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Microplastics in Spanish Table Salt OPEN

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Marine debris is widely recognized as a global environmental problem. One of its main components, microplastics, has been found in several sea salt samples from diferent countries, indicating that sea products are irremediably contaminated by microplastics. Previous studies show very confusing results, reporting amounts of microparticles (MPs) in salt ranging from zero to 680MPs/kg, with no mention of the possible causes of such diferences. Several errors in the experimental procedures used were found and are reported in the present work. Likewise, 21 diferent samples of commercial table salt from Spain have been analyzed for MPs content and nature. The samples comprise sea salts and well salts, before and after packing. The microplastic content found was of 50–280MPs/kg salt, being polyethylene-terephthalate (PET) the most frequently found polymer, followed by polypropylene (PP) and polyethylene (PE), with no signifcant diferences among all the samples. The results indicate that even though the micro-particles might originate from multiple sources, there is a background presence of microplastics in the environment.

Marine debris (MDs) are widely recognized as a global environmental problem^{1-[3](#page-5-1)}. They produce a wide range of negative efects, not only environmental, but also economical, safety, health and cultural impact.

The materials which are most commonly found in marine debris comprise glass, metal, paper and plastic^{[4](#page-5-2)}; being plastic debris the most abundant in the marine environment^{4-[8](#page-5-3)}. They represent between 60 and 80% of the total marine debris⁹.

Plastics are relatively recent, they have existed only for around one century¹⁰, but since the development of the plastics industry, plastic items have spread around the world. Recent studies estimate that between 4.8 and 12.7 million tons of plastic waste ends up in the world's oceans every year², making them the largest plastic dumps.

Plastics are harmful to the marine environment, mainly because of their resistance to degradation. Once in the environment, and especially in the marine environment, the decomposition of plastic items occurs in an exceed-ingly long time, usually estimated between hundreds and thousands of years^{[11](#page-5-7)}. Furthermore, during this time, plastics are fragmented into small pieces, becoming plastic micro-particles (with a diameter of less than 5 mm)¹².

Many products of human consumption, such as salt, fsh… come from the sea. Marine pollution can afect these products, and microplastics can reach our organism through them. Recent studies¹³ show the presence of plastic microfber in deep-sea organisms, showing that the deep sea is already worryingly exposed to human waste.

There are several types of table salt according to their origin: sea salt, lake salt, rock salt and river or well salt. Sea salt and lake salt are obtained by evaporation, rock salt comes from the mining of a mineral rock called halite, and river or well salt is obtained from wells in non-coastal zones. Sea salt and rock salt are the most sold and consumed in Spain.

In the production of sea or lake salt, saltwater is pumped into evaporation ponds, where it is concentrated by the action of sun and wind. Afer that, the salt condenses and crystallizes on the surface of the crystallizers, where the salt is cut and collected by means of a controlled collection process. Subsequently, salt undergoes diferent physical processes before its packing in diferent containers for their multiple uses and diferent applications. Figure [1](#page-1-0) shows a scheme of the salt production process. For manufacturing river or well salt, the saline water is pumped from below ground into salt lakes, where it concentrates similarly to sea salt.

Recent studies showed that microplastics (fragments, fibers and pellets) are present in seas all over the world $14-16$ $14-16$, although only microfibers (not microbeads) where found in deep-sea organisms^{[13](#page-5-9)}. This lead to the hypothesis that sea salt might contain microplastics in it, because it is directly produced from seawater¹⁷, pointing out the importance of monitoring the presence of such contaminants in sea salts.

In the work of Yang *et al*. the amounts of microparticles (MPs, comprising fragments, fbers and pellets) found in diferent Chinese salts are in the range 7–680MPs/kg. On the contrary, an study recently published in Scientific Reports by Karami et al.¹⁸ concluded that the amount of MPs in salt of different origins is nil, in the range $0-10$ MPs/kg, without a comparison with previous studies. Surprisingly, the authors of the paper¹⁸ are

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Figure 1. Scheme of the salt manufacturing process.

trying to retain microfbers of a size of between 10 and 200 μm, approximately, in a flter whose pore size is 150 μm. Tis result is that the amount of fbers detected is extremely low, as most are expected to pass through the flter. Previous studies by Yang *et al*. [17](#page-5-12) mention amounts of fbers around 700 microparticles MP/kg (using a 5μm pore sized filter), while the amounts detected in this case are 0-10 MP/kg. The authors specifically cite the work of Yang *et al*., but do not even compare their results nor mention of the diferences found. By contrast, Karami *et al*. perform a validation of the experimental method with model samples, but in this case they used a flter whose pore size is much smaller, 8μm.

With the aim of designing a veritable method to isolate MPs and check it with samples from the Spanish salt market, in the present work several types of salt produced in diferent points of Spain have been studied. Marine and well salt have been collected to compare both types. Also the effect of the packaging process is discussed. The abundance and nature of the microplastics found have been analyzed by means of stereo microscopy and Fourier Transform Infrared Spectroscopy (FT-IR).

Results and Discussion

Number of microfibers. As a representative example, a photograph of one of the filters is shown in Fig. [2.](#page-2-0) The size of plastic fibers ranged from $30 \mu m$ to $3.5 \mu m$ in all table salt samples. No fibers smaller than $30 \mu m$ were found. The most common colors found were black, red, blue, white and transparent.

The number of particles/kg in the different types of salt is summarized in Table [1](#page-3-0) (analysis was triplicated and confdence intervals are given, data were analyzed using the fxed-efects model of analysis of variance). As can be seen, diferent brands presented no signifcant diferences in the abundance of microplastics in marine salts. There is also no significant difference between the amount of fibers found in well salt samples (ranging from 115 to 185 particles/ kg) and sea salt samples (from 50 to 280 particles/ kg).

From the data presented in Table [1,](#page-3-0) when comparing the samples 1 (before grinding) and 2 (afer grinding), both from the same manufacturer, it can be concluded that the grinding process does not infuence the amount of microplastics found in these table salts. For the well salt samples, no important diferences were found in relation to the grinding process either, i.e. between the sample 17 (before grinding) and 18 (afer grinding).

In the study, packed and un-packed salt of the same well have also been analyzed (samples 18 and 19), in order to assess the infuence of the packing process. Table [1](#page-3-0) shows that the origin of the microplastic is not related to the packaging process, since a similar microplastic content was found before and afer this process.

Figure 2. Optical microscope image of a filter after the filtration of a salt sample (The target used was $3.0 \times$).

Microfiber characterization. The identification of the fibers found was done by Fourier Transform Infrared Spectroscopy (FT-IR), which is one of the most popular methods used to confrm the composition of microplastics¹⁹.

Using this technique, several types of microplastics were identifed, being the most common ones polyethylene terephthalate (PET), polyethylene (PE) and polypropylene (PP), with a presence, respect to the total of fbers analyzed, of 83.3%, 3.3% and 6.7%, respectively. The remaining 7% corresponds to other particles that have not been identifed. Figure [3a–c](#page-4-0) shows the spectrum of some of the main types of fbers analyzed. In each case, two spectra are shown for fbers found in diferent samples, in order to compare them. As can be seen, the spectra are very similar, and the identifcation is defnite.

The abundance of the different plastics is logical, since the most commonly used plastics in the world are polyethylene, polypropylene and polyethylene-terephthalate; therefore, they are the most frequently found in the marine environment too^{3, [20](#page-5-15)[–22](#page-5-16)}. However, the relative abundance of PET is very high, accounting for 83.3% of the analyzed fibers. Yang *et al.* suggested that this might be due to the high density of PET (1.30 g/cm³) in comparison to PE (0.94 g/cm³) and PP (0.90 g/cm³), which causes it to remain with the salt during the crystallization process in the salt production.

PET is a material which is widely used in packaging, either fexible (as in plastic flms) or rigid (as in bottles). PET is also the most used polyester in the textile sector, hence it is in the form of fbers in the sea and in the global environment.

Although experimental methods are not exactly the same, the amount of fbers found in the present study should be compared with the work of Yang *et al*., in which the table salt came from supermarkets throughout China²³. Table [2](#page-4-1) shows the comparison of some data from both studies. The results found by Yang *et al.* indicate that the number of microplastics found in Chinese salts is much higher than the one in the present study, almost double in the case of sea salts; these authors also pointed out remarkable increase in the amount of fbers found in salt which come from the marine environment. In this way, Yang *et al*. indicate that the abundance of microplastics in sea salt was signifcantly higher than in lake and rock/well salts. However, this result is not found in the present work, being the microplastics content very similar in salt coming from marine and non-marine environments. Tis should be related with the pollution level and the plastic presence in the Spanish and Chinese seas. Eriksen *et al*. [24](#page-6-0) report an estimate of the total number of plastic particles in the world's ocean, indicating a very high plastic pollution in north Pacifc compared to Mediterranean sea. Yang *et al*. [23](#page-5-17) indicate that in China the sources of sea salts were from coastal waters in locations where the population density is very high, not being this true in the present study. In the present case, the data indicate that there is not a clear source of these micro-particles, but there is a background presence of the microplastics in the environment. In line with this, microplastic pollution has also been detected in honey and sugar samples²⁵ and other^{[26](#page-6-2)}, as well as in marine organisms^{[27](#page-6-3),2}

As mentioned before, the work published by Karami *et al*. [18](#page-5-13) found amount of MPs in salts in the range 0–10MPs/kg, but an experimental error made this result unessential.

Microplastics in food and risk to human health. Marine debris produces a wide variety of negative environmental, economic^{[29](#page-6-5)}, safety, health^{[30](#page-6-6)} and cultural impacts¹¹. The presence of microplastics in the sea water has been revealed as hazardous. In literature, three possible toxic efects of plastic particle have been indicated: frst due to the plastic particles themselves, second to the release of persistent organic pollutant adsorbed to the plastics³¹, and third to the leaching of additives of the plastics²⁶. Persistent organic pollutants (POPs) such as polychlorinated biphenyls (PCBs) and organochlorine pesticides are present in aquatic systems worldwide. Coastal environments contain keystone species and are a source of food supply³². Plastics might

Table 1. Number of microplastics found in the diferent types of table salt analyzed. .

absorb these contaminants from the seawater and transfer them to the sea products (fsh, salt,..). So the presence of plastics in the sea salt might pose a threat to food safety. Diferent studies have addressed the sorption capacity of plastic debris for POPs^{33[–36](#page-6-10)}. For this, authors such as Seltenrich and Bouwmeester highlighted in their research^{[26,](#page-6-2) [37](#page-6-11)} the importance of investigating the risk of the transmission of these microplastics from the food chain to humans.

According to the World Health Organization (WHO), the maximum salt intake in an adult should not exceed 5 g per day. In this way, an average consumer of Spanish salt would ingest a maximum of approximately 510 plastic particles each year. However, microplastics can reach the human organism through other type of food from sea, such as fish and mussels. There are several studies on ingestion of plastic by fish in the field^{38–41}. Foekema *et al*. concluded in a recently research that more than 80% of the fsh that ingested plastic contained only one particle, suggesting that microplastics do not seem to remain over a long period of time inside the gastrointestinal tract of fish^{[42](#page-6-14)}. However, the presence of microplastics is higher in mussels^{[43](#page-6-15), 44}, reaching concentrations of up to 178 microfibres per mussel⁴⁵. So if you compare our result with the intake of a person's microplastics through another type of food from the sea, such as fsh and mussels, it can see that 510 particles/year is not a big quantity of microplastics.

Method and Materials

Twenty-one diferent table salt samples were collected for the present study, all of them from Spanish salt producers, during September 2016–June 2017. An average package with a weight of 1 kg was chosen. All samples were collected from supermarkets, and all of them corresponded to salts manufactured in saltworks along Spain. Figure [4](#page-5-18) details the origin of each of the salt samples. Note that some of the samples (samples 17 to 21) are coming from salt producers not located in the coast, i.e., these samples are well salts. As described in Table [1,](#page-3-0) samples of diferent particle size were used. Also samples were taken afer and before packaging, in order to check for possible diferences. Mediterranean Sea and Atlantic Ocean are the coastal areas of study.

As remarked before, two brand of well salt were also analyzed, but collecting separate samples at diferent stages of the manufacturing process: before grinding, before packing and once packed (see scheme in Fig. [1](#page-1-0)), corresponding to samples 17, 18 and 19, respectively. This was done in order to elucidate the possible origin of the microplastics. Note that the well salt is also concentrated in salt lakes by evaporation, and collected by vehicles, but its origin is not marine.

Approximately 200 g of each type of salt were dissolved in 1 L of distilled water. The density of this mixture was determined (by using a pycnometer) and give an average value of 1.27 g/cm³. Then, the solution was placed in a centrifuge at 1900rpm for 1h, in order to separate the possible sand contained by the salt from the saline solution. Although the density of PET is quite similar to that of the saline solution (ca. 1.3 g/cm^3), the fact that the plastics are in form of microfibers highly decreases their apparent density⁴⁶ so all MPs are expected to remain at the upper part of the solution. Tat solution was immediately fltered through a piece of 5 μm pore size, 47 mm cellulose nitrate filter paper using a vacuum system 23 23 23 .

Figure 3. FTIR spectra of fbers found in the table salt samples: (**a**) Two PET fbers; (**b**) Two PP fbers; (**c**) A PE fber.

Table 2. Comparison of the results of the present study with Yang *et al*. (2015) study on Chinese salts. * Cellophane (CP).

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Once the solution had been fltered, the flter paper was placed into a clean Petri dish with a cover and was dried at room temperature, to later examine the total number of particles under a microscope. This procedure was repeated in triplicate for each sample, using salt from the same package.

The filters were observed under a Leica S6 D Stereozoom CLS150X microscope (range of magnification $0.63\times -4.0\times$), under polarized light. The amount of microplastics was determined by manually counting the particles contained in the flters.

In order to rule out the possibility of contamination during the sample handling process, a blank measurement was performed. The procedural blank only contained 6 particles of microplastics per filter.

Some fibers of the fibers found on the filters were randomly selected for analysis using FT-IR^{[21,](#page-5-19) 23} in a Microscope FTIR JASCO IRT-5200 with 16x Cassegrain lens and MCT detector (7000 – 600 cm⁻¹). All samples were measured in transmission. The spectrum range was set to 850–4000 cm⁻¹. The spectral resolution was 4 cm[−]¹ for all samples and the aperture size varied widely depending on the size of the fbers. All the spectra

Figure 4. Map of Spain showing the sampling points. This Figure was done using Photoshop CS6 and was created modifying an image freely available at the following web: [http://d-maps.com/carte.php?num_](http://d-maps.com/carte.php?num_car=2190&lang=es) $car = 2190\&$ lang $=$ es. The maps on this page are free for any use, even commercial.

obtained were compared to the library to identify the polymer type (NICODOM IR Libraries). The spectrum analysis followed the method of Woodall *et al*. [28](#page-6-4).

Data Availability. The datasets generated during and/or analyzed during the current study are available from the corresponding author on reasonable request.

References

- 1. Cózar, A. *et al*. Plastic debris in the open ocean. *Proceedings of the National Academy of Sciences* **111**, 10239–10244 (2014).
- 2. Jambeck, J. R. *et al*. Plastic waste inputs from land into the ocean. *Science* **347**, 768–771 (2015).
- 3. Iñiguez, M. E., Conesa, J. A. & Fullana, A. Marine debris occurrence and treatment: A review. *Renewable and Sustainable Energy Reviews* **64**, 394–402 (2016).
- 4. OSPAR. OSPAR Pilot Project on Monitoring Marine Beach Litter: Monitoring of marine litter on beaches in the OSPAR region. *OSPAR Commission, London* (2007).
- 5. Tompson, R. C., Moore, C. J., vom Saal, F. S. & Swan, S. H. Plastics, the environment and human health: current consensus and future trends. *Philosophical Transactions of the Royal Society of London B: Biological Sciences* **364**, 2153–2166 (2009).
- 6. UNEP-CAR/RCU. Marine Litter in the Wider Caribbean Region: A Regional Overview. *United Nations Environment Programme, Nairobi, p. 81*. (2008).
- 7. UNEP. Marine Litter an Analytical Overview. United Nations Environment Programme, Kenya, p. 47. (2005).
- 8. UNEP. Marine Litter: A Global Challenge. *Nairobi: UNEP*, 232 (2009).
- 9. Gregory, M. R. & Ryan, P. G. In *Marine Debris Springer Series on Environmental Management* (eds JamesM Coe & DonaldB Rogers) Ch. 6, 49–66 (Springer New York, 1997).
- 10. Gorman, M. Environmental Hazards—Marine Pollution. *ABC-CLIO Inc, Santa Barbara*. (1993).
- 11. Barnes, D. K. A., Galgani, F., Thompson, R. C. & Barlaz, M. Accumulation and fragmentation of plastic debris in global environments. *Philosophical Transactions of the Royal Society of London B: Biological Sciences* **364**, 1985–1998 (2009).
- 12. Arthur, C., Baker, J. & Bamford, H. Proceedings of the International Research Workshop on the Occurrence, Efects and Fate of Microplastic Marine Debris. *NOAA Technical Memorandum NOS-OR&R-30* (2009).
- 13. Taylor, M. L., Gwinnett, C., Robinson, L. F. & Woodall, L. C. Plastic microfbre ingestion by deep-sea organisms. *Scientifc Reports* **6**, 33997, doi[:10.1038/srep33997](http://dx.doi.org/10.1038/srep33997) (2016).
- 14. Gajšt, T., Bizjak, T., Palatinus, A., Liubartseva, S. & Kržan, A. Sea surface microplastics in Slovenian part of the Northern Adriatic. *Marine Pollution Bulletin* **113**, 392–399, doi[:10.1016/j.marpolbul.2016.10.031](http://dx.doi.org/10.1016/j.marpolbul.2016.10.031) (2016).
- 15. Russell, M. & Webster, L. In *Fate and Impact of Microplastics in Marine Ecosystems* 46–47 (Elsevier, 2017).
- 16. Isobe, A., Uchiyama-Matsumoto, K., Uchida, K. & Tokai, T. Microplastics in the Southern Ocean. *Marine Pollution Bulletin* **114**, 623–626, doi:[10.1016/j.marpolbul.2016.09.037](http://dx.doi.org/10.1016/j.marpolbul.2016.09.037) (2017).
- 17. Yang, D. *et al*. Microplastic Pollution in Table Salts from China. *Environmental Science & Technology*, doi[:10.1021/acs.est.5b03163](http://dx.doi.org/10.1021/acs.est.5b03163) (2015).
- 18. Karami, A. et al. The presence of microplastics in commercial salts from different countries. *Scientific Reports* 7, 46173, doi:[10.1038/](http://dx.doi.org/10.1038/srep46173) [srep46173](http://dx.doi.org/10.1038/srep46173),<https://www.nature.com/articles/srep46173#supplementary-information>(2017).
- 19. Song, Y. K. *et al*. A comparison of microscopic and spectroscopic identification methods for analysis of microplastics in environmental samples. *Marine Pollution Bulletin* **93**, 202–209 (2015).
- 20. Heo, N. *et al*. Distribution of small plastic debris in cross-section and high strandline on Heungnam beach, South Korea. *Ocean Sci. J.* **48**, 225–233, doi:[10.1007/s12601-013-0019-9](http://dx.doi.org/10.1007/s12601-013-0019-9) (2013).
- 21. Hidalgo-Ruz, V., Gutow, L., Tompson, R. C. & Tiel, M. Microplastics in the Marine Environment: A Review of the Methods Used for Identifcation and Quantifcation. *Environmental Science & Technology* **46**, 3060–3075, doi:[10.1021/es2031505](http://dx.doi.org/10.1021/es2031505) (2012).
- 22. Martins, J. & Sobral, P. Plastic marine debris on the Portuguese coastline: A matter of size? *Marine Pollution Bulletin* **62**, 2649–2653, doi[:10.1016/j.marpolbul.2011.09.028](http://dx.doi.org/10.1016/j.marpolbul.2011.09.028) (2011).
- 23. Yang, D. *et al*. Microplastic pollution in table salts from China. *Environmental Science & Technology* **49**, 13622–13627 (2015).
- 24. Eriksen, M., Lebreton, L. C. M., Carson, H. S., Tiel, M. & Moore, C. J. Plastic Marine Pollution Global Dataset. *PLoS ONE* **9**, 1–15, doi[:10.6084/m9.fgshare.1015289.v1](http://dx.doi.org/10.6084/m9.figshare.1015289.v1) (2014).
- 25. Liebezeit, G. & Liebezeit, E. Non-pollen particulates in honey and sugar. *Food Additives and Contaminants - Part A Chemistry, Analysis, Control, Exposure and Risk Assessment* **30**, 2136–2140, doi[:10.1080/19440049.2013.843025](http://dx.doi.org/10.1080/19440049.2013.843025) (2013).
- 26. Bouwmeester, H., Hollman, P. C. & Peters, R. J. Potential health impact of environmentally released micro-and nanoplastics in the human food production chain: experiences from nanotoxicology. *Environmental Science & Technology* **49**, 8932–8947 (2015).
- 27. Van Cauwenberghe, L., Vanreusel, A., Mees, J. & Janssen, C. R. Microplastic pollution in deep-sea sediments. *Environ. Pollut.* **182**, 495–499, doi:[10.1016/j.envpol.2013.08.013](http://dx.doi.org/10.1016/j.envpol.2013.08.013) (2013).
- 28. Woodall, L. C. et al. The deep sea is a major sink for microplastic debris. *Royal Society Open Science* 1, doi[:10.1098/rsos.140317](http://dx.doi.org/10.1098/rsos.140317) (2014).
- 29. Legat, A., French, V. & McDonough, N. An economic perspective on oceans and human health. *Marine Biological Association of the United Kingdom. Journal of the Marine Biological Association of the United Kingdom* **96**, 13–17, doi[:10.1017/S0025315415001319](http://dx.doi.org/10.1017/S0025315415001319) (2016).
- 30. Depledge, M. H. *et al*. Changing Views of the Interconnections Between the Oceans and Human Health in Europe. *Microbial Ecology* **65**, 852–859, doi:[10.1007/s00248-012-0173-0](http://dx.doi.org/10.1007/s00248-012-0173-0) (2013).
- 31. Iñiguez, M. E., Conesa, J. A. & Fullana, A. Pollutant content in marine debris and characterization by thermal decomposition. *Marine Pollution Bulletin*, doi:[10.1016/j.marpolbul.2017.02.022](http://dx.doi.org/10.1016/j.marpolbul.2017.02.022) (2017).
- 32. Madeira, D., Narciso, L., Diniz, M. S. & Vinagre, C. Synergy of environmental variables alters the thermal window and heat shock response: An experimental test with the crab Pachygrapsus marmoratus. *Marine Environmental Research* **98**, 21–28, doi[:10.1016/j.](http://dx.doi.org/10.1016/j.marenvres.2014.03.011) [marenvres.2014.03.011](http://dx.doi.org/10.1016/j.marenvres.2014.03.011) (2014).
- 33. Lee, H., Shim, W. J. & Kwon, J.-H. Sorption capacity of plastic debris for hydrophobic organic chemicals. *Sci. Total Environ.* **470–471**, 1545–1552, doi:[10.1016/j.scitotenv.2013.08.023](http://dx.doi.org/10.1016/j.scitotenv.2013.08.023) (2014).
- 34. Rochman, C. M., Manzano, C., Hentschel, B. T., Simonich, S. L. M. & Hoh, E. Polystyrene Plastic: A Source and Sink for Polycyclic Aromatic Hydrocarbons in the Marine Environment. *Environmental Science & Technology* **47**, 13976–13984, doi[:10.1021/es403605f](http://dx.doi.org/10.1021/es403605f) (2013).
- 35. Fisner, M., Taniguchi, S., Moreira, F., Bícego, M. C. & Turra, A. Polycyclic aromatic hydrocarbons (PAHs) in plastic pellets: Variability in the concentration and composition at diferent sediment depths in a sandy beach. *Marine Pollution Bulletin* **70**, 219–226, doi:[10.1016/j.marpolbul.2013.03.008](http://dx.doi.org/10.1016/j.marpolbul.2013.03.008) (2013).
- 36. Krüger, O., Kalbe, U., Meißner, K. & Sobottka, S. Sorption efects interfering with the analysis of polycyclic aromatic hydrocarbons (PAH) in aqueous samples. *Talanta* **122**, 151–156, doi[:10.1016/j.talanta.2014.01.038](http://dx.doi.org/10.1016/j.talanta.2014.01.038) (2014).
- 37. Seltenrich, N. New link in the food chain? Marine plastic pollution and seafood safety. *Environmental health perspectives* **123**, A34–A41 (2015).
- 38. Boerger, C. M., Lattin, G. L., Moore, S. L. & Moore, C. J. Plastic ingestion by planktivorous fshes in the North Pacifc Central Gyre. *Marine Pollution Bulletin* **60**, 2275–2278 (2010).
- 39. Possatto, F. E., Barletta, M., Costa, M. F., do Sul, J. A. I. & Dantas, D. V. Plastic debris ingestion by marine catfsh: an unexpected fsheries impact. *Marine Pollution Bulletin* **62**, 1098–1102 (2011).
- 40. Lusher, A., McHugh, M. & Tompson, R. Occurrence of microplastics in the gastrointestinal tract of pelagic and demersal fsh from the English Channel. *Marine Pollution Bulletin* **67**, 94–99 (2013).
- 41. Neves, D., Sobral, P., Ferreira, J. L. & Pereira, T. Ingestion of microplastics by commercial fsh of the Portuguese coast. *Marine Pollution Bulletin* **101**, 119–126, doi:[10.1016/j.marpolbul.2015.11.008](http://dx.doi.org/10.1016/j.marpolbul.2015.11.008) (2015).
- 42. Foekema, E. M. *et al*. Plastic in north sea fsh. *Environmental Science & Technology* **47**, 8818–8824 (2013).
- 43. Van Cauwenberghe, L. & Janssen, C. R. Microplastics in bivalves cultured for human consumption. *Environmental Pollution* **193**, 65–70 (2014).
- 44. Van Cauwenberghe, L., Claessens, M., Vandegehuchte, M. B. & Janssen, C. R. Microplastics are taken up by mussels (Mytilus edulis) and lugworms (Arenicola marina) living in natural habitats. *Environmental Pollution* **199**, 10–17 (2015).
- 45. Mathalon, A. & Hill, P. Microplastic fbers in the intertidal ecosystem surrounding Halifax Harbor, Nova Scotia. *Marine Pollution Bulletin* **81**, 69–79 (2014).
- 46. Branca, D. L. Uniformity of low density fibrous structures and the effects of manufacturing processes on apparent density 1446161 thesis, State University of New York College of Environmental Science and Forestry, (2007).

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Author Contributions

Maria E. Iñiguez did the experimental work under the supervision of Juan A. Conesa and Andres Fullana. Maria Iñiguez wrote the main manuscript text and prepared Figures 2 to 5. Juan A. Conesa prepared Figures 1 and 6. All authors reviewed the manuscript.

Additional Information

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