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8	Environmental Deterioration of Biodegradable, Oxo-biodegradable, Compostable
9	and Conventional Plastic Carrier Bags in the Sea, Soil and Open-Air Over a Three-
10	<u>Year Period</u>
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18 19	Acknowledgements:
20 21 22 23 24 25 26	The authors would technicians from the Ecology Research and Plymouth Microscopy Centre throughout the
27 28	Graphical Abstract:
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31	A carrier bag labelled as biodegradable after 3
32	years in the marine environment
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## 44 <u>Abstract</u>

There is clear evidence that discarded single-use carrier bags are accumulating in the environment. As a result, various plastic formulations have been developed which state they deteriorate faster and/or have fewer impacts on the environment because their persistence is shorter. This study examined biodegradable, oxo-biodegradable, compostable and high-density polyethylene (i.e. a conventional plastic carrier bag) materials over a 3 year period. These materials were exposed in 3 natural environments; open-air, buried in soil and submersed in seawater, as well as in controlled laboratory conditions. In the marine environment, the compostable bag completely disappeared within 3 months. However, the same compostable bag type was still present in the soil environment after 27 months but could no longer hold weight without tearing. After 9 months exposure in the open-air, all bag materials had disintegrated into fragments. Collectively, our results showed that none of the bags could be relied upon to show any substantial deterioration over a 3 year period in all of the environments. It is therefore not clear that the oxo-biodegradable or biodegradable formulations provide sufficiently advanced rates of deterioration to be advantageous in the context of reducing marine litter, compared to conventional bags. Keywords: Plastics, Carrier bags, Biodegradability, Waste Management 

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78	1. Introduction
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80	Plastics are lightweight, strong, durable and corrosion-resistant materials which have
81	become an integral part of daily life worldwide <sup>1</sup> . The versatility of plastic, together
82	with its low cost, has resulted in annual worldwide production exceeding 335 million
83	tonnes <sup>2</sup> .
84	
85	Approximately 50% of plastics are discarded after a single-use <sup>3,4</sup> . This creates a
86	major waste management problem, with plastics accounting for approximately 8–
87	10% of all the waste generated in the U.K. <sup>3,5</sup> . Considerable quantities of end of life
88	plastics also escape to the environment as litter, and single-use items constitute a
89	large proportion of the litter found in marine and terrestrial environments.
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91	There is evidence that plastic debris can harm maritime industries, tourism and
92	human wellbeing <sup>6–8</sup> . In the marine environment the accumulation of plastic debris
93	has been identified as a major global issue by the United Nations Environment
94	Assembly and in the G7 Leader's declaration 2015 9-11.
95	
96	Plastic debris is widespread in terrestrial and freshwater environments. However,
97	much of the existing information about the presence of plastics in these
98	environments is focused on sources and transportation pathways to the oceans.
99	Given that the majority of all plastics will be used and disposed of on land, terrestrial
100	environments will themselves be subject to extensive pollution by plastics of all
101	sizes, based on large amounts of anthropogenic litter from both point (e.g. landfill)
102	and diffuse (e.g. general littering) sources. As such it is highly likely that soils may
103	also act as long-term sinks for plastic debris <sup>12,13</sup> .

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Since their introduction in the 1970s, plastic carrier bags have become widespread in daily life worldwide <sup>14</sup>. They are typically considered as single-use items and are commonly made from polyethylene <sup>15</sup>. These bags are an iconic symbol of our 'throw-away' society and their waste is often viewed as a very visible nuisance. In 2010, it was estimated that 98.6 billion plastic carrier bags were placed on the European Union (EU) market and about 100 billion plastic bags have been placed additionally every year since <sup>16</sup>.

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116 Plastic carrier bags are often supplied free or for a low charge and used in high 117 volumes. Consumption figures vary greatly between countries, with annual use per capita exceeding 450 bags in some EU countries <sup>16</sup>. Interventions to reduce the use 118 of plastic bags have been varied in range and scope. Governments in many nations 119 120 have strategies to either ban the sale of lightweight bags, charge customers for their use and/or generate taxes from stores who sell them <sup>17</sup>. Several countries have 121 122 already included bans or taxes, which have resulted in substantial reductions in use 123 <sup>18</sup>. However, there is no consistency between countries. 124

There are concerns that littering of plastic carrier bags presents a substantial source of contamination in the oceans. They have been found to be one of the most common items in the intertidal <sup>19,20</sup> and subtidal benthos <sup>21</sup>. Even if properly discarded, lightweight bags can unintentionally be transferred away from landfill sites

129 or other areas by wind or heavy rain 5.

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131 The presence of carrier bags in the marine environment can have a number of

132 effects; for example, previous research by Bugoni et al. (2001), found that out of 50

133 stranded dead sea turtles, plastic carrier bags were the main debris ingested.

Additionally, Green et al. (2015) found that within 9 weeks in the marine

135 environment, plastic carrier bags can create anoxic conditions within the sediment,

136 and that their presence can significantly lower abundances of infaunal invertebrates.

137 This indicates carrier bags can rapidly alter marine assemblages and the ecosystem

- services they provide <sup>23</sup>. Additionally, Hodgson et al. (2018) used laboratory
  experiments with carrier bags and showed that amphipods can shred plastic carrier
  bags, generating numerous microplastic fragments.
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The hydrophobicity and long carbon chain molecular structure of polyethylene, which
is widely used for plastic bags, makes it resistant to biodegradation under normal
conditions. The timeframe for the complete mineralisation is unknown, creating a
major waste management issue.

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Awareness of the accumulation of end of life plastic and its impact on the environment has, in part led, to interest in the development of degradable polymers. Biodegradable, oxo-biodegradable and compostable plastics are often regarded as potential solutions to the accumulation of plastic litter and waste. Some of these products are marketed accompanied by statements indicating they can be *'recycled back into nature much more quickly than ordinary plastic*<sup>'25</sup> or *'plant-based alternatives to plastic*<sup>'26</sup>.

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These materials are widely used for the production of carrier bags and some are also used to make a variety of other items, including single-use cutlery, water bottles and straws.

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Biodegradation takes place through the action of enzymes and/or chemical deterioration associated with living organisms, bacteria, fungi and algae. This occurs in two steps; the first is the fragmentation of the polymers into sections of lower molecular mass by means of either abiotic reactions (i.e. oxidation, photodegradation, hydrolysis), or biotic reactions (i.e. degradation by microorganisms). This is followed by bio-assimilation of the polymer fragments by microorganisms and its mineralisation <sup>27</sup>.

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A material may be labelled as 'biodegradable' if it conforms to certain national or regional standards <sup>28,29</sup>. Such standards could include: ISO, European Norm – EN and American Society for Testing and Materials (ASTM) International. Some standards are appropriate for conditions that occur in an industrial composter, in which temperatures are expected to reach 70 °C. Other standards, focus on laboratory-based biodegradation tests using measurements of oxygen demand or CO<sub>2</sub> evolution; for example, ISO 19679:2016 (2016) tests for the aerobic biodegradation of plastics at the interface between seawater and sandy marine sediment. Oxo-biodegradable plastics (oxo-plastics) are reported to contain an additive (pro-oxidant) which is intended to break the molecular chain within the polymer which will then lead to its biodegradation <sup>31</sup>. However, there is typically no clearly defined timeframe given for the breakdown of oxo-/biodegradable plastics <sup>32</sup>.

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179 In this context the term 'Composting' relates to enhanced biodegradation under 180 managed conditions, predominantly characterised by forced aeration and natural heat production resulting from biological activity decomposing the material. The resulting 181 182 output material, *compost*, contains nutrients and can be used as a soil improver <sup>33</sup>. Therefore, compostable plastics should biodegrade in a managed composting process 183 184 through the action of naturally occurring micro-organisms and typically do so in relation to a specified timeframe <sup>32</sup>. However, this can only occur it there is a specific waste 185 186 stream dedicated to compostable waste.

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There is a lack of clear evidence that biodegradable, oxo-biodegradable and compostable materials offer an environmental advantage over conventional plastics, and the potential for fragmentation into microplastics causes additional concern <sup>28,34</sup>. To date, studies focusing on the deterioration of different types of degradable plastics in the environment give varying results and are shorter in timeframe.

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The EU is proposing a process to restrict the use of oxo-plastics <sup>33</sup>, because of the lack of consistent evidence about rates of deterioration in the environment, allegedly misleading claims to consumers and risks that labelling products as biodegradable may inadvertently promote littering behaviour.

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The present study describes the deterioration in different natural environments of bags, which were stated to have biodegradable, oxo-biodegradable or compostable

204 properties. We do not specifically attempt to quantify biodegradation performance in

205 relation to any specific standard such as degradability in a commercial composting

206 facility. Rather we assess whether or not there has been any meaningful 207 deterioration in the context of reducing marine litter; for example, had the bag 208 remained intact or deteriorated into visible fragments? A conventional polyethylene 209 plastic carrier bag was also examined for comparison. All bags were available at the 210 point of sale in U.K. high-street retailers. These materials were exposed in various 211 environments that discarded carrier bags could encounter; in open-air, buried in soil 212 and submersed in the marine environment. This is the first research where plastic 213 deterioration has been examined simultaneously across these three natural 214 environments, together with controlled conditions in the laboratory. Five different 215 plastic carrier bag formulations were considered, and their deterioration was 216 evaluated over a 3-year period. Deterioration was considered in terms of visible loss 217 in surface area, as well as approaches to detect more subtle changes in tensile 218 stress, surface texture and chemical structure.

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## 220 2. Methodology

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## 222 2.1 Sample Preparation

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Five different types of plastic carrier bag were compared (Table 1): these included two types of oxo-biodegradable bag (labelled here as Oxobio1 and Oxiobio2), one biodegradable bag, one compostable bag, and a high-density polyethylene (HDPE) carrier bag (labelled in this research as a conventional carrier bag), which was not stated to have any particular deterioration/compostable properties. Deterioration in this study is used to describe the process of a becoming a lower quality or condition.

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Designated	Degradation	Information stated on websites linked to the product	Disposal/anti-littering
label for	properties (as		information (as stated on
testing	stated on bag)		bag)
Oxobio1	Degradable Plastics (D <sub>2</sub> W trademark,	Oxo-biodegradable (https://www.symphonyenvironmental.com/d2w/)	No information
Oxobio2	Planet safe plastic; incorporating EPI's totally degradable plastic additives (EPI trademark, logo)	Oxo-biodegradable (http://www.epi-global.com/en)	No information
Biodegradable	Biodegradable bag (exo plastics logo, sustainable bioplastic; Biodegradable ISO 14855)	No claims about biodegradability on exo plastics website (https://www.exoplastics.com/) ISO 14855 is an international standard covering aerobic biodegradability of plastic materials under controlled composting conditions	Recyclable (no numerical category for recycling type stated)
Compostable	Completely compostable, recycle me with food	Plant-based compostable foodservice packaging         (https://www.vegware.com/about/info_1.html)         Compostable packaging is designed to be recycled together with food waste.         https://www.vegware.com/close-the-loop/info_50.html         EN13432 is the packaging waste directive and standards for compostability	Reuse me first for shopping, and then as a food waste caddy liner! This completely compostable bag complies with standard EN13432.         Suitable for industrial food waste recycling – visit www.foodwastenetwork.org.uk. Recycling category '7 - other'
Conventional	High Density Polyethylene; No degradation properties stated	http://www.bpf.co.uk/topics/standards_for_compostability.aspx No manufacturer given	Reuse at home or recycle. Recycling category '2 – HDPE'

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Table 1. Information on the tested carrier bags and the properties as stated on the
 manufacture's website. All bags were opaque and obtained based on their

239 prevalence in retail stores in and around Plymouth, U.K.

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242 The bags were chosen as they were all opaque and were obtained based on their prevalence in retail stores in and around Plymouth, U.K. Sixteen samples of each 243 244 bag were obtained. In order to obtain a representative sample of each bag type, a 245 maximum of two bags were sourced from any one store on any single occasion. 246 Where repeat visits to the same store were necessary to obtain sufficient 247 independent samples, these visits were separated by at least 2 weeks. Hence our 248 experiment was designed to contain a range of products and production batches so 249 as to be as representative as possible. Since the specific retail stores from which the carrier bags were obtained is not of particular relevance, bags will only be describedbased on their formulation (Table 1).

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253 Each carrier bag type was cut into strips; 15 x 25 mm. The strip samples were taken 254 from the main body of the carrier bag (not the handles or the sides), to provide areas 255 of similar structure. A strip of each plastic carrier bag type was then placed into a 256 pouch made of high-density polyethylene (HDPE) mesh and sewn secure using 257 nylon fishing twine. Each pouch structure (150 x 200 mm) was sewn together to 258 provide 5 equally spaced separated compartments. These compartments were then 259 used to house an individual strip of each bag type (Fig.S1). The HDPE mesh (1 mm 260 x 1 mm) allowed exposure to external environments and each compartment was 261 sewn so as to allow the bag samples to move relatively freely. Each pouch structure 262 was attached to a permanent panel to aid removal.

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These permanent panels were placed in one of four different conditions; buried in 264 265 soil, exposed outdoors in the air, submerged in the marine environment and placed 266 in a blacked-out box in the laboratory as a control. The buried samples were situated 267 at the University of Plymouth's Skardon Garden (50°22'38.4"N, -4°08'11.9"W) and 268 were buried to a depth of approximately 0.25 m (Fig.S1a). The samples that were 269 exposed in open-air were also situated in Skardon Garden and were placed on a 270 south facing wall (Fig.S1b). Samples placed in the marine environment were 271 submerged on a beam at Queen Anne's Battery Marina (50°36'48.4"N, -272 4°12'96.5"W) at a depth of approximately 1 m (Fig.S1c). 3 kg weights were 273 connected on each side of the beam to maintain depth. Control samples were placed 274 in a blacked-out box (kept at room temperature) in a laboratory at the University of 275 Plymouth.

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All samples were deployed on the 10<sup>th</sup> July 2015. There were 3 subsequent
sampling dates; 6<sup>th</sup> April 2016 (9 months), 6<sup>th</sup> Jan 2017 (18 months), 6<sup>th</sup> October
2017 (27 months). Additionally, whole bags of each material were also deployed in
polypropylene mesh in each environment at the same time and used for visual
inspection over the 3-year period (23<sup>rd</sup> August 2018).

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283 Over this period, the samples would have been exposed to sea (8.8 °C - 18.8 °C; United Kingdom Sea Temperatures, 2019) and air (1.5 °C - 21.5 °C; Met Office, 284 285 2016) temperatures, typical of those in a temperate environment. The soil type in the 286 South West of the U.K. is freely draining and slightly acidic <sup>35</sup>. 287 Before deployment, 4 subsample strips from each carrier bag type were tested to 288 289 provide a comparison starting point. After deployment, four replicate samples of 290 strips from differing bag replicates were collected from each environment on each 291 sampling date. Samples were removed from the mesh structure, gently cleaned 292 using distilled water, air dried (30 °C) and tested (see below) within 48 hours. 293 294 295 2.2 Visual Inspection 296 297 298 The first step on each sampling date was to visually inspect the samples to check for 299 surface area loss, holes or disintegration. Random samples of each carrier bag type 300 were then also visualised by scanning electron microscopy (JEOL, 7001F) prior to 301 deployment, and then from each environment at 27 months. 302 303 Measurements of tensile stress and molecular structure using Fourier transform 304 infra-red spectroscopy (FTIR) were made in order to detect any more subtle 305 changes. 306 307 308 2.3 Tensile Stress Testing 309 310 The thickness of each strip was measured using an electronic micrometer (Sealey; 311 AK9635D). Each strip was measured at 10,50,100 and 140 mm from a central point. 312 This produced 4 reference points for each sample and the mean was then 313 calculated. The maximum load (N) for each strip was then measured using a tensile 314 testing machine at a rate of 100 mm min<sup>-1</sup> (Instron, system ID 3345 k1669 - USA, force transducer model 2519-104, capacity 500 N). Then, the maximum tensile 315 stress of each strip was calculated using the following equations: 316

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i) 
$$A = bh$$
 ii)  $\sigma = \frac{F}{A}$ 

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where *b* is the width (25 mm), *h* is the height (mean thickness) and *F* (maximum

321 load, N) is the force for each extracted strip. For each strip Eq. (i) allowed calculation

of the cross-sectional area (A) and Eq. (ii) allowed calculation of the tensile stress ( $\sigma$ ,

323 MPa). The maximum tensile stress of a material is also termed as its ultimate

324 strength (and referred to as the rate of disintegration within this research).

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326 Normality of the data was confirmed by using QQ plots to examine distribution. One-327 way analysis of variance (ANOVA) was used to compare the maximum tensile stress 328 difference between the different bag types before being exposed in any environment. 329 The effects of bag type, environment and time on the maximum tensile stress was 330 then examined. This was compared using the percentage change of tensile stress 331 from 0 to 9 months and 9 to 27 months using a three-way ANOVA; the three factors 332 were (bag type, environment, time). Time had two levels (0-9 and 9-27 months), bag 333 type had 5 levels (Oxobio1, Oxobio2, biodegradable, compostable and conventional) 334 and environment consisted of 4 levels (control, open-air, marine, soil). Post-hoc 335 Tukey tests were then used to identify the significant effects. Any samples which 336 were too brittle to test or were no longer visible were omitted from the analysis. All 337 statistical tests were performed in R ver. 3.4.1 (R Core Team 2017). 338

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339 <u>2.4 Molecular Composition Analysis (FTIR) and Image Analysis</u>

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In order to assess any subtle deterioration effects on the molecular composition of the materials, samples were analysed by FTIR microscopy in transmission mode with a Hyperion 1000 microscope coupled to a Vertex 70 spectrometer (Bruker). For each sample, the spectra was recorded with 32 scans in the region of 4000 to 600 cm. Prior to FTIR, samples were cleaned with absolute ethanol to remove any residues. The spectra obtained were compared against a spectral database of synthetic polymers (BPAD polymer & synthetic fibres ATR).

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- 350 **<u>3. Results</u>**
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- 352 Prior to exposure in different environments, the maximum tensile stress and

353 thickness of the bags were measured. Oxobio2 had the highest tensile stress and

thickness (28.82 ± 1.55 MPa and 0.04 mm), the compostable bag had the lowest

tensile stress (10.47 ± 1.23 MPa) and the biodegradable, conventional and Oxobio1

bag had the lowest thickness (0.02 mm) (Table S1). All bag types had relatively

- 357 consistent thickness.
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359 Before commencing the experiment there were significant differences in mean

maximum tensile stress [F (4, 15) = 12.94,  $p = \langle 0.01 \rangle$ ] between the carrier bag

361 materials (Table S2). Post-hoc Tukey's HSD tests showed that the compostable bag

had a significantly lower maximum tensile stress when compared against all other

363 bag types. All other comparisons were not significant.

364

After the various exposure periods, all pouch structures were successfully recovered
 from all environments. The strips and whole bags were then analysed visually.
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For plastic bag strips in both the control and soil environment, no surface area loss was measurable over the period of 27 months. Within the marine environment, a microbial biofilm was visible on the surface of all bags after 1-month. However, the compostable bag samples (including whole bags) were no longer visible by the 1<sup>st</sup> sampling date of 9 months.

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374 After 9 months, in the open-air environment all bag types (including conventional 375 polyethylene) were too brittle to test and had or were disintegrating into pieces. Most 376 of the pieces were in the microplastic size range (<5 mm); therefore, they could not be examined for tensile stress. The whole bags were also found to have 377 378 disintegrated into microplastic pieces. Substantial quantities of the fragments that 379 formed were visible to the naked eye on the ground beneath the test rig and in the 380 pouches. While disintegration into microplastic was apparent it was not clear whether 381 this fragmentation could have altered the potential for the plastic to biodegrade and 382 more work would be needed to establish this together with the associated timescale.

Scanning electron images were obtained before environmental exposure and then
again after 27 months. Minor changes were noticeable within the open-air
environment for sample fragments from both conventional and compostable bag
types. Cracks and holes were present in the conventional bag material suggesting
deterioration (Fig. S2;1b). For the compostable material, solid deposits that looked
like filamentous bacteria were visible on the surface; however, no cracks or holes
were present nearby (Fig. S2;2b).

After 3 years, photographs were taken of the whole bags from both the soil and
marine environment (Fig. 1). As a qualitative assessment of functionality, the bags
were loaded with typical groceries from a local supermarket (weight 2.25 kg).
Oxobio1, Oxobio2, biodegradable and conventional were still functional and retained
the items with no breakages. However, the compostable bag type (which was only
present in the soil environment for 27 months) was unable to hold any weight without
tearing.



Figure 1. Oxo-biodegradable bags (oxobio2) which had either been submerged in
the marine environment (left) or buried in soil (right) for over three years. Each bag is
holding 2.25 kg of typical groceries.

The maximum tensile stress of all plastic types decreased in all environments over
time, but at different rates (Fig. 2). This testing involved destructive sampling of 262
samples, with each measurement taken from a previously untested strip.



Figure 2. Mean maximum tensile stress of plastic carrier bag samples, shown as maximum stress before breakage displayed (mean + S.D.) over a 27-month exposure period in 4 different environments (control, marine, soil, open-air). Open-air is labelled as air in this graph. If bag type is not shown in relation to an environment, it denotes complete disintegration / fragmentation and hence samples were not testable. 

As the compostable bag samples had completely disappeared from the mesh in the marine environment this gave an unbalanced data set, and so this bag type was examined using a separate analysis just considering the remaining environments and sampling dates. Additionally, all bag types after 9 months in the open-air environment could not be tested due to being too brittle; these were subsequently omitted from statistical testing from 9-27 months.

436

437 From the perspective of litter and potential interactions with biota, most of the bag 438 samples remained intact. However, subtle changes in tensile stress were apparent in 439 all the bag materials indicating some degree of deterioration; the factors time, bag 440 type and environment showed significant differences between 0 - 9 months exposure 441 (Table S3). Post hoc comparisons found that Oxobio1 lost strength at a significantly 442 faster rate than the other bags between 0 - 9 months (p = <0.01). There were also differences between the two Oxo-biodegradable samples; Oxobio1 lost strength 443 444 significantly faster than Oxobio2 (p = 0.01). Additionally, bags exposed in the open-445 air environment lost strength more rapidly when compared to the other 446 environments: control (p = <0.01), marine (p = <0.01) and soil (p = <0.01). 447

A second ANOVA was conducted which included the compostable bag type. This bag type needed a separate analysis as all its samples had completely disappeared within the marine environment after 3 months. This ANOVA showed that the compostable bag material had a significant difference in tensile stress to Oxobio1 (p= <0.01). The compostable bag material exposed in the open-air environment also lost its strength more rapidly when compared to bags exposed to both control (p = <0.01) and soil (p = <0.01) environments.

455

Exposure from 9 - 27 months produced similar relative changes in tensile stress patterns as 0 - 9 months. However, Oxobio1 and the Conventional bag type were also found to differ in tensile stress (p = <0.01). As samples in the open-air were too brittle to test after 9 months, the only significant difference between environments was between soil and control (p = <0.01). Furthermore, the specific order of tensile stress between the bags was largely unchanged throughout 27 months, whereas environment type seemed to have a greater effect (Table S4). 464 When comparing bag types (and ignoring any samples that had deteriorated to such an extent tensile stress could not be tested), Oxobio1 had the greatest loss in tensile 465 466 stress over 27 months for all environments; soil (75% loss), marine (60% loss) and 467 control (29% loss). Conventional plastic had the least reduction in tensile stress for 468 both soil (34% loss) and the marine environment (14% loss). Compostable plastic had the lowest change in tensile stress within the control environment (11% loss), 469 470 but samples within open-air and marine environment showed total disintegration 471 (Table S5).

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473 Subtle changes in chemical composition were indicated by FTIR analysis. Some

474 samples developed a small poorly defined carbonyl stretch at a wave number of

475 approximately 1715 cm<sup>-1</sup>; this is indicative of oxidation which is a sign of

476 deterioration and was more evident for samples exposed in the open-air. However,

- 477 this varied between materials and environments, with no clear pattern being evident.
- 478

## 479 <u>4.0 Discussion</u>

480

481 Here we report the deterioration of several plastic carrier bag materials after 482 exposure in the marine, soil, open-air and control environment over a period of 3 483 years. All bags were obtained from mainstream retail shops and 4 of the materials 484 were promoted as having some level of enhanced degradability or composability 485 presumably in relation to conventional polyethylene. Apart from the compostable bag 486 material deployed in the marine environment, fragments or whole samples of each 487 bag material type were present in all environments after 27 months and some of the 488 whole bag samples were still functional as plastic bags after 3 years in the natural 489 environment.

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497 Over a 27-month period, little change in the chemical structure of any of the samples was revealed. Additionally, some subtle, but statistically testable and 498 499 significant, changes in tensile stress were apparent overtime, but the extent of these 500 changes varied among materials and environments. The rates of degradation of 501 plastics in different environments will strongly depend on the local conditions to which they are exposed <sup>36</sup>. Physical and chemical changes in polymers can be 502 503 caused by environmental factors including light (photo-oxidation), heat (photo-504 thermal oxidation), mechanical abrasion, moisture, chemical conditions or biological activity (fungi, bacteria, yeasts, algae, and their enzymes) <sup>24,37,38</sup>. For example, on 505 506 the compostable bag samples in the open-air environment solid deposits that looked 507 like filamentous bacteria were visible on the surface of the material (Fig. S2;2b). 508

509 The tensile stress of bags exposed to sunlight outdoors (labelled as open-air) decreased faster than in the other environments. Between 9 -18 months all of the 510 511 samples exposed in the open air had fragmented and could no longer hold their 512 original shape because they were too brittle. The faster rate of fragmentation in air 513 may be due to greater levels of ultraviolet (UV) radiation and oxygen, in combination with higher temperatures than in the other environments <sup>39,40</sup>. The amount of 514 515 exposure to UV would be decreased if plastics are buried in soil, landfill, or 516 submerged in the marine environment and this may explain the slower rates of 517 deterioration observed in these conditions during our study.

518

519 Samples buried in soil were found to lose tensile stress significantly faster than 520 samples in the control environment possibly because of increased moisture content 521 in the soil. Understanding the degradation of different plastic types in terrestrial 522 environments is important as substantial quantities of plastic will end up in landfills 523 <sup>41</sup>. Further, in the absence of a specific waste management pathway, for example to 524 a commercial composter, all of these materials will, unless littered be sent as 525 residual waste to landfill or incineration. When plastic accumulates within the soil, it becomes part of a complex mixture of organic matter and mineral substituents. It has 526 527 been suggested that within this environment microplastics could negatively impact 528 organisms including earthworms <sup>42,43</sup>.

529

530 Our research showed that all carrier bag materials tested appeared intact after they 531 were buried in soil conditions after 27 months. However, more subtle changes were 532 detectable with a 25 – 69% reduction in tensile stress between the different bag 533 types. These results are perhaps more realistic than the previous studies due to 534 being exposed for a longer time period and being exposed to naturally fluctuating soil 535 moisture or air temperature <sup>34,44</sup>.

536

Samples exposed in soil and open-air were, overall, found to lose tensile stress 537 538 significantly quicker than in the marine environment. However, there were no 539 significant differences between the marine environment and control samples 540 suggesting that deterioration in the marine environment was slow. Reduced 541 deterioration in seawater has been observed previously. Rutkowska et al., (2002) 542 exposed polyethylene (PE) for 20 months in 2 m water depth in the Baltic sea and reported that there was no biodegradation. Pegram and Andrady (1989) studied the 543 544 weathering of several plastics typically found in beach debris using floating marine 545 exposure tests over a 6-month period. They measured the rate of deterioration from 546 the changes in tensile elongation at break (and, in some instances, by the force to 547 rupture) and found it to be much slower (2%) for samples exposed in the sea 548 compared to samples exposed in open-air (95%). In the current experiment, after 9 549 months, conventional polyethylene was found to lose 31% in tensile stress in open-550 air, but only 2% in the marine environment.

551

552 Colonisation by micro- and macro-marine organisms (a process described as 553 fouling), occurs in natural environments and will vary according to conditions (e.g. 554 temperature). This fouling process may affect plastic in a variety of ways <sup>47</sup>. Firstly, the biofilm may 'shield' the plastic from UV light <sup>34</sup> thus reducing the rate of photo-555 degradation. Within the marine environment, fouling can also make plastics 556 negatively buoyant causing buoyant items to sink <sup>48</sup>; hence further reducing 557 558 irradiance. In the current experiment, all samples in the marine environment readily 559 acquired a coating of biofilm.

560

All samples of the compostable bag (Compost), including the whole bag, completely
deteriorated within a 3-month period in the marine environment. Similarly, research
by O'Brine and Thompson (2010) also found that a compostable bag type had 100%

- surface area loss between 16 and 24 weeks when deployed in the marine
- 565 environment. This suggests that deterioration of compostable bags can be relatively
- rapid in seawater. However, more work would be needed to establish what the
- 567 breakdown products of this deterioration are, such as microplastics or nanoplastics,
- 568 and to consider any potential environmental consequences.
- 569
- 570 From the perspective of the remaining bag types, it might have been expected that 571 the two oxo-biodegradable materials would degrade faster than both the
- 572 biodegradable and conventional bag types as these bags have pro-oxidants which
- are incorporated into the polymer chains to accelerate photo- and thermo-oxidation
- <sup>44</sup>. However, throughout the 27 months of this experiment, Oxobio1 was the only bag
- 575 type to lose tensile stress significantly faster compared to biodegradable,
- 576 conventional and Oxobio2 bag types.
- 577

Koutny et al., (2006) studied the biodegradability of high-density polyethylene film 578 579 (HDPE) and low-density polyethylene film (LDPE) containing pro-oxidants and 580 antioxidants. These were tested against microbial strains (Koutny et al., 2006; Larkin 581 et al., 2005). After an abiotic pre-treatment consisting of photooxidation and 582 unnaturally high thermo-oxidation (60 °C) which was intended to mimic around 3 583 years of outdoor weathering, the samples were inoculated, incubated up to 200 days (27 °C) and their metabolic activities were followed. An initial phase of fast microbial 584 585 growth was observed, and the authors suggest this was probably caused by 586 utilization of low molecular extractable compounds. This was followed by a long 587 period of stabilized metabolic activity. Analysis performed at the end of incubation 588 indicated that any biodegradation had probably only affected the surface layer of the 589 materials.

590

591 The current study showed that, oxo-biodegradable, degradable and conventional 592 carrier bag materials did not degrade quickly in any of the natural environments 593 examined and, in some cases, formulations merely disintegrated into small pieces 594 (such as those in the open-air environment). There are considerable concerns about 595 the accumulation of microplastics in the environment and it remains to be 596 established whether fragmentation into microplastics presents greater environmental 597 risks than the original intact items of litter. From the perspective of cleansing, fragments are certainly considerably harder, if not impossible, to remove from theenvironment compared to intact items.

600

It is of importance to understand the actual environmental degradability performance of materials which are claimed to have enhanced degradation properties as these could make consumers more relaxed about discarding, or even littering them, rather than reusing and recycling. Due to the growing interest in products which indicate enhanced environmental outcomes, we should be careful that such products do not inadvertently encourage littering or compromise alternative approaches to waste reduction such as recycling.

608

609 Designing products specifically to degrade in the environment is very challenging 610 because of the natural variability between environment types, as illustrated by the present study. In addition, formulations that are designed to be less durable may 611 612 compromise recyclability since they decrease the durability of the recyclate. It is also 613 important to set the benefits of the various formulations into a wider context since 614 reducing the diversity of polymers that are widely used is likely to facilitate greater 615 recycling. To gain the maximum benefit from materials with enhanced rates of 616 degradability, it is essential to have clear definitions and product labelling to indicate 617 appropriate usage and disposal <sup>7</sup>.

618

619 If products are designed or marketed with the intent to make a valuable contribution 620 in reducing the impacts of plastic litter in the natural environment then it is imperative 621 to have appropriate standard tests against which to assess performance. These 622 standards would need to incorporate the variability of natural environmental 623 conditions (e.g. temperature/pH/light) and an appropriate time scale of deterioration 624 such that it is clear items are deteriorating sufficiently rapidly to make a difference and not leave any potentially harmful degradation products (chemicals of fragments). 625 626 In addition to appropriate standards and tests, the relevant receiving environment in 627 which breakdown is expected to occur also needs to be stated. 628

629 Clearly there may be drivers for the design of products with modified degradability

other than deterioration in the natural environment, but in order for any these

631 potential benefits to be realised it is essential that such products have a high

632 probability of actually reaching the appropriate waste stream. This will require availability of a dedicated waste stream, the appropriate infrastructure such as an 633 634 industrial composting facility and sufficient understanding amongst consumers to 635 correctly separate their waste accordingly. Some nations actively promote the use of 636 carrier bags with biodegradable, degradable or compostable formulations, for 637 example using fiscal measures or other legislation. This includes some nations with 638 relatively poor waste management infrastructure where the likelihood of these 639 products reaching an appropriate waste stream seems low. Given the findings of this 640 study, the benefits of such policy measures are unclear.

641

642 In conclusion, the current experiment has shown that biodegradable, oxo-

643 biodegradable and conventional plastic formulations persist and remain functional in 644 the soil and the marine environment for over 3 years. The compostable bag was the 645 only material that completely disappeared from the experimental test rig within the 646 marine environment and did this within a 3-month period, but this product remained 647 intact in soil. Hence the current study indicated that over a 3-year period, none of the 648 materials examined could be relied upon to deteriorate sufficiently enough to reduce 649 the negative effects of littering on biota or aesthetics across all three environments. 650 Moreover, it was not clear that materials which claimed to have enhanced 651 degradation consistently deteriorated faster than conventional polyethylene. 652 Deterioration was influenced by the receiving environment, but this was not 653 consistent among material types. Hence, we suggest that statements about the 654 degradation of products should be clearly linked to appropriate standards, made in 655 conjunction with statements on the receiving environment (air, soil, water) and 656 timescale to which those claims relate. Since degradable and compostable materials 657 are typically not compatible with widely available recycling infrastructure, it is also important that the users are informed of the appropriate disposal route which in most 658 circumstances will be disposal to the residual waste stream. It is only by providing 659 accurate, unambiguous and complete guidance to the user regarding disposal that 660 the potential benefits of these novel materials can be realised without the negative 661 662 consequences that could result in inappropriate disposal as well as unintended 663 environmental consequences. For many applications in which plastic carrier bags 664 are used, perhaps durability in the form of a bag that *can* and *is* reused many times 665 presents a better alternative to degradability.

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667		References			
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