



The Potential Role of Marine Fungi in Plastic Degradation – A Review

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Plastic debris has been accumulating in the marine realm since the start of plastic mass production in the 1950s. Due to the adverse effects on ocean life, the fate of plastics in the marine environment is an increasingly important environmental issue. Microbial degradation, in addition to weathering, has been identified as a potentially relevant breakdown route for marine plastic debris. Although many studies have focused on microbial colonization and the potential role of microorganisms in breaking down marine plastic debris, little is known about fungi-plastic interactions. Marine fungi are a generally understudied group of microorganisms but the ability of terrestrial and lacustrine fungal taxa to metabolize recalcitrant compounds, pollutants, and some plastic types (e.g., lignin, solvents, pesticides, polyaromatic hydrocarbons, polyurethane, and polyethylene) indicates that marine fungi could be important degraders of complex organic matter in the marine realm, too. Indeed, recent studies demonstrated that some fungal strains from the ocean, such as *Zalerion maritimum* have the ability to degrade polyethylene. This mini-review summarizes the available information on plastic-fungi interactions in marine environments. We address (i) the currently known diversity of fungi colonizing marine plastic debris and provide (ii) an overview of methods applied to investigate the role of fungi in plastic degradation, highlighting their advantages and drawbacks. We also highlight (iii) the underestimated role of fungi as plastic degraders in marine habitats.

Keywords: fungi, marine plastic debris, marine pollution, plastic degradation, biodegradation

OCEAN PLASTIC POLLUTION

Plastics are man-made materials of mostly petrochemical origin – so-called conventional plastics that are commonly considered as non-biodegradable (Wayman and Niemann, 2021). Most conventional plastics comprise a carbon-carbon backbone, e.g., polyethylene (PE), polypropylene (PP) and polystyrene (PS), while other plastics feature heteroatoms, e.g., polyethylene terephthalate

(PET), polyurethane (PU) or polyamides (PA, Nylon). Conventional plastics are used in nearly all industrial sectors, most importantly packaging, construction and transport. The global annual production of plastics reached 359 Mt in 2018 (PlasticsEurope, 2020). Recently, biopolymer alternatives, based on renewable sources (e.g., starch and cellulose) have been developed, but their global market share is still small. In 2020, the global production of bioplastics amounted to approximately 2 Mt (Bioplastics Market Data update, 2020). Among the plastic types produced from renewable carbon sources, approximately half are biodegradable, while the second half is non-biodegradable.

A substantial fraction of plastic waste is mismanaged (Geyer et al., 2017), and it has been estimated that this fraction represented an average of 80 Mt (60 to 99 Mt) accumulated in total (Lebreton and Andrady, 2019). Mismanaged plastic waste ends up in all ecosystem compartments, such as soils (Chae and An, 2018), lakes (Mason et al., 2016) and rivers (Lebreton et al., 2017). Thus, freshwater bodies function as a transport vector for plastic ultimately reaching the ocean though the magnitude of this transport mechanism is discussed controversially (Weiss et al., 2021). In addition, plastic litter can be airborne, thus, the atmospheric deposition of plastic litter is a potentially important source for the ocean's plastic budget (Liss, 2020). Indeed, plastic litter has emerged as a major pollution issue in marine environments (Eriksen et al., 2014; UNEP, 2014; van Sebille et al., 2015; Wayman and Niemann, 2021). Plastic debris forms the most abundant component of marine litter, accounting for up to 95% of waste found on shorelines, ocean surface water and the seafloor (Galgani et al., 2015). In fact, sedimented plastic litter might become a stratigraphic marker horizon of the Anthropocene epoch (Zalasiewicz et al., 2016). It has been estimated that 0.1% (Cózar et al., 2015) to up to 4.6% (Jambeck et al., 2015) of the global plastic production enters the ocean. As a result, the total accumulated plastic waste in the ocean might have reached 320 Mt in the year 2015 (Wayman and Niemann, 2021).

Plastic floating at the ocean surface is typically dominated by PE and PP, roughly in accordance with global production figures (Erni-Cassola et al., 2019). However, plastic debris in the ocean varies in size: macroplastics (>5 mm), microplastics (1 μm – 5 mm) and nanoplastics (<1 μm) (Wayman and Niemann, 2021). Several studies have tried to quantify the different size fractions to estimate the total concentration of plastics and the fate of it in marine environments. For example, in the Mediterranean Sea, the density of floating microplastic was about 2.5×10^5 items per km^2 (Cózar et al., 2015). The abundance of floating plastic in the ocean is usually determined by surface trawling with nets (typically > 300 μm mesh size). Consequently, these methods discriminate against smaller size classes, which could account for an important fraction of the floating plastic budget (Poulain et al., 2019). Furthermore, surface trawls do not account for submerged and sedimented plastics debris (Woodall et al., 2014). Finally, the distribution of floating plastic is not homogenous, as large quantities concentrate in subtropical ocean gyres and enclosed basins, making balanced sampling efforts over large areas difficult (Lindeque et al., 2020).

MICROBIAL BIOFILMS ON MARINE PLASTIC DEBRIS

Rapid microbial colonization occurs on any available material that ends up in the ocean, whether it is of natural or synthetic origin (such as plastic debris). From a microbial perspective, transiting from pelagic to a particle-attached lifestyle provides advantages; e.g., better access to nutrients, and protection against UV exposure and grazing (de Carvalho, 2018). Several studies have investigated the composition of microbial communities living on plastic materials in different marine environments (Zettler et al., 2013; Amaral-Zettler et al., 2015; Eich et al., 2015; Oberbeckmann et al., 2016; Debroas et al., 2017; Dussud et al., 2018; Ogonowski et al., 2018; Miao et al., 2019; Dudek et al., 2020; Krause et al., 2020; Vaksmaa et al., 2021b). Some of these reported a difference in community structure when comparing microbes in biofilms on plastics to those in the surrounding seawater or those attached to natural surfaces. A detailed overview of this matter is compiled in Wright et al. (2020). However, most studies applied amplicon sequencing of the 16S rRNA gene, and focused on bacterial and archaeal communities in biofilms attached to the plastic debris. In contrast, fungal community composition on plastic in the marine environment has, until now, been investigated seldomly (Table 1).

A BRIEF INTRODUCTION TO THE FUNGAL KINGDOM

The fungal kingdom constitutes a major lineage within the domain Eukarya and diverged from a common ancestor with the animals >800 million years ago (Parfrey et al., 2011; Chang et al., 2015). Fungi exhibit highly diverse lifestyles and can cope with different redox conditions. The majority of fungi seemingly prefer oxic environments, however some fungal species inhabit oxygen minimum zones (Stief et al., 2014). Furthermore, anaerobic fungi (mainly studied in gut microbiomes) have been identified (Gruninger et al., 2014; Mura et al., 2019). Morphologically, fungi may be unicellular (e.g., yeasts and cells with a flagellum such as zoospores), filamentous (e.g., molds and mushrooms) or dimorphic (i.e., they exists in two forms, yeast-like single cells or hyphae forming) (Boyce and Andrianopoulos, 2015). They can be found as free-living organisms, in mutualistic symbiotic associations (such as those forming mycorrhizas, lichens, in gut microbiomes) (Watkinson, 2016) or as parasitic pathogens of several plants and animals, including humans (Szabo and Bushnell, 2001). Fungi are ubiquitous and occur throughout terrestrial, freshwater and marine environments (Sutherland, 1916; Raghukumar, 2017; Walker et al., 2017; Gladfelter et al., 2019).

Taxonomically, nineteen major fungal phyla are recognized: Aphelidiomycota, Ascomycota, Basidiobolomycota, Basidiomycota, Blastocladiomycota, Calcarisporiellomycota, Caulochytriomycota, Chytridiomycota, Entomophthoromycota, Entorrhizomycota, Glomeromycota, Kickxellomycota, Monoblepharomycota, Mortierellomycota, Mucoromycota, Neocallimastigomycota, Olpidiomycota, Rozellomycota, and

TABLE 1 | Studies on the diversity of fungal communities on microplastics in marine environments.

Source	Polymer	Location	Primers	Phyla detected
Oberbeckmann et al., 2016	PET bottles	North Sea	1391F – 1795R	Ascomycota, Basidiomycota and Chytridiomycota
Kettner et al., 2017, 2019	PE, PS	Baltic Sea and Warnow river	Eu565F – Eu918R	Ascomycota, Basidiomycota, Chytridiomycota and Rozellomycota
De Tender et al., 2017	PE	North Sea (coast and offshore)	fITS7bis (adapted) – ITS4NGSr	Ascomycota, Basidiomycota, Chytridiomycota, Glomeromycota, and Mucoromycota
Kirstein et al., 2018	HDPE, LDPE, PP, PS and PET	North Sea (flow-through system)	Eu565F – Eu918R	Chytridiomycota
Lacerda et al., 2020	PE, Nylon, PU, PP and PS	Western south Atlantic and Antarctic Peninsula	1391F – EukB, TAReuk454FWD1 – TAReukREV3, ITS1f – ITS4 and gITS7 – ITS4	Aphelidomycota, Ascomycota, Basidiomycota, Blastocladiomycota, Chytridiomycota, Rozellomycota, Mucoromycota and Zoopagomycota

All studies applied next generation sequencing by using the Illumina MiSeq platform.

Zoopagomycota (Wijayawardene et al., 2020). However, the phylogeny and taxonomy of the fungal kingdom has been a matter of debate (Hibbett et al., 2007; Spatafora et al., 2016, 2017). Some fungal phyla are rarely sampled and our understanding of the deeply branching groups is insufficient (Naranjo-Ortiz and Gabaldón, 2019). Lately, the number of available fungal genomes has been increasing. But these new data showed that complex inter-species relationships occur, such as introgression and hybridization (Gabaldón, 2020), that blur the definition of a species. The estimated total number of fungal species could range from 2 to 4 million (Hawksworth and Lücking, 2017), but only ~100000 have been described (Wu et al., 2019) of which only ~1100 species originate from marine environments (Amend et al., 2019).

Recently, an online database, www.marinefungi.org, containing marine fungal species was created (Jones et al., 2019). Many fungal species are found in both, terrestrial and marine environments (Gladfelter et al., 2019). It has been proposed that early diverging fungi, Chytridiomycota and Rozellomycota, originate from aquatic environments (Berbee et al., 2017, 2020). These phyla also have members found in the present-day ocean. On the other hand, fungi from the Ascomycota and Basidiomycota phyla, commonly present in marine environments, are proposed to originate from terrestrial ancestors. Because of this difference in origin (terrestrial vs. aquatic), a clear definition of “marine fungus” is challenging. The recent definition suggests that a marine fungus is “any fungus that is recovered repeatedly from marine habitats and: (1) is able to grow and/or sporulate (on substrata) in marine environments; or (2) forms symbiotic relationships with other marine organisms; or (3) is shown to adapt and evolve at the genetic level or is metabolically active in marine environments” (Pang et al., 2016). Marine fungi are found ubiquitously in the ocean (Comeau et al., 2016; Tisthammer et al., 2016; Raghukumar, 2017) including coastal environments (Taylor and Cunliffe, 2016; Picard, 2017; Banos et al., 2020), mangroves (Hyde and Lee, 1995; Alias et al., 2010; Pang et al., 2010; Lee et al., 2019), the water column (Richards et al., 2015; Tisthammer et al., 2016; Morales et al., 2019) and sediments (Kohlmeyer and Kohlmeyer, 1979a; Khudyakova et al., 2000;

Mouton et al., 2012; Tisthammer et al., 2016). Marine fungi have also been detected in extreme marine habitats such as the deep sea biosphere (Orsi et al., 2013; Rédou et al., 2015), the Arctic (Rämä et al., 2017; Hassett et al., 2019) and oxygen minimum zones (Cathrine and Raghukumar, 2009; Jebaraj et al., 2010; Manohar et al., 2015).

Metabolically, fungi participate in different biogeochemical processes and occupy a plethora of ecological niches. Fungi play a major role in decomposing recalcitrant substrates, making them an integral part of food web structures, contributing to carbon cycling and nutrient regeneration in, at least, terrestrial environments. For example, saprophytic fungi accelerate carbon and nitrogen cycling and symbiotic fungi such as mycorrhiza networks, enhance primary production in symbionts and hosts (Lindahl et al., 2007; Walder et al., 2012). Similar to terrestrial systems, parasitic fungi in aquatic environments were found to have a great impact on pelagic food webs and thus biogeochemical cycling (Jobard et al., 2010; Sime-Ngando, 2012). Also some symbiotic interactions are known from the marine environment, for example, with phytoplankton seaweeds and sponges (Rasconi et al., 2011; Richards et al., 2012; Webster and Taylor, 2012; Du et al., 2019). Similar to terrestrial systems, fungi in coastal and surface marine environments were found to degrade wood (lignin, cellulose and hemicellulose) (Bucher et al., 2004) and remains of marine animals (Kohlmeyer and Kohlmeyer, 1979b,c; Raghukumar, 2017). Marine fungi were furthermore found to degrade complex components of algae such as agar (in laboratory conditions; Balabanova et al., 2018) and fungi dominate the microbial composition of bathypelagic marine snow where they might play the role as saprotrophs (Bochdansky et al., 2017). Fungi contribute to different nitrogen cycle processes such as nitrification (Falih and Wainwright, 1995) and denitrification (Shoun et al., 1992; Stief et al., 2014; Maeda et al., 2015) in both, terrestrial and the marine environment (Cathrine and Raghukumar, 2009). Marine fungi mobilize metals by excreting siderophores and act as bio-sorbents for some metals, thereby alleviating metal toxicity (Taboski et al., 2005; Vala, 2010). These processes influence the cycles of several elements among them: Fe, Mn, Hg, Ni, Zn, Ag, Cu, Cd, and Pb (Gadd, 2004). Finally, some

fungi break down rocks and minerals to harvest nutrients (Ortega-Morales et al., 2016).

LIFE ON PLASTIC IS FANTASTIC: FUNGAL COLONIZATION OF MARINE PLASTIC DEBRIS

Fungi in Marine Environments as Part of Marine Plastic Debris Associated Biofilms

Marine fungi are generally understudied, which is also reflected by the relatively limited number of studies targeting fungi on marine plastic debris (MPD) (Jacquin et al., 2019). Oberbeckmann et al. (2016) conducted a seasonal comparison of microbial communities including both bacteria and fungi on submerged PET bottles, glass slides, and seawater in the North Sea. Fungal communities were represented by Ascomycota, Basidiomycota and Chytridiomycota. The study showed that the PET-attached eukaryotic communities (fungi among them) varied significantly with season and location. Kettner et al. (2017, 2019) conducted exposure experiments with PE and PS in the Baltic Sea and the river Warnow and investigated colonization of these plastic surfaces by fungi. The fungal genus *Chytridium*, as well as fungi-like Rhinosporideaceae, *Rhizidiomyces*, and *Pythium* taxa, had a high read count on both plastic types at both locations. However, a substantial number of sequences was assigned as unclassified fungi. The fungal community composition was significantly influenced by location but not polymer type. Furthermore, the alpha diversity was significantly lower for PE and PS compared to the surrounding water and wood particles. De Tender et al. (2017) conducted a 44-week experiment during which PE plastic sheets and dolly ropes were weighed down close to the sediment at a harbor and an offshore location in the North Sea. Similar to the Baltic Sea studies by Kettner et al. (2017, 2019), they found that the Ascomycota were highly abundant followed by a smaller fraction of Basidiomycota and Mucoromycota. Furthermore, a minor fraction of members of Ascomycota from the Lecanoromycetes class (*Physconia*, *Candelariella*, *Caloplaca*) were identified. Analyses on the beta diversity of the fungal community composition showed statistically significant effects of sample type (natural substrate vs. plastic polymers), environment, and exposure time. Some of the detected species were previously identified as potential PE degraders in terrestrial environments: *Cladosporium cladosporioides* (Bonhomme et al., 2003) and *Fusarium redolens* (Albertsson, 1980; Albertsson and Karlsson, 1990).

Recently, Lacerda et al. (2020) studied the fungal diversity associated with plastics in the surface waters of the Western South Atlantic and the Antarctic Peninsula by investigating three different molecular marker genes for fungal identification: ITS2 (Internal transcribed spacer), the variable regions V4 and V9 of the 18S rRNA gene sequence. To date, this is the only study aiming to resolve fungal diversity on plastic using multiple marker genes. Across the tested marker genes, a total of 64 different fungal orders were associated with plastics. The primers

targeting the 18S rRNA gene (V4 and V9 regions) were able to detect a higher number of Chytridiomycota. Some taxa that were totally omitted by the ITS2 marker were detected, such as Rozellomycota, Zoopagomycota, Aphelidomycota, and Blastocladiomycota. Across all samples, the genus *Aspergillus* was the most abundant. Some of its OTUs were assigned at the species level, identifying *A. vitricola*, *A. restrictus*, and *A. wentii*. None of the identified strains were previously reported as plastic degraders, although other *Aspergillus* species have been shown to be able to oxidize different plastic types (Table 2). Lacerda and colleagues also reported on fungal taxa, such as Aphelidomycota, Zoopagomycota, Mucoromycota, and Blastocladiomycota, that had previously not been detected to colonize plastic in the marine environment.

In contrast to environmental studies, Kirstein et al. (2018) studied the microbial composition formed on microplastics under controlled laboratory conditions. Their set-up consisted of a flow-through system with North Sea water in which HDPE (High Density Polyethylene), LDPE (Low Density Polyethylene), PP, PS and PET were incubated in the dark. The 18S rRNA gene sequencing analysis of the eukaryotic community of biofilms revealed that the highest fungal read abundances belonged to Chytridiomycota (up to 3% of sequences on PET). This can be explained by this taxa's dominance throughout aquatic environments (Comeau et al., 2016) or the compatibility of biofilm presence on PET and chytrid's parasitic lifestyle.

Plastic Biodegradation Potential of Fungi

Biodegradation is the degradation of compounds and substrates mediated by living organisms, most commonly microorganisms. The soluble products of biodegradation (typically low molecular weight compounds) are absorbed or assimilated by the microorganisms. The biodegradation can be partial or complete. Complete biodegradation results in the formation of CO₂ and is also referred to as biomineralization. On the other hand, degradation of organic matter without a terminal electron acceptor, conditions that are countered in some reduced environments, leads to the formation of CH₄ and/or other short-chain hydrocarbons. In natural environments, biodegradation is mediated by enzymes or by other compounds (such as acids and peroxides), secreted by microorganisms.

Fungi are able to degrade synthetic compounds; e.g., persistent organic pollutants (POPs) (Singleton, 2001), polycyclic aromatic hydrocarbons (PAHs) (Cerniglia and Sutherland, 2001), benzene, toluene, ethylbenzene and xylenes (BTEX compounds) (Buswell, 2001) and pesticides (Pinto et al., 2012). The metabolic versatility of fungi and ability to degrade complex compounds indicates that biodegradation of plastics in the environment could be a potential metabolic trait of some fungi (Vaksmas et al., 2021a). To date, some plastic degrading fungi have indeed been identified mostly from the Ascomycete phylum to which also *Aspergillus*, *Fusarium*, and *Penicillium* belong (Table 2 *Fusarium sp.* and *F. oxysporum* isolated from soil provoked weight loss of Nylon, PE, and PU) (Tachibana et al., 2010; Raghavendra et al., 2016). *F. solani* from a collection hydrolyzed PET to terephthalic acid (TPA) via a cutinase (Ronkvist et al., 2009). Furthermore, several strains belonging to *Penicillium* are considered as potential

TABLE 2 | Selection of fungal strains that showed plastic degradation potential.

Strains	References	Environment of isolation	Polymer	Degradation assessment technique	Incubation time experiment	Main observed results
<i>Alternaria alternata</i>	Ameen et al., 2015	Mangrove	LDPE	Weight loss, SEM, enzyme activity assays, quantification of CO ₂	28 days	Increased biomass in culture with LDPE, CO ₂ emission and increased production of laccase, MnP and lignin peroxidase
<i>Aspergillus caespitosus</i>	Ameen et al., 2015	Mangrove	LDPE	Weight loss, SEM, enzyme activity assays, quantification of CO ₂	28 days	Increased biomass in culture with LDPE, CO ₂ emission and increased production of laccase and MnP
<i>Aspergillus flavus</i>	Alshehrei, 2017	Seawater	PE	Weight loss, tensile strength, SEM, FTIR	30 days	16.2% weight loss of polyethylene
<i>Aspergillus flavus</i>	Deepika and Madhuri, 2015	Soil from waste disposal site	PE	Halo test, weight loss	180 days	Reduction of 16% in molecular weight
<i>Aspergillus flavus</i>	Zhang et al., 2020	Guts of wax moth <i>Galleria mellonella</i>	HDPE	Halo test, HT-GPC, FTIR	28 days	Decreased Mn. Carbonyls groups FTIR and two laccase-like multicopper oxidases (LMCOs) genes, afla_006190 and afla_053930, displayed up-regulation
<i>Aspergillus flavus</i> ITCC no. 6051	Mathur and Prasad, 2012	Soil from waste disposal site	PU	Weight loss, SEM, FTIR, and thermogravimetric analysis, enzyme activity assay	30 days	Weight loss of PU, detection of PU degradation via FTIR and production of esterase
<i>Aspergillus flavus</i> VRKPT2	Sangeetha Devi et al., 2015	Coastal area of gulf of Mannar	HDPE	SEM, FTIR, weight loss, total protein content measurement	30 days	Weight loss of 8.5 ± 0.1%
<i>Aspergillus fumigatus</i>	Alshehrei, 2017	Seawater	PE	Weight loss, tensile strength, SEM, FTIR	30 days	20.5% weight loss of polyethylene
<i>Aspergillus fumigatus</i>	Zahra et al., 2010	Landfill soil	LDPE	UV treated LDPE, SEM, FTIR	100 days	Molecular weight decreases and use of UV LDPE as sole Carbon source and detection of structural changes by FTIR
<i>Aspergillus fumigatus</i>	Raghavendra et al., 2016	Soil from waste disposal site	PU and LDPE UV irradiated for 50 h	Halo test, weight loss	90 days	Weight loss for PE and PU after 90 days and loss of tensile strength
<i>Aspergillus glaucus</i>	Kathiresan, 2003	Mangrove soil	PE	Weight loss	30 days	Weight loss of 28.8 ± 2.4%
<i>Aspergillus niger</i>	Alshehrei, 2017	Seawater	PE	Weight loss, tensile strength, SEM, FTIR	30 days	19.5% weight loss of PE
<i>Aspergillus niger</i>	Raghavendra et al., 2016	Soil from waste disposal site	PU and LDPE UV irradiated for 50 h	Halo test, weight loss	90 days	Weight loss for PE and PU after 90 days and loss of tensile strength
<i>Aspergillus niger</i>	Deepika and Madhuri, 2015	Soil from waste disposal site	PE	Halo test, weight loss	180 days	Reduction of 26% in Mn
<i>Aspergillus niger</i>	Kathiresan, 2003	Mangrove soil	PE	Weight loss	30 days	Weight loss of 17.4 ± 2%
<i>Aspergillus niger</i> (ITCC no. 6052)	Mathur et al., 2011	Soil from waste disposal site	HDPE	Weight loss, tensile strength, SEM, FTIR	30 days	Reduction of 3.44% in Mn and 61% reduction in tensile strength
<i>Aspergillus nomius</i>	Munir et al., 2018	Soil from waste disposal site	LDPE	Weight loss, tensile strength	45 days	Weight loss of 6.63% and tensile strength reduction of 40%
<i>Aspergillus oryzae</i>	Muhonja et al., 2018	Soil from waste disposal site	LDPE	Weight loss, FTIR analysis	112 weeks	Weight loss of 36.4 ± 5.53% and degradation products detection by FTIR
<i>Aspergillus sp.</i>	Pramila and Ramesh, 2011	Seawater	LDPE	SEM, Quantification of CO ₂	7 to 17 days	Visual signs of degradation via SEM and about 4g ^{-L} of CO ₂ produced
<i>Aspergillus sp.</i>	Osman et al., 2018	Soil from waste disposal site	PU	Weight loss, quantification of CO ₂ , SEM, FTIR, DSC	28 days	15–20% of weight loss. Change in melting temperature and detection of degradation products via FTIR
<i>Aspergillus sp.</i> in co-culture with <i>Lysinibacillus xylanilyticus</i> XDB9	Esmaeili et al., 2013	Landfill soils	UV and non-UV irradiated LDPE	Tensile strength, SEM, FTIR, CO ₂ measurements	126 days	Carbon dioxide measurements: biodegradation 7.6 and 8.6% of mineralization for the non-UV irradiated and UV irradiated LDPE respectively after 126 days vs. 29.5 and 15.8% in presence of co-culture
<i>Aspergillus sydowii</i>	Sangale et al., 2019	Mangrove dumpsite	PE	Weight loss, tensile strength, SEM, FTIR	60 days	Cracks and holes visible by SEM, FTIR analysis, weight loss of 37.94 ± 3.06% (pH = 7) and tensile strength reduction
<i>Aspergillus terreus</i>	Sangale et al., 2019	Mangrove dumpsite	PE	Weight loss, tensile strength, SEM, FTIR	60 days	Cracks and holes visible by SEM, FTIR analysis, weight loss of 41.82 ± 5.47% (pH = 9.5) and tensile strength reduction
<i>Aspergillus terreus</i>	Alshehrei, 2017	Seawater	PE	Weight loss, tensile strength, SEM, FTIR	30 days	21.8% weight loss of polyethylene
<i>Aspergillus terreus</i>	Ameen et al., 2015	Mangrove	LDPE	Weight loss, SEM, enzyme activity assays, quantification of CO ₂	28 days	Increased biomass in culture with LDPE, CO ₂ emission and increased production of laccase, MnP and lignin peroxidase
<i>Aspergillus terreus</i>	Zahra et al., 2010	Soil from waste disposal site	LDPE	UV treated LDPE, SEM, FTIR	100 days	Molecular weight decreases and use of UV LDPE as sole carbon source and detection of structural changes by FTIR

(Continued)

TABLE 2 | (Continued)

Strains	References	Environment of isolation	Polymer	Degradation assessment technique	Incubation time experiment	Main observed results
<i>Aspergillus terreus</i> MF12	Balasubramanian et al., 2014	Soil with PE wastes	HDPE was pretreated by physical (heat and UV), chemical (citric acid and KMnO ₄ /HCl), and biological (microbial) treatments in different combinations	SEM, GC-MS, weight loss and FTIR	30 days	Highest degradation rates for UV treated PE
<i>Aspergillus tubingensis</i>	Khan et al., 2017	PU buried in soil	PU	SEM, tensile strength and ATR-FTIR	20 days	SEM surface cracking, erosion, pore formation or loss in tensile strength and ATR-FTIR detected degradation products
<i>Aspergillus tubingensis</i> VRKPT1	Sangeetha Devi et al., 2015	Coastal area of gulf of Mannar	HDPE	SEM, FTIR, weight loss, total protein content (alkaline hydrolysis treatment)	30 days	Weight loss of 6 ± 0.2%
<i>Bjerkandera adusta</i>	Friedrich et al., 2007	Collection	Nylon-6	Halo test, SEM, DSC, HPLC, Enzyme activity assay	60 days	Decrease in number Mn and production of MnP in presence of Nylon
<i>Cladosporium cladosporioides</i>	Brunner et al., 2018	Shoreline of lake Zurich	PU	Halo test	several days	Halos indicating potential plastic usage
<i>Cladosporium pseudocladosporioides</i> strain T1.PL.1	Álvarez-Barragán et al., 2016	Soil	PU (Impranil)	Halo test, SEM, FTIR, GCMS, enzyme activity assay	14 days	detection of PU degradation via FTIR and GCMS. Production of esterase in presence of PU
<i>Eupenicillium hirayamae</i>	Ameen et al., 2015	Mangrove	LDPE	Weight loss, SEM, enzyme activity assays, quantification of CO ₂	28 days	Increased biomass in culture with LDPE, CO ₂ emission and increased production of laccase and MnP
<i>Fusarium oxysporum</i>	Raghavendra et al., 2016	Soil from waste disposal site	PU and LDPE UV irradiated for 50 h	Halo test, weight loss	90 days	Weight loss for PE and PU after 90 days and loss of tensile strength
<i>Fusarium oxysporum</i>	Nimchua et al., 2007	Terrestrial environment	PET	Enzyme activity assay and increase of hydrophilicity detection	7 days	Increase of hydrophobicity and release of TPA from PET in presence of esterase
<i>Fusarium solani</i>	Ronkvist et al., 2009	Collection	PET (low crystallinity)	Weight loss, SEM, DSC and HPLC	4 days	Degradation of PET into TPA via cutinase (named FsC) and 5% of film weight loss
<i>Fusarium sp.</i>	Tachibana et al., 2010	Nylon 4 films buried in composted soil	Nylon-4	Weight loss, Biochemical oxygen demand (BOD), NMR, MALDI-TOF and SEM	35 days	Decreased average weight and SEM evidence
<i>Gloeophyllum trabeum</i>	Krueger et al., 2015	Collection	PS (polystyrene sulfonate)	Size exclusion chromatography (SEC)	20 days	Depolymerization of up to 50% reduction in Mn
<i>Lasioidiplodia crassispota</i>	Raghavendra et al., 2016	Soil from waste disposal site	PU and LDPE UV irradiated for 50 h	Halo test, weight loss	90 days	Weight loss for PE and PU after 90 days and loss of tensile strength
<i>Lasioidiplodia theobromae</i>	Sheik et al., 2015	Terrestrial environment	LDPE and PP (Gamma irradiated)	Weight loss, DSC, SEM, FTIR, enzyme activity assay	90 days	production laccase, weight loss in PP and PE and detection of degradation products via FTIR
<i>Leptosphaeria sp.</i>	Brunner et al., 2018	Shoreline of lake Zurich	PE, PU	Halo test	several days	Halos indicating potential plastic usage
<i>Paecilomyces variotii</i>	Ameen et al., 2015	Mangrove	LDPE	Weight loss, SEM, Enzyme Activity Assays, Estimation of CO ₂ Evolution	28 days	Increased biomass in culture with LDPE, CO ₂ emission and increased production of laccase, MnP and lignin peroxidase
<i>Penicillium chrysogenum</i>	Ojha et al., 2017	Soil from waste disposal site	LDPE and HDPE	SEM, AFM, and FTIR	90 days	Visual with SEM and detection of degradation compounds with FTIR after 60 days
<i>Penicillium citrinum</i>	Liebmingner et al., 2007	Landfill soil	PET	Increase of hydrophilicity detection (rising height and drop dissipation measurements)	1 day	Polyesterase hydrolyzed PET showed by rising height (5.1 cm) and drop dissipation measurements (55 s)
<i>Penicillium griseofulvum</i>	Brunner et al., 2018	Shoreline of lake Zurich	PE, PU	Halos test	several days	Halos indicating potential plastic usage
<i>Penicillium oxalicum</i>	Ojha et al., 2017	Soil from waste disposal site	LDPE and HDPE sheets	SEM, AFM, and FTIR	90 days	Visual with SEM and detection of degradation compounds with FTIR after 60 days
<i>Penicillium simplicissimum</i>	Sowmya et al., 2015	Soil from waste disposal site	UV treated PE, non uv autoclaved, non uv surface sterilized	Halo test, SEM, FTIR, NMR spectroscopy, Enzyme activity assay	90 days	Weight loss for UV treated polyethylene was 38%. FTIR detected degradation products. NMR indicated degradation. Both laccase and MnP when treated with PE disk showed weight loss and morphological changes in FTIR spectrum.
<i>Penicillium simplicissimum</i> YK	Yamada-Onodera et al., 2001	Soil	PE	HT-GPC, FTIR, visual growth assessment	90 days	Lower molecular weight and FTIR detection of degradation products
<i>Penicillium sp.</i>	Alshehrei, 2017	Seawater	PE	Weight loss, tensile strength, SEM, FTIR	30 days	43.4% weight loss of PE
<i>Penicillium sp.</i>	Raghavendra et al., 2016	Soil from waste disposal site	PU and LDPE UV irradiated for 50 h	Halo test, weight loss	90 days	Weight loss for PE and PU after 90 days and loss of tensile strength

(Continued)

TABLE 2 | (Continued)

Strains	References	Environment of isolation	Polymer	Degradation assessment technique	Incubation time experiment	Main observed results
<i>Penicillium sp.</i>	Magnin et al., 2019	Waste from terrestrial environment	PU	Weight loss, FTIR, SEM	60 days	SEM showed visual signs of degradation. Detection of PU degradation via FTIR
<i>Phialophora alba</i>	Ameen et al., 2015	Mangrove	LDPE	Weight loss, SEM, Enzyme Activity Assays, Estimation of CO ₂ Evolution	28 days	Increased biomass in culture with LDPE, CO ₂ emission and increased production of laccase and MnP
strain IZU-154	Deguchi et al., 1997	Collection	Nylon-6	NMR	20 days	NMR showed formation of four end groups, CHO, NHCHO, CH ₃ , and CONH ₂ indicating degradation
strain IZU-154	Iiyoshi et al., 1998	Collection	PE	Tensile strength, enzyme activity assay, HT-GPC	12 days	Decrease of tensile strength in presence of strain. Production of MnP. Loss in Mn with MnP treatment
<i>Thermomyces (formerly Humicola) insolens</i>	Ronkvist et al., 2009	Collection	PET (low crystallinity)	Weight loss, SEM, DSC and HPLC	6 days	Degradation of PET into TPA via cutinase (named HiC) and 97% weight loss
<i>Trichoderma harzianum</i>	Raghavendra et al., 2016	Soil from waste disposal site	PU and LDPE UV irradiated for 50 h	Halo test, weight loss	90 days	Weight loss for PE and PU after 90 days and loss of tensile strength
<i>Trichoderma harzianum</i>	Sowmya et al., 2014	Dumpsite soil	PE	SEM, FTIR, NMR analyses and enzyme assay	90 days	Weight loss of UV PE was 40%. Enzymes causing degradation were identified as laccase and MnP
<i>Trichoderma viride</i>	Munir et al., 2018	Soil from waste disposal site	LDPE	Weight loss, tensile strength	45 days	Weight loss of 5.13% and tensile strength reduction of 58%
<i>Xepiclopsis graminea</i>	Brunner et al., 2018	Shoreline of lake Zurich	PE, PU	Halos test	several days	Halos indicating potential plastic usage
<i>Zalerion maritimum</i> (ATCC 34329)	Paço et al., 2017	From collection but marine strain	PE	FTIR-ATR, NMR	28 days	FTIR analysis (carbonyl index), mass loss of 56.7 ± 2.9% of the plastic

PE, polyethylene; LDPE, low density polyethylene; HDPE, high density polyethylene; PP, polypropylene; PS, polystyrene; PU, polyurethane; Nylon, polyamides; PET, polyethylene terephthalate; TPA, terephthalic acid; SEM, scanning electron microscopy; FTIR, Fourier-transform infrared spectroscopy; NMR, nuclear magnetic resonance; DSC, differential scanning calorimetry; HPLC, high-performance liquid chromatography; HT-GPC, high temperature gel permeation chromatography; GC-MS, gas chromatography-mass spectrometry; MALDI-TOF, matrix-assisted laser desorption/ionization – time-of-flight mass spectrometry; Mn, average molecular weight; MnP, manganese peroxidase.

plastic degraders. *P. chrysogenum*, *P. oxalicum*, *P. simplicissimum* isolated from soil (Yamada-Onodera et al., 2001; Sowmya et al., 2012; Ojha et al., 2017) and *Penicillium sp.* isolated from seawater (Alshehrei, 2017) showed potential to degrade PE. Similar to *Penicillium*, different species belonging to the genus *Aspergillus* were found to be potential plastic degraders as well (Table 2). *A. flavus* isolated from soil (Deepika and Madhuri, 2015), wax moth gut (Zhang et al., 2020) and a marine environment (Sangeetha Devi et al., 2015; Alshehrei, 2017) exhibited the potential to degrade PE and *A. niger* isolated from soil (Deepika and Madhuri, 2015; Raghavendra et al., 2016) and seawater (Alshehrei, 2017) showed potential to degrade PE and PU. Also, *A. terreus* isolated from soil (Zahra et al., 2010), mangrove sediments (Ameen et al., 2015; Sangale et al., 2019) and seawater (Alshehrei, 2017) were potentially able to degrade PE. In addition to these species, several other types of *Aspergillus* strains were considered as potential PE and PU degraders. Many of the plastic degrading species that were isolated from terrestrial environments (Table 2) are also found in marine habitats. However, it has not been confirmed if all of the plastic degrading strains found in terrestrial environments perform equally well in the marine realm.

The enzymes utilized by plastic degrading fungi in the environment are typically not constrained. However, fungi produce a wide range of enzymes that have the potential to break down the chemical bonds of the plastic polymers (Figure 1). Amongst these are manganese peroxidase (MnP) and lignin peroxidase (LiP), which are commonly associated

with lignin degradation (Xu et al., 2013). These enzymes catalyze oxidation-reduction reactions, involving free radicals, transforming several compounds into oxidized or polymerized products (Wei and Zimmermann, 2017). Peroxidase are also used in industrial applications for degrading recalcitrant organic pollutants, PAHs, industrial dyes and chlorophenols (Qin et al., 2014). Lignin peroxidase is characterized by a high redox potential, and enables oxidation of non-phenolic aromatic compounds. Another enzyme that might be involved in plastic degradation is laccase, a multicopper oxidase, which is a well classified lignin-modifying enzyme (Mehra et al., 2018) and mediates the oxidation of the polymers' carbon backbone (Amobonye et al., 2021).

Elevated laccase, manganese peroxidase and lignin peroxidase activities were observed during PE degradation by a fungal consortium in a mangrove (Ameen et al., 2015). It appears likely that these enzymes also play a role in potential plastic degradation in the ocean. Marine adapted fungi have the ability to regulate the expression of their enzymes according to salinity. Cultivated in marine conditions, *Peniophora sp.*, for example, showed multigene transcription of ligninolytic laccase enzymes (Otero et al., 2017). Fungi isolated from marine environments can also produce enzymes that allow them to grow in liquid media with sole carbon sources such as agar, alginate, carrageenans, laminarians, and ulvans, i.e., polymers which are common in the marine realm (Wang et al., 2016). However, whether these compounds are also utilized by marine fungi *in situ*, i.e., in the ocean, still needs to be shown. Similarly to several

respectively, after 45 days (Munir et al., 2018). *Cladosporium tenuissimum* caused 25.9 and 65.3% of weight loss of PE-PU foams with and without flame retardants, respectively (Álvarez-Barragán et al., 2016). The marine fungus *Zalerion maritimum* exposed to PE microplastics caused mass loss of $56.7 \pm 2.9\%$ of the plastic, corresponding to 43% of removal after 2 weeks of exposure (Paço et al., 2017).

Although determining the weight loss of plastic is a straightforward method, it requires a long monitoring time, ranging from months to years to yield measurable gravimetric changes, while short term studies often yield inconclusive results (Lee et al., 1991). Gravimetric measurements require post incubation treatments, i.e., the removal of the biofilm from the incubated plastic. This can easily cause measurement artifacts; e.g., the incomplete removal of biofilm or accidental removal of a polymer respectively leads to an under or overestimation of weight loss. Weight loss measurements indicate that plastic polymers indeed disintegrate over time, however, this type of measurement does not reveal whether the polymer is broken down physiochemically (e.g., plastics release lower molecular weight compounds as a result of photooxidation; Wayman and Niemann, 2021) or if the plastic polymer was hydrolyzed and metabolized by microbes. Some plastics (e.g., polyvinyl chloride) contain high quantities of additives which, if soluble, can be released, and thus bias weight loss measurements. Also, biomass figures (i.e., cell numbers, culture dry weight) are typically not reported for

culture-based studies, making these investigations quantitatively non-repeatable, thus, severely limiting comparability. Hence, gravimetric measurements are insufficient to identify plastic degrading organisms and microbial kinetics, but should be accompanied by other methods.

Determining microbial biomass growth on plastic as the sole carbon source (e.g., monitoring cell numbers) has been used to infer the biodegradability of a specific polymer. Together with monitoring the weight loss of the polymer, some studies have measured an increase in fungal biomass as an indicator for the activity of fungi (Paço et al., 2017), and interpreted that the increase in fungal biomass occurred at the expense of carbon originating from the plastics.

Clearance Zone Formation

The growth of fungi at the expense of plastic can be monitored visually with degradation assays based on plates coated with agar and solubilized plastic. Inoculation at the surface and utilization of the plastic leads to the formation of clearance zones, referred to as 'halos.' Appearance of 'halos' on plates with plastics as the sole carbon source is used as an indicator for plastic degradation. The fungal species *Cladosporium cladosporioides*, *Xepiculopsis graminea*, and *Penicillium griseofulvum* and *Leptosphaeria* sp., isolated from plastic debris from the lake Zurich, formed clearance zones on PU. However, none of these strains were able to degrade polyethylene (Brunner et al., 2018). Attempts to use polyethylene and clearance zone formation

TABLE 3 | Most common methods used to assess plastic degradation advantages and disadvantages.

Methods	Principle	Advantages	Disadvantages
Gravimetric measurements and growth of biomass	Monitoring strains biomass growth with plastic as sole carbon source and plastic weight changes	Easily performed measurement and inexpensive	Not accurate and weight loss can be due to other processes than biodegradation
Clearance zone formation	Visual assessment based on the appearance of clearance zones on agar plates containing solubilized plastic	Easy assessment and cheap. Great screening technique before further studies.	The plastic is not the sole carbon source (presence of agar) so the strain growth cannot be automatically imputed to the biodegradation of plastic
Scanning Electron Microscopy (SEM)	Microscopy technique allowing visual assessment of strain growth and physical changes within the polymer	High resolution allowing a clear visual assessment	Changes in polymer physical structure and strain growth are not sufficient to prove plastic biodegradation
Atomic Force Microscopy (AFM)	Microscopy technique allowing collection of data on the surface roughness	Precise quantitative and qualitative data on the surface roughness	Changes in polymer surface roughness are not sufficient to prove plastic biodegradation
Fourier-Transform Infrared Spectroscopy (FTIR)	Spectroscopy technique allowing the obtention of an infrared spectrum of physio-chemical properties of a sample	Reliable detection and semi-quantification of changes of the polymer configuration	Sensitive to biofilm attachment and chemical treatments. Only proves degradation of the plastic
Assays with ^{13}C or ^{14}C labeled polymers	Tracing of isotopically labeled carbon from the plastic polymers	Precise quantification of plastic degradation products and traces incorporation in microbial biomass	High costs
Respiratory assays	Measurements of O_2 consumption and/or CO_2 production of strains	Can provide quantitative information on degradation	Respiration cannot be automatically imputed to biodegradation of plastic
Tensile resistance	Detection of physio-chemical properties changes (tensile resistance strength, thermal stability, glass transition, molecular weight). The decrease detected in these properties indicates structural alteration of the plastic polymer	Easy measurements that can indicate plastic polymer degradation	Changes in these properties do not automatically indicate biodegradation of plastic as these methods are sensitive to other processes of degradation
Thermo Gravimetric Analysis (TGA)			
Differential scanning calorimetric analysis (DSC)			
High-Temperature Gel Permeation Chromatography (HT-GPC)			

have been successful for *Aspergillus niger* and *Aspergillus flavus* (Deepika and Madhuri, 2015).

Clearance zone tests allow for rapid visual observations. However, the technique has the drawback that it requires dissolving the polymer in an organic solvent that can be applied to the petri dish at relatively cold temperatures to avoid agar melting. Furthermore, remnants of the solvents (rather than the plastic) may act as a carbon source for the fungi, leading to false-positive results. In addition, bias associated with clearance zone formation is relatively large, as clearance zones are not uniform in size or shape. Finally, fungi with hyphae may grow through the plastic, thus gain access to the agar below, which can also lead to false-positive results. As agar also represents a carbon source, this test alone does not prove microbial plastic degradation.

Scanning Electron Microscopy

Scanning electron microscopy (SEM) is used to create a surface image by directing a high-intensity electron beam at the surface and scanning over this surface. SEM allows high magnification, thus offers high resolution at the nanometer range. SEM-based observations are used to examine and evaluate the colonization of plastic films or particles by microorganisms and to simultaneously visualize cracks, pits and deformations on the plastic surface (Zettler et al., 2013; Vaksmaa et al., 2021a), which in return can indicate if the polymer is degraded. SEM has been applied in several studies to investigate fungi on plastics, for example, to visualize the growth of *C. tenuissimum* and *C. pseudocladosporioides* hyphae within PE-PU foams (Álvarez-Barragán et al., 2016). Furthermore, Paço et al. (2017) visualized the attachment of *Z. maritimum* on PE. With SEM, also surface roughness of single plastic fragments can be visualized. Floating marine plastics often feature signs of abrasion, cracking and ongoing fragmentation (Zettler et al., 2013; Vaksmaa et al., 2021b). Also, plastics exposed to the marine environment developed such signs during the incubation. For example, Welden and Cowie (2017) exposed PE, PP and Nylon to marine sediment for 12 months and found breaks and increased fraying on nylon ropes, surface scratching and roughening on PE filament rope, cracks, fissures and finer surface fibers scaling off from PP.

Scanning electron microscopy is a rapid technique, and allows to visualize surface attachment and morphological microstructures. Observations by SEM without chemical fixation can be achieved by applying FIB-SEM (Focused Ion Beam milling combined with Scanning Electron Microscopy). However, SEM does not allow phylogenetic identification of the microbes (unless the strain-specific morphological characteristics allow this) and the formation and attachment of biofilms is not necessarily an indication for biodegradation. Furthermore, though SEM is a valuable tool to visualize surface defects (e.g., cracks) of the plastic, it does not allow to scale in Z-direction (i.e., to measure the depth of cracks). Alternatively, this may be achieved by atomic force microscopy (AFM). For example, an increase in surface roughness and formation of cracks and grooves on PE films were investigated with AFM after exposure to two different *Penicillium* strains (Ojha et al., 2017).

Fourier-Transform Infrared Spectroscopy

Fourier-transform infrared spectroscopy (FTIR) is a technique used to obtain an infrared spectrum of absorption, emission, and photoconductivity of a material allowing to determine the chemical identity of most polymers. Furthermore, the FTIR spectrum enables detection and semi-quantification of changes of the original polymer configuration, for example, the introduction of carbonyl groups during polymer oxidation (Xu et al., 2019; Almond et al., 2020). The degree of carbonylation can be enumerated by determining the carbonyl index (calculated from the ratio between the integrated band absorbance of the carbonyl and that of the methylene peaks). Carbonyl indexes, as a measure of degradation, has been applied for a variety of polymers, such as PU (Filip, 1979; Álvarez-Barragán et al., 2016), PE (Paço et al., 2017), PS (Tian et al., 2017), and PP (Sheik et al., 2015). PE degradation has been evaluated by FTIR in co-cultures of bacteria and fungi, *Lysinibacillus xylanilyticus* and *A. niger*, isolated from soil (Esmaili et al., 2013). For example, *Penicillium variabile* CCF3219 strain decreased the carbonyl peaks of pre-oxidized 14C-βPS after 16 weeks of incubation and ozonation pre-treatment enhanced subsequent biodegradation (Tian et al., 2017). An additional advantage of FTIR is that it can be used for small plastic particles (the diffraction limit in IR spectroscopy is ~10–20 μm).

Fourier-transform infrared spectroscopy is a straightforward and reliable technique, however, it often yields non-quantitative results that are difficult to compare between studies. Also, the attachment of biofilm on the plastic surfaces affects the plastics optical properties in the IR range because proteinic and polysaccharide contents of the biomass will change the polymer's IR spectrum (Bonhomme et al., 2003). Samples must hence be pretreated to remove biofilms, e.g., using hydrogen peroxide (Löder and Gerdtts, 2015) or sodium dodecyl sulfate (Zhang et al., 2020). These chemicals have been described as the least aggressive. Nevertheless, chemical treatments can modify the molecular structure of plastic surfaces, which can introduce biases.

Assays With ¹³C or ¹⁴C Labeled Polymers

Isotopically labeled plastics can be traced into biodegradation products sensitively and quantitatively (Lanctôt et al., 2018; Taipale et al., 2019). The first studies involving isotopically labeled plastics were conducted with the radio isotope ¹⁴C. The fungus *Fusarium redolens* liberated ¹⁴C-CO₂, originating from pulverized ¹⁴C-labeled HDPE (Albertsson, 1978, 1980; Albertsson et al., 1978). ¹⁴C labeled ¹⁴C-CO₂ formation was also evaluated after photoirradiation of ¹⁴C-αPS and ¹⁴C-βPS, exposed to garden soil and activated sludge, showing higher ¹⁴C-CO₂ formation in treatments with photo oxidized polymer. The authors quantified degradation rates and showed that complete degradation of the ¹⁴C-αPS polymer in garden soil would require 20 to 80 years and in activated sludge from 11 to 24 years (Guillet et al., 1974). ¹⁴C labeled polystyrene was also used to test PS degradation capabilities of 17 different fungal species in a 14 days incubation experiment during

which 0 to 0.24% of the ^{14}C -PS was degraded (Kaplan et al., 1979). Because radioactivity can be measured and quantified extremely sensitively, radio isotope probing allows determining extremely low plastic degradation rates. Nevertheless, the use of radio-chemicals entails the necessity for specialized laboratory facilities and trained personnel, and the radio-labeled base materials for synthesizing polymers are extremely expensive or not available at all. This makes the use of radio isotopes mostly impractical. However, using plastics labeled with stable isotopes (e.g., containing a high degree of ^{13}C or ^2H) provides a good alternative. Similar to radio isotope probing, stable isotope probing (SIP) offers the advantages to allow tracing plastic derived matter into degradation products (Sander et al., 2019; Taipale et al., 2019), though the ubiquitous presence of ^{13}C and ^2H in most types of matter makes SIP assays less sensitive compared to approaches with radio isotopes. Nevertheless, no study using ^{13}C or ^2H labeled conventional plastics (PE, PP, PET, PS, PU, and nylon) for investigating interactions of marine fungi and plastics has been published yet. On the other hand, Zumstein et al. (2018) used ^{13}C -PBAT (Polybutylene adipate terephthalate) to study its microbial degradation in soil.

Other Methods

Additional methods to evaluate plastic degradation include the following techniques: (i) Respiratory measurements of O_2 consumption or CO_2 production, which can provide quantitative information of degradation, but cannot discriminate between different respiration pathways. (ii) Tensile resistance alterations of plastic can be used as a measure of the strength/integrity of the plastics, which will decrease as a function of degradation (yet, also physicochemically induced degradation reduces tensile strength). (iii) Thermo Gravimetric Analysis (TGA) characterizes the thermal stability of a polymer, which can potentially indicate its degradation (similar to tensile strength). (iv) Differential scanning calorimetric (DSC) analysis assesses the thermal properties of synthetic polymers, such as glass transition Temperature (T_g).

Lower T_g temperatures are often related to a decrease in the stability, indicating degradation (Lucas et al., 2008). (v) High-Temperature Gel Permeation Chromatography (HT-GPC) provides information on the molecular weight (Mn%) and molecular weight distribution of the polymer. A decrease in Mn% is evidence of chain cleavage that can be related to microbial degradation. Nevertheless, just as alterations of tensile resistance, glass transition Temperature, thermal stability and the molecular weight will also change in response to physicochemical processes. Finally, none of the other methods described in this section is able to unambiguously prove the occurrence of complete microbial biodegradation, from initial depolymerization to mineralization and biomass assimilation.

LIMITATIONS OF STUDYING FUNGAL COMMUNITIES

In comparison to bacteria, fungi in the marine realm are understudied and often overlooked. It seems likely that fungi are relevant as saprotrophs in general and may act as plastic

degraders, yet this needs to be demonstrated in future studies. However, in comparison to investigating bacterial communities, taxonomic and physiological characterization of (marine) fungi is not as straightforward and standardized methods are generally lacking. Indeed, molecular studies on fungi still encounter classic difficulties and biases related to molecular techniques such as polymerase chain reaction (PCR) bias, library preparation bias, sequencing bias, bioinformatics biases and unequal sequencing depth. Perhaps the most hindering factors in molecular studies of fungi are: (i) nucleic acid extraction method bias (ii) marker gene bias when using 18S rRNA or ITS spacer and (iii) primer bias. In addition, also (iv) culture-based methods are hindered by the fact that identification based on morphological features (alone) is difficult, growth conditions are hard to determine and fungi have complex life cycles.

- (i) Fungal genomic DNA extraction is less straightforward in comparison to extracting DNA from bacteria. Fungal cell walls are made of chitin making them more robust than a peptidoglycan bacterial cell wall (Fredricks et al., 2005; Shin, 2018). Breaking down fungal cells requires further steps such as the addition of lysing agents (for example, adapting the lysis buffer or adding enzymes such as cutinases) and/or mechanical disruption (e.g., increasing bead-beating steps or introducing freeze-thaw cycles) to increase fungal DNA yields, while maintaining the integrity of the DNA. Furthermore, different fungal strains may require different extraction steps to be added, complicating DNA extraction from the whole fungal community in environmental samples.
- (ii) Selection of a reliable marker gene (gene section) is influenced not only by the number of targeted taxa, but also the feasibility of down-stream analysis as this depends on the availability of data stored in publicly available databases. For prokaryotes, several and up to date databases exist for 16S rRNA gene sequences (and are publicly accessible), while far less information is available for 18S rRNA sequences. One of the most commonly used database is SILVA, but while the SILVA ref 138.1 release (Quast et al., 2013) contains 2,052,220 16S rRNA gene sequences (1,983,022 bacterial and 69,198 archaeal sequences), it only contains 172,520 18S rRNA gene sequences, of which 30,386 were classified as fungi. SILVA also, hosts a repository of 9329 representative 18S rRNA gene sequences covering all of the fungal kingdom and includes a manually curated alignment, and reference phylogenetic tree (Yarza et al., 2017). An alternative database for the identification of fungi is the ITS sequence database UNITE, but the most recent release (Nilsson et al., 2018) contains a similarly low number of 30,555 fungal sequences. Reich and Labes (2017) reviewed the available molecular ecology tools including their advantages and drawbacks when applied to the community studies of marine fungi. Even when the available number of 18S rRNA gene sequences, ITS sequences and fungal genomes has increased substantially, these numbers stand in stark contrast to the estimated several million fungal species (Nilsson et al., 2015). About 50% of the described fungal

species still lack any DNA sequence information in public databases (Xu, 2016). In order to fill the taxonomic marker-related gaps in public databases, it has been suggested to use third-generation sequencing and apply ribosomal tandem repeat sequencing to cover all ribosomal markers for fungi (Wurzbacher et al., 2019).

- (iii) Different nuclear ribosomal DNA marker genes are used for the identification of fungal species. However, environmental studies on fungal communities, apply most commonly the ITS and the 18S rRNA marker gene. The ITS region has been proposed as a universal barcode for fungal DNA as it may allow for a better resolution of fungal taxonomy than the 18S rRNA gene overall (Schoch et al., 2012). ITS barcoded sequencing has been successfully applied for both unicellular as well as for filamentous fungi (Vu et al., 2016, 2019). Many ITS targeting primers have been developed in recent years (Martin and Rygiewicz, 2005; Manter and Vivanco, 2007; Toju et al., 2012). However, choosing adequate thresholds for taxonomic assignments in ITS processing pipelines is delicate as the average intraspecific ITS variability is fluctuating (Smith et al., 2007; Simon and Weiß, 2008). For example, 0.2% for *A. fumigatus*, 3.1% for *F. solani* and up to 24.2% for the Ascomycota *Xylaria hypoxylon* (Nilsson et al., 2008). Vu et al. (2016, 2019) suggested that a threshold of 98.41% should be applied for ITS to distinguish between yeast species and a threshold of 99.6% should be applied to discriminate between filamentous fungi. Moreover, within an individual, ITS polymorphism (Alper et al., 2011) and ITS hybrid forms (Sriswasdi et al., 2019) were reported. Consequently, even when using fungal specific primers (ITS), intraspecific variability might not lead to species identification. Use of ITS shows even more limitations when dealing with marine fungi. Indeed, the ITS marker has a higher divergence rate compared to the 18S rRNA marker gene in marine fungal communities where early diverging fungi are abundant, and thus results into low classification success rates for ITS (Nilsson et al., 2019). For example, De Tender et al. (2017) could not assign 28 to 99% of the fungal reads acquired in their study when using ITS. To overcome this issue, it has been suggested to combine multiple marker genes for the molecular identification of the members of marine fungal communities. Banos et al. (2018) suggested to apply a fungi-specific 18S rRNA primers according to different environments, conditions or goals. This raises a new issue when using fungal specific ITS/18S primers, as different sets may overestimate some taxa and, in some cases, not target any fungus at all. By using a long read sequencing approach multiple marker genes can be retrieved at the same time, which allows for better taxonomic resolution (Heeger et al., 2018). Studies focusing on fungi colonizing plastic debris are scarce and the use of different marker genes and short read sequencing technology (Table 1) makes comparison between different datasets difficult. Unfortunately, no credible hypotheses on the presence or absence of a core fungal community living on plastic can be formulated yet.

- (iv) Culture-based methods are time-consuming and the culture media will have a selective effect on fungal growth. There is no specific medium for marine fungi available to date, but only adaptations of media selective for terrestrial fungi. The morphological diversity associated with different developmental stages of the same species, complicates the identification of fungal isolates (Xu, 2016). Nevertheless, only culturing of strains isolated from environmental samples enables in-depth investigation of their physiological and metabolic capabilities. Overy et al. (2019) published a toolkit of best practices for the culturing and isolation of marine fungi. We thus argue that combining molecular and isolation/culturing effort is the best way to evaluate interactions between fungal communities and plastics in marine environments.

CONCLUSION-OUTLOOK

This mini-review summarizes the current knowledge on marine fungi – PMD interactions; i.e., the ability of fungi to colonize plastics and specific strains known to degrade plastics as well as the methodological advances and difficulties in studying fungi – PMD interactions. Investigating the interaction of marine fungi and PMD is an emerging and exciting field of research when considering the high potential for plastic degradation of several fungal strains. Yet, for a well-constrained appraisal of marine fungi and their role as plastic degraders, several knowledge gaps still need to be filled. Firstly, a more general and fundamental understanding of fungi in the marine environment needs to be achieved by addressing fungal prevalence and diversity in the ocean. By using new molecular markers, the available sequence databases could be expanded for future classification of fungi detected on plastic polymers and to classify currently unclassified fungi. To address the biodegradation potential of fungi, comparable detection methods should be used to enable comparison between different strains, polymers and studies. Also, we suggest applying several complementary techniques for assessing biodegradation, particularly if the used techniques might cause false positives. Once fungal species degrading plastic in the marine environment are identified, future research should address the enzymatic potential of these fungi, which might then serve biotechnological applications for plastic waste bioremediation.

AUTHOR CONTRIBUTIONS

EZ wrote the manuscript with contributions from all co-authors. HN supervised the project. All authors contributed to the article and approved the submitted version.

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