Biomass composition of the golden tide pelagic seaweeds *Sargassum fluitans* and *S. natans* (morphotypes I and VIII) to inform valorisation pathways.

Doleasha Davis[a,b]*, Rachael Simister[b]*, Sanjay Campbell[a,b], Melissa Marston[a], Suranjana Bose[c] Simon J. McQueen-Mason[b], Leonardo D. Gomez[b], Winklet A. Gallimore[a], and Thierry Tonon[b#].

[a] Department of Chemistry, University of the West Indies, Mona Campus, Mona, Kingston 7, Jamaica.

[b] Department of Biology, Centre for Novel Agricultural Products, University of York, Heslington, York YO10 5DD, United Kingdom.

[c] Green Chemistry Centre of Excellence, Department of Chemistry, University of York, Heslington, York YO10 5DD, United Kingdom.

* Equal contribution.

Corresponding author: thierry.tonon@york.ac.uk

Abstract

Massive strandings of the pelagic brown algae *Sargassum* have occurred in the Caribbean, and to a lesser extent, in western Africa, almost every year since 2011. These events have major environmental, health, and economic impacts in the affected countries. Once on the shore, *Sargassum* is mechanically harvested and disposed of in landfills. Existing commercial applications of other brown algae indicate that the pelagic *Sargassum* could constitute a valuable feedstock for potential valorisation. However, limited data on the composition of this *Sargassum* biomass was available to inform on possible application through pyrolysis or enzymatic fractionation of this feedstock. To fill this gap, we conducted a detailed comparative biochemical and elemental analysis of three pelagic *Sargassum* morphotypes identified so far as forming Atlantic blooms: *Sargassum natans* I (SnI), *S. fluitans* III (Sf), and *S. natans* VIII (SnVIII). Our results showed that SnVIII accumulated a lower quantity of metals and metalloids compared to SnI and Sf, but it contained higher amounts of phenolics and non-cellulosic polysaccharides. SnVIII also had more of the carbon storage compound mannitol. No differences in the

content and composition of the cell wall polysaccharide alginate were identified among the three morphotypes. In addition, enzymatic saccharification of SnI produced more sugars compared to SnVIII and Sf. Due to high content of arsenic, the use of pelagic *Sargassum* is not recommended for nutritional purposes. In addition, low yields of alginate extracted from this biomass, compared with brown algae used for industrial production, limit its use as viable source of commercial alginates. Further work is needed to establish routes for future valorisation of pelagic *Sargassum* biomass.

Keywords

Sargassum; the Caribbean; western African; composition analysis; biomass valorisation; seaweed strandings.

1. Introduction

Sargassum fluitans and *S. natans* are species of surface dwelling (pelagic) brown seaweeds that have inundated the shores of the Caribbean, and, to a lesser extent, the western African shoreline, since 2011 (Smetacek and Zingone, 2013; Langin, 2018; Milledge et al., 2020). Large populations of *Sargassum* have proliferated almost every year since then, forming what Wang et al. (2019) described as the "Great Atlantic *Sargassum* Belt (GASB)". This GASB was estimated to be 8,850 km long and to contain over 20 million metric tons of biomass in June 2018. Such massive inundation events are known as golden tides due to the golden brown colour of *Sargassum*. These seaweeds threaten coastal environments because they begin to rot shortly after reaching shallow waters and beaches, removing oxygen from the surrounding water, killing fish and other marine organisms. *Sargassum* also raises human health concerns due to large amounts of toxic gases, including hydrogen sulphide and ammonia, are produced when the seaweeds start decomposing on the seashore (Resiere et al., 2018). Exposure to high concentrations of hydrogen sulphide can lead to pulmonary, neurological, and cardiovascular lesions. In addition, *Sargassum* has negative impacts on the fishing and tourism industries in the Caribbean, as well as in western Africa (Adet et al., 2017; Ofori et al. 2020).

Once on the beach, *Sargassum* is mechanically harvested and brought to a landfill. *Sargassum* clean-up costs for the Caribbean region were estimated at USD $210 million for the year 2018 by the Caribbean Regional Fisheries Mechanism, causing severe impacts on local economies, and highlighting the need for more information on the massive stranding events. At present, efforts to forecast and estimate biomass volumes are based mainly on the analysis of satellite data (Ody et al., 2019; Wang et al., 2019). Seaweeds causing these massive algal blooms in the Atlantic have been identified as being *S. fluitans* III (Sf), *S. natans* I (SnI), and *S. natans* VIII (SnVIII) morphotypes (Schell et al., 2015; Amaral-Zettler et al., 2017; Govindarajan et al., 2019). Recent reports suggest a combination of factors, including winds, currents and sources of nutrients, to explain the establishment, and recurrence, of pelagic *Sargassum* blooms (Putman et al., 2018; Oviatt et al., 2019; Wang et al., 2019; Johns et al., 2020).

Sargassum seaweeds are very abundant in tropical and subtropical regions, forming large floating rafts and inhabiting rocky reefs. Because these algae account for large biomass, Gouvêa et al.

(2020) suggested their contribution is relevant for global carbon stocks and consequently for mitigating climate change as CO_2 remover. There is also interest in exploiting the bioremediation potential of *Sargassum* sp. to tackle pollution in coastal environments (Saldariagga- Hernandez et al., 2020). Brown seaweeds are harvested from wild populations or farmed to provide valuable products, including texturing agents for the food industry, biofuels, fertilisers, animal feed, nutraceuticals, and cosmeceuticals (Kraan, 2013). A biorefinery approach, valorising different fractions of the biomass, has been put forward for *S. muticum* to obtain high value/low volume and high volume/low value products (Balboa et al., 2015; Milledge et al., 2016; Pérez-Larrán et al., 2019). The prospect for valorisation of *S. fluitans* and *S. natans* biomass has not been unnoticed, and Milledge and Harvey (2016) have reviewed the potential uses and obstacles for exploitation of pelagic *Sargassum*. In line with this, Thompson et al. (2020) have recently investigated the feasibility of using this biomass as a feedstock for the production of fertiliser and electricity in Barbados, one of the Caribbean nations affected by golden tides. However, considering results obtained by Milledge et al. (2020), exploitation of pelagic *Sargassum* biomass alone for the production of biogas may be challenging because of the limited production of methane from such feedstock.

The exploitation of seaweed biomass and definition of valorisation pathways should be informed by thorough knowledge of the biomass composition. Previous compositional analysis of pelagic *Sargassum* collected outside the Sargasso Sea was conducted on mixtures of *S. natans* and *S. fluitans*. Samples from the Nigerian coast were used to determine crude protein, crude fat, fibre, moisture, ash, carbohydrate, minerals and phytochemical content (Oyesiku and Egunyomi, 2014). Biomass harvested along the coast of the western region of Ghana were analysed for nutritional (nitrogen, phosphate, ammonia, nitrate and potassium) and toxicological (copper, zinc, iron, lead, cadmium, mercury, arsenic and chloride) parameters (Addico and deGraft-Johnson, 2016). Algae collected from the Southern North Atlantic were considered for biochemical analysis (C: N ratio, fatty acids, and stable isotopes) (Baker et al., 2018). Sembera et al. (2018) assessed compost quality of *Sargassum* harvested from the shoreline of a Texas beach. Other studies have dealt with individual morphotypes. Rhein-Knudsen et al. (2017) and Mohammed et al. (2018, 2019) used *S. natans* from Ghana and Trinidad and Tobago respectively, and Rosado-Espinosa et al. (2020) *S .fluitans* from the

Yucatan coast (Mexico), for extraction and characterization of the cell wall polysaccharide alginate. More recently, Rodríguez-Martínez et al. (2020) described variations in the elemental concentrations between Sf, SnI and SnVIII harvested on the Mexican Caribbean coast. Milledge et al. (2020) investigated differences in methane potential related to moisture, ash, salt, carbon, hydrogen, nitrogen, sulphur, phenolic, lipid, amino acid, metal and metalloid contents among the three pelagic morphotypes collected from the Caribbean islands of Turks and Caicos. However, no information such as pyrolysis analysis, content of the antioxidant phlorotannins, and quantification of individual monosaccharides were available to define pathways for valorisation. In addition, data was missing on the potential to fractionate *Sargassum* biomass using enzymes.

In this context, the main objectives of our study were to produce an in-depth characterisation of the pelagic *Sargassum* morphotypes, and to investigate the influence of enzymes to facilitate biomass fractionation, to inform pathways for the valorisation of this seaweed biomass and the benefits of affected countries. To this aim, we report a detailed comparative biochemical and elemental characterization, and the monosaccharide profiles obtained after enzymatic treatments, of the individual Sf, SnI, and SnVIII morphotypes, and of a bulk mixture (containing mostly Sf) collected from the south coast region of Jamaica in February 2019. Our results point to differences between morphotypes and establish a compositional library that will contribute to define possible valorisation routes for pelagic *Sargassum* biomass.

2. Materials and Methods

2.1. Study sites, field collection, and preparation of seaweed samples

Sargassum biomass was collected on the 6th of February 2019 from three different sites in the vicinity of Port Royal, Jamaica. These sites were labelled for analysis purposes as follows: Site A, Fort Rocky (17°56'12.0"N 76°49'08.4"W); Site B, Port Royal (17°56'09.2"N 76°50'17.2"W); site C, Lime Cay (17°55'06.1"N 76°49'12.8"W) (Figure 1A). Wet algae were manually collected fresh from inshore water, before they reached the shore and begun to dry. After harvesting, algae were bagged and brought

126 to the laboratory the same day of collection. After washing with tap water to remove natural solid
127 contaminants, biomass from each of the three sampling sites was separated into three different
128 morphotypes according to criteria previously described (Schell et al., 2015; Amaral-Zettler et al., 2017):
129 *S. fluitans* III (Sf), *S. natans* I (SnI), and *S. natans* VIII (SnVIII) (Figure 1B). A bulk sample of
130 unprocessed algae (Sfm), observed to be composed mainly of Sf (estimated by visual observation to be
131 10 times more than the *S. natans* biomass), was also considered for the three sites as it represents raw
132 biomass that could be used for subsequent processing without any prior separation. The resulting twelve
133 samples were kept for two days in the freezer, before drying at University of the West Indies, Mona
134 Campus (Jamaica). For this, samples were placed on drying trays, exposed in direct sunlight during the
135 days for approximately 6-8 hours daily (estimated temperatures of 29 - 31 °C), and frequently rotated
136 to ensure thorough drying. They were stored at room temperature during the nights of the three days of
137 the drying for the morphotype samples, and of the seven days for the bulk samples (longer process due
138 to difference in volume of samples). After drying, approximately half of the dried biomass for each
139 sample was sent to York (UK), where it was milled for 50 sec at 300 MHz with a tissueLyser II (Qiagen)
140 using a 20 mm stainless steel grinding ball in a 10 ml grinding jar (Qiagen).
141
142 2.2. Thermogravimetric analysis
143 Seaweed samples and calcium carbonate powder (Sigma) were analysed using a NETZSCH
144 STA 409 instrument. The heating rate was controlled at 10 °C min^{-1} from 25 to 800 °C. Nitrogen was
145 used as the carrier gas at a flow rate of 100 ml/min.
146
147 2.3. Analysis of elemental composition by inductively coupled plasma mass spectrometry (ICP-MS)
148 Three technical replicates were prepared for the three morphotypes collected at site A.
149 Approximately 0.2 g of sample was weighed accurately into a digestion vessel, and 8 ml of concentrated
150 HNO_3 and 2 ml of 30% H_2O_2 were added. The digestion vessels were sealed and placed into a
151 microwave (Milestone Ethos Up). A thermocouple was placed into the first digestion vessel to monitor
152 the temperature of the liquid inside. The microwave was programmed to heat the contents of the
153 digestion vessels to 200 °C over a period of 30 min. Once at the desired temperature, contents were

kept at 200 °C for a period of 15 min. After this, the digestion vessels were cooled down before diluting to 100 ml with distilled water. Ten ml of each sample were used for subsequent analysis. An environmental stock calibration fluid (ICP-MS calibration, Agilent part number 5183-4688), with a known concentration of many common elements found in the environment, was used to produce low (Ag, Al, As, Ba, Be, Cd, Co, Cr, Cu, Mn, Mo, Ni, Pb, Sb, Se, Th, Tl, U, V, Zn: 10,000 ppb) and high (Ca, Fe, K, Mg, Na: 1,000,000 ppb) concentration calibration standards. Algal samples and calibration solutions were run on an Agilent 7700x ICP-MS equipped with a helium collision cell.

2.4. Quantification of phenolic and phlorotannin contents

For each sample, 0.25 g of dried algal powder was loaded into a glass flask. Acetone-H_2O mixture (70% /30 % v/v, 17.5 ml) was added, and the flasks were covered with aluminium foil for extraction at room temperature in the dark for 24 h. The extracts were centrifuged at 4,000 rpm at 4°C for 8 min, and the supernatants concentrated by evaporation using a rotary evaporator. The dried extracts were thereafter re-suspended in 3-4 ml of methanol.

The phenolic content in each sample was determined by the Folin-Ciocalteu (FC) method (Singleton et al., 1999), using phloroglucinol methanolic solutions (0-0.250 mg/ml) as standards. The methanolic extract of the brown algae (100 µl) was mixed with 800 µl of the 10% 2N FC reagent, and 800 µl of 1M Na_2CO_3 were added. The resulting mixture was incubated at 40 °C for 15 min and then at room temperature for one hour. Post incubation, the absorbance was read at 650 nm on a Sunrise microplate reader spectrophotometer (Tecan) and phenolic contents were expressed as phloroglucinol equivalents (PGE) in mg/g of biomass dried weight (DW).

The phlorotannin content of each fraction was determined by the 2,4-dimethoxybenzaldehyde (DMBA) assay, which reacts specifically with m-diphenolics (1,3- and 1,3,5-substituted phenols), and is more specific than the FC reagent that reacts also with mono- and o-diphenolics (Stern et al., 1996). Phloroglucinol (0-0.06 mg/ml) methanolic solutions were used as standards. An aliquot of the methanolic solution of the dry extract (50 µl) was mixed in a capped tube with 50 µl of MeOH and 1.5 ml of working solution. This solution was obtained by mixing equal volumes of DMBA (3%, m/v) and HCl (3%, v/v), both prepared in glacial acetic acid and mixed prior to use. Samples were incubated for

one hour at room temperature in the dark. The absorbance was read at 515 nm on a visible spectrophotometer (Varian 50 Bio), and the concentrations of the phlorotannins were expressed as phloroglucinol equivalents (PGE) in mg/g of algal biomass DW.

2.5. Determination of monosaccharide composition of the non-cellulosic fraction of the biomass

Approximately five mg of each sample were weighed in triplicate in screw capped tubes. Samples were partially hydrolysed by adding 0.5 ml of 2 M trifluoroacetic acid (TFA). The vials were flushed with dry argon, mixed and heated at 100 °C for four hours, mixing periodically. The vials were then cooled to room temperature and dried in a centrifugal evaporator with fume extraction. Five hundred µl of 2-propanol were added to the samples, and vortexed before drying in centrifugal evaporator. This was then repeated once. Finally, the samples were resuspended in 200 µl of deionised water, mixed, centrifuged at 11,600 rpm for 5 min. The supernatant was filtered with 0.45 µm PTFE filters into HPLC vials, and analysed by high-performance anion-exchange chromatography on a Dionex Carbopac PA-10 column using integrated amperometry detection as described in Jones et al. (2003). To enable quantification, a standard sugar mixture containing arabinose, fucose, galactose, glucose, mannose, rhamnose, xylose, galacturonic acid, glucuronic acid, guluronic acid, mannuronic acid, and mannitol, was prepared and treated as indicated above for algal samples.

2.6. Production of alcohol insoluble residues (AIRs)

Sn I site C, Sf site C, and Sn VIII site B were selected because they represented the samples with the largest amounts of biomass available for each morphotype. Approximately half of the dried biomass from each sample was weighed, and the accurate mass recorded to calculate the % of soluble material). *Sargassum* powder was mixed with 75% ethanol, placed in boiling bath for 5 min, and span down for 5 min at room temperature. The supernatants were discarded, and pellets used to repeat wash with 75% ethanol twice. After this, a fourth wash was done with 96% ethanol under identical conditions. The resulting pellets were then washed in acetone, dried at 50 ^{o}C for 2 days, and weighed. The dried samples were kept for enzyme digestion analysis.

210 2.7. Enzymatic digestion

211 AIRs were used for enzymatic digestions with enzyme at 1 mg/ml final concentration, or in
212 absence of enzyme, replaced by buffer, as control. All samples were incubated for 24 hours at the
213 indicated optimum temperature for each enzyme: (1) 50 mg of biomass were re-suspended in 1.5 ml of
214 25 mM Na Acetate Buffer pH 4.5, and 215 µl of Cellic2® CTec2 (Novozymes) added before incubation
215 at 50 °C; (2) 50 mg of biomass were re-suspended in 1.5 ml of 10 mM potassium phosphate buffer, pH
216 6.5, 1 mM $CaCl_2$, 0.05% NaN_3, and 150 µl of amyloglucosidase (Sigma) added before incubation at 25
217 °C; (3) 50 mg of biomass were re-suspended in 1.5 ml of 100 mM potassium phosphate buffer pH 7,
218 and 100 µl of pronase (Roche) added before incubation at 40 °C; (4) 50 mg of biomass were re-
219 suspended in 1.5 ml of laminarinase buffer pH 6.2 (HEPES 50 mM, NaCl 25 mM, $CaCl_2$ 3.5 mM), and
220 50 µl of laminarinase (NZYTech, Portugal) added before incubation at 90 °C; (5) 50 mg of biomass
221 were re-suspended in 1.5 ml of alginate lyase buffer pH 9 (HEPES 50 mM, NaCl 25mM, $CaCl_2$ 3.5
222 mM), and 50 µl of alginate lyase (NZYTech, Portugal) added before incubation at 30 °C. After
223 digestion, samples were centrifuged for five minutes at 13,000 rpm, 100 µl of the supernatant was dried
224 down, and 0.5 ml of 2 M TFA was added for monosaccharide composition analysis as described in
225 section 2.5.
226
227 2.8. Data analysis

228 To assess potential differences in the biochemical and elemental composition between the three
229 pelagic *Sargassum* morphotypes, statistical analysis was conducted using SigmaPlot version 14.0. For
230 thermogravimetric analysis, as well as for comparison of monosaccharide composition, and of phenolic
231 and phlorotannin contents, values for samples corresponding to the same morphotype, or to the bulk,
232 and harvested at the three sites of collection, were pooled together. However, bulk values were not
233 considered for the statistical tests to focus on comparison between the three morphotypes. Data were
234 first tested for normality and homogeneity of variance using the Shapiro-Wilk test and Brown-Forsythe
235 test, respectively. When these tests were passed, one-way ANOVA was performed, followed by a post-
236 hoc Holm-Sidak test for all pairwise multiple comparisons. When the normality test (Shapiro-Wilk)
237 failed, Kruskal-Wallis one-way analysis of variance on ranks was applied, followed by a post hoc Tukey

238 test for all pairwise multiple comparisons. In addition, T-test done in Excel was used to assess the
239 impact of enzymatic treatment on monosaccharide composition of alcohol insoluble residues. The
240 significance level was set at p-value ≤ 0.05 for all the data analysis.
241
242 **3. Results and Discussion**
243
244 3.1. Thermogravimetric (TG) analysis of *S. natans* and *S. fluitans* morphotypes
245 The first weight loss in the TG profiles corresponded to evaporation of water at 100 °C, and the
246 moisture content represented approximately 7-8 weight % in the three morphotypes and the bulk
247 sample, with no significant differences between Sf, SnI, and SnVIII (Figure 2, and Supplementary Table
248 S1). After the start of pyrolysis, i.e. once vaporization of all moisture has happened, the main weight
249 loss was observed between 200 and 400 °C. Based on previous analysis of different brown algal biomass
250 (Ross et al., 2009; Bae et al., 2011; Kim et al., 2012 and 2013), and of polysaccharides and
251 carbohydrates of these organisms (Anastasakis et al., 2011), this can be attributed to the decomposition
252 of carbohydrates (between 200-300 °C) and of proteins (300-400 °C). These mass losses ranged
253 between 28.86 ± 0.40 (Sf) and 33.73 ± 1.47 (SnVIII) weight %, with significant differences observed
254 between SnVIII and Sf ($p = 0.017$), and SnVIII and SnI ($p = 0.015$). A third mass loss step was identified
255 in most of the samples, occurring from 600 °C, and was suggested to be due to calcium carbonate, i.e.
256 the mineral part of the exoskeleton of encrusting bryozoan. Indeed, variable quantities of white material
257 in the dried *Sargassum* samples before analysis were observed, and the occurrence of bryozoans on the
258 surface of pelagic *Sargassum* has been previously reported in the literature (Weis, 1968; Taylor and
259 Monks, 1997). We confirmed the presence of calcium carbonate in the algal samples by analysing in
260 parallel calcium carbonate standard powder. Calcium carbonate accounted for 6.30 ± 0.47 (SnVIII) to
261 10.75 ± 1.10 (Sf) weight %, with a significant difference only between SnVIII and Sf ($p = 0.032$).
262 When thermal decomposition was carried out up to 800 °C to assess pyrolysis of pelagic
263 *Sargassun* samples, the content of char ranged between 35.16 ± 6.05 (SnI) and 39.63 ± 1.34 (Sf) weight
264 %, without any significant differences between the three morphotypes. It was slightly below (34.12 ±
265 3.46) in the bulk samples (Figure 2). Such values were in the lower range compared to those measured

by TG analysis of *Sargassum* sp. from Vietnam (average of 46.17 weight %; Kim et al., 2013) or from the Red Sea (46.15 weight %; Ali and Bahadar, 2017). One potential use of the char produced by pyrolysis from pelagic *Sargassum* could be as soil enhancer, as previously suggested for other seaweeds (Bird and Benson, 1987), including *Sargassum* sp. (Roberts et al., 2015). This possible route for valorisation was discussed in details by Milledge and Harvey (2016), who indicated that the use of solar drying prior to pyrolysis for biochar production could potentially balance the insufficient energy within the *Sargassum* feedstock for drying. However, the high concentration of salt, and the accumulation by pelagic *Sargassum* morphotypes of high level of toxic elements, e.g. arsenic, should be considered cautiously.

3.2. Determination of the elemental composition of pelagic *Sargassum* morphotypes

For this analysis, samples collected from site A were used, and eighteen elements were quantified, representing about 10% of the biomass DW (Table 1). The total amounts determined were similar between SnI and Sf, while being statistically different, and lower, for SnVIII (SnVIII vs. SnI: $p = 0.004$; SnVIII vs. Sf: $p = 0.002$; Supplementary Table S2). Significant variations were observed for the macroelements Na, Mg, and Ca (all p-values ≤ 0.001 for SnVIII vs. SnI and SnVIII vs. Sf). For the microelements, amounts of Fe and Mn were significantly different among the three morphotypes (all p-values ≤ 0.05). Among metalloids, arsenic content was significantly higher in SnI (64.91 ± 0.61 µg/g DW), compared with SnVIII (60.30 ± 0.34 µg/g DW; $p = 0.031$) and Sf (58.32 ± 2.29 µg/g DW; $p = 0.009$). These values were in the wide range of levels previously reported for *Sargassum* species (20–231 µg/g DW; Milledge et al., 2016). However, they are higher than those determined by Milledge et al. (2020) from the same three morphotypes collected in Turks and Caicos (Atlantic Ocean) (21-30 µg/g DW). Arsenic contents measured in Jamaican samples were above the maximum level permitted for seaweed meal and feed materials derived from seaweed in Europe (40 µg/g DW; Official Journal of the European Union, 2019), and exceeded limits recommended for agricultural soils in different countries (15-50 µg/g DW; Rodríguez-Martínez et al., 2020). This has been previously observed in *S. fluitans* and *S. natans* biomass harvested in Nigeria (Oyesiku and Egunyomi, 2014), Ghana (Addico and deGraft-Johnson, 2016), Dominican Republic (Fernández et al., 2017), and more recently for the three

294 pelagic *Sargassum* morphotypes collected along the Caribbean coast of Mexico (Rodríguez-Martínez
295 et al., 2020). In this latter study, it was clearly suggested that the use of pelagic *Sargassum* for nutritional
296 purposes should be avoided. In this context, the elemental concentration represents an important
297 constraint to be taken into account when considering *S. fluitans* and *S. natans* for incorporation into
298 animal feed and/or human diet, as well as for the production of compost and fertiliser. Importantly,
299 organic and inorganic forms of metals and metalloids, such as for arsenic, are known to have varying
300 degrees of toxicity (Gong et al. 2002); this underlines the need to investigate the speciation of these
301 elements in *S. fluitans* and *S. natans* biomass before implementing valorisation pathways.
302
303 3.3. Determination of phenolic and phlorotannin content
304 After combining results from samples collected for each morphotype at the different collection
305 sites, the phenolic content ranged between 1.20 ± 0.43 (Sf) and 3.11 ± 0.74 (SnVIII) mg/g of biomass
306 DW. Phlorotannins are phloroglucinol-based phenolic compounds produced by brown macroalgae, and
307 have been recognized for their bioactive properties and commercial potential (Ford et al., 2019). Their
308 contents was comprised between 0.39 ± 0.21 (Sf) and 0.91 ± 0.32 (SnVIII) mg/g of biomass DW (Table
309 2). The values determined for the bulk samples were in the same ranges as those for Sf. These phenolic
310 contents are similar to those determined by Milledge et al. (2020) in pelagic *Sargassum* collected in
311 Turks and Caicos. The percentage (w/w) of phenolic compounds in the Jamaican samples ranged
312 between 0.12-0.43 % of the biomass DW, and between 0.04 and 0.09 % of the biomass DW for the
313 phlorotannins. These values were lower than those determined by Oyesiku and Egunyomi (2014), i.e.
314 phenolics representing 0.8% of the biomass DW and tannins 1.22%, using a dried mixture of *S. natans*
315 and *S. fluitans* collected in Nigeria and different analytical approaches.
316 When comparing the three morphotypes, significant variations in the phenolic content were
317 observed between SnVIII and Sf ($p < 0.001$), and SnVIII and SnI ($p = 0.01$), but not between SnI and
318 Sf (Supplementary Table S3). For the phlorotannins, the only statistically supported difference occurred
319 between SnVIII and Sf ($p = 0.002$). In line with our results, Milledge et al. (2020), using the FC method,
320 observed significant variations in the phenolic content between the three *Sargassum* morphotypes.
321 These observations, based on seaweeds collected in very distant locations, support the fact that *S. natans*

and *S. fluitans* contains different levels of phenolic compounds. In the context of exploiting pelagic *Sargassum* biomass, it will be interesting to assess if there are seasonal changes in their phenolic and phlorotannin content, since such variations have been previously reported in *S. muticum* harvested in the Isle of Wight (Gorham and Lewey, 1984), and in western Brittany (Plouguerné et al., 2006). In addition, due to the increasingly recognised nutritional and health benefits of brown algal polyphenols (Fernando et al., 2016), it will be important to explore further the chemical structure and reactivity of pelagic *Sargassum* phlorotannins for possible use in nutraceuticals, functional foods, cosmetic, and pharmaceutical applications.

3.5. Analysis of the non-cellulosic fraction of pelagic *Sargassum* morphotypes.

The total monosaccharide content across the three morphotypes ranged between 142.76 ± 32.95 (Sf) and 183.94 ± 27.46 (SnVIII) µg of monosaccharides per mg biomass DW (Figure 3), and was found to be slightly higher in the bulk samples (193.98 ± 55.05). Statistical difference was only found between content in SnVIII and Sf ($p = 0.027$) (Supplementary Table S4).

The most abundant monosaccharides investigated were mannuronic (M) and guluronic (G) acids (65-67 % of the total monosaccharides, Table 3). These are the monomers forming alginate, the main brown algal cell wall polysaccharides, which represented 9-12 % of the morphotype DW. These values were in the lower range compared to alginate content determined for representative *Sargassum* species from different locations (9.3-49.9 % DW; Rosado-Espinosa et al. (2020). Guluronic and mannuronic acid accounted for 34-35 % and 65-66 % of the alginates respectively, with M/G ratios ranging from 1.87 ± 0.12 (SnI) to 1.97 ± 0.39 (SnVIII). No statistically supported differences were observed in the alginate content and the M/G ratio between the three morphotypes. Previous analysis using *S. natans* (no information given on the morphotype) harvested on the Ghanaian coast in January 2015 shown that alginate represented 30% of the seaweed biomass (Rhein-Knudsen et al., 2017), contained 53% of G and 32% of M, with an M/G ratio equal to 0.6. Differences in alginate content and composition observed between previous and current studies can be related to the distinct sites of collection (Gulf of Guinea and Caribbean Sea), to adaptation to changing growth conditions through the years, and the use of different experimental procedures. Lower yields of alginate observed in the

three pelagic *Sargassum* morphotypes (9-12%), compared to 12-45% from the brown seaweeds used for industrial production (Peteiro 2018), are likely to limit their use as a viable source of commercial alginate. This polysaccharide is one of the most versatile polymers, historically used in a wide ranges of industries (food, feed, textile printing, papermaking and pharmaceutical), and with more recent applications in the biomedical and bioengineering fields (Peteiro et al. 2018). Interestingly, recent work using *S. natans* harvested in the Caribbean suggested that sodium alginate extracted from this species could be used as an alternative for packaging and encapsulation (Mohammed et al. 2018), and calcium alginate could be considered as a successful biosorbent of heavy metals ions (Mohammed et al. 2019). These observations warrant further analysis of the structure and properties of alginates extracted from pelagic *Sargassum*.

Two other uronic acids were identified in pelagic *Sargassum* samples. Very low amounts of galacturonic acid were quantified, while glucuronic acid content ranged between 6.82 ± 2.00 (Sf) and 9.44 ± 1.58 µg/mg of biomass DW (SnVIII). Significant variations were observed only for glucuronic acid between SnVIII and Sf ($p = 0.008$) (Supplementary Table S4).

Apart from uronic acids, the most abundant monosaccharide was fucose, accounting for 15.46 ± 1.47 (Sf) to 16.83 ± 0.94 (SnVIII) µg/mg of biomass DW. Variations in its content were statistically supported only between Sf and SnVIII ($p = 0.046$). Fucose is the main precursor of the brown algal sulphated cell wall polysaccharides fucoidans and fucans (Deniaud-Bouët et al., 2014). Galactose corresponded to the second most abundant sugar quantified in the algal samples, ranging from 10.44 ± 2.44 (Sf) to 12.50 ± 0.81 (SnI) µg/mg of biomass DW, and its content did not show significant variation when comparing the three morphotypes. Similar observations were made for glucose (from 4.25 ± 1.53 (Sf) to 4.92 ± 0.40 (SnI) µg/mg of biomass DW), and xylose (from 4.19 ± 0.33 (SnI) to 4.61 ± 0.99 (SnVIII) µg/mg of biomass DW). In contrast, quantities of mannose, from 3.53 ± 1.37 (Sf) to 4.82 ± 1.10 (SnVIII) µg/mg of biomass DW, were significantly different between SnVIII and Sf ($p = 0.04$), as for fucose. Lower quantities of rhamnose (from 1.16 ± 0.0.43 (SnVIII) to 1.48 ± 0.31 (SnI) µg/mg of biomass DW) and arabinose (from 1.13 ± 0.17 (SnI) to 1.19 ± 0.16 (Sf) µg/mg of biomass DW) were determined, without any differences across three morphotypes. Higher content of fucose in SnI, SnVIII, and Sf compared to other sugars was not unexpected based on previous reports for *S. muticum*. Several

studies have described differential fractionation methods for valorisation of this seaweed, and analysis of sugar content (fucose, galactose, xylose, glucose and mannose) in extracts produced after different treatments shown that fucose was the most abundant sugar in the majority of the fractions obtained (Balboa et al., 2015; Álvarez-Viñas et al., 2019; Pérez-Larrán et al., 2019).

SnVIII contained approximately four times more mannitol than the other morphotypes, which content ranged from 1.77 ± 0.80 (Sf) to 7.24 ± 1.13 (SnVIII) µg/mg of biomass DW. This represented less than 1% of the biomass DW of these algae. Differences in mannitol content were supported statistically between SnVIII and Sf, and SnVIII and SnI, with $p < 0.001$. Mannitol, one form of carbon storage used by brown algae, is therefore the only monosaccharide investigated for which significant changes were observed between the two *S. natans* morphotypes. In line with this, a great variability in the mannitol content has been reported within the genus *Sargassum* (1-34% biomass DW; Zubia et al., 2008), depending on season, site of collection, and species.

3.6. Production of alcohol insoluble residues (AIRs) and release of monosaccharides by different enzymatic treatments

AIRs were prepared, and mass loss between morphotypes was similar: 15.7% for SnI, 17.7% for Sf, and 22.4% for SnVIII (Supplementary Figure S1). This was within the range of soluble content observed in land plant biomass (Templeton et al., 2016). A study evaluating the potential of plant feedstock for sustainable production of biorenewables production showed soluble contents ranging between 3.39 and 28.29% of biomass (Lima et al., 2014). AIRs were subsequently used for individual enzymatic treatment using enzymes known to hydrolyse land plant cell walls (Cellic® CTec2) and starch (amyloglucosidase), brown seaweed cell wall component alginates (alginate lyase), and carbon storage polysaccharide (laminarinase), as well as a protease (pronase). The buffer only controls extracted a small amount of monosaccharides (Figure 4). This was not unexpected due to the length of the extraction and the high solubility of some compounds in macroalgae. Most of the enzymatic treatments released significantly higher amounts of monosaccharides compared to the control conditions, with the exception of Sf incubated in presence of alginate lyase and of SnVIII in presence of laminarinase (Supplementary Table S5). The highest amount of monosaccharides released after

406 hydrolysis was observed for SnI when compared with the other two morphotypes. The enzymes Cellic®
407 Ctec2 and amyloglucosidase released the highest quantities of monosaccharides among the five
408 enzymes tested and across all the samples investigated, while the enzymes acting on brown algal
409 polysaccharides and pronase generally gave a lower yield, with the exception of SnI. Glucose, mannose,
410 and galactose were the most abundant monosaccharides released through enzyme hydrolysis, mainly
411 by the Cellic® CTec2 and amyloglucosidase. The low efficiency of commercial laminarinase and
412 alginate lyase may be due to a lack of specificity towards carbon storage and cell wall polysaccharides
413 of the pelagic *Sargassum* biomass for which structure has not been characterised yet. In addition, it is
414 anticipated that sequential enzyme combinations, with and without acid pre-treatment, will release a
415 higher amount of monosaccharides from pelagic *Sargassum* for subsequent applications. Such
416 experiments have been conducted using *Sargassum* sp. harvested in different countries. Borines et al.
417 (2013) subjected *Sargassum* biomass harvested from Philippines to acid hydrolysis, and then to enzyme
418 saccharification in the presence of cellulase and cellobiase for the production of ethanol. Similarly,
419 Saravanan et al. (2018) reported ethanol production based on algal hydrolysate produced by acid and
420 enzyme (cellulase and pectinase) treatment considering *Sargassum* from India. In the same vein, Azizi
421 et al. (2017) considered *Sargassum* sp. from Persian Gulf for acid hydrolysis and enzyme
422 saccharification (cellulase and cellobiase) to produce algal hydrolysate subsequently used for microbial
423 production of polyhydroxyalkanoates, some of the most encouraging alternatives to conventional
424 plastics. These examples pave the way for further experiments on enzymatic pre-treatments of pelagic
425 *Sargassum* biomass for the production of ethanol and bioplastics, and may contribute to extend the
426 portfolio of potential applications for this feedstock.
427

428 **4. Conclusions and future considerations**

429 The three pelagic *Sargassum* morphotypes investigated in this study present differences in their
430 biochemical and elemental composition. This might be related to different source regions and dispersal
431 patterns for SnI, SnVIII, and Sf, as recently suggested (Govindarajan et al., 2019). This will affect the
432 properties of the *Sargassum* feedstock depending on the predominant morphotype during inundation
433 events. Composition may also vary with the season and after storage of the collected biomass. Char

produced by pyrolysis can potentially be used as a soil enhancer. However, high concentrations of toxic arsenic may hamper this application. Accumulation of this metalloid in high concentrations can also render the pelagic *Sargassum* unusable for nutritional purposes, despite the presence of phlorotannins which have dietary and health benefits. In addition, *S. natans* and *S. fluitans* may not be suitable as a viable source of commercial alginates because yields of extraction are low compared to those of brown algae currently used for industrial production of alginate. However, both species may represent an interesting source of carbohydrates for microbial production of ethanol and bioplastics. Further investigation will help to define the best routes for exploitation of pelagic *Sargassum* harvested after inundation events. Valorisation of this seaweed biomass, informed by biological knowledge, will contribute to the sustainable management of the *Sargassum* crisis in the affected countries.

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Legends of figures

Figure 1. Sampling sites (A) and morphological identification (B) of *S. fluitans* and *S. natans* morphotypes. For *S. fluitans*, characteristic oblong to spherical air bladders with wings but without spines (A1), broad and medium length lanceolate blades with serrated edges (B1), and lateral branches with small spines (C1) were observed. For *S. natans* I, spherical air bladders without wings but with spines (A2), narrow and long linear blades with serrated edges (B2), and lateral branches with spines were present, except for Site C (C2). *S. natans* VIII featured spherical air bladders without wings and without spines (A3), broad and medium length to long lanceolate blades with serrated edges (B3), and lateral branches without spines (C3); presence of hydroid colonies (D3) was also observed.

462

463 Figure 2. TG plots of *S. natans* I (SnI), *S. fluitans* III (Sf), *S. natans* VIII (SnVIII), and bulk samples
464 (Sfm) collected at three different sites (A, B, and C). Values in the inserted table corresponded to
465 moisture, organic matter, calcium carbonate, and char content of each morphotype calculated by
466 averaging data from weight loss curves obtained for the three sites of collection.

467

468 Figure 3. Monosaccharide composition in the non-cellulosic fraction of *S. natans* I (SnI), *S. fluitans* III
469 (Sf), *S. natans* VIII (SnVIII), and bulk samples (Sfm).

470

471 Figure 4. Quantification of monosaccharides released by individual enzymatic treatments of *S. natans*
472 I (SnI) site C, *S. fluitans* (Sf) site C, and *S. natans* VIII (SnVIII) site B. NE: no enzyme control; ED:
473 enzyme digestion.

474

475 **List of supplementary material**

476 Supplementary Figure S1. Mass loss in *S. natans* I (SnI) site C, *S. fluitans* (Sf) site C, and *S. natans*
477 VIII (SnVIII) site B samples estimated by preparation of alcohol insoluble residues (AIRs).

478

479 Supplementary Table S1. Moisture, organic matter, calcium carbonate, and char content in *S. natans* I
480 (SnI), *S. fluitans* III (Sf), *S. natans* VIII (SnVIII), and bulk samples (Sfm) collected at three different
481 sites (A, B, and C). Results are expressed in weight %.

482

483 Supplementary Table S2. Element composition analysis by ICP-MS of pelagic *Sargassum*. Results are
484 expressed as µg/kg of biomass DW.

485

486 Supplementary Table S3. Determination of phenolic and phlorotannin contents in pelagic *Sargassum*.
487 Results are expressed as mg/g of biomass DW.

488

Supplementary Table S4. Monosaccharide composition of pelagic *Sargassum*. Results are expressed as μg of monosaccharides/mg biomass DW.

Supplementary Table S5. Quantification of monosaccharides released after different enzymatic treatments of pelagic *Sargassum* biomass. Results are expressed as μg of monosaccharides/mg biomass DW.

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Table 1. Element contents determined by ICP-MS in pelagic morphotypes *S. natans* I (SnI), *S. natans* VIII (SnVIII), and *S. fluitans* III (Sf) collected at site A. Results (mean ± SD) are expressed as µg/g of biomass DW.

Elements	SnI	SnVIII	Sf
Na	11,441.00 ± 237.24	14,436.18 ± 575.76	11,310.71 ± 406.27
Mg	8,456.26 ± 300.36	6,193.47 ± 146.48	8,684.03 ± 292.54
Al	335.69 ± 18.70	187.70 ± 31.79	427.57 ± 54.94
K	28,701.30 ± 527.46	32,865.84 ± 1003.03	30,503.78 ± 1225.51
Ca	56,138.23 ± 1864.90	36,435.64 ± 690.72	57,726.79 ± 1813.97
V	2.37 ± 0.06	2.28 ± 0.18	4.21 ± 0.43
Cr	3.18 ± 0.99	1.50 ± 0.54	9.18 ± 0.37
Mn	39.62± 0.36	13.03 ± 0.48	22.92 ± 0.66
Fe	634.79 ± 18.18	237.07 ± 44.26	832.97 ± 101.84
Co	0.91 ± 0.07	0.47 ± 0.03	0.89 ± 0.06
Ni	4.21 ± 0.16	3.87 ± 0.10	3.52 ± 0.08
Cu	4.29 ± 0.16	2.78 ± 0.14	4.47 ± 0.20
Zn	14.71 ± 1.98	6.35 ± 0.62	7.2 ± 1.20
As	64.91 ± 0.61	60.30 ± 0.34	58.32 ± 2.29
Cd	0.77 ± 0.43	0.40 ± 0.02	0.57 ±0.02
Ba	22.17 ± 0.67	19.21 ± 0.65	23.21 ± 0.42
Pb	2.47 ± 1.79	0.33 ± 0.13	1.11 ± 0.47
U	0.80 ± 0.08	0.79 ± 0.01	0.83 ± 0.04
Total	105,867.69 ± 2926.44	90,467.20 ± 2410.74	109,622.32 ± 3618.09

Table 2. Determination of the phenolic (FC method) and phlorotannin (DMBA method) contents in pelagic morphotypes *S. natans* I (SnI), *S. natans* VIII (SnVIII), *S. fluitans* III (Sf), and in the bulk samples (Sfm). Results (mean ± SD) are expressed as mg PGE/g of biomass DW.

	SnI	SnVIII	Sf	Sfm
FC	2.13 ± 0.46	3.11 ± 0.74	1.20 ± 0.43	1.42 ± 0.48
DMBA	0.62 ± 0.11	0.91 ± 0.32	0.39 ± 0.21	0.34 ± 0.14

Table 3. Alginate content (mean ± SD), determined by quantification of mannuronic (M) and guluronic (G) acid monomers, in pelagic morphotypes *S. natans* I (SnI), *S. natans* VIII (SnVIII), *S. fluitans* III (Sf), and in the bulk samples (Sfm).

Samples	Alginate (% dry weight)	Alginate (% total monosaccharides)	M (% alginate)	G (% alginate)	M/G ratio
SnI	11.13 ± 2.02	66.85 ± 2.45	65.05 ± 1.47	34.95 ± 1.47	1.87 ± 0.12
SnVIII	12.18 ± 2.10	66.09 ± 2.59	65.86 ± 3.58	34.14 ± 3.58	1.97 ± 0.39
Sf	9.36 ± 2.51	65.15 ± 4.83	65.49 ± 3.80	34.51 ± 3.80	1.94 ± 0.42
Sfm	13.50 ± 4.61	68.51 ± 4.29	64.61 ± 1.47	35.39 ± 1.47	1.83 ± 0.11

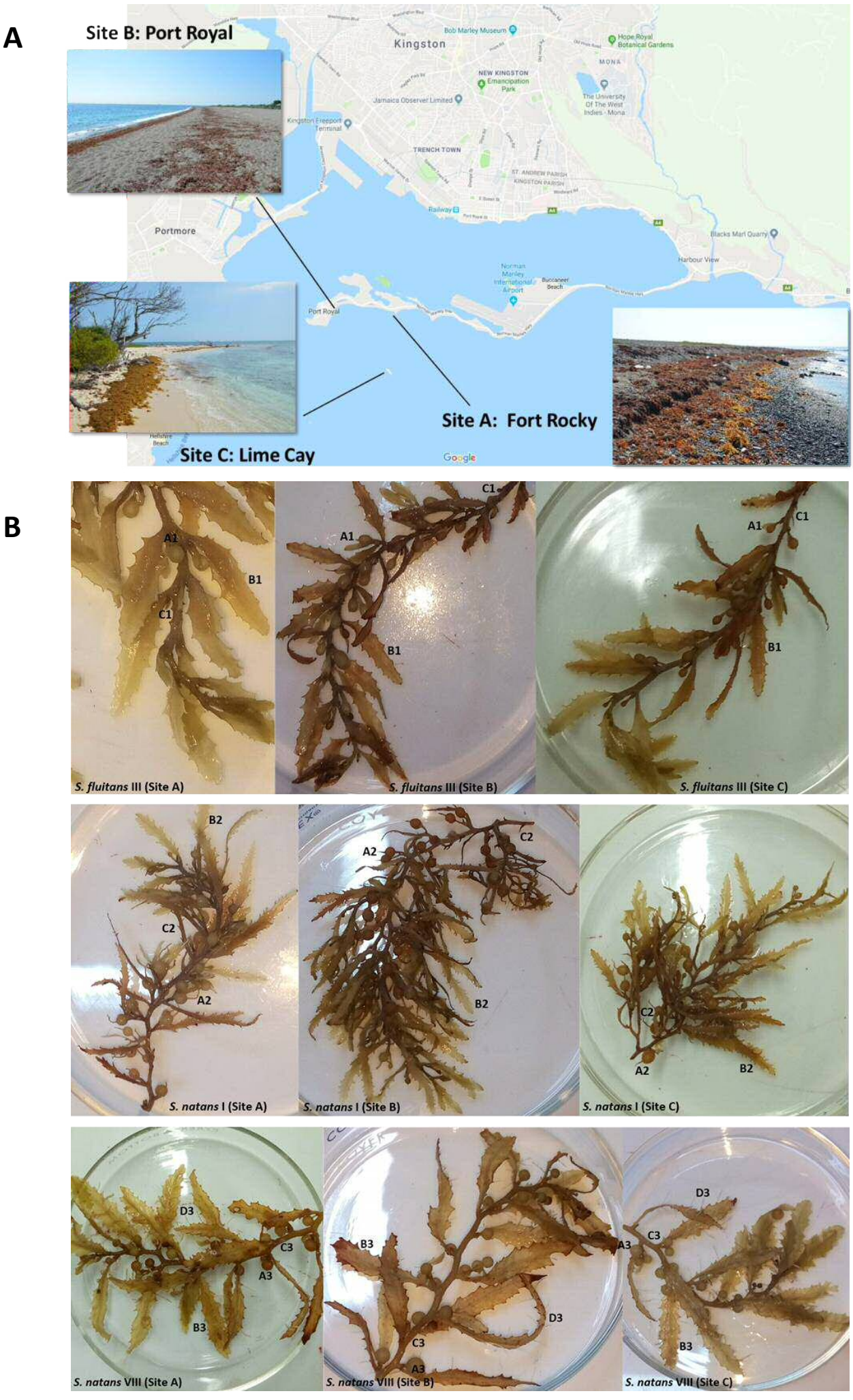

Figure 1. Sampling sites (A) and morphological identification (B) of *S. fluitans* and *S. natans* morphotypes. For *S. fluitans*, characteristic oblong to spherical air bladders with wings but without spines (A1), broad and medium length lanceolate blades with serrated edges (B1), and lateral branches with small spines (C1) were observed. For *S. natans* I, spherical air bladders without wings but with spines (A2), narrow and long linear blades with serrated edges (B2), and lateral branches with spines were present, except for Site C (C2). *S. natans* VIII featured spherical air bladders without wings and without spines (A3), broad and medium length to long lanceolate blades with serrated edges (B3), and lateral branches without spines (C3); presence of hydroid colonies (D3) was also observed.

Samples	Moisture content (weight %)	Organic matter content (weight %)	Calcium carbonate content (weight %)	Char content (weight %)
SnI	7.61 ± 0.31	29.11 ± 1.32	9.49 ± 1.75	35.16 ± 6.05
SnVIII	7.91 ± 0.95	33.73 ± 1.47	6.30 ± 0.47	36.60 ± 3.24
Sf	7.25 ± 0.42	28.86 ± 0.40	10.75 ± 1.10	39.63 ± 1.34
Sfm	8.19 ± 0.71	30.61 ± 0.75	8.56 ± 1.53	34.12 ± 3.46

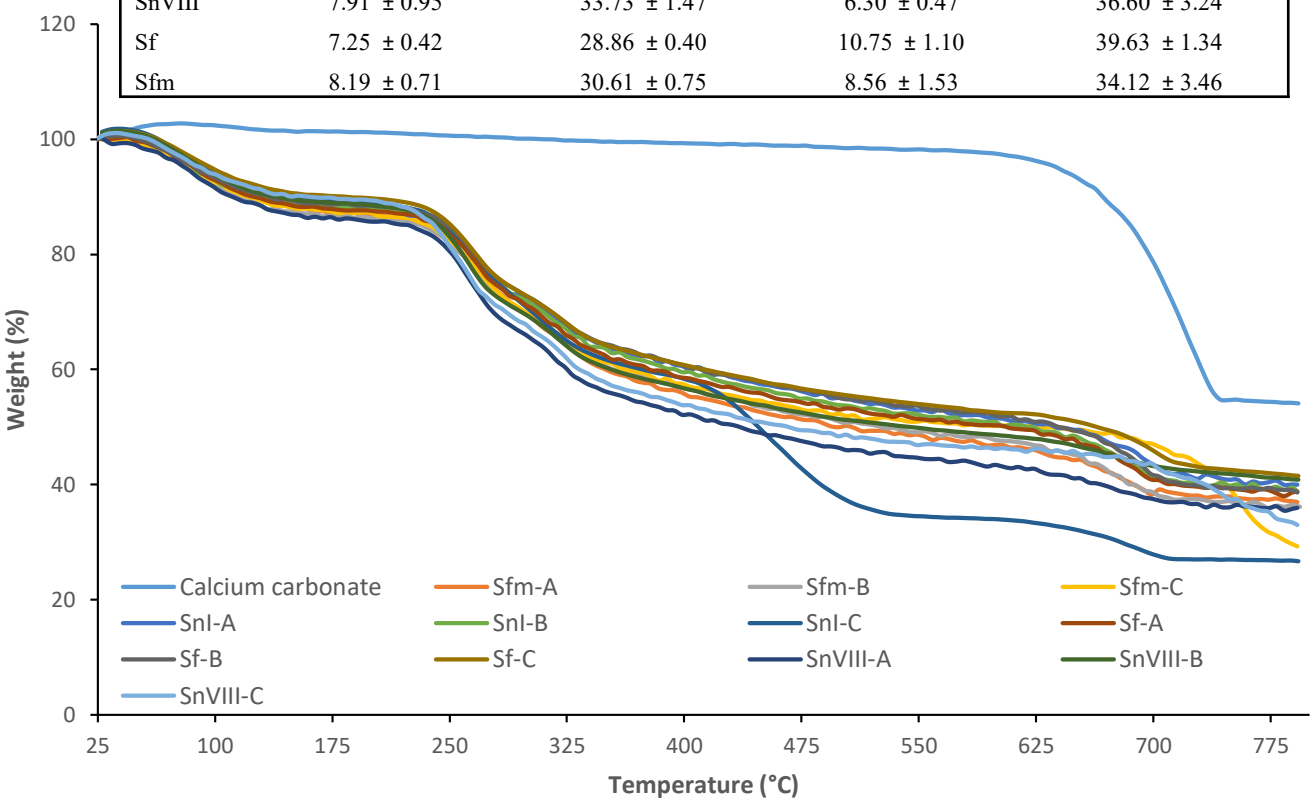

Figure 2. TG plots of *S. natans* I (SnI), *S. fluitans* III (Sf), *S. natans* VIII (SnVIII), and bulk samples (Sfm) collected at three different sites (A, B, and C). Values in the inserted table corresponded to moisture, organic matter, calcium carbonate, and char content of each morphotype calculated by averaging data from weight loss curves obtained for the three sites of collection.

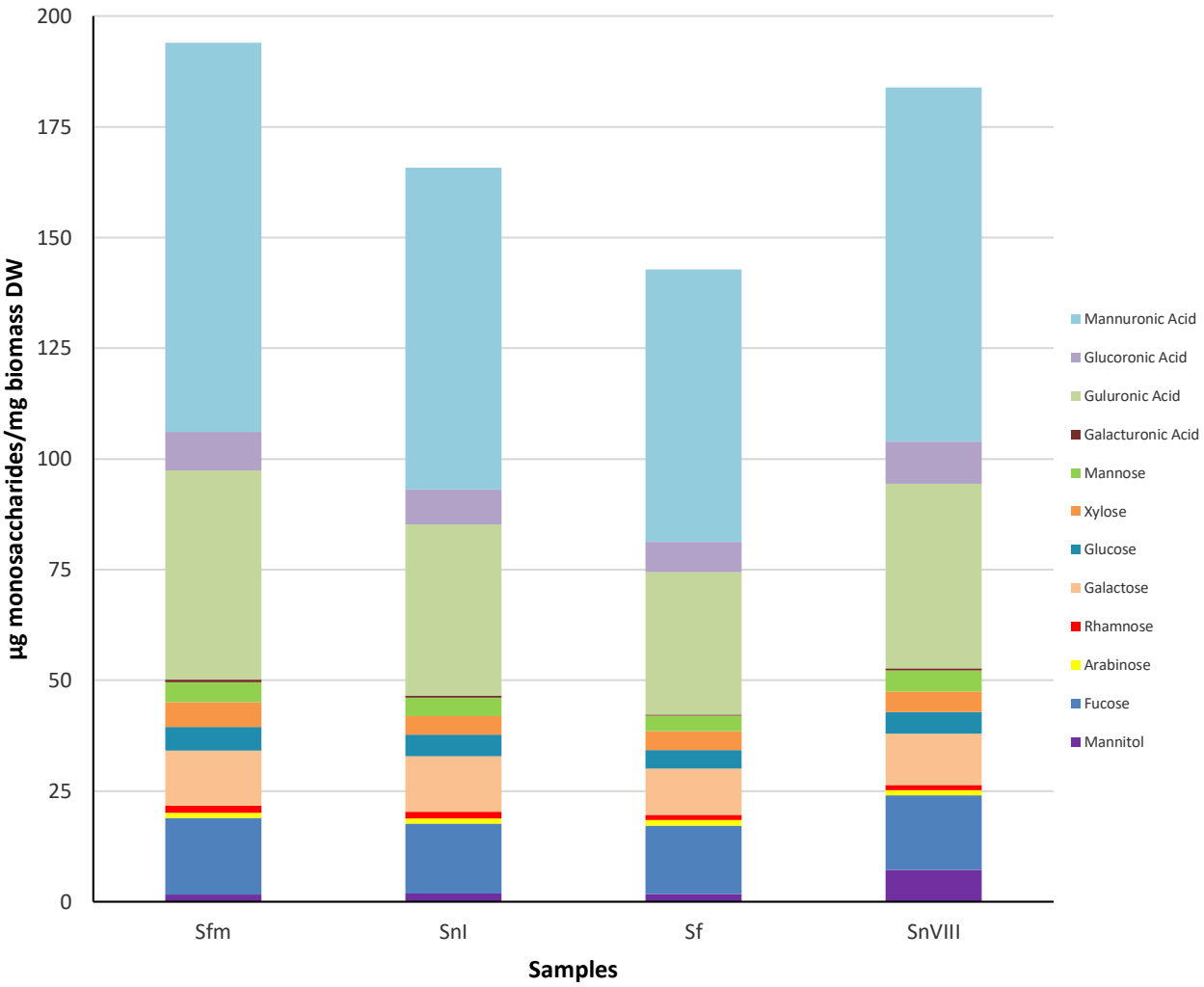

Figure 3. Monosaccharide composition in the non-cellulosic fraction of *S. natans* I (SnI), *S. fluitans* III (Sf), *S. natans* VIII (SnVIII), and bulk samples (Sfm).

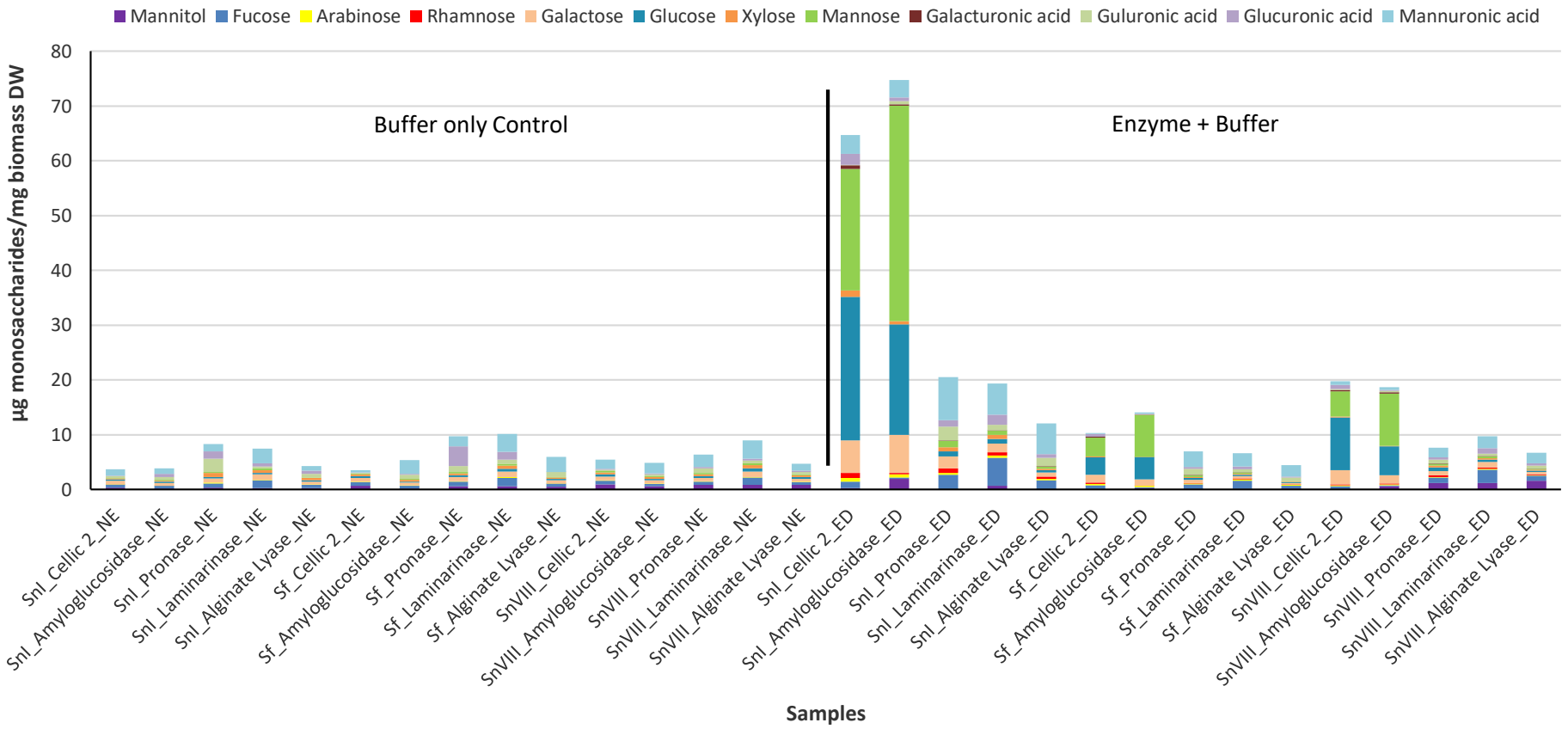

Figure 4. Quantification of monosaccharides released by individual enzymatic treatments of *S. natans* I (SnI) site C, *S. fluitans* (Sf) site C, and *S. natans* VIII (SnVIII) site B. NE: no enzyme control; ED: enzyme digestion.