



1 Hyperspectral reflectance dataset of pristine, weathered and biofouled

2 plastics

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14 **Abstract**

This work presents a hyperspectral reflectance dataset of macroplastic samples acquired using 15 Analytical Spectral Devices (ASD) FieldSpec 4. Samples analysed consisted of pristine, artificially 16 weathered and biofouled plastic items and plastic debris samples collected in the docks of the Port of 17 Antwerp and in the river Scheldt near Temse Bridge (Belgium). The hyperspectral signal of each sample 18 was measured in controlled dry conditions in an optical calibration facility at the Flemish Institute for 19 Technological Research, and, for a subset of plastics, under wet and submerged conditions in a silo tank 20 21 at Flanders Hydraulics. The wet and submerged hyperspectral signals were measured in a mesocosm 22 setting that mimicked environmentally relevant concentrations of freshwater microalgae and of 23 suspended sediment. The ASD was equipped with an 8° field of view at the calibration facility, and a 1° 24 field of view was used in the mesocosm setting. The dataset obtained complies with the Findability, Accessibility, Interoperability, and Reuse (FAIR) principles and is available in the open-access repository 25 Marine Data Archive (https://doi.org/10.14284/530, Leone et al., 2021). 26





27 **1 Introduction**

The spectral reflectance measurements, collected in the framework of the Plastic Flux for Innovation and Business Opportunities in Flanders (PLUXIN) project, contribute to the current knowledge on the detection of plastics using remote sensing techniques.

In recent years, focus has been placed on the detection of plastic litter using remote sensing techniques such as optical sensors on satellites, aircrafts, and drones (Garaba and Dierssen, 2018; Martínez-Vicente et al., 2019). With the increasing demand of these technologies, it is crucial to generate knowledge on the diagnostic spectral properties of not only pristine, but also weathered and biofouled plastics (Moshtaghi et al., 2021) that are representative of the variety of environmental plastics.

Currently the spectral reflectance of dry plastics is known, and is already applied in the field of material recycling (Moroni et al., 2015), but it is restricted to items assessed for dry measurements. To be able to identify plastic litter in aquatic environments such as rivers, harbours, and oceans, we require the acquisition of the spectral features when plastics are either wet or submerged (Moshtaghi et al., 2021), as water absorbs the light in both the near (NIR) and shortwave infrared (SWIR). In addition, other water constituents such as sediment or algae, could further impact the reflected signal of plastic items (Moshtaghi et al., 2021).

To date, only a limited number of high-quality datasets consisting of hyperspectral measurements of wet and submerged plastic litter, have been published in open-access repositories (e.g., Garaba and Dierssen, 2019, Garaba and Dierssen, 2020; Knaeps et al., 2021). The dataset described in the current paper aims at complementing the existing datasets by adding new information about the hyperspectral reflectance of pristine plastic items, harvested plastic litter, and artificially weathered and biofouled plastic samples. In addition, the dataset reports the optical features of plastics acquired in various water turbidity conditions obtained by adding sediment or algae to the water, at selected concentrations.

50 The dry spectral reflectance of plastic specimens, consisting of different polymers, was measured using 51 an Analytical Spectral Device (ASD) FieldSpec 4. For a selection of these samples, the spectral 52 reflectance was also collected in wet and submerged settings.

53 The presented data can be used to generate insights on the spectral properties of plastic litter and how 54 these features change when plastics are exposed to natural agents or different depths. The dataset was

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compiled following the FAIR (Findable, Accessible, Interoperable, Reusable; go-fair.org) principles and
 it is publicly available at https://doi.org/10.14284/530.

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58 2. Data collection

Data collection consisted in measuring the spectral reflectance of different plastic specimens using the ASD. The measurements were conducted in two different settings: at an optical calibration facility located at the Flemish Institute for Technological Research (Mol, Belgium), and at a silo tank at the Flanders Hydraulics Research facility (Antwerp, Belgium). The spectral reflectance of each plastic specimen was collected at least 5 times ("pseudo-replicates") within 1 minute. These pseudo-replicates are measurements taken by slightly changing the position of the plastic sample. Therefore, very homogeneous plastics will have less variation in the pseudo-replicates' reflectance.

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67 **2.1 Plastic specimens**

We aimed to collect the spectral reflectance of plastic items as representative as possible of commonly 68 69 found plastics in the environment (LI et al., 2016), and to provide additional information on their 70 reflectance when exposed to natural agents such as sunlight (UV radiation) or biofilm growth. To do so, 71 we have selected and analysed a total of 10 polymer types (Table 1). The selected polymers were Polyethylene (PE), Polypropylene (PP), Polystyrene (PS), and two types of Polyethylene terephthalate 72 (PET), crystalline and amorphous, Fluorocarbon, Thermoplastic elastomer, Polyvinyl chloride (PVC), 73 Nylon 6 (PA6) and Paraffin. In addition, we have further artificially weathered and biofouled a selection 74 of 6 plastic polymer types. The polymer discrimination was based on the available information from the 75 supplier or marked on the plastic itself (e.g., plastic bottle or bags with an identifiable polymer tag). In 76 77 addition, for a set of samples, we confirmed the polymer type using a micro-Fourier Transform Infrared Spectroscopy (µFTIR, PerkinElmer, FTIR spectrometer Frontier). The percentage of matching score with 78 79 a library was recorded and it is reported in the dataset.

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Table 1. Overview of the plastic specimens analysed in the current work. Polymer type refers to the available information from the supplier or product, unless stated otherwise. Selected polymer types were confirmed by uFTIR. Pictures in this table are taken by Giulia Leone and Mattias Bossaer.





Туре	Plastic specimen	Condition of the assessment	Picture	Polymer type	Available measurements (d: dry, w: wet, s: submerged)	Dimensions (cm)
Commercial plastics	Sheet	Dry		PE (confirmed by μFTIR)	d, w, s	6 x 6
	Sheet	Dry		PP (confirmed by µFTIR)	d, w, s	6 x 6
	Sheet	Dry		PET amorphous (confirmed by µFTIR)	d, w, s	6 x 6
	Sheet	Dry		PET crystalline (confirmed by µFTIR)	d, w, s	6 x 6
	Sheet	Dry		PS (confirmed by µFTIR)	d, w, s	6 x 6





Weathered plastics from shops	Sheet	Dry	PVC (Thermoplastic elastomer under µFTIR)	d	2 x 4
	Sheet	Dry	PP (confirmed by µFTIR)	d	2 x 4
	Sheet	Dry	PS (confirmed by µFTIR)	d	2 x 4
Biofilm plastics from shops	Sheet	Dry	PVC (Thermoplastic elastomer under µFTIR)	d	2 x 10
	Sheet	Dry	PP (confirmed by µFTIR)	d	2 x 10
	Sheet	Dry	PS (confirmed by µFTIR)	d	2 x 10





Plastic from shops	Sheet	Dry	ND	PVC (Thermoplastic elastomer under µFTIR)	d	2 x 10
	Sheet	Dry	ND	PP (confirmed by µFTIR)	d	2 x 10
	Sheet	Dry	ND	PS (confirmed by µFTIR)	d	2 x 10
	Spar bag	Dry		LDPE (paraffin under µFTIR)	d	ND
	Jumbo bag	Dry	Halo	ND	d	ND
	Bottle	Dry		ND	d	ND





Biofilm commercial plastics	Sheet	Wet rough	PP (confirmed by µFTIR)	d, w, s	6 x 6
	Sheet	Wet smooth	PP (confirmed by µFTIR)	d, w, s	6 x 6
	Sheet	Wet rough	PE (confirmed by µFTIR)	d, w, s	6 x 6
	Sheet	Wet smooth	PE (confirmed by µFTIR)	d, w, s	6 x 6
	Sheet	Wet rough	PS (confirmed by µFTIR)	d, w, s	6 x 6
	Sheet	Wet smooth	PS (confirmed by µFTIR)	d, w, s	6 x 6





Sheet	Wet rough	PET Amorphous (confirmed by µFTIR)	d, w, s	6 x 6
Sheet	Wet smooth	PET amorphous (confirmed by µFTIR)	d, w, s	6 x 6
Sheet	Wet rough	PET Crystalline (confirmed by µFTIR)	d, w, s	6 x 6
Sheet	Wet smooth	PET crystalline (confirmed by µFTIR)	d, w, s	6 x 6
Sheet	Dry rough	PS (confirmed by µFTIR)	d	<mark>6 x 6</mark>
Sheet	Dry smooth	PS (confirmed by µFTIR)	d	6 x 6
Sheet	Dry rough	PE (confirmed by μFTIR)	d	6 x 6





Sheet	Dry smooth	PE (confirmed by µFTIR)	d	6 x 6
Sheet	Dry rough	PET amorphous (confirmed by µFTIR)	d	6 x 6
Sheet	Dry smooth	PET amorphous (confirmed by µFTIR)	d	6 x 6
Sheet	Dry rough	PP (confirmed by µFTIR)	d	6 x 6
Sheet	Dry smooth	PP (confirmed by µFTIR)	d	6 x 6
Sheet	Dry rough	PET crystalline (confirmed by µFTIR)	d	6x6
Sheet	Dry smooth	PET crystalline (confirmed by µFTIR)	d	6x6





Artificially weathered	Sheet	Dry	PS (confirmed by µFTIR)	d, w, s	2 x 4
	Sheet	Seawater	PS (confirmed by µFTIR)	d, w, s	2 x 4
	Sheet	Dry	PE (confirmed by µFTIR)	d, w, s	2 x 4
	Sheet	Seawater	PE (confirmed by µFTIR)	d, w, s	2 x 4
	Sheet	Dry	PP (confirmed by µFTIR)	d, w, s	2 x 4
	Sheet	Seawater	PP (confirmed by µFTIR)	d, w, s	2 x 4
	Sheet	Dry	PET amorphous (confirmed by µFTIR)	d, w, s	2 x 4





	Sheet	Seawater	PET amorphous (confirmed by µFTIR)	d, w, s	2 x 4
	Sheet	Dry	PET crystalline (confirmed by µFTIR)	d, w, s	2 x 4
	Sheet	Seawater	PET crystalline (confirmed by µFTIR)	d, w, s	2 x 4
Real samples	Green Buoy	Real	PP (confirmed by µFTIR)	d, w, s	ND
	Milk bottle	Real	HDPE (confirmed by µFTIR)	d, w, s	ND
	Foam	Real	PS	d	ND





Toy placemat	Real	5	PP – PE (confirmed by µFTIR)	d, w, s	ND
Green Bottle	Real		PET (confirmed by µFTIR)	d, w, s	ND
Yellow sheet	Real		HDPE (confirmed by µFTIR)	d	ND
Bag dog food	Real	The second secon	PE (confirmed by µFTIR)	d	ND
Rope	Real	Contraction of the second	PP (confirmed by µFTIR)	d	ND
Shoe	Real		Fluorocarbon (confirmed by µFTIR)	d	ND





	Bubble wrap	Real	PE (confirmed by µFTIR)	d	ND
	Foam	Real	PE (confirmed by µFTIR)	d	ND
	Red sheet	Real	PP (confirmed by µFTIR)	d	ND
	Cup	Real	PP (confirmed by µFTIR)	d	ND
The Ocean Cleanup samples	HDPE	Pristine	PE	d	30x30
	Extruded PS	Pristine	Extruded PS	d	30x30







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86 2.2.1 Pristine plastic specimens

To enhance the physio-chemical properties of an end plastic product, plastic manufacturers often incorporate additive substances into the pristine polymer (Garaba et al., 2021). Additives included in plastics, such as coloured plastics, may be influencing the spectra (Garaba et al., 2021). Thus, to avoid confounding measurements, sheets of 6 x 6 cm were obtained from a commercial supplier (Carat,





Germany, https://www.carat-lab.com) as pure pristine plastic polymers without any additional additive
for the following polymers: PP, PE, PS, PET amorphous, PET crystalline. In addition, no colour or stains
were added to the commercial plastics, and as such, these selected samples were transparent or white.
The other pristine plastic samples were either purchased from a local shop (Oostende, Belgium), or
obtained from other institutes (Marine Remote Sensing Group, University of the Aegean in Plastic Litter
Projects (2021 and 2022), The Ocean Cleanup).

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98 2.1.2 Weathered plastics

To mimic the effect of solar radiation on plastic items in the environment, pristine plastics without additives were exposed to UV radiation in an Atlas SunTest CPS+ weathering chamber, simulating one solar year in central Europe (Gewert et al., 2018) (Table 2). Prior to the treatment, plastic specimens were cut, using a hot knife, into 2 x 10 cm sheets in order to fit them into closed Quartz cuvettes. Different treatments for weathering conditions were tested: i.e., dry UV exposure, seawater UV exposure (35 psu) and dark controls (dry and wet), and each treatment consisted of a set of three independent replicates for each polymer type (Table 4).

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107 **Table 2.** Parameters of the UV chamber, used in artificial weathering experiments

 UV	Black Stand Temperature	Chamber Temperature	Test
irradiation	(BST)	(CHT)	durations
W / m ²	°C	°C	h
60	50	30 - 35	

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109 **2.1.3 Biofouled plastics**

110 To test the effects of biofilm attached to the surface of plastic items on their spectral reflectance, we induced biofilm growth on the two surfaces of pristine samples. Since surface roughness can affect 111 biofilm growth and its survival (Rodriguez, D., B. Einarsson, 2012), one of the two surfaces of the 112 commercial plastic specimens, was manually treated for approximately 10 seconds with sandpaper 113 (grain size 80) to create a rougher surface compared to the smooth and untouched one. An aquarium 114 was filled with unfiltered seawater collected from the port of Ostend (Belgium), which was renewed every 115 116 two weeks, kept at 20 °C, and aerated with an air pump. Plastic specimens were suspended in the aquarium with paper clippers and rope (Fig. 1) to allow the biofilm to form on all of the surfaces. 117







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- 119 **Figure 1.** Aquarium set-up for biofilm growth on pristine plastic specimens. Picture taken by Giulia Leone
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121 **2.1.4 Field plastic items**

- 122 Plastic specimens were collected from the Port of Antwerp and in the Scheldt River, near Temse Bridge
- 123 (Belgium) (Fig.2). The plastic coming from the river Scheldt were collected by a plastic collector installed
- by the Belgian company Dredging, Environmental and Marine Engineering NV (DEME) Environmental
- 125 Contractors (DEC) on behalf of Vlaamse Waterweg (the Flemish authority responsible for waterways in
- 126 Flanders, Belgium).







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Figure 2. Example of field litter items collected in 2020 from the river Scheldt, near Temse Bridge (Belgium) by the plastic collector installed by the Belgian company DEME Environmental Contractors (DEC) on behalf of Vlaamse Waterweg (the Flemish authority responsible for waterways in Flanders, Belgium). Picture taken by Giulia Leone

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132 **2.2 Spectral measurements of plastic items**

The spectral reflectance measurements of plastic specimens were performed using an Analytical Spectral Device (ASD) Field-Spec4 (Table 3). The reflectance was automatically derived and normalized to a 99% Labsphere Spectralon® Lambertian panel. In the silo tank, the Spectralon panel was positioned on the holder of the adjustable arm, at the same distance from the ASD field of view as the dry plastic specimen.

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- **Table 3.** Hyperspectral-radiometer specifications used in the two locations of the study, calibration facility and water silo tank
 ASD: Analytical Spectral Device; VNIR: visible-near infrared; SWIR: shortwave infrared
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	ASD specifications in the calibration facility	ASD specifications in the tank
Spectral range (nm)	350 - 2500	350 - 2500
Spectral resolution (nm)	VNIR: ca. 3 nm SWIR: 10 - 12 nm	VNIR: ca. 3 nm SWIR: 10 - 12 nm
Scans per measurements	30	10
Replicate measurements	5	5
Foreoptic field of view	8°	1°

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143 **3 Experimental set-ups**

Following the procedure described by Knaeps et al., 2021, we measured the spectral reflectance of the different plastic samples at two different facilities: 1) Measurements done in dry conditions were performed at the optical calibration laboratory of the Flemish Institute for Technological Research (VITO), Belgium; 2) the series of wet and submerged measurements were acquired in a mesocosm setting, with experiments being performed in a conical shaped silo tank at the Flanders Hydraulics Research facility in Antwerp, Belgium.

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151 **3.1 Laboratory set-up**

All the plastic specimens were measured in the optical calibration facility (VITO, Belgium). This laboratory
 consisted of a dark room where a desk equipped with two halogen tungsten lamps and a holder for the
 ASD field of view, allowed us to collect the measurements.

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156 **3.1 Silo tank set-up**

The water silo tank (2 top diameter x 3 depth m), at the Flanders Hydraulics Research (Belgium), was 157 equipped with a controlled mixer with a double pitch blade impeller that permitted the mixing of the water 158 159 to obtain suspensions of sediment or algae. No information about the rpm was available on the metal impeller at the water tank. We attached a tailor-made aluminium frame to the water tank for the mounting 160 of spectroradiometer detector and light, together with a plexiglass sample holder to lower the plastic 161 162 specimens in the tank and measure their spectral reflectance at different water depths. The plexiglass 163 was cut to fit the plastic samples which were held by black plastic paper clips. In the current study, we 164 acquired data using a single lamp attached to the frame, with an angle of 40 degrees. To reduce 165 undesired stray light, a dark environment was created with black plastic cover around the experimental 166 set-up and in addition black plastic bags were held against the tank by means of wooden rings with 167 weights and waterproof tape. In the tank, samples attached to the plexiglass sample holder were first measured at dry conditions, just above the water level and then carefully lowered in the water at fixed 168 169 depths. The water level above the sample was measured with a ruler and the chosen depths were: 1 cm, 2 cm, 4 cm and 8 cm. After the submersion, the plastic was measured again above the water level 170 as wetted sample. The same steps were performed in clear and turbid water. 171

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3.1.1 Water with added sediment

174 Natural sediment was collected using a manual Van Veen grab by the Flanders Marine Institute (VLIZ) 175 during a sampling campaign as part of the PLUXIN project, in Nieuwpoort (Belgium). The sediment was transported to the laboratory (VLIZ), placed into a metal container, and dried in the oven for 4 days at 60 176 °C, and was afterwards manually crushed using a mortar, and stored in the dark until further use. To add 177 the crushed sediment into the tank, before any measurements, the tank metal impeller was activated for 178 179 a one and a half minute to allow the sediment to stay in suspension in the water. This action was repeated approximately every 30 minutes to maintain the sediment in suspension. The actual concentration of 180 sediment added in the tank was measured by Flanders Hydraulics. The suspended sediment 181 182 concentration (SSC) was determined gravimetrically, after filtration of the sample through a filter with a pore size of 0.45 µm and drying at a temperature of 105 °C. The results showed a low concentration of 183 sediment of 4 mg / L and a higher concentration of 16 mg / L. 184





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186 **3.1.2 Water with added microalgae**

187 To evaluated how the readings of spectral reflectance of plastic specimens can be affected by turbidity in the water due to microalgae suspension, measurements were performed using two concentrations of 188 189 freshwater Pseudokirchneriella subcapitata. The concentrations were selected to mimic the natural conditions of spring and late summer seasonal concentration of green algae for a Western European 190 River (Ibelings et al., 1998). The nominal concentrations of microalgae were 1500 cells / mL and 3000 191 192 cell / mL. The microalgae P. subcapitata was cultured at the Research unit "Health" at VITO, and the stock solution of 8,18 * 10⁶ algae / mL was kept for 11 days in a cold room and in dark before being used 193 in the experimental set-up. After poring 450 mL of stock solution, before any measurements, the metal 194 195 impeller was activated for a one and a half minute to allow the algae to suspend into the water. This 196 action was repeated approximately every 30 minutes to maintain the algae in suspension. To obtain a 197 higher concentration of algae that would mimic a nominal concentration of 3000 cells / mL, other 450 mL of stock solution were then added into the tank. 198

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200 **4. Data description**

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202 In this study we created a dataset with the spectral reflectance of ten plastic polymers undergone to two treatments (i.e., artificial weathering and artificial biofouling) in addition to field collected samples and 203 204 pristine plastics (Table 4). All data included in the presented dataset were curated before submission and consist of the raw reflectance data of the samples. A metadata section, included in the dataset, 205 explains the data, adding additional and essential information. For instance, information on the polymer 206 type, µFTIR results with the corresponding library matching score, origin of the sample, date and time of 207 208 collection. The sample code was created following the naming convention of Knaeps et al., 2021. To 209 ensures that end users can correctly interpret each section of the metadata, a README file is available 210 together with the dataset.

We suggest that users of this dataset perform splice correction on the data using Python (Fig. 3) or any suitable software. This processing step will lead to the obtention of spectra that do not present radiometric steps at the joints of the detectors. In addition, pseudo-replicates of each plastic sample were





- taken, and therefore we advise that users calculate the mean of these measurements to obtain a single
- spectrum for each observed item (Fig. 4).
- 216
- 217 **Table 4.** Overview of the polymer and treatment performed during the study (SSC: suspended sediment concentration)

Treatment	Polymer tested	Origin/Supplier	State of samples	Condition of the water in the tank
 Pristine Artificially dry weathered Artificially seawater weathered Artificially biofouled on pristine surface Artificially biofouled on rough surface Field 	 PS PE PP PET amorphous PET crystalline PVC Extruded polystyrene Thermoplastic elastomer Fluorocarbon Paraffin 	 Carat The Ocean Cleanup Shop Port of Antwerp Scheldt VITO Marine Remote Sensing Group 	 Dry Wet Submerged 	 Clear water, no turbidity Algae: 1.Nominal conc. 3000 cells/ml 2.Nominal conc. 1500 cells/ml Sediment: 1.SSC 4mg/L 2.SSC 16mg/L





Figure 3. Spectrum of polypropylene (PP) collected in the field without (a) and with splice correction (b). The circle indicates the jump around 1000nm due to the connection between the different ASD cameras.

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Figure 4. Spectra of field sample. Spectrum (a) shows the pseudo-replicates of the low homogeneous field samples. Spectrum (b) is the mean of all the pseudo-replicates.

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5. Data availability

The data are available in the open access repository Marine Data Archive at https://doi.org/10.14284/530 (Leone et al., 2021)

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232 6. Conclusions

The use of remote sensing technologies can be used in the detection, observation and monitoring of 233 marine plastic pollution. However, due to a lack of knowledge of the optical features of environmental 234 plastics, small steps can be made in designing algorithms to appropriately detecting plastic pollution. 235 236 The presented hyperspectral dataset is a step forward in the knowledge of the optical features of plastic litter when exposed to natural agents such as UV radiations or the growth of biofilm. We anticipate that 237 this dataset will contribute to the definition of optical spectral bands and assist in the development of 238 algorithms for the observation, monitoring and discrimination of plastics in a (semi-) operational 239 240 environment.

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243 Funding

cSBO Plastic Flux for Innovation and Business Opportunities in Flanders (PLUXIN, HBC.2019.2904) project, financed by Flanders Innovation & Entrepreneurship (VLAIO) and supported by The Blue Cluster, a Flemish members organisation for sustainable entrepreneurship in blue growth (Belgium).



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Since November 2021, Giulia Leone is supported by the Research Foundation of Flanders (FWO), as a
 PhD grant strategic basic research, application number 1S13522N (Belgium).

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250 Credit author statement

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258

259 **Competing interests**

260 The authors declare that they have no conflict of interest.

261

262 Acknowledgement

We would like to thank Flanders Hydraulics for the use of the silo tank and for the sediment concentration measurements, Hilda Witters and Guy Geukens, of the research Unit Health at VITO (Belgium) for culturing the freshwater algae. Special thanks to the Marine Remote Sensing Group, University of the Aegean (Greece) and The Ocean Cleanup (The Netherlands) for sharing plastic specimens with us.

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