

Journal of Hazardous Materials

Are biodegradable plastics an environmental rip off?

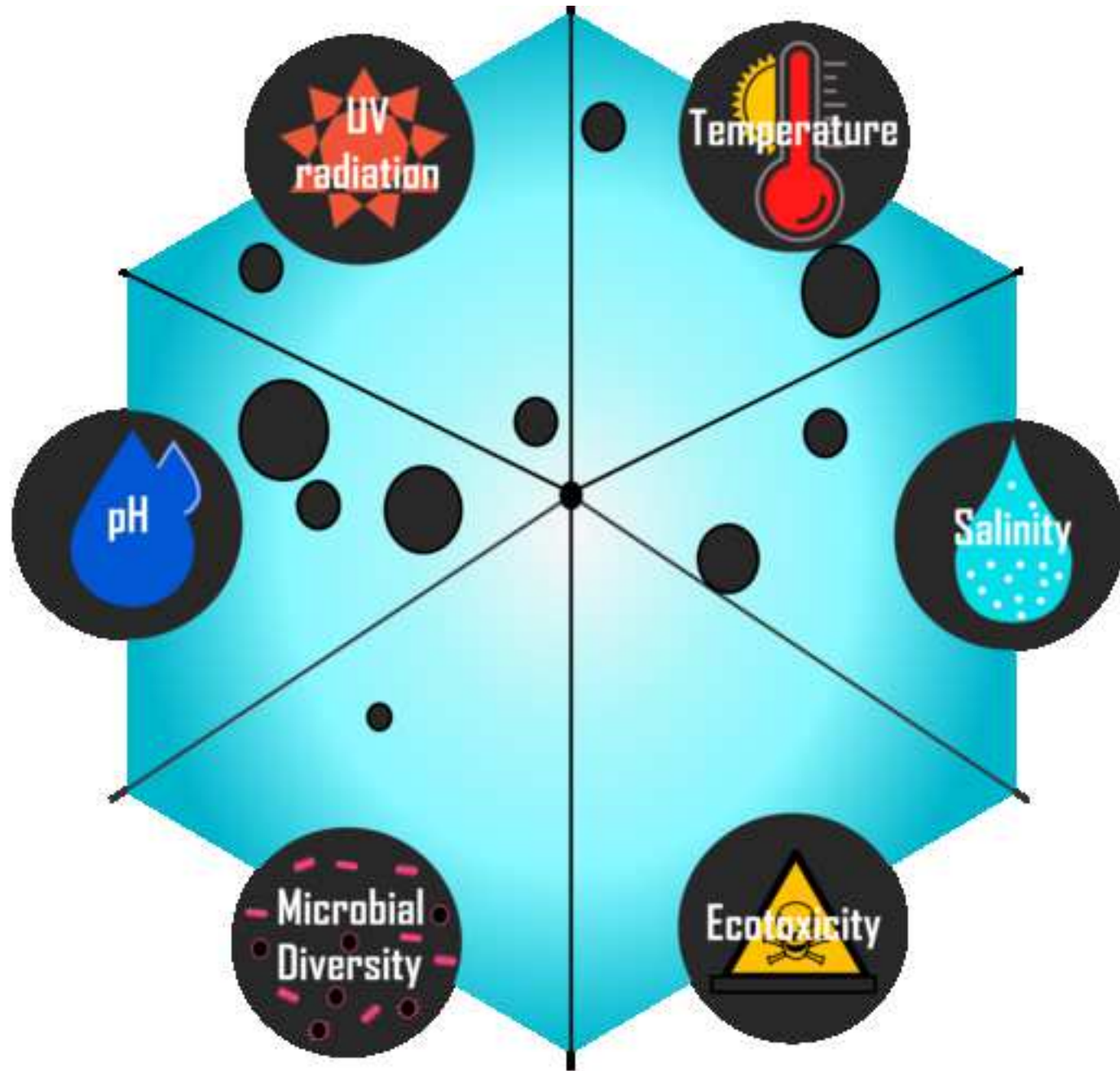
--Manuscript Draft--

Manuscript Number:	HAZMAT-D-21-01752
Article Type:	Research Paper
Keywords:	bioplastic; technical standards; pollution; marine debris; contamination
Abstract:	<p>While the use of supposedly biodegradable polymers has been recognized as a global strategy to minimize plastic pollution, the technical standards (TS) used to attest biodegradability of alternative polymers may not be in compliance with most environmental parameters observed in marine and coastal ecosystems worldwide. Indeed, through a careful assessment of the TS currently in use, this study evidenced that these guidelines cover only a fraction of the diversity of biogeochemical parameters seen in nature and largely disregard the deep-sea. Thus, these are not able to ensure degradation in natural environments. This is alarming, considering that a relevant parcel of plastic debris ends up in the oceans and reaches abyssal and hadal zones. Moreover, aquatic environments present ranges of microbial activity, pH, temperature, salinity, UV radiation and pressure that are not covered by any TS. Therefore, from a scientific perspective, claims of biodegradability placed on labels seeking to influence purchasing decisions can be considered an environmental rip off. Therefore, there is an urgent need to revise such TS, which must consider the real fate of plastic debris alongside microplastic formation and ecotoxicology effects. Furthermore, certification should provide information on time scale, degradation rates and, preferably, be globally harmonized.</p>

Statement of novelty

While biodegradable polymers are recognized as strategy to minimize plastic pollution, standards used to attest biodegradability are not in compliance with environmental parameters in aquatic systems. These guidelines cover only a fraction of the biogeochemical parameters seen in nature largely disregarding deep-sea. Thus, they are not able to ensure degradation in natural conditions. Therefore, claims of biodegradability can be considered an environmental rip off. Was proposed a revision of such standards considering the fate of plastic alongside microplastic formation and ecotoxicology effects. Furthermore, the certification of bioplastics should provide information on time scale, degradation rates and be globally harmonized.

**Parameters controlling
Bioplastic degradation in natural environments**



● Range covered by Technical standards

Highlights

- Certification of biodegradability do not reflect degradation in natural environments.
- Ranges of biogeochemical factors observed on ocean are not covered by any guide.
- New biodegradability certification schemes should be globally adopted.
- Certification of biodegradable plastics must be tested under real ocean conditions.
- Microplastics formation should be monitored in degradation experiments.

Are biodegradable plastics an environmental rip off?

João S. C. Viera^a, Mônica R.C. Marques^b, Monick Cruz Nazareth^b, Paula Christine Jimenez^a, Carlos Sanz-Lazaro^c and Ítalo Braga Castro^{a*}

^a Instituto do Mar, Universidade Federal de São Paulo (IMAR-UNIFESP), Rua Maria Máximo, 11030-100 Santos, SP, Brazil

^b Programa de Pós-Graduação em Química do Instituto de Química, Universidade do Estado do Rio de Janeiro (UERJ), Rua São Francisco Xavier, 524 Pavilhão Haroldo Lisboa da Cunha, 20559-900, RJ, Brazil

^c Department of Ecology, University of Alicante, PO Box 99, E-03080, Alicante, Spain

*Corresponding author: Ítalo Braga Castro - ibcastro@unifesp.br

Abstract

While the use of supposedly biodegradable polymers has been recognized as a global strategy to minimize plastic pollution, the technical standards (TS) used to attest biodegradability of alternative polymers may not be in compliance with most environmental parameters observed in marine and coastal ecosystems worldwide. Indeed, through a careful assessment of the TS currently in use, this study evidenced that these guidelines cover only a fraction of the diversity of biogeochemical parameters seen in nature and largely disregard the deep-sea. Thus, these are not able to ensure degradation in natural environments. This is alarming, considering that a relevant parcel of plastic debris ends up in the oceans and reaches abyssal and hadal zones. Moreover, aquatic environments present ranges of microbial activity, pH, temperature, salinity, UV radiation and pressure that are not covered by any TS. Therefore, from a scientific perspective, claims of biodegradability placed on labels seeking to influence purchasing decisions can be considered an environmental rip off. Therefore, there is an urgent need to revise such TS, which must consider the real fate of plastic debris alongside microplastic formation and ecotoxicology effects. Furthermore, certification should provide information on time scale, degradation rates and, preferably, be globally harmonized.

Keywords: bioplastic; technical standards; pollution; marine debris; contamination

32 **Introduction**

33 The occurrence of plastic debris in ocean and coastal areas has increased
34 dramatically due to a growing demand for synthetic polymers and poor management after
35 these polymers become residues. Studies indicate that in water surface of the North
36 Central Pacific Gyre there are over 334,000 plastic fragments per square kilometer [1],
37 while estimates point out 71.5 to 116 billion large (> 5mm) plastic debris on the ocean
38 floor. It is worth mentioning that such values disregards microplastics, which are also
39 widely distributed in sediments. In addition, up to 70% of plastic debris discarded in the
40 oceans reaches the deep sea [2].

41 In this sense, the superficial fouling processes undergone by floating plastics
42 may lead to sinking of such residues which reach benthic regions. In fact, plastic amounts
43 in the seabed are several orders of magnitude higher than in the water column [3]. Further,
44 plastic residues have been found beyond 1,000 km off the coast [4] and microplastics are
45 abundant in the hadal sediments of the Marianas trench, ranging from 200 to 2,200
46 particles per liter [5]. After sedimentation, the plastic deposited on the ocean floor can
47 still be resuspended and scattered by mechanical processes. According to Tosin et al., [6],
48 evidences indicates that different types of plastics circulate through various marine and
49 coastal environments due to oceanographic processes such as tides, currents and wind
50 action. Many of these factors are also responsible for the transportation and global
51 distribution of plastic debris. Therefore, along this cycle, plastic waste is exposed to a
52 multitude of environmental conditions which influence their transformation processes
53 [7,8].

54 Given the global issue of plastic pollution, scientific community and industry
55 have proposed and adopted biodegradable materials as a strategy to replace traditional
56 polymers Such materials, are supposedly able to be decomposed into CO₂ and water or

57 CH₄ by means of living organisms under a time frame comparable to average periods for
58 disintegration of biological substances [9]. In addition, these materials were designed to
59 suffer rapid degradation under environmental conditions. Simultaneously, technical
60 standards (TS) have been developed to certify biodegradability properties of commercial
61 products [10]. In fact, standardization can be critical in determining the success of
62 emergent technologies and plays a vital role in supporting key technological trends,
63 improving international trade and reducing costs and trade barriers [11]. Moreover, labels
64 certifying that a product is biodegradable may positively influence buying decisions by
65 consumers willing to pay extra for environmental benefits [12]. To meet certification
66 purposes there are internationally recognized standardization bodies such as the
67 International Standards Organization (ISO). Additionally, the American Society of
68 Methods and Materials (ASTM), the European Standardization Committee (EN) and the
69 Organization for Economic Cooperation and Development (OECD), operating at regional
70 levels, increments the list. Furthermore, several other national institutions as French
71 standardization association (AFNOR), have issued standards with protocols for
72 certification of commercial plastic products. Although most of these certification bodies
73 operate voluntarily with support of the industry, the TS produced often become part of
74 the legal framework of governments, as in the European Union [13].

75 The TS used to analyze polymers biodegradability under controlled laboratory
76 conditions often evaluate their conversion rates into carbon dioxide, water and biomass
77 [14,15] considering a particular time frame. These TS establish minimum requirements,
78 measure CO₂ emissions, or are based on visual evidence of degradation and loss of
79 polymer mass [16]. During these experiments, polymer samples may be exposed to an
80 artificially inoculated medium, which can be enriched with nutrients, and are kept under
81 specific ranges of temperature and pH. Thus, the conditions are designed to assess

82 biodegradability in optimal but artificial conditions, whereas the goal should be to
83 simulate conditions of natural environments [17]. Considering the variety of natural
84 environments experienced, seasonally and geographically, by plastic residues after
85 irregular disposal, the degradation kinetics will depend on a complex combination of
86 biotic and abiotic factors over which persists doubts if the tests established by TS are able
87 ~~of reproduce~~ [18,19]. This situation is worrisome since, based on the obtained
88 certifications, commercial products receive labels suggesting biodegradability under
89 naturally existing conditions [10]. Based on these scenarios, the present study aimed to
90 qualitatively assess whether the TS used to attest biodegradability of plastic polymers are
91 in compliance with the majority of environmental parameters observed in marine and
92 coastal ecosystems around the world.

93 **Material and Methods**

94 Several TS issued by different organizations to assess biodegradability of plastic
95 polymers based on different parameters were chosen to quantify the degradation rates
96 [20]. To carry out this study, TS for biodegradability of plastic products were obtained
97 from the main publishing organizations in the world, as shown in Table 1. Subsequently,
98 each standard was carefully reviewed and grouped according to the environment types
99 for which the tests were designed (aqueous, soil and marine environments). The intervals
100 of the physical-chemical parameters (temperature, pH, light incidence and inoculum) and
101 the operational procedures including, exposure period and biodegradation rates described
102 by each TS, were tabulated. Subsequently, the obtained data were compared to the
103 biogeochemical parameters that mainly characterized marine and coastal environments
104 of the world considering coastal and oceanic zones, as well deep-sea environments.

105 **Results**

106 A total of 17 TS issued by ASTM, ISO, EN, AFNOR and OECD were analyzed
107 (Table 2). As far as we could verify, this set represents nearly all the technical guidelines
108 developed worldwide to assess the biodegradability of plastic materials [10,21].
109 Considering the categories, were evaluated 4, 6 and 7 TS for soil, water and marine
110 environments respectively.



111 *Biodegradability of plastic polymers in soils*

112 Soils present heterogeneous characteristics and their properties are affected by
113 temperature, water content, chemical composition and pH. These parameters, separately
114 or combined, create different conditions that exert a strong influence on polymer
115 biodegradability [22,23]. Soils are usually where plastic waste is initially discarded [24].
116 In addition, soil covers made by plastic films has been globally employed as an efficient
117 strategy to improve soil properties contributing to the growth of crops [25]. Thus,
118 considering the impacts resulting from the use of conventional plastics, biodegradable
119 polymer films have become ecological alternatives to polyethylene. Based on widespread
120 use for agriculture purpose, specific standards for soil degradation of polymers have been
121 developed, especially in the European Union [26]. Such actions, seeking to ban or reduce
122 certain microplastics sources [27], are requiring certification of biodegradable plastic
123 films for agriculture use [28].

124 The four TS directed at evaluating biodegradability of plastics in soil analyzed
125 herein were designed to do so considering ideal and controlled conditions, and were
126 issued by ASTM, CEN, AFINOR and ISO. These standards recommend exposure periods
127 ranging from 2 weeks to 12 months, with temperatures between 20 and 37°C, while soil
128 pH conditions between 4.5 and 8 are also considered by AFNORNFU52-002 [29],
129 ASTMD5247 [30], ASTMD5988 [31], ISO17556 [32]. The aerobic biodegradability
130 rates considered acceptable by these standards are measured through oxygen demand or

131 CO₂ emissions and have been established in values between 60% and 90%, respectively.
132 Considering the inoculum composition, only ASTM D5988 [33] and ISO 17556 [34]
133 recommend the use of forest and field soils. In addition, none of the documents mentions
134 light exposure during the tests.

135 *Biodegradability of plastic polymers in freshwater environments*

136 High levels of microplastics have been reported for freshwater environments and
137 estuaries. Nevertheless, specific biodegradability TS for plastic items deposited in open
138 freshwater ecosystems have not been issued [18]. In contrast, several standards and
139 experimental methods have been developed to measure the degree and rate of aerobic
140 biodegradation of plastic materials in wastewater. The six TS analyzed that contemplate
141 this matter, published by CEN and ISO, recommend exposure of plastic polymers to
142 activated sludge from a sewage treatment plant and controlled digestion systems.
143 Moreover, temperature ranges should be set between 20 and 58°C and biodegradability
144 rates must be measured by comparing biochemical to theoretical oxygen demands. TS
145 issued by ISO [35,36] adopt experimental periods varying from 45 days to 6 months and
146 specify the test should be performed under dark or diffused light conditions. All TS
147 indicate pH values between 6 and 8.

148 *Biodegradability of plastic polymers in marine environments*

149 Seven TS, published by ASTM, ISO and OECD, describing methods for
150 assessing biodegradability of plastic polymers were considered in this study. The ISO
151 18830 [37] and ISO 19679 [38] TS assesses biodegradability for plastic materials in the
152 water-sediment interface, simulating sublittoral-like conditions. On the other hand,
153 ASTM D7991 [39] and ISO 22404 [40] evaluate biodegradation considering conditions
154 occurring in intertidal zones. ISO 16221 [41] and OCDE 306 [42] depict methods that
155 weigh on biodegradability by aerobic microorganisms in static aqueous systems

156 dissolving a determined amount of the material, incubated in the test medium. According
157 to ASTM D6691 [43], aerobic biodegradation is assayed under laboratory conditions that
158 simulate the pelagic zone by a defined microbial consortium, with the plastic sample
159 suspended in a synthetic sea salt solution. In all cases, aerobic biodegradation is appraised
160 by measurements of oxygen and CO₂ emissions under temperatures ranging from 15 to
161 30° C. Tests should span from 60 days up to 2 years, in which biodegradation rates must
162 reach between 60 and 70%. In TS in which it is stated, pH should be adjusted within 7
163 and 8.5, while OECD306 [44] does not include this requirement.

164 Most of these TS imply the use of sediments and filtered seawater incorporating
165 their natural microbiota; still this may be replaced by artificial seawater. In such cases,
166 ASTM D6691 [43] specifies an inoculum containing a minimum of ten microorganisms
167 including a defined taxa list. Additionally, samples of natural seawater must contain
168 inorganic nutrients such as ammonium chloride and monopotassium phosphate and be
169 obtained in uncontaminated areas. OECD306 [42] instructs the pretreatment of natural
170 seawater to remove coarse particles and to add mineral nutrients such as nitrogen and
171 phosphorus, however in concentrations higher than what are generally found in natural
172 seawater. Likewise, salinity must be adjusted to 32 ppt according to OECD306 [42],
173 ISO18830 [45] and ISO19679 [38], while ASTM D6691 [43] requires that to be 34 ppt.
174 ISO22404 [40] and ISO16221 [41] and ASTM D7991 [39] do not specify salinity to run
175 biodegradability tests.

176 **Discussion**

177 The TS issued by different institutions showed a lack of uniformity with physical
178 and chemical properties observed in actual environments, which are known to influence
179 biodegradation processes [46,47]. In Europe, the certification of biodegradability of
180 materials created by Vinçotte and managed by TUV (OK Biodegradable) is the most

181 broadly used and encompasses the following modalities: WATER, SOIL and MARINE.
182 Regarding the MARINE modality, it has been based on the technical standard ASTM
183 6691, which measures the biodegradability of the material in the water column. However,
184 oceans are composed by different environmental compartments (water column and
185 seabed) resulting in dissimilar metabolic capacities to degrade materials. In this context,
186 material degradation in the oceans, rather it is biodegradable plastic or algae, is mainly
187 mediated by microorganisms. Sediments may shelter nearly 1,000 times more
188 microorganisms than the water column [48], while various microbial groups known for
189 their function as decomposers feed on organic matter and largely lodge in marine
190 sediments [49]. Therefore, degradation processes occur in sediment at higher degrees than
191 they do in the water column. Indeed, biodegradable plastics have been shown to have a
192 10 times greater degradation rate in the sediment than in the water column [7]. This is
193 especially relevant given that 70% of plastic debris that reach the oceans end up in the
194 sediment.

195 Physico-chemical parameters in oceans and coastal zones can vary dramatically
196 in time and space. In this sense, studies have shown that pH, salinity, dissolved oxygen
197 and temperature can differ substantially among samples collected within few centimeters
198 or minutes apart [50,51]. Common oceanographic processes such as tides, currents,
199 continental discharges and euryhaline circulation are crucial agents in such dynamics.
200 Furthermore, storms, tsunami and other extreme events are also important factors to be
201 considered when assessing the physico-chemical parameters of water and marine
202 sediments [52]. Hence, fragments of biodegradable plastics discarded inappropriately will
203 be subjected to a wide variety of biogeochemical conditions.

204 Temperature is a parameter that may present pronounced horizontal and vertical
205 variations, as small oscillations in tidal cycles can induce changes in the order of 10°C

206 over a single day. Sudden variations in temperature are also observed in coastal and
207 oceanic areas where seasonal upwelling are typical, such as in South Africa, when these
208 processes alter surface water temperatures, causing declines from 20°C to less than 17°C
209 [53]. On the West Coast of North America, the recorded variations are of 3°C, reaching
210 9°C near the coasts of Oregon and California [54]. In addition, latitudinal variations in
211 surface temperatures around 28°C (in the equatorial zones) and -1.9°C (at the poles) are
212 often seen in Earth's oceans and seas [55].

213 Regarding ocean stratification, the warmest surface layers are separated from the
214 deep cold ocean by thermoclines. Below these bands of abrupt variations – deep-sea
215 regions represent 90% of oceanic areas, with a mean depth of 3,800m [56] –, the water
216 column extends to the seafloor where the temperature averages 3.5°C, while varying
217 between 1°C and 5°C [57]. Under such perspective, temperature is a crucial factor in
218 chemical reactions and may strongly influence metabolic activity of the microorganisms.
219 In fact, a temperature rise of 10°C can produce a two-fold increase in the metabolic
220 capacity of the sediment [58]. Moreover, experiments carried out using Poly-3-
221 hydroxybutyrate (P3HB) immersed in seawater, showed that at 27°C the biodegradation
222 rates were almost twice as high as those observed in water at 10°C. Temperature
223 influences biochemical reactions and taxonomic composition of microbial communities,
224 controlling reproduction, growth and distribution of decomposing microorganisms, thus
225 this is a key environmental parameter for biodegradability [59].

226 The examination conducted herein disclosed the temperatures adopted by some
227 TS, like ASTM 6691 that requires experiments to be carried out at 30°C, are generally far
228 higher than the mean occurring temperature in the oceans. In this sense, considering the
229 effects of this parameter on chemical reactions of any nature, the time allotted until full
230 degradation of a material could significantly vary depending on the temperature selected.

231 In other cases, like in ISO 19679, temperature is not accurately specified, allowing ranges
232 from 15 to 28°C, which are excessively large and can result in marked dissimilarities in
233 biodegradability outcomes, while being still very far from the conditions observed in the
234 deep-sea. Furthermore, in the deep-sea, abiotic characteristics are mostly uniform,
235 however hydrostatic pressure is a prevalent physical variable, which in the hadal systems
236 can reach between 600 to 1100 atm. An environment enduring increased pressure along
237 with reduced temperature is expected to impact the activity of enzymes in organisms
238 therein [60]. Experiments using fungi isolated from the deep-sea revealed their reduced
239 polymer degradation capacity under increased hydrostatic pressure, while no degradation
240 could be observed above 296 atm [61].

241 Salinity is another environmental factor that affects polymer degradation rates
242 as this further plays an essential role in the selection of microbial communities and over
243 their metabolic activity [62]. In surface extracts from the oceans, salinity varies between
244 32 and 37 ups, with higher values found in semi-closed seas, where evaporation far
245 exceeds precipitation, such as the Mediterranean Sea (37 to 39) and the Red Sea (40 to
246 41) [63]. On the other hand, runoff leads to salinity reduction (27 to 30) in coastal waters.
247 In estuarine systems, salinity may vary between 0 and 30 [64], whereas in the deep ocean
248 it tends to be more stable, averaging 34.8 [65]. Reasonably, only TS designed for marine
249 environments recommend adjustments in this parameter (32 – 34), whereas those
250 regarding non-saline circumstances, such as soil and freshwater environments, do not
251 consider such parameter.

252 The pH in aquatic environments is fundamentally controlled by addition or
253 removal of CO₂ due to physical and biological processes. This important parameter can
254 be modified by up to one unit due to microbial activity and phytoplankton density. In
255 addition, increases in partial pressure of CO₂ experienced after the industrial revolution

256 have contributed to gradual decreases in the pH of global aquatic systems [66]. Especially
257 in coastal areas, pH is subjected to daily and seasonal fluctuations. Thus, higher values
258 are often observed during the summer owing to changes in water temperature and
259 increased photosynthesis rates. Therefore, pH in coastal waters routinely vary between
260 7.5 and 8.5 [67], while lower values are usually observed in the eulittoral (5.5 to 6.8) and
261 sublittoral (7.8 and 7.9). In contrast, in surface waters of open oceans, pH values are
262 generally found between 7.9 and 8.3 [68]. In the deep ocean, vertical pH profiles reveal
263 average values between 7.5 and 7.6 [68]. When confronted with the analyzed TS, pH
264 values of 7, 6-8 and 7-8.5 are recommended for aqueous medium, soil and marine
265 environments, respectively. Indeed, microorganisms from different groups are known to
266 demand certain conditions to perform in degradation processes. For instance, fungi
267 usually tolerate wider pH ranges compared to bacteria [69]. Studies carried out with
268 butylene polyadipate-terephthalate (PBAT) and its biocomposites have shown this
269 material to better degrade in activated sludge compound than in natural sea water (*in situ*).
270 This study also showed that the low temperature and alkalinity of seawater stimulated the
271 activity of psychrotrophic bacteria, while acidity in the sludge favored fungi activity [70].
272 On the counter side, polylactic acid (PLA), a known biodegradable polymer, underwent
273 faster degradation in alkaline solutions influenced by the high concentration of hydroxide
274 ions [71].

275 The penetration and distribution of light radiation in water bodies is dependent
276 on depth and turbidity. In coastal regions, where the amount of particulate matter in
277 suspension is high, the photic zone can vary from a few to 60 m depth. On the other hand,
278 in pelagic environments, availability of light can reach 200m. Beyond this measure,
279 dysphotic and aphotic zones show an evident decrease in incidence of light [72].
280 Therefore, the deepest portions of the ocean, which comprises 90% of its volume, remains

281 in complete darkness. Degradation rates of plastic polymers are intensified in surface
282 environments as photodegradation take up a significant share of the process [73,74].
283 Moreover, UV radiation also plays an essential role in the vertical distribution of
284 microbial diversity [75]. In this regard, it is important to highlight that fungal and bacterial
285 species produce enzymes that mediate biodegradation of bioplastics which are, directly
286 or indirectly, dependent of light radiation [8,76]. Thus, light conditions should be a
287 controlled variable in TS testing of biodegradability, and if darkness is not applied, light
288 sources should be able to simulate the whole radiation spectrum that the oceans receive.

289 Although there are a wide variety of microorganisms and enzymes able to
290 biodegrade polymers, these are not capable of universally degrading all types of plastics.
291 Further, biodegradation rates are also dependent of polymer morphology and their
292 physical and chemical properties. In this regard, the surface area, chemical structure,
293 molecular mass, elasticity and crystalline structure are crucial features [77]. In addition,
294 environmental parameters, such as humidity, temperature, pH, salinity and hydrostatic
295 pressure, along with the availability and type of nutrients and presence of xenobiotics
296 greatly influence the dynamics of microbial systems [78]. A recent study assessing
297 degradation rates of polyhydroxybutyrate (PHB) and polybutylene sebacate-co-
298 terephthalate (PBSeT) in three different marine environments showed sample
299 disintegration under intertidal, pelagic and benthic conditions. However, significant
300 differences in degradation rates were observed and related to the distinct abiotic and biotic
301 conditions, which strongly influence microbiota diversity and function [79]. Another
302 study, which evaluated degradation of polyhydroxyalkanoate (PHA) in marine
303 environments under different climatic zones, showed disintegration rates related to light
304 exposure, temperature and oxygen [80]. Therefore, considering that microbiota is
305 connected to habitat [81], formulated inoculums are unlikely to approximate accurate

306 microbial consortia of marine environments. In this regard, ASTM D6691 [82] indorses
307 the use of either local sea water for an inoculum or one consisting of a minimum of ten
308 organisms, then listing the following species, claiming to have been identified by Gram
309 staining and biochemical evidences: *Alteromonas haloplanktis*, *Xanthomonas campestris*,
310 *Vibrio alginolyticus*, *Vibrio proteolyticus*, *Actinomycece* sp., *Bacillus megaterium*,
311 *Bacillus* sp., *Zooster* sp. and *Pseudomonas* sp. It must be noted, herein, that this list is
312 constant since, at least, the 2009 version of this document, which seemingly reflects the
313 bacteria identified in a certain environmental sample used to run the modeled experiment,
314 and not a reproduceable consortium of microorganisms which would, thus, allow
315 standardization of this protocol. Moreover, to the best our knowledge, *Zooster* sp. and
316 *Actinomycece* sp. are not valid microorganism taxa. In such a scenario, one cannot
317 overlook the title held by the ASTM D6691 [82] (*Standard test method for determining*
318 *aerobic biodegradation of plastic materials in the marine environment by a defined*
319 *microbial consortium or natural sea water inoculum*), for which the guidelines provide
320 fragile, under sought and inconsistent instructions on how to certify for biodegradability
321 in the marine environment.

322 Analogously, the tests provided by ASTM D5247- 92, which employ soil-
323 specific microorganisms, limit the plastic samples as the only carbon source, although, in
324 natural environments, the tested polymer may not be the preferred substrate in the
325 presence of more conventional nutrients. Furthermore, the ISO 16221-01, ASTM D6691-
326 17 and OECD 306-92, which add mineral nutrients to the inoculum, can stimulate growth
327 and microbial activity in a different way than what in fact occurs in natural environments,
328 while studies have indicated that nutrient addiction affects the degradation process (Tosin
329 et al., 2012b). Furthermore, the microorganisms used in experimental conditions
330 generally differ quantitatively and qualitatively from the environmental microbiota [83].

331 On the other hand, as it is reasonable to agree that achieving an environmentally accurate
332 microbial inoculum to be used under experimental conditions is not feasible, the means
333 by which TS assess and ponder microorganisms in degradation processes is still far from
334 allowing their adequate use in certification schemes.

335 Biodegradable plastics may contain toxic substances that can be released during
336 the degradation process [84]. Thus, TS should include assessments of the potential
337 toxicity derived from the biodegradation process of the tested materials. In this regard,
338 TÜV's OK Biodegradable, based on ASTM D6691 [82] includes a standard for
339 ecotoxicity tests using *Daphnia sp.* [85]. The certification scheme from the US
340 Environmental Protection Agency also indicates further protocols for ecotoxicity
341 assessments using other organisms, such as fish, algae and cyanobacteria, but does not
342 clarify if these tests must be performed in order to certify the materials. Moreover, the
343 toxicity induced during biodegradation process, could also be related to formation of
344 microplastics, which are becoming of substantial concern since those could be the most
345 harmful portion of plastics debris. Due to their reduced size (< 5 mm), these materials can
346 be ingested and prompt immune and neurotoxicity disorders [86]. Moreover, microplastic
347 also act as carriers for other pollutants and/or additives adsorbed to their surface, which
348 can then become bioavailable during degradation [87]. In this sense, there is no technical
349 standard for certification of biodegradability of plastic polymers that considers
350 microplastic formation as an intermediate or end product.

351 **Final remarks**

352 Biodegradability is a widely misused term that has been distorted generally
353 aiming to give a specific product an added “green” value that, in many cases, is not
354 accurate. At least in part, this issue is derived from the lack consensus on biodegradability
355 meaning. Indeed, decomposition under natural conditions may present different time

356 frames depending on the specific environmental variables. Thus, a biodegradability
357 concept must always be linked to a specific environment [9]. Biologically diverse marine
358 environments cover about 70% of the Earth's surface [88], offering 300 times more
359 habitable space than terrestrial and freshwater environments. In addition, these areas
360 portray a remarkable diversity of physical and chemical conditions [64]. The TS currently
361 in use contemplate only a fraction of this diversity and largely disregard the deep-sea,
362 where 32% of the accumulated debris are plastic, most of which (89%) are from single-
363 use utensils [4]. These environments have specific ranges of microbial activity, pH,
364 temperature, salinity and pressure that are not fully covered by any technical standard.
365 Even the TS designed to simulate marine environments have established experimental
366 conditions that are far from real environmental parameters.

367 Despite this, labels claiming biodegradability of these materials are currently
368 issued based on tests carried out using TS which do not reflect their degradation in natural
369 environments. As such labels wield a positive influence on consumers' purchasing
370 decision, they certainly portray an environmental rip off. Therefore, ideal TS for attesting
371 degradation should consider deep-sea environmental conditions, which could provide a
372 more appropriate assessment on material biodegradability for these ecosystems.
373 Considering the above, we propose three key points that new biodegradability
374 certification schemes should consider: (1) simulation of deep-sea environmental
375 conditions as much as possible, taking into consideration temperatures close to 0°C and
376 absence of light, among other factors; (2) using toxicity tests to assess potential toxicity
377 through degradation and (3) monitoring of microplastic formation. Furthermore, it is
378 important that new TS make truly clear to the users the meaning of the certification,
379 providing information on time scale and degradation rates. Finally, as plastic waste is
380 subject to uncontrollable cross-border movements, worldwide harmony among

381 certification schemes is a real necessity in a globalized and interconnected planet.

382 **Acknowledgments:**

383 This research was supported by São Paulo Research Foundation (FAPESP n. 2019/13750-
384 4). I.B. Castro (PQ 302713/2018-2) and M.R.C. Marques (PQ 304295/2018-3) were
385 recipient of research productivity fellowship from the Conselho Nacional de
386 Desenvolvimento Científico e Tecnológico (CNPq). M.C. Nazareth was sponsored by
387 Coordenação de Aperfeiçoamento de Pessoal de Nível Superior (CAPES / Finance code:
388 001).

389 **Table Captions**

390 Table 1: Technical standards currently used to assess the biodegradability of plastic
391 polymers.

392
393 Table 2: Main parameters used to assess biodegradability of plastic materials according
394 to several technical standards.

395

396 **References**

- 397 [1] W.C. Li, H.F. Tse, L. Fok, Plastic waste in the marine environment: A review of
398 sources, occurrence and effects, *Science of The Total Environment*. 566–567
399 (2016) 333–349. <https://doi.org/10.1016/j.scitotenv.2016.05.084>.
- 400 [2] F. Galgani, G. Hanke, T. Maes, Global Distribution, Composition and Abundance
401 of Marine Litter, in: M. Bergmann, L. Gutow, M. Klages (Eds.), *Marine*
402 *Anthropogenic Litter*, Springer International Publishing, Cham, 2015: pp. 29–56.
403 https://doi.org/10.1007/978-3-319-16510-3_2.
- 404 [3] G. Erni-Cassola, V. Zadjelovic, M.I. Gibson, J.A. Christie-Oleza, Distribution of
405 plastic polymer types in the marine environment; A meta-analysis, *Journal of*
406 *Hazardous Materials*. 369 (2019) 691–698.
407 <https://doi.org/10.1016/j.jhazmat.2019.02.067>.
- 408 [4] S. Chiba, H. Saito, R. Fletcher, T. Yogi, M. Kayo, S. Miyagi, M. Ogido, K.
409 Fujikura, Human footprint in the abyss: 30 year records of deep-sea plastic debris,
410 *Marine Policy*. 96 (2018) 204–212. <https://doi.org/10.1016/j.marpol.2018.03.022>.
- 411 [5] X. Peng, M. Chen, S. Chen, S. Dasgupta, H. Xu, K. Ta, M. Du, J. Li, Z. Guo, S.
412 Bai, Microplastics contaminate the deepest part of the world's ocean, *Geochemical*
413 *Perspectives Letters*. 9 (2018) 1–5. <https://doi.org/10.7185/geochemlet.1829>.
- 414 [6] M. Tosin, M. Weber, M. Siotto, C. Lott, F. Degli-Innocenti, Laboratory Test
415 Methods to Determine the Degradation of Plastics in Marine Environmental
416 Conditions, *Frontiers in Microbiology*. 3 (2012) 225.
417 <https://doi.org/10.3389/fmicb.2012.00225>.

- 418 [7] A. Beltrán-Sanahuja, N. Casado-Coy, L. Simó-Cabrera, C. Sanz-Lázaro,
419 Monitoring polymer degradation under different conditions in the marine
420 environment, *Environmental Pollution*. 259 (2020) 113836.
421 <https://doi.org/10.1016/j.envpol.2019.113836>.
- 422 [8] S.M. Emadian, T.T. Onay, B. Demirel, Biodegradation of bioplastics in natural
423 environments, *Waste Management*. 59 (2017) 526–536.
424 <https://doi.org/10.1016/j.wasman.2016.10.006>.
- 425 [9] A.-C. Albertsson, M. Hakkarainen, Designed to degrade, *Science*. 358 (2017) 872.
426 <https://doi.org/10.1126/science.aap8115>.
- 427 [10] J.C. Philp, A. Bartsev, R.J. Ritchie, M.-A. Baucher, K. Guy, Bioplastics science
428 from a policy vantage point, *New Biotechnology*. 30 (2013) 635–646.
- 429 [11] P.M. Wiegmann, H.J. de Vries, K. Blind, Multi-mode standardisation: A critical
430 review and a research agenda, *Research Policy*. 46 (2017) 1370–1386.
431 <https://doi.org/10.1016/j.respol.2017.06.002>.
- 432 [12] J.S.C. Viera, M.R.C. Marques, M.C. Nazareth, P.C. Jimenez, Í.B. Castro, On
433 replacing single-use plastic with so-called biodegradable ones: The case with
434 straws, *Environmental Science & Policy*. 106 (2020) 177–181.
435 <https://doi.org/10.1016/j.envsci.2020.02.007>.
- 436 [13] UNE, Report on the standardization landscape and applicable standards, 2017.
437 [https://circpack.eu/fileadmin/user_upload/DLV7.4_Standardization_map_UNE_V](https://circpack.eu/fileadmin/user_upload/DLV7.4_Standardization_map_UNE_VF.pdf)
438 [F.pdf](https://circpack.eu/fileadmin/user_upload/DLV7.4_Standardization_map_UNE_VF.pdf).
- 439 [14] EN 13432, Requisitos para embalagens recuperáveis através de compostagem e
440 biodegradação – esquema de teste e critérios de avaliação para aceitação final da
441 embalagem., 2000.
- 442 [15] EN 14046, Embalagem – avaliação da biodegradabilidade aeróbica final dos
443 materiais de embalagem sob condições controladas de compostagem – método por
444 análise de dióxido de carbono liberado., 2003.
- 445 [16] S. Chinaglia, M. Tosin, F. Degli-Innocenti, Biodegradation rate of biodegradable
446 plastics at molecular level, *Polymer Degradation and Stability*. 147 (2018) 237–
447 244.
- 448 [17] A. Krzan, S. Hemjinda, S. Miertus, A. Corti, E. Chiellini, Standardization and
449 certification in the area of environmentally degradable plastics, *Polymer*
450 *Degradation and Stability*. 91 (2006) 2819–2833.
451 <https://doi.org/10.1016/j.polymdegradstab.2006.04.034>.
- 452 [18] J.P. Harrison, C. Boardman, K. O’Callaghan, A.-M. Delort, J. Song,
453 Biodegradability standards for carrier bags and plastic films in aquatic
454 environments: a critical review, *Royal Society Open Science*. 5 (2018) 171792.
- 455 [19] B. Laycock, M. Nikolić, J.M. Colwell, E. Gauthier, P. Halley, S. Bottle, G. George,
456 Lifetime prediction of biodegradable polymers, *Progress in Polymer Science*. 71
457 (2017) 144–189. <https://doi.org/10.1016/j.progpolymsci.2017.02.004>.
- 458 [20] R. Jayasekara, I. Harding, I. Bowater, G. Lonergan, Biodegradability of a selected
459 range of polymers and polymer blends and standard methods for assessment of
460 biodegradation, *Journal of Polymers and the Environment*. 13 (2005) 231–251.
- 461 [21] A. Ammala, S. Bateman, K. Dean, E. Petinakis, P. Sangwan, S. Wong, Q. Yuan, L.
462 Yu, C. Patrick, K.H. Leong, An overview of degradable and biodegradable
463 polyolefins, *Prog. Polym. Sci.* 36 (2011) 1015.
- 464 [22] A. Hoshino, H. Sawada, M. Yokota, M. Tsuji, K. Fukuda, M. Kimura, Influence of
465 weather conditions and soil properties on degradation of biodegradable plastics in
466 soil, *Soil Science and Plant Nutrition*. 47 (2001) 35–43.
467 <https://doi.org/10.1080/00380768.2001.10408366>.

- 468 [23] F. Maréchal, Biodegradable Plastics, in: E. Chiellini, R. Solaro (Eds.),
469 Biodegradable Polymers and Plastics, Springer US, Boston, MA, 2003: pp. 67–71.
- 470 [24] Y. Chae, Y.-J. An, Current research trends on plastic pollution and ecological
471 impacts on the soil ecosystem: A review, *Environmental Pollution*. 240 (2018)
472 387–395.
- 473 [25] X. Zhang, S. You, Y. Tian, J. Li, Comparison of plastic film, biodegradable paper
474 and bio-based film mulching for summer tomato production: Soil properties, plant
475 growth, fruit yield and fruit quality, *Scientia Horticulturae*. 249 (2019) 38–48.
476 <https://doi.org/10.1016/j.scienta.2019.01.037>.
- 477 [26] F. Touchaleaume, L. Martin-Closas, H. Angellier-Coussy, A. Chevillard, G. Cesar,
478 N. Gontard, E. Gastaldi, Performance and environmental impact of biodegradable
479 polymers as agricultural mulching films, *Chemosphere*. 144 (2016) 433–439.
480 <https://doi.org/10.1016/j.chemosphere.2015.09.006>.
- 481 [27] J. Šerá, L. Serbruyns, B. De Wilde, M. Koutný, Accelerated biodegradation testing
482 of slowly degradable polyesters in soil, *Polymer Degradation and Stability*. 171
483 (2020) 109031. <https://doi.org/10.1016/j.polymdegradstab.2019.109031>.
- 484 [28] EN 17033, Plastics - Biodegradable mulch films for use in agriculture and
485 horticulture - Requirements and test methods, 2018.
- 486 [29] AFNOR NF U52-001, Materiais biodegradáveis., (2005).
- 487 [30] ASTM D5247, Método de teste padrão para determinação da biodegradabilidade
488 aeróbica de plásticos degradáveis por microrganismos específicos., 1992.
- 489 [31] ASTM D5988, Método de teste-padrão para determinar a biodegradação aeróbica
490 no solo de materiais plásticos ou material plástico residual após a compostagem.,
491 2018.
- 492 [32] ISO 17556, Plásticos – determinação do grau de desintegração de materiais sob
493 condições de compostagem em um teste de escala piloto., 2019.
- 494 [33] ASTM D5988-18, Test Method for Determining Aerobic Biodegradation of Plastic
495 Materials in Soil, ASTM International, 2018. <https://doi.org/10.1520/D5988-18>.
- 496 [34] ISO 17556, Plastics — Determination of the ultimate aerobic biodegradability of
497 plastic materials in soil by measuring the oxygen demand in a respirometer or the
498 amount of carbon dioxide evolved, ISO. (2019).
499 [https://www.iso.org/cms/render/live/en/sites/isoorg/contents/data/standard/07/49/7](https://www.iso.org/cms/render/live/en/sites/isoorg/contents/data/standard/07/49/74993.html)
500 [4993.html](https://www.iso.org/cms/render/live/en/sites/isoorg/contents/data/standard/07/49/74993.html) (accessed November 17, 2020).
- 501 [35] ISO 14851, Determinação da biodegradabilidade aeróbica final de materiais
502 plásticos em meio aquoso – método através da medição da demanda de oxigênio
503 em um respirômetro fechado, 2019.
- 504 [36] ISO 14852, Determinação da biodegradabilidade aeróbia final de materiais
505 plásticos em meio aquoso – método por análise de dióxido de carbono., 2018.
- 506 [37] ISO18830, Determinação da biodegradação aeróbica de materiais plásticos não
507 flutuantes em uma interface água do mar / sedimentos arenosos - Método através
508 da medição da demanda de oxigênio no respirômetro fechado, 2016.
- 509 [38] ISO19679, Determinação da biodegradação aeróbica de materiais plásticos não
510 flutuantes em uma interface água do mar / sedimentos - Método por análise de
511 dióxido de carbono evoluído., 2020.
- 512 [39] ASTM D7991, Método de teste padrão para determinação da biodegradação
513 aeróbica de plásticos enterrados em sedimentos marinhos arenosos em condições
514 laboratoriais controladas, 2015.
- 515 [40] ISO 22404, Plásticos - Determinação da biodegradação aeróbica de materiais não
516 flutuantes expostos a sedimentos marinhos., 2019.
- 517 [41] ISO 16221, Qualidade da água – orientação para determinação da

- 518 biodegradabilidade no ambiente marinho., 2001.
- 519 [42] OCDE 306, Biodegradabilidade em água do mar., 1992.
- 520 [43] ASTM D6691, Método de Teste Padrão para Determinação da Biodegradação
521 Aeróbica de Materiais Plásticos no Ambiente Marinho por um Consórcio
522 Microbiano Definido ou Inóculo Natural de Água do Mar., 2017.
- 523 [44] OECD, 306, OECD Guideline for Testing of Chemicals (306): Biodegradability in
524 Seawater, n.d. [https://www.oecd-ilibrary.org/environment/test-no-306-](https://www.oecd-ilibrary.org/environment/test-no-306-biodegradability-in-seawater_9789264070486-en)
525 [biodegradability-in-seawater_9789264070486-en](https://www.oecd-ilibrary.org/environment/test-no-306-biodegradability-in-seawater_9789264070486-en).
- 526 [45] ISO18830, Determinação da biodegradação aeróbica de materiais plásticos não
527 flutuantes em uma interface água do mar / sedimentos arenosos - Método através
528 da medição da demanda de oxigênio no respirômetro fechado., 2016.
- 529 [46] E. Balestri, V. Menicagli, F. Vallerini, C. Lardicci, Biodegradable plastic bags on
530 the seafloor: A future threat for seagrass meadows?, *Science of The Total*
531 *Environment*. 605–606 (2017) 755–763.
532 <https://doi.org/10.1016/j.scitotenv.2017.06.249>.
- 533 [47] J.-M. Restrepo-Flórez, A. Bassi, M.R. Thompson, Microbial degradation and
534 deterioration of polyethylene – A review, *International Biodeterioration &*
535 *Biodegradation*. 88 (2014) 83–90. <https://doi.org/10.1016/j.ibiod.2013.12.014>.
- 536 [48] B.C. Sander, J. Kalf, Factors controlling bacterial production in marine and
537 freshwater sediments, *Microbial Ecology*. 26 (1993) 79–99.
538 <https://doi.org/10.1007/BF00177045>.
- 539 [49] X. Luo, X. Xiang, G. Huang, X. Song, P. Wang, K. Fu, Bacterial Abundance and
540 Physicochemical Characteristics of Water and Sediment Associated with
541 Hydroelectric Dam on the Lancang River China, *International Journal of*
542 *Environmental Research and Public Health*. 16 (2019) 2031.
543 <https://doi.org/10.3390/ijerph16112031>.
- 544 [50] C.A. Frieder, S.H. Nam, T.R. Martz, L.A. Levin, High temporal and spatial
545 variability of dissolved oxygen and pH in a nearshore California kelp forest.,
546 *Biogeosciences*. 9 (2012).
- 547 [51] O. Guadayol, N.J. Silbiger, M.J. Donahue, F.I. Thomas, Patterns in temporal
548 variability of temperature, oxygen and pH along an environmental gradient in a
549 coral reef, *PloS One*. 9 (2014) e85213.
- 550 [52] Í.B. Castro, Improper environmental sampling design bias assessments of coastal
551 contamination, *Trends in Environmental Analytical Chemistry*. 24 (2019) e00068.
- 552 [53] J.R.E. Lutjeharms, J.M. Meeuwis, The extent and variability of South-East
553 Atlantic upwelling, *South African Journal of Marine Science*. 5 (1987) 51–62.
554 <https://doi.org/10.2989/025776187784522621>.
- 555 [54] G. Rehder, R.W. Collier, K. Heeschen, P.M. Kosro, J. Barth, E. Suess, Enhanced
556 marine CH₄ emissions to the atmosphere off Oregon caused by coastal upwelling,
557 *Global Biogeochemical Cycles*. 16 (2002) 2–1.
558 <https://doi.org/10.1029/2000GB001391>.
- 559 [55] Open-Bio, Work Package 5 In situ, biodegradation deliverable review of curret
560 methods and standards relevant to marine degradation, 2015. www.OpenUBio.eu
561 (accessed July 17, 2019).
- 562 [56] E. Ramirez-Llodra, A. Brandt, R. Danovaro, B. De Mol, E. Escobar, C.R. German,
563 L.A. Levin, P. Martinez Arbizu, L. Menot, P. Buhl-Mortensen, B.E.
564 Narayanaswamy, C.R. Smith, D.P. Tittensor, P.A. Tyler, A. Vanreusel, M.
565 Vecchione, Deep, diverse and definitely different: unique attributes of the world's
566 largest ecosystem, *Biogeosciences*. 7 (2010) 2851–2899.
567 <https://doi.org/10.5194/bg-7-2851-2010>.

- 568 [57] R. Chester, T. Jickells, Descriptive oceanography: water -column parameters., in:
569 Marine Geochemistry, John Wiley & Sons, Ltd, Chichester, UK, 2012: pp. 125–
570 153. <https://doi.org/10.1002/9781118349083.ch7>.
- 571 [58] C. Sanz-Lázaro, T. Valdemarsen, A. Marin, M. Holmer, Effect of temperature on
572 biogeochemistry of marine organic \square enriched systems: Implications in a global
573 warming scenario, *Ecological Applications*. 21 (2011) 2664–2677.
- 574 [59] A. Pischedda, M. Tosin, F. Degli-Innocenti, Biodegradation of plastics in soil: The
575 effect of temperature, *Polymer Degradation and Stability*. 170 (2019) 109017.
576 <https://doi.org/10.1016/j.polymdegradstab.2019.109017>.
- 577 [60] J.D. Gage, P.A. Tyker, The physical environment of the deep-sea, in: *Deep-Sea*
578 *Biology: A Natural History of Organisms at the Deep-Sea Floor*, Cambridge
579 University Press, 1991: p. 18 a 28.
- 580 [61] K.E. Gonda, D. Jendrossek, H.P. Molitoris, Fungal degradation of the
581 thermoplastic polymer poly- β -hydroxybutyric acid (PHB) under simulated deep
582 sea pressure, in: G. Liebezeit, S. Dittmann, I. Kröncke (Eds.), *Life at Interfaces*
583 *and Under Extreme Conditions*, Springer Netherlands, Dordrecht, 2000: pp. 173–
584 183.
- 585 [62] T. Artham, M. Doble, Biodegradation of Aliphatic and Aromatic Polycarbonates,
586 *Macromolecular Bioscience*. 8 (2008) 14–24.
587 <https://doi.org/10.1002/mabi.200700106>.
- 588 [63] P.C. Fiedler, Ocean Environments, in: B. Würsig, J.G.M. Thewissen, K.M. Kovacs
589 (Eds.), *Encyclopedia of Marine Mammals (Third Edition)*, Academic Press, 2018:
590 pp. 649–654. <https://doi.org/10.1016/B978-0-12-804327-1.00014-5>.
- 591 [64] C.M. Lalli, T.R. Parsons, *Biological Oceanography: An Introduction*, The open
592 university, 1997.
- 593 [65] F.J. Millero, *Chemical Oceanography*, 4th ed., Taylor & Francis Group, 2016.
- 594 [66] B. Hönisch, A. Ridgwell, D.N. Schmidt, E. Thomas, S.J. Gibbs, A. Sluijs, R.
595 Zeebe, L. Kump, R.C. Martindale, S.E. Greene, W. Kiessling, J. Ries, J.C. Zachos,
596 D.L. Royer, S. Barker, T.M. Marchitto, R. Moyer, C. Pelejero, P. Ziveri, G.L.
597 Foster, B. Williams, The Geological Record of Ocean Acidification, *Science*. 335
598 (2012) 1058. <https://doi.org/10.1126/science.1208277>.
- 599 [67] P. Kerrison, J.M. Hall-Spencer, D.J. Suggett, L.J. Hepburn, M. Steinke,
600 Assessment of pH variability at a coastal CO₂ vent for ocean acidification studies,
601 *Estuarine, Coastal and Shelf Science*. 94 (2011) 129–137.
602 <https://doi.org/10.1016/j.ecss.2011.05.025>.
- 603 [68] A.G. Dickson, The measurement of sea water pH, *Marine Chemistry*. 44 (1993)
604 131–142. [https://doi.org/10.1016/0304-4203\(93\)90198-W](https://doi.org/10.1016/0304-4203(93)90198-W).
- 605 [69] H.-C. Flemming, Relevance of biofilms for the biodeterioration of surfaces of
606 polymeric materials*, *Polymer Degradation and Stability*. 59 (1998) 309–315.
607 [https://doi.org/10.1016/S0141-3910\(97\)00189-4](https://doi.org/10.1016/S0141-3910(97)00189-4).
- 608 [70] K. Krasowska, J. Brzeska, M. Rutkowska, H. Janik, M.S. Sreekala, K. Goda, S.
609 Thomas, Environmental degradation of ramie fibre reinforced biocomposites.,
610 *Polish Journal of Environmental Studies*. (2010) 937–945.
- 611 [71] M. Karamanlioglu, R. Preziosi, G.D. Robson, Abiotic and biotic environmental
612 degradation of the bioplastic polymer poly(lactic acid): A review, *Polymer*
613 *Degradation and Stability*. 137 (2017) 122–130.
614 <https://doi.org/10.1016/j.polymdegradstab.2017.01.009>.
- 615 [72] A. Mitra, S. Zaman, *Basics of Marine and Estuarine Ecology*, Springer New York,
616 india, 2016.
- 617 [73] M. Bergmann, L. Gutow, M. Klages, *Marine anthropogenic litter*, Springer, 2015.

- 618 [74] N.F.A. Biber, A. Foggo, R.C. Thompson, Characterising the deterioration of
619 different plastics in air and seawater, *Marine Pollution Bulletin*. 141 (2019) 595–
620 602. <https://doi.org/10.1016/j.marpolbul.2019.02.068>.
- 621 [75] V.I. Rich, R.M. Maier, Chapter 6 - Aquatic Environments, in: I.L. Pepper, C.P.
622 Gerba, T.J. Gentry (Eds.), *Environmental Microbiology* (Third Edition), Academic
623 Press, San Diego, 2015: pp. 111–138. <https://doi.org/10.1016/B978-0-12-394626-3.00006-5>.
- 624 [76] J. Jacquin, J. Cheng, C. Odobel, C. Pandin, P. Conan, M. Pujo-Pay, V. Barbe, A.-L.
625 Meistertzheim, J.-F. Ghiglione, *Microbial Ecotoxicology of Marine Plastic Debris:*
626 *A Review on Colonization and Biodegradation by the “Plastisphere,”* *Frontiers in*
627 *Microbiology*. 10 (2019) 865. <https://doi.org/10.3389/fmicb.2019.00865>.
- 628 [77] Y. Tokiwa, P.B. Calabria, U.C. Ugwu, S. Aiba, *Biodegradability of Plastics,*
629 *International Journal of Molecular Sciences*. 10 (2009).
630 <https://doi.org/10.3390/ijms10093722>.
- 631 [78] S.K. Kale, A.G. Deshmukh, M.S. Dudhare, V.B. Patil, *Microbial degradation of*
632 *plastic: a review,* *Journal of Biochemical Technology*. 6 (2015) 952–961.
- 633 [79] D. Briassoulis, A. Pikasi, Chr. Briassoulis, A. Mistriotis, *Disintegration behaviour*
634 *of bio-based plastics in coastal zone marine environments: A field experiment*
635 *under natural conditions,* *Science of The Total Environment*. 688 (2019) 208–223.
636 <https://doi.org/10.1016/j.scitotenv.2019.06.129>.
- 637 [80] C. Lott, A. Eich, B. Unger, D. Makarow, G. Battagliarin, K. Schlegel, M.T. Lasut,
638 M. Weber, *Field and mesocosm methods to test biodegradable plastic film under*
639 *marine conditions,* *BioRxiv*. (2020) 2020.01.31.928606.
640 <https://doi.org/10.1101/2020.01.31.928606>.
- 641 [81] J. Li, W. Huang, R. Jiang, X. Han, D. Zhang, C. Zhang, *Are bacterial communities*
642 *associated with microplastics influenced by marine habitats?,* *Science of The Total*
643 *Environment*. 733 (2020) 139400. <https://doi.org/10.1016/j.scitotenv.2020.139400>.
- 644 [82] ASTMD6691, *Método de Teste Padrão para Determinação da Biodegradação*
645 *Aeróbica de Materiais Plásticos no Ambiente Marinho por um Consórcio*
646 *Microbiano Definido ou Inóculo Natural de Água do Mar,* 2017.
- 647 [83] T.P. Haider, C. Völker, J. Kramm, K. Landfester, F.R. Wurm, *Plastics of the*
648 *Future? The Impact of Biodegradable Polymers on the Environment and on*
649 *Society,* *Angewandte Chemie International Edition*. 58 (2019) 50–62.
650 <https://doi.org/10.1002/anie.201805766>.
- 651 [84] F. Degli-Innocenti, G. Bellia, M. Tosin, A. Kapanen, M. Itävaara, *Detection of*
652 *toxicity released by biodegradable plastics after composting in activated*
653 *vermiculite,* *Polymer Degradation and Stability*. 73 (2001) 101–106.
654 [https://doi.org/10.1016/S0141-3910\(01\)00075-1](https://doi.org/10.1016/S0141-3910(01)00075-1).
- 655 [85] OECD 202, *Daphnia sp. Acute Immobilisation Test,* 2004.
- 656 [86] H. Ma, S. Pu, S. Liu, Y. Bai, S. Mandal, B. Xing, *Microplastics in aquatic*
657 *environments: Toxicity to trigger ecological consequences,* *Environmental*
658 *Pollution*. 261 (2020) 114089. <https://doi.org/10.1016/j.envpol.2020.114089>.
- 659 [87] M. Smith, D.C. Love, C.M. Rochman, R.A. Neff, *Microplastics in Seafood and the*
660 *Implications for Human Health,* *Current Environmental Health Reports*. 5 (2018)
661 375–386. <https://doi.org/10.1007/s40572-018-0206-z>.
- 662 [88] A.M.P. Walag, *Bioactivities of extracts from different marine organisms around the*
663 *world (2000 to present),* *Clin Oncol*. 2 (2017) 355–361.
- 664
- 665

Table 1: Technical standards currently used to assess the biodegradability of plastic polymers.

Standard	Description
ASTM D5511-2018	Standard test method for determining anaerobic biodegradation of plastic materials under high-solids anaerobic-digestion conditions
ASTM D5988-2003	Standard test method for determining aerobic biodegradation of plastic materials in soil
ASTM D6691-2017	Standard test method for determining aerobic biodegradation of plastic materials in the marine environment by a defined microbial consortium or natural seawater inoculum
ASTM D7991-2015	Standard test method for determining aerobic biodegradation of plastics buried in sandy marine sediment under controlled laboratory conditions
EN 14047:2002	Packaging. Determination of the ultimate aerobic biodegradability of packaging materials in an aqueous medium. Method by analysis of evolved carbon dioxide
EN 14048:2002	Packaging. Determination of the ultimate aerobic biodegradability of packaging materials in an aqueous medium. Method by measuring the oxygen demand in a closed respirometer
EN 17033:2018	Plastics. Biodegradable mulch films for use in agriculture and horticulture. Requirements and test methods
ISO 13975:2019	Plastics — Determination of the ultimate anaerobic biodegradation of plastic materials in controlled slurry digestion systems — Method by measurement of biogas production
ISO 14851:1999	Determination of the ultimate aerobic biodegradability of plastic materials in an aqueous medium — Method by measuring the oxygen demand in a closed respirometer
ISO 14852:2018	Determination of the ultimate aerobic biodegradability of plastic materials in an aqueous medium — Method by analysis of evolved carbon dioxide — Technical Corrigendum
ISO 14853:2016	Plastics — Determination of the ultimate anaerobic biodegradation of plastic materials in an aqueous system — Method by measurement of biogas production
ISO 17556:2003	Plastics — Determination of the ultimate aerobic biodegradability in soil by measuring the oxygen demand in a respirometer or the amount of carbon dioxide evolved
ISO 16221:2001	Water quality — Guidance for determination of biodegradability in the marine environment
ISO 18830: 2016	Plastics — Determination of aerobic biodegradation of non-floating plastic materials in a seawater/sandy sediment interface — Method by measuring the oxygen demand in closed respirometer
ISO 19679: 2020	Plastics — Determination of aerobic biodegradation of non-floating plastic materials in a seawater/sediment interface — Method by analysis of evolved carbon dioxide
ISO 22404:2019	Plastics — Determination of the aerobic biodegradation of non-floating materials exposed to marine sediment — Method by analysis of evolved carbon dioxide
AFNOR NF U52-001	Biodegradable materials for use in agriculture and horticulture - Mulching products - Requirements and test methods
OCDE 306	Biodegradability in Seawater

Table 2: Main parameters used to assess biodegradability of plastic materials according to several technical standards.

Medium	Certification body	Technical Standard	pH	Exposure time	Inoculum	Measurements	Temperature	% Biodegradation
Soil	ASTM	D5988-18	4.5 - 8	2 weeks to - 4months	Bacteria / fungi / soil	CO ₂ emission, mass loss, tensile strength	20 - 37°C	70%
	ISO	17556:03	6 - 8	6 - 24 months	Field or forest soil	O ₂ uptake and CO ₂ emissions	20 - 28°C	60%
	AFNOR	NF U52-001	6 - 8	12 months	N/C	CO ₂ emissions	28°C	60%
	EN	17033-12	N/C	24 months	Field or forest soil	CO ₂ emissions	20–28 ° C	90%
Aqueous	EN	14047:02 14048:02	7.7	45 - 55 days	Minerals	O ₂ uptake and CO ₂ emissions	20 - 58°C	70%
	ISO	14851-99 14852-99 13975-12 14853-16	6 - 8	3 - 6 months	Soil / sludge / compost	ratio between biochemical oxygen demand and theoretical oxygen demand	20 - 25° C	60 a 70%
	ASTM	D6691-17 D7991-15	7 - 8	3 months to 2 years	Sediments / seawater	CO conversion to CO ₂	15 - 30°C	60 a 70%
Seawater	ISO	18830:16 19679:01 16221:01 22404:19	7 – 8.5	60 days to 24 months	Sediments / seawater	O ₂ uptake and CO ₂ emissions	15–28° C	60 a 70%
	OCDE	306:92	N/C	60 days	Treated seawater	Chemical oxygen demand	15-20°C	> 60%