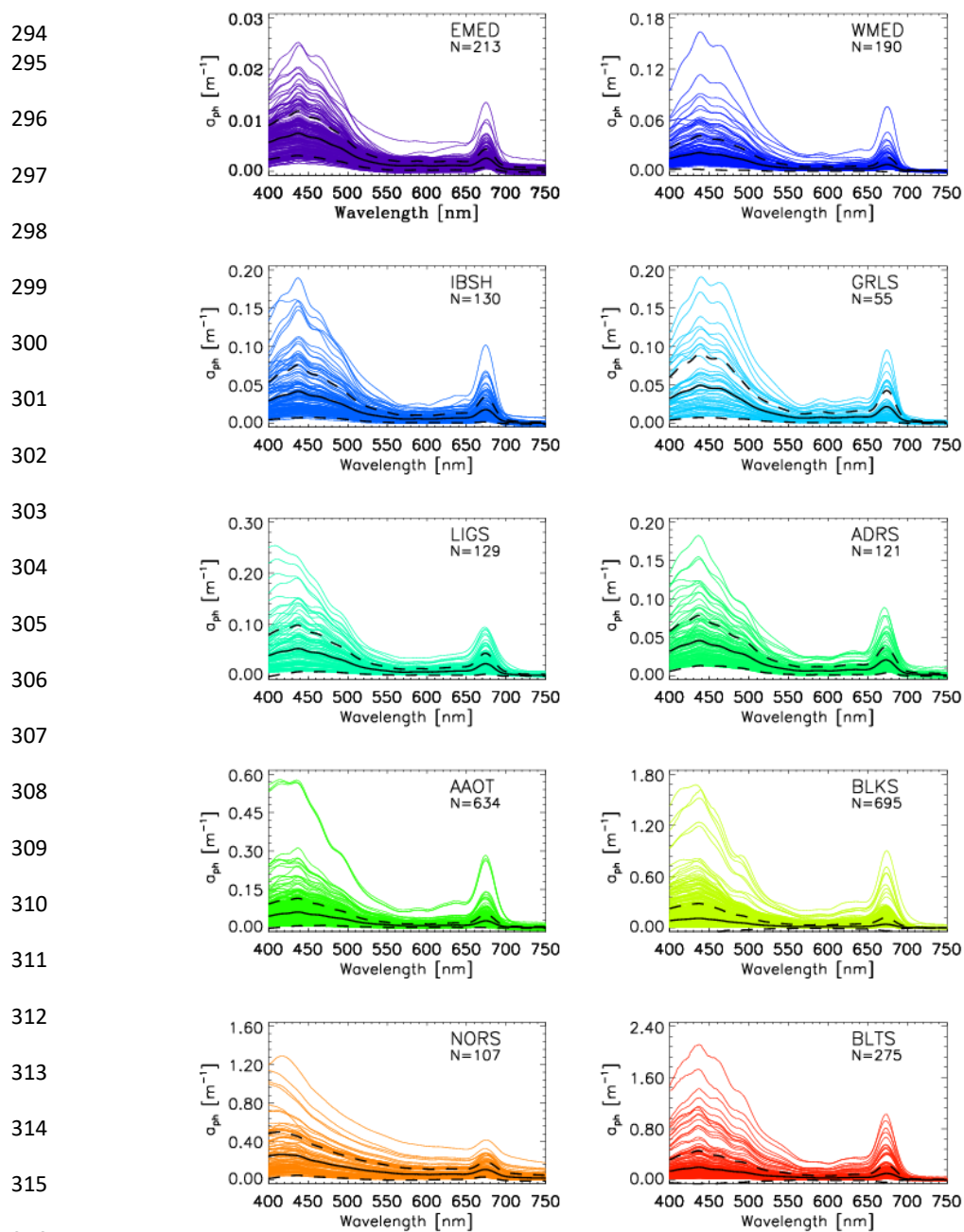
266 where A_{dt} and A_{ys} indicate the absorption coefficients fitted at 412 nm, S_{dt} and S_{ys} the slope of
267 the exponential function, and, B_{dt} and B_{ys} account for the background.

268 With reference to Zibordi and Berthon (2024), the slopes S_{dt} exhibit mean values varying
269 from 0.009 nm^{-1} for the Eastern Mediterranean Sea (EMED) up to 0.013 for the North Sea
270 (NORS). Similarly, the slopes S_{ys} show mean values ranging from 0.012 nm^{-1} for the Eastern
271 Mediterranean Sea (EMED) up to 0.019 nm^{-1} for the Baltic Sea (BLTS).

272 Ternary plots of a_{ys} , a_{dt} and a_{ph} at 442 nm as a percent of the total absorption (*i.e.*,
273 $a_{ys}(442)+a_{dt}(442)+a_{ph}(442)$) are presented in Fig. 4. Consistent with the equivalent plots
274 presented in Zibordi and Berthon (2024), very few cases (most refer to the Black Sea for values
275 greater than 60%) suggest dominance of absorption by particles with a_{ph} and a_{dt} values clustering
276 near the upper and lower right vertices, respectively. Indeed, most of the cases show
277 predominance of absorption by CDOM with a_{ys} values clustered nearby the lower left vertex.
278 While this is naturally expected for the Black Sea (BLKS) optically complex waters and for all
279 the Baltic Sea (BLTS) highly absorbing ones, peculiar is the pronounced clustering observed for
280 the Eastern Mediterranean Sea (EMED) oligotrophic waters, and also for some geographic areas
281 of the Western Mediterranean Sea (WMED) oligotrophic/mesotrophic ones. This outcome for
282 either EMED and WMED waters, conflicts with the Case-1 water classification (IOCCG 2000).
283 Still, equivalent findings were reached by Pérez et al. (2016), further confirming the presence of
284 a significant CDOM background in the Mediterranean Sea waters

285 Minor differences between Fig. 4 and the equivalent one published in Zibordi and Berthon
286 (2024) are explained by the diverse number of spectra (see § 4).

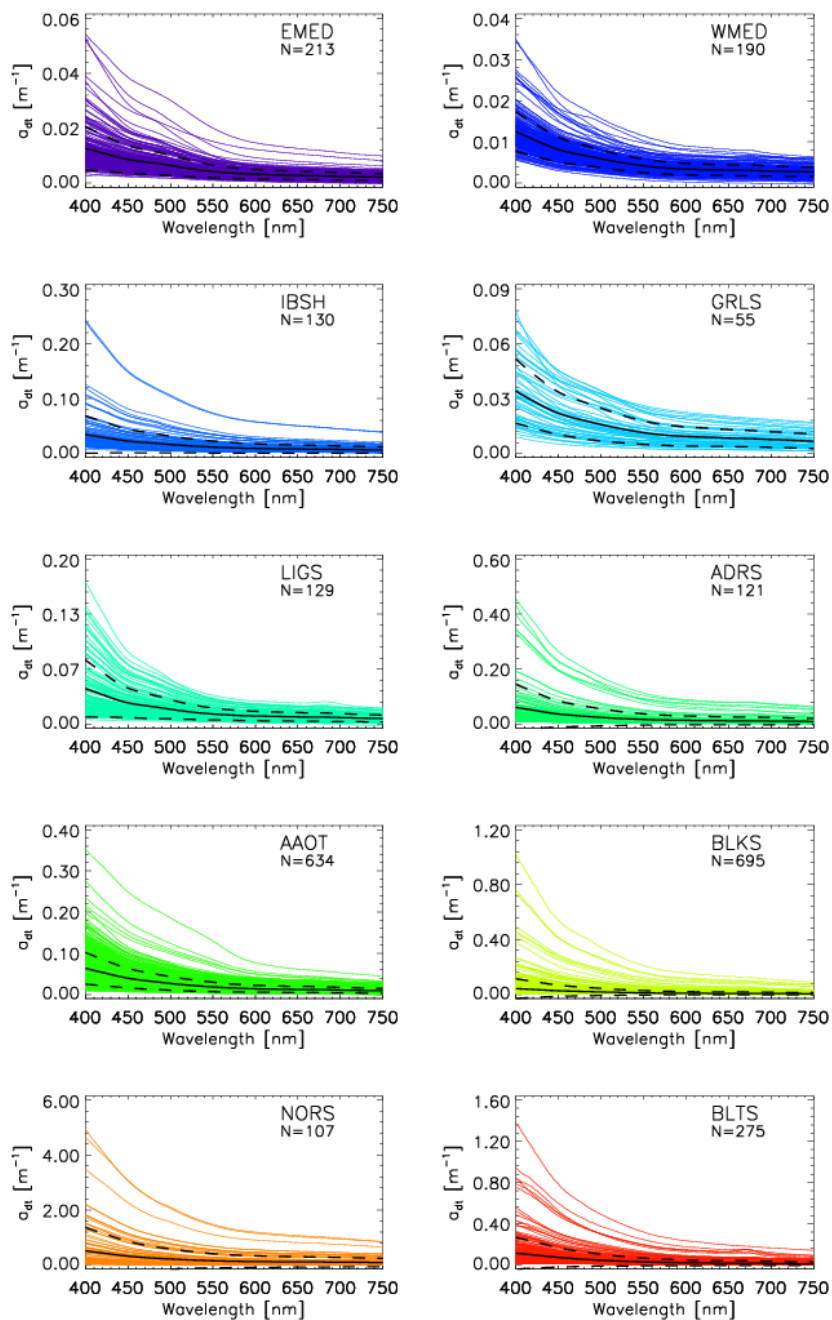
287 As an applicative example, scatter plots of $a_{ph}(442)$ versus $a_{ph}(676)$ in m^{-1} are displayed
288 in Fig. 5 to investigate the impact of regional pigments. As expected, data show decreasing
289 values of the average $a_{ph}(442)/a_{ph}(676)$ ratio from oligotrophic (*i.e.*, EMED and WMED) to more
290 eutrophic waters (*i.e.*, NORS and BLTS), with the exception of the high value characterizing the
291 Black Sea (BKLS). This suggests a higher mean cell size (and consequently an increased
292 packaging effect) in conjunction with a lower proportion of photoprotective pigments in
293 eutrophic environments, which lead to a lower ratio with respect to oligotrophic environments.



317 Figure 1. Spectra of $a_{ph}(\lambda)$ for the various marine regions. N is the number of spectra. The
318 continuous thick black lines indicate mean values while the dashed lines indicate ± 1 standard
319 deviation.



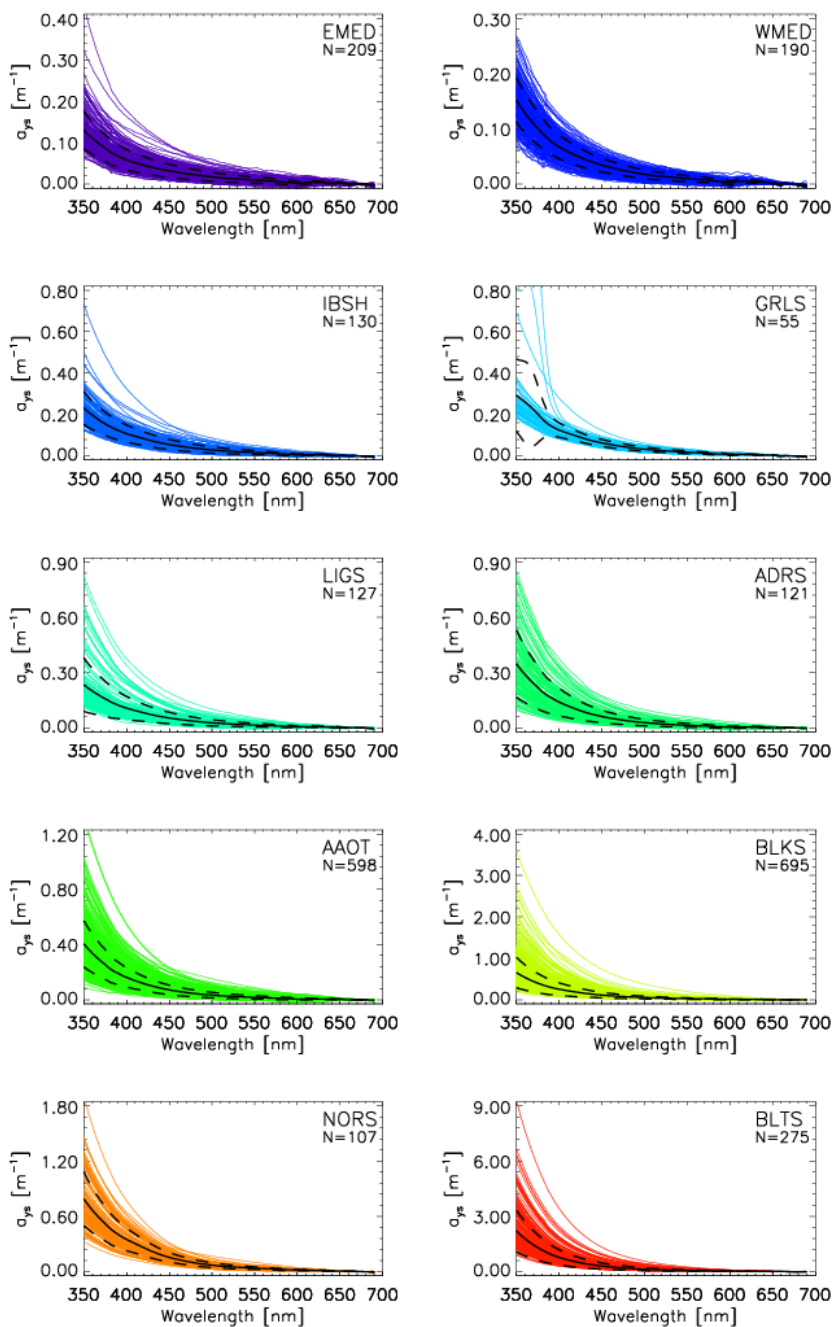
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361 Figure 2. Spectra of $a_{dt}(\lambda)$ for the various marine regions. N is the number of spectra. The
362 continuous thick black lines indicate mean values while the dashed lines indicate ± 1 standard
363 deviation.



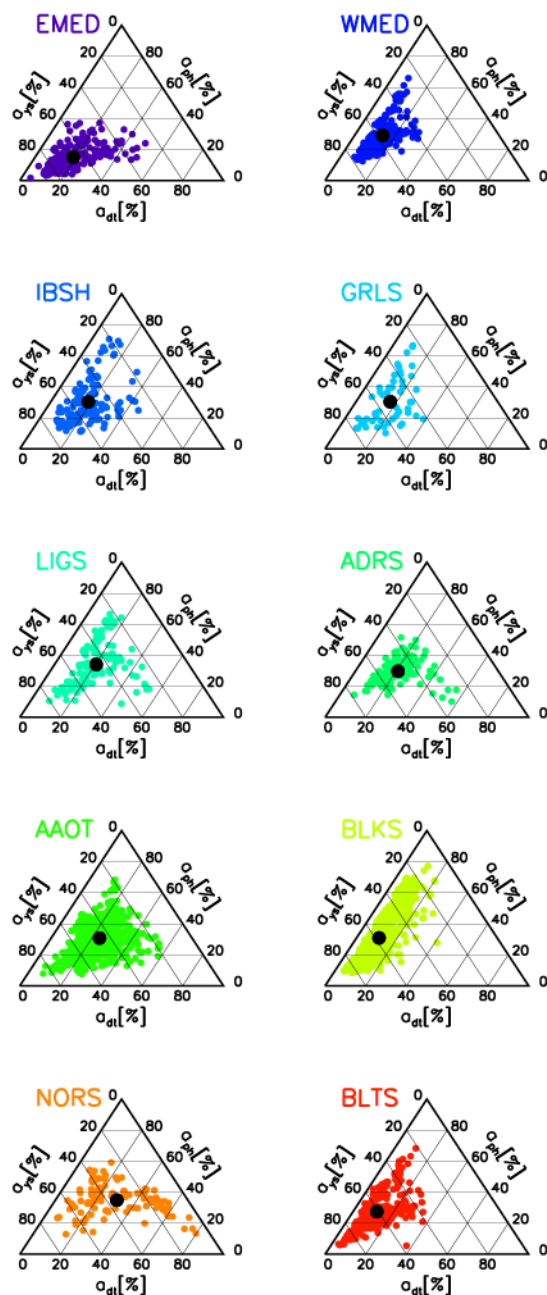
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406 Figure 3. Spectra of $a_{ys}(\lambda)$ for the various marine regions. N is the number of spectra. The
407 continuous black lines indicate mean values while the dashed lines indicate ± 1 standard
408 deviation.



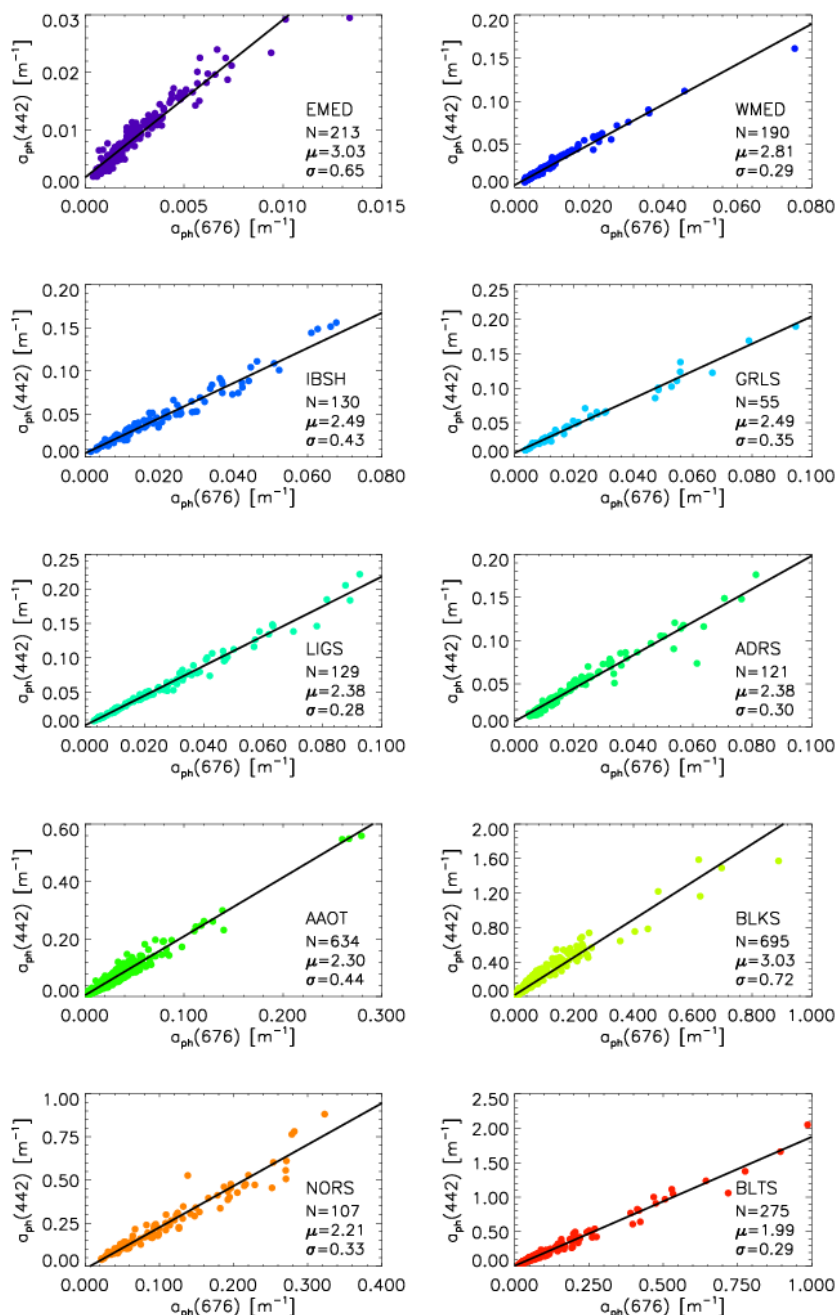
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452 Figure 4. Ternary plots of the absorption coefficients a_{ys} , a_{dt} and a_{ph} expressed in percent of the
453 total (except sea water) absorption (*i.e.*, with respect to $a_{ys}+a_{dt}+a_{ph}$) at the 442 nm center-
454 wavelength. The filled black circles indicates the mean of the plotted values.



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497 Figure 5. Scatter plots of $a_{ph}(442)$ versus $a_{ph}(676)$ in m^{-1} for the various geographic regions. N is
 498 the number of spectra, while μ and σ are the mean and standard deviation of the ratio
 499 $a_{ph}(442)/a_{ph}(676)$, respectively. Correlations coefficients (not shown) exhibit values within 0.96
 500 and 0.99. The continuous thick black lines indicate the linear regression fits.



501

502 **5. Summary and conclusions**

503 The CoASTS and BiOMaP measurement programs, established by the JRC Marine Optical
504 Laboratory in collaboration with various European institutions, from 1995 up to 2022 produced a
505 time-series and geographically distributed bio-optical data at the AAOT site in the northern
506 Adriatic Sea and in European Seas, respectively. The measurements from the two programs
507 included the same quantities rigorously obtained by applying standardized instruments,
508 community measurement methods, extended quality control schemes and consolidated
509 processing codes.

510 Complementary to a previous work that introduced the CoASTS-BiOMaP dataset
511 comprising the near surface multispectral bio-optical data (see Zibordi and Berthon 2024), this
512 work describes the CoASTS and BiOMaP dataset of hyperspectral absorption coefficients by
513 pigmented and non-pigmented particles as well those by colored dissolved organic matter, both
514 obtained applying spectrophotometric measurements to field samples. This unique dataset has
515 relevance for bio-optical investigations supporting the exploitation of advanced hyperspectral
516 satellite ocean color sensor data.

517

518 **6. Author contributions**

519 Jean-François Berthon and Giuseppe Zibordi equally contributed to the generation of the data set
520 and to the writing of the manuscript.

521

522 **7. Competing interests**

523 Both authors declare no competing interest.

524

525 **8. Data availability**

526 Interested researchers can download the CoASTS-BiOMaP data set at
527 <https://doi.pangaea.de/10.1594/PANGAEA.993510> (Berthon and Zibordi 2026).

528

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533

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539 Administration (NASA), the European Space Agency (ESA), the Romanian Space Agency
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541 Oceanology of the Polish Academy of Sciences, the Finnish Environment Institute, the Italian
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599 **Appendix A: Acronyms**

600	AAOT	Acqua Alta Oceanographic Tower
601	ADRS	Adriatic Sea
602	BiOMaP	Bio-Optical mapping of Marine Properties
603	BLKS	Black Sea
604	BLTS	Baltic Sea
605	CDOM	Colored Dissolved Organic Matter
606	CoASTS	Coastal-Atmosphere and Sea Time-Series
607	EMED	Eastern Mediterranean Sea
608	GRLS	Greenland Sea
609	IBS	Iberian Shelf
610	LIGS	Ligurian Sea
611	MADP	Mean Absolute Percent Differences
612	JRC	Joint Research Center
613	NORS	North Sea
614	SPM	Suspended Particulate Matter
615	T-R	Transmission and Reflection spectrophotometric method
616	WMED	Western Mediterranean Sea