What do we know about how the terrestrial multicellular soil fauna reacts to microplastic?

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Final responses to all referees plus marked-up version
Dear Referee #1

First I would like to express my sincere thanks to you for thoroughly reviewing our manuscript and for your very helpful and precise suggestions. In the following I will answer your points. Our corrections are marked-up with yellow numbers within the corrected manuscript at the end of this document.

Best regards,
Frederick Büks

Abstract

[1] Lines 20-21: “Most of the studies applied MP concentrations similar to amounts in slightly to very heavily polluted soils.” This sentence makes the reader expect that generally, the concentrations in the experimental environments are mostly the same as expected in the environment, but is this really the case? I would suggest showing the percentage of experiments with high microplastic exposure that is not representative of most soils.

Thanks a lot for this point. We now write: “About 58% of the studies thereby use inappropriate concentrations or units, but 42% applied MP concentrations similar to amounts in slightly to very heavily polluted soils.”

Introduction

[2] Line 53: Instead of “microbial decay”, I’d suggest “processing by soil organisms”, since this includes any process relevant for the generation of smaller plastic particles.
Done.

[3] Line 61: I’d suggest changing the sentence to “intensive use of plastic mulching and sewage sludge”, for the former, Huang et al. (2020) show an increase in microplastic by approx. 1 order of magnitude between fields with 5 and 24 continuous years of plastic mulching.
Done.

[4] Line 95: Suggest changing “feed on” to “inadvertently ingest”, otherwise it sounds like the organisms are actually able to metabolize the microplastics.
Done and reference added.

Search pattern

[5] The cut-off dates (time period that was considered) of the search should be mentioned somewhere.
Information added to this chapter (see the answer to referee #3)

[6] Figure 1: This figure shows the phylogenetic tree of edaphic fauna, rather than “edaphic tree of faunal life”.
Thank you. And done.

Data collection

[7] Line 113-122: I’ve been having some difficulties understanding the search methodology and table 8 (table 8 should be moved at the appropriate place to become table 1).
We moved the table to line 124 and mentioned that it contains the number of found studies. All table numbers were adjusted within the text.

It would be great if the authors could re-word this, specifying:

What does it mean that some combinations would have caused too much search effort?
It means e.g. that searching for a taxon only in combination with “PET” gives results for PET bottles for cultivation and experiments and also the “use” as pets, if the search is not case sensitive. We now tried to clarify this in our text.
“Organism-plastic” is not a type-shape combination.
Oh, yes, that's right. Corrected.

What exactly does the number of studies in table 8 mean? The number of articles or single experiments (sometimes more than one taxon or plastic type is used in one article)?
The number counts for how often type-shape combinations were used in all reviewed experimental setups independently of organism.

Some articles are included that studied the uptake of macroplastics by organisms, mainly termites and ant species. It is reasonable to include these studies, but it should be mentioned more prominently, in the abstract and aims of the review, that macroplastics are included. Where macroplastics were used in the reviewed studies, the size was explicitly mentioned in the article text, so we do not see a necessity for elaborating the text. We did add a mention of macroplastics to the abstract.

Maybe also in the synthesis, a sentence about the proportions of experiments using macro-, micro-, and nanoplastic would be a helpful piece of information.
Now mentioned in “4.2 Limitations of previous studies”

Tables 1-7: What does N/A mean in the tables? In some cases I assume “not analysed” (e.g., passive transport), but in other cases it should mean “not mentioned” (e.g., aging, coating, etc.) or “not observed” (e.g., measured adverse effects). I think this needs to be specified. Usually, N/A refers to “not applicable”, but this doesn’t fit in the tables.
In this work it means “(data) not available”. We marked it at the tables.

Synthesis

Lines 549-550: Could you cite the studies that imitated weathering in the described way?
We did so. Tsunoda et al. (2010) artificially aged their plastic by soaking in hot water at 90°C for 21 days, and then it was sanded/scratched with medium-grade paper prior to the test. Gebhard and Forster (2018) incubated particles in seawater for 4 weeks to stimulate the formation of biofilms.

[9] Lines 555-557: This is true, but it should be acknowledged that these additives are mainly present in commercial plastics, and therefore, mentioning of additives is not expected for “clean” microbeads specifically synthesized for the experiments. Nevertheless, the disadvantages of using these microbeads has been clearly discussed earlier in this section.
Done.

Conclusions

[10] Line 620-621: I am a little concerned about describing the results as “alarming”. Is it really? The following sentences actually refute this rather strong statement.
Replaced with “considerable”.

[11] Lines 624-629: I would suggest changing the sentence to: “To elucidate [...], the most exact reproduction of plastic concentrations and properties [...].” However, the difficulty here is that very scarce data of limited quality is available on concentrations of microplastic in soils, so a range of concentrations need to be used for future experiments in order to match the “real world” concentrations in soil, while expecting a decrease in uncertainty in analytic results in the future. Especially in the lower size ranges (<100μm) quantification is currently challenging. Therefore, little is known about size distributions occurring in soils. It might be worth mentioning this dilemma in a sentence.
Done.

[12] Technical corrections:
All done.
Dear PD Dr. Werner Kratz (referee #2)

Thank you very much for your review. In the following I will try to answer your comments at my best. Our corrections are marked-up with green numbers within the corrected manuscript at the end of this document.

Best regards,
Frederick Büks

[1] Line 53: Is that only “microbial” decay?
We agree, we will change this to “processing by soil organisms” as it is actually micro- as well as macroorganisms.

[2] Figure 1: The taxonomic group “further Panarthropods” is placed centrally, the other groups are not.
Done.

[3] Table 7: The last three experiments within this table were conducted by feeding the mice with a MP suspension. You might write “(food)” behind the concentration data as in the other tables.
Thanks a lot. Done.

[4] Line 507: “Preferably” instead of “preferrably”.
Done.

[5] Table 8: Could you explain the meaning of the numbers within the table. Are these the numbers of experiments with the named type-shape combinations?
Yes. Please see the answer to referee #1 (Table 8 is now Table 1).

[6] Lines 549-507: Is that proved that carboxylation of microspheres decreases hydrophobicity in an appreciable extent?
We ask the manufacturers of Polyciences Europe GmbH, a leading producer of PS microspheres, and they said no. We added this important information to the review.
Dear Referee #3

Thank you very much for your critical review of our manuscript. It has helped us to see some points which still need clarification. In the following, we want to explain how we propose to adjust our article based on the reviewer's comments and also explain why in some cases we do not agree with the reviewer's proposed changes. Our corrections are marked-up with purple numbers within the corrected manuscript at the end of this document.

(1) First and foremost, please have the manuscript edited by a professional (!) native (!) biologist (!). The English of your text is largely understandable, but rough. Apart from annoying typos, I found sentences the meaning of which I only understood when trying to translate them to German (my native language). So, your text will heavily benefit from thorough native editing.

Rereading our article we did indeed see that some typos had escaped our notice. We are slightly surprised by the request of the reviewer to have the manuscript edited by a “professional (!) native (!) biologist (!)”. We rephrased some stiff sentences and corrected grammatical errors. If a proofreading is indeed wished, we will have a scientific translator (English native speaker) correct the article.

(2) Then, the text lacks conciseness, it is overly long. For example, I suggest to omit all biological/ecological details you provide when introducing a taxon. This is per se interesting, but not to the point here (except when the reader needs background to understand microplastic effects). Then, figure 1 does not contribute to the understanding of your presentation, omit it. And I do not think it necessary to present taxa for which there is no information available, especially if the taxa are of minor or no importance in soil (e.g. line 183ff, 205, 220, 227, 450ff) or if the literature is not on edaphic species (435ff). As a reviewer, you are of course required to address blind spots of research (thus pointing out important taxa that are missing in literature), but you need to better balance completeness with a concise presentation.

Your suggestion to omit the ecological presentation of some key taxa is understandable. If we would expect all readers to be well acquainted with the soil fauna, we would definitely go along with this. However, SOIL is a multi-disciplinary journal connecting a broad spectrum of soil scientists. Therefore, we think it is helpful to provide a short overview of information on the soil fauna, such as ecological functionalities (marker function, transport, degradation, habitat and food selection), which might influence how they cope with microplastics. We have critically gone through the article and here we summarize which parts we will shorten.

- Proposal: We shortened the introduction of the springtail section, as it is indeed oversized.
- Proposal: We would agree with moving it to the supplements in order to save space, in case this is wished.
- Proposal: We shortened the chapter about Onychophora.

Potamopyrgus antipodarum in fact is a benthic snail.
- Proposal: We use this benthic species to show more clearly how inconsistent the few results for benthic and terrestrial snails are.

(3) I miss a convincing argumentation why you focus on multicellular animals (but then, you provide many details about bacteria, fungi, algae, plant roots in 72ff... omit this). A good line of reasoning could be that you follow up on the Rillig and Bonkowski (2018) paper. The aim of this review is to depict the influence of microplastic contamination in soils to the soil fauna. But, to present a holistic view on the food web, we refer to microorganisms, plant roots and biofilms within the introduction section. Being large fields of knowledge on their own, these organisms are not part of the focus in this review, however they are food sources for meso- and macroorganisms and, thus, worthy of mention. Given that we only use 22 lines to describe these other parts of the phylogenetic tree of soil life, we think this is merited and wish to leave this part in the review.

Unfortunately, we do not understand how Rillig and Bonkowski (2018), a paper on soil protozoa, matches your point. We have read this paper and do mention it elsewhere in the review.

(4) Please provide details of your literature search (123ff). When did you search? Which time span did you cover? Which search strings? Please consider the literature on meta-analyses how to properly specify these technical aspects.
The search was applied between June 2019 and January 2020, repeated in the first week of January 2020 and covers publications until January 2020. The search strings result from combinations of taxon, plastic type and particle shape shown in Table 1 (formerly Table 8).

- [5] Proposal: Information added to section 2.

(5) Thank you very much for the positive note.

(6) line 636f: Please reconsider including your supervisor as a co-author. What “supervision” means is nowhere clearly defined, however, co-authorship is only justified for significant contributions to the manuscript. Honorary authorship violates the principle of scientific honesty. We understand this point completely and agree that it is not good practice to include scientists who have not contributed significantly to a paper. We also acknowledge that supervision is a very broad term and would like to specify the contribution of Martin Kaupenjohann to the paper. [6] Martin Kaupenjohann was involved in the development of the idea and concept for this paper. During the literature reading and writing phase he has supported the work with frequent discussions of the contents of the article. And finally he has critically revised the manuscript.

Best regards,

Dr. Frederick Büks
Dr. Loes van Schaik
Prof. Dr. Martin Kaupenjohann
Dear Dr. Maha Deeb (referee #4)

Thank you very much for the repeated check and your friendly report.

Best regards,

Dr. Frederick Büks
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Abstract. The ubiquitous accumulation of microplastic (MP) particles across all global ecosystems comes along with the uptake into soil food webs. In this review, we analyzed studies on passive translocation, active ingestion, bioaccumulation and adverse effects within the phylogenetic tree of multicellular soil faunal life. The representativity of these studies for natural soil ecosystems was assessed using data on the type of plastic, shape, composition, concentration and time of exposure.

Available studies cover a wide range of soil organisms, with emphasis on earthworms, nematodes, springtails, beetles and lugworms, each focused on well-known model organisms. About 58% of the studies thereby used inappropriate concentrations or units, but 42% applied MP concentrations similar to amounts in slightly to very heavily polluted soils. In many cases, however, polystyrene microspheres have been used, a combination of plastic type and shape, that is easily available, but does not represent the main plastic input into soil ecosystems. In turn, MP fibers are strongly underrepresented compared to their high abundance within contaminated soils. A few studies also examined the comminution of macroplastic by the soil fauna. Further properties of plastic such as aging, coating and additives were insufficiently documented. Despite these limitations, there is a recurring pattern of active intake followed by a population shift within the gut microbiome and adverse effects on motility, growth, metabolism, reproduction and mortality in various combinations, especially at high concentrations and small particle sizes.

For the improvement of future studies, we identified problems of past experiments and recommend that coming studies take into account the type, shape, grade of aging, specific concentrations of MP fractions and long-term incubation, in natural and contaminated soils.
1 Introduction

Imagine a compact plastic cube of nearly 2 km side length and a weight of 7300000000 tons, with major percentages by weight of 36 % polyethylene (PE), 21 % polypropylene (PP), 12 % polyvinyl chloride (PVC) and 10 % of each polyethylene terephthalate (PET), polyurethane (PU) and polystyrene (PS). That is the accumulated global non-fiber production of the six main plastic types until 2015. It accounts to 87 % of the all-time plastic production, which evolved exponentially, since the early 1950s, from some megatons (Mt) to 8300 Mt in 2015, with only 260 Mt annual output in 2009 increased to 380 Mt in 2015 (Thompson et al., 2009; Geyer et al., 2017). Of this ever produced plastic, 6300 Mt became waste until 2015, of which only 21 % were recycled or incinerated, whereas 5000 Mt ended up in landfills and nature (Geyer et al., 2017). As a corollary of production, use and disposal, a certain part of plastic waste is constantly released into the environment through various pathways, but our knowledge about rates of mass flow into global ecosystems is very limited. Based on waste generation in coastal countries, Jambeck et al. (2015) calculated the global plastic input to marine ecosystems to be roughly 4.8 to 12.7 Mt in 2010. Such data on soils are lacking, but Nizzetto et al. (2016) estimated that the load of microplastic (MP) to agricultural sites in Europe is in the same order of magnitude as that in marine environments.

By littering, plastic mulching, the application of sewage sludge, digestates and composts as well as windblown dispersal (Bertling et al., 2018; Weithmann et al., 2018; Zhang et al., 2019; Wang et al., 2019a), plastic from our technosphere arrives in soil ecosystems in various forms as large and small fragments, fibers and particles. Exposed to UV radiation, mechanical stress and processing by soil organisms, plastic items become weathered and prone to a successive comminution towards the size range of MP with increased surface, charge and biofilm cover (Kale et al., 2015; Andrady, 2017). However, the resistance of plastic to metabolization causes a constant accumulation in soils as long as the release rate from human processes is above the very slow rate of degradation.

Due to a lack of monitoring programs, data on MP concentrations in terrestrial soils are rare, and those using w/w concentrations represent only a small part compared to item concentrations. In soils with only slightly contaminated conditions, amounts seem to average about 1 mg kg\(^{-1}\) soil dry weight (and approx. 200 items kg\(^{-1}\) dry soil) (Rezaei et al., 2019). On sites with industrial activity or intensive use of plastic mulching and sewage sludge in agriculture, concentrations can be increased by 2 to 4 orders of magnitude (Fuller and Gautam, 2016; Zhang and Liu, 2018; Huang et al., 2020). Semisubhydric soils such as beaches, mudflats, mangroves or lagoons, that are additionally contaminated from the aquatic side, contain MP of the order of 10 to 100 items kg\(^{-1}\) dry soil and single extreme samplings contained several thousand items (Nor and Obbard, 2014; Naji et al., 2017; Garcés-Ordóñez et al., 2019; Li et al., 2018a). More informative data using mg kg\(^{-1}\) are only available for beaches and coastal deconstruction yards in municipal neighborhoods and amount to 0.5 and
70 mg kg⁻¹ dry soil, 0.00005 and 0.007 % w/w, respectively (Reddy et al., 2006; Claessens et al., 2011). All these concentration data represent a wide range of particle sizes between 0 and 5000 µm with different materials, shapes and degrees of aging.

Plastic particles can possibly enter and accumulate in the food web by either direct uptake from soil or by consumption of other soil biota contaminated by adhesion or ingestion (Huerta Lwanga et al., 2017a). There is evidence, that MP is incorporated even by plants and unicellular organisms at the base of the food web. Bacteria, for example, that are reasonably assumed to avoid MP uptake due to their minor size and the prevalent lack of phagocytosis, were shown to take up inorganic nanoparticles of a few nanometers (Kumar et al., 2011). Although the physiochemical properties of weathered nanoparticulate plastics might differ from these, also their uptake seems likely.

A similar argument can be made for fungi and soil algae, but studies on incorporation are lacking, whereas the transfer into a freshwater food web by adhesion of nanoplastic on algae has been shown by Chae et al. (2018). The uptake of MP into plant roots is also inhibited (Rillig et al., 2019), but occurred for nanoplastics that permeate into the plant tissue (Li et al., 2019). Also the integration into root tissue after adsorption to the rhizodermis has yet to be studied.

In contrast, protozoa feature phagocytosis for the active ingestion of particles. Diverse soil, freshwater and marine ciliates ingest PS/latex beads of 0.1 to 14.4 µm in laboratory experiments, with preferences to their natural prey size (Fenchel, 1980; Jonsson, 1986; Lavin et al., 1990). Soil amoebas act similarly, but additionally select according to food quality (Weisman and Korn, 1967; Vogel et al., 1980; Bowers and Olszewski, 1983; Avery et al., 1995; Elloway et al., 2006).

Finally, many soil microbiota live protected within biofilms. Plastic particles were shown to be a potential surface for the formation of those biofilms (Lobelle and Cunliffe, 2011), which are a food source for grazing primary consumers. inadvertent ingestion might also transfer occluded or abrased MP to higher trophic levels.

But what about the larger organisms that feed on all these, free plastic particles, contaminated microorganisms, biofilms and one another? Recent work discussed the effects of MP on soil biota (Chae and An, 2018) or called for intensified research on certain taxonomic groups (Rillig and Bonkowski, 2018). Thus, we were motivated to give on our part a review with focus on the most-produced plastics and their passive translocation, ingestion, bioaccumulation and adverse effects on the multicellular soil fauna. The types, sizes and shapes of plastic used in former laboratory studies were compared with the available knowledge on plastic in the environment, and recommendations are given for future research.

This analysis aims to support the assessment of the influence of MP on the ecosystem services provided by diverse soil organisms.
Within the tree of life, edaphic branches were identified comprising taxa that permanently inhabit the soil, are both-sided part of the soil food web and/or the burrowing macro- and megafauna or have active subterranean larval stages. The resulting tree of soil life based on the NCBI taxonomy database (Fig. 1) was drawn using the software phyloT and shows the leading taxonomic rank, which is mainly the family, but in exceptions – e.g. if one species represents the only soil-born between many aquatic – a lower rank.

Figure 1: [2] Tree of [6] edaphic fauna. Taxonomic ranks, that were examined in this qualitative study, are placed at the outer rim of the diagram. The length of the connecting line between two taxa represents the grade of phylogenetic relationship.
A pattern of search terms was established (Table 1), consisting of „taxon“ (Linné’s binominal nomenclature, common name, plural-sensitive search), „plastic type“ (plastic, microplastic, nanoplastic, PE or polyethylene, PP or polypropylene, PVC or polyvinyl chloride, PS or polystyrene, PU or polyurethane, PET or polyethylene terephthalate and latex) and „common shapes“ (fragments, particles, fibers, microfibers, beads, microbeads, microspheres). Some type-shape combinations caused problems, as they led to a very large amount of unuseful, off-topic papers – e.g. using any taxon combined with PET, papers with the use of PET bottles in experimental set-ups were selected or also studies on pets. Those combinations of search terms were excluded from this pattern. Further plastic types and shapes occuring within the found studies were also included in the review. Data on microspheres and microbeads were pooled, as both names describe one and the same.

Table 1: Types and shapes of microplastic particles in edaphon studies within this review. (X) symbolizes combinations excluded from the search pattern. The number counts for how often type-shape combinations were used in all reviewed experimental setups independently of organism. Empty fields stand for zero results. Microbeads and microspheres are often mixed up terms and, thus, counted together.

| Organism: | Linné’s systematic names OR common name |
|-----------|----------------------------------------|
| plastic   | X                                      |
| microplastic | X                                      |
| nanoplastic | X                                      |
| PE OR polyethylene | X 4 10 1 1 4 7 |
| PP OR polypropylene | X 1                                      |
| PVC OR polyvinyl chloride | X 4 6 1                                      |
| PS OR polystyrene | X 6 3 24 4                                      |
| PU OR polyurethane | X                                      |
| PET OR polyethylene terephthalate | X 3 2 6 |
| latex | X                                      |
| other | X 6 3 1 1 |
| N/A | 1 1 2 3 |

The search was applied between June 2019 and January 2020 within the Web of Science Core Collection Database, repeated in the first week of January 2020 and covers publications until January 2020. The search strings result from combinations of taxon, plastic type and particle shape shown in Table 1. Based on the search pattern, data on passive transport,
ingestion, bioaccumulation and adverse effects were collected for each edaphic group. Studies that only use uncommon, local, outdated, weird or nicknames are excluded by the search pattern. Studies testing injection to tissues, lymph or blood were excluded, as they do not represent natural ways to incorporate MPs. Data on inhalation by the megafauna in fact represent a natural way of uptake, but were also excluded as they are exclusively related to above-ground organisms, that only occur on the outer edge of the food-web. Also running debates on phylogenetic classifications are not part of this work and the taxonomists will be able to adjust the branches accordingly to their purpose.

The data of related taxonomic groups were pooled and evaluated for their environmental representativity based on exposure time, plastic concentrations and properties used. From this synthesis recommendations for a structured experimental design were derived for application in future studies.
3 Data collection

3.1 Insects

Within the Panarthropoda, the insects comprise the highest taxonomic diversity. And, regarding MPs, they represent an unevenly studied taxonomic group.

Within the Insecta, the Coleoptera (beetles) are an extraordinarily diverse and abundant taxon. Studies on plastic uptake into adult individuals mainly focus on the subfamily of Scarabaeinae (dung beetles). Comprehensive experiments with latex microbeads showed, that many species only ingest [12] smaller particles with maximum diameters of about 10 to 83 µm and retain them within the gut – with a slightly positive dependency on body size. Larger particles were rejected by a filtering mechanism within the mouth region and not ground with the mandibles (Holter, 2000; Holter et al., 2002; Holter and Scholtz, 2005). Beside those on Nematods, these data comprise by far the most detailed information about [12] size-dependent uptake of MP particles compared to other edaphic taxa. This gives a good foundation for future studies on adverse concentrations. In addition, several studies with plastic as predominant food source could show chewing, ingestion and intestinal degradation of different PS and PE foams in feeding experiments with Tenebrio sp. larvae (mealworms). These experiments also pointed out an alteration of the gut microbiome, but no adverse effects on reproduction and survival, with only one case of a non-significant tendency to higher mortality after 1 month of exposure (Yang et al., 2015; Brandon et al., 2018; Yang et al., 2018; Peng et al., 2019).

The Isoptera (termites), recently categorized as part of the order Blattodea, are the oldest social insects with a tribal history of about 130 million years (Korb, 2008). Especially in arid ecosystems with a lack of earthworms they play an important role in homogenization of soils, but also in sorting of soil mineral particles for building mounds as well as decomposition and distribution of organic matter (De Bruyn and Conacher, 1990). Tsunoda et al. (2010) and Lenz et al. (2012) could show, that different termite species are picky feeders and erode PE, but avoid other plastic cable sheathing. This suggests the excretion of ground MP particles by termites, but metabolic impacts are unknown. In contrast to termites, data on other Blattodea (e.g. cockroaches) were not found.

The suborder Apocrita comprises some flying insects, that inhabit burrows within the soil, such as ground-dwelling wasps within the Vespidae superfamily, mining bees within the Apoidea superfamily and the Spheciformes. They mostly do not prey and feed on subterrestrial organisms, but may move MP particles into the ground, as implied by a report of Allasino et al. (2019) on [12] solitary bees, which built nests fully made of plastic fragments. The Apocrita also contain the Formicidae (ants). Some ant species are considered an important factor for seed dispersal, a behavior, that could also be shown for artificial plastic seeds with ~2 mm diameter (Hughes and Westoby, 1992; Angotti et al., 2018). Robins and
Robins (2011) found that this also includes differently shaped cultural objects: *Rhytidoponera metallica*, a representative of ground-nesting, omnivore ants, is capable not only of a remarkable bioturbation, but also of an active, apparently random burying of anthropogenic plastic artifacts >1 mm. Seeds are used as a food source, thus, the ingestion of plastic bites is conceivable, but not documented. The uptake of latex microspheres ≥0.88 µm with liquids by larvae of *Solenopsis invicta* seems to be prevented by filtration within the mouth and the particles are released as larger aggregates, whereas other species ingest by far larger particles up to 150 µm (Glancey et al., 1981). However, also here data on adverse effects are missing.

Further insects with edaphic adult stages, e.g. *Dermaptera* (earwigs), *Heteroptera* (true bugs) and *Zygentoma* (silverfish, fishmoth, firebrat) or soil- or litter-dwelling larvae such as *Embioptera* (webspinners, footspinners), *Thysanoptera* (thrips), *Psocoptera* (booklice, barklice, barkflies), *Neuroptera* (lacewings), *Raphidioptera* (snakeflies) or *Zoraptera* (angel insects) were not yet researched with focus on soil MP.

Regarding insects, mainly studies on translocation and uptake of MP were carried out. In contrast, work on bioaccumulation is completely lacking and adverse effects are sparsely tested using *Tenebrio sp.* larvae. Such studies could provide information whether or not the input of MP in soil ecosystems is one of many factors causing the global decline of the entomofauna (Oliveira et al., 2019; Sánchez-Bayo and Wyckhuys, 2019).
Table 2: Microplastic studies on Coleoptera, Blattodea (Blattod.), Apoidea (A.) and Formicidae (mb=microbeads, fr=fragments, ms=microspheres, b=beads, [8] N/A=information not available). Concentrations refer to mg kg\(^{-1}\) dry soil, if not specially marked.

| organism | experimental environment | plastic type | aging | coating | additives | shape | size span [µm] | concentrations | exposure time | passive transport | active uptake | bioaccum. dynamics | measured adverse effects | reference |
|----------|--------------------------|--------------|-------|---------|-----------|-------|---------------|---------------|---------------|------------------|---------------|---------------------|-----------------------------|-----------|
| Aphodius erraticus | vial | latex | N/A | N/A | N/A | mb | 5 | 2.39 | N/A | 45 min | N/A | no | N/A | Holter (2000) |
| Aphodius rufipes | Petri dish | latex | N/A | N/A | N/A | | | | | | | | | |
| Aphodius ater | Petri dish | latex | N/A | N/A | N/A | | | | | | | | | |
| Aphodius fimetarius | Petri dish | latex | N/A | N/A | N/A | | | | | | | | | |
| Aphodius contaminatus | Petri dish | latex | N/A | N/A | N/A | | | | | | | | | |
| Aphodius fossor | Petri dish | latex | N/A | N/A | N/A | | | | | | | | | |
| diverse dung beetles | Petri dish | latex | N/A | N/A | N/A | | | | | | | | | |
| Tenebrio molitor larvae | container | PS | N/A | N/A | no | foam | N/A | 100% w/w (food) | 31 d | N/A | yes | biodegrad. | N/A | Yang et al. (2015) |
| Tenebrio molitor larvae | container | PS | N/A | N/A | no | foam | N/A | 86..100% w/w (food) | 31 d | N/A | yes | biodegrad. | microbiome | Brandon et al. (2018) |
| Tenebrio molitor larvae | container | PS | N/A | N/A | no | foam | N/A | 86..100% w/w (food) | 31 d | N/A | yes | biodegrad. | no | Yang et al. (2018) |
| Tenebrio obscurus larvae | container | PS | N/A | N/A | no | foam | N/A | 86..100% w/w (food) | 31 d | N/A | yes | biodegrad. | microbiome | Peng et al. (2019) |
| Captotermes formosanus | mesocosm | LD-PE | N/A | N/A | no | flame retardant | foam | 8..27 cm\(^3\) | 50..100% w/w (food) | 32 d | N/A | yes | biodegrad. | N/A | Tsunoda et al. (2010) |
| diverse termites | Petri dish | latex | N/A | N/A | fluorescence | ms | 0.9..4.5 | 2.5% w/w (food) | direct | N/A | filtration | N/A | Glancey et al. (1982) |
| Solenopsis invicta | Petri dish | latex | N/A | N/A | fluorescence | ms | 0.9..4.5 | 2.5% w/w (food) | direct | N/A | filtration | N/A | Glancey et al. (1982) |
| Rhytidoponera metallica | in situ | MD-PE | N/A | N/A | anti-oxidant stabilizer | cable sheet | 4 cm, ∅ 0.8 cm | N/A | 42 d | N/A | yes | no | N/A | N/A | Lenz et al. (2012) |
| Aphaenogaster longiceps | in situ | MD-PE | N/A | N/A | anti-oxidant stabilizer | cable sheet | 30 cm, ∅ 1.4 cm | N/A | 6 yr. | N/A | yes | no | N/A | N/A | Lenz et al. (2012) |
| Pheidole sp. | in situ | MD-PE | N/A | N/A | anti-oxidant stabilizer | cable sheet | 30 cm, ∅ 1.4 cm | N/A | 6 yr. | N/A | yes | no | N/A | N/A | Lenz et al. (2012) |
| Formicidae | in situ | MD-PE | N/A | N/A | anti-oxidant stabilizer | cable sheet | 30 cm, ∅ 1.4 cm | N/A | 6 yr. | N/A | yes | no | N/A | N/A | Lenz et al. (2012) |
| Rhytidoponera metallica | in situ | MD-PE | N/A | N/A | anti-oxidant stabilizer | cable sheet | 30 cm, ∅ 1.4 cm | N/A | 6 yr. | N/A | yes | no | N/A | N/A | Lenz et al. (2012) |
3.2 Other panarthropods

Apart from the insects, *Acari* (mites) comprise many abundant soil-living taxa, that feed on litter, fungi and fauna as predators and parasites and are bioindicators, as they are sensitive to changes in the soil physiochemical environment (Gulvik, 2007). Experiments indicated, that mites passively transport MP due to pushing and dragging after attachment to their cuticle, as shown with 80 to 250 µm sized PVC particles in a Petri dish experiment without soil (Zhu et al., 2018a). The population within manure pats slightly declines when exposed to mm-sized unweathered PE and PS particles at concentrations of 5 % v/v and declines strongly at ≥60 % v/v (Stamatiadis and Dindal, 1990). This could probably be an effect of moisture deficiency due to a reduced water holding capacity in an unnaturally enriched substrate, but not necessarily through plastic intake. In contrast, no data was found on their arachnid, preying relatives, the order of *Pseudoscorpiones* (false scorpions).

Just as many other highly abundant and diverse representatives of the soil mesofauna, the *Oniscideae* (woodlice) contribute to the decomposition of litter by chewing and passage through their digestive system (Warburg, 1987) and react strongly to environmental pollution, thus, they are potentially used as bioindicators (van Gestel et al., 2018). They practice a strict selection of natural food sources (Hassall and Rushton, 1984). This is also demonstrated for starch and cellulose based plastic films (4 cm²), which were consumed and digested in experiments with the model organism *Porcellio scaber*, in contrast to PHB (polyhydroxybutyrate) films, that reduces the feeding rate (Wood and Zimmer, 2014). Smaller PE particles (137±51 µm and 183±93 µm) embedded into food pellets (0.4 % w/w) were taken up easily by *Porcellio scaber*, and the smaller fraction caused a slight and non-significant reduction of body mass after 14 days of exposure, but not of feeding, defecation or energy reserves (Kokalj et al., 2018).

Other panarthropodean groups are even less studied in terms of MP. We did not find literature on the subphylum of *Myriapoda* containing the classes of *Diplopoda* (millipedes), *Chilopoda* (centipedes), *Pauropoda* and *Symphyla* (pseudocentipedes or symphilids), which are important litter-feeders and predators within various soil ecosystems.

The situation is nearly similar with the phylum of *Tardigrada* (water-bears or tardigrades), that has many ecologically relevant and well studied species feeding on microorganisms and detritus particles. Sparse field research in semisubhydric environments showed no uptake of MP fibers by tardigrada (Gusmão et al., 2016), but comprehensive data on terrestrial soils are lacking. Similarly, the related phylum of *Onychophora* (velvet worms), primordial invertebrates that are mainly native in litter and soils with high water holding capacity under pleistocene-like forest vegetation within tropical and moderate regions (Monge-Nájera, 1994). Another branch within the panarthropoda, the phylum of *Onychophora* (velvet worms), comprises primordial invertebrates that are mainly native in litter and soils with high water...
holding capacity under pleistocene-like forest vegetation within tropical and moderate regions (Monge-Nájera, 1994). As predators, they most likely take up plastic debris appearing within or on their prey, but no studies on MP are available, most likely due to their remote habitats, low-abundance and little scientific focus.

The phylum of Collembola (springtails) \[1\] together with the Diplura and Protura (Westheide and Rieger, 1996; Pass et al., 2011), an abundant, diverse and ubiquitous soil-borne phylum with a broad spectrum of food sources (Hopkin, 1997), also represent an intensively studied group within the Arthropoda. Together with the Diplura (which mainly live in tropic and subtropic regions in litter and humid topsoil and feed on fungal hyphae, POM and prey) (Westheide and Rieger, 1996) and the Protura (Pass et al., 2011), the Collembola build an intensively studied morphological group, that \[12\] exhibits similar ecological functions, such as distribution and decomposition of organic matter as well as the control of fungal abundance (Hopkin, 1997). Springtails provide up to 27 % of the soil biomass and up to 33 % of the total soil respiration (with higher shares in colder ecosystems) (Petersen, 1994) with up to 100000 individuals per square meter (Hopkin, 1997). Thus, their well-being plays an important role for ecosystem functioning.

In a Petri dish experiment without soil, Maaß et al. (2017) showed the passive transport of urea-formaldehyde particles <400 µm and undefined PET fragments by two Collembola species (Folsomia candida and Proisotoma minuta) due to attachment, but found no ingestion. Within a soil matrix, trials of Kim and An (2019) indicated hindrance of collembolan migration by larger PS particles (44±39, 282±131 and 676±479 µm) at concentrations of 1000 mg kg\(^{-1}\) corresponding to highly contaminated soils. In addition, they found suppressed mobility due to the attachment of even smaller PS microbeads (0.47 to 0.53 µm) at concentrations of 8 mg kg\(^{-1}\) dry soil, which is equivalent to values found in nature. Small particles <50 µm were moved, while larger particles were most likely \[12\] cast off. When F. candida encounters two of its predators, the mites Damaeus exspinosus and Hypoaspis aculeifer, the dispersal of 80 to 250 µm PVC particles is enhanced as shown by Zhu et al. (2018a) in a Petri dish experiment. Without proving the ingestion or the minimal effective MP concentration, Zhu et al. (2018b) published an alteration of the gut microbiome and adverse effects on growth and reproduction of F. candida by 80 to 250 µm PVC particles mixed in soil at concentrations of 1000 mg kg\(^{-1}\) dry soil. These data were not considered robust (van Gestel and Selonen, 2018), but match with a later study that found inhibited reproduction at \(\geq\)1000 mg kg\(^{-1}\) and avoidance behavior as well as microbiome alteration at \(\geq\)5000 mg kg\(^{-1}\) (Ju et al., 2019). Such concentrations can occur in highly contaminated soils (Fuller and Gautam, 2016). However, documentations on the active uptake, gnawing and grinding of MP by springtails proposed by Rillig (2012) is still lacking and also studies on Diplura and Protura were not found.
Table 3: Microplastic studies on Acari, Oniscidea (Onisc.), Tardigrada (T.) and Collembola (fr=fragments, p=particles, mf=microfibers, mb=microbeads, ms=microspheres, s=semisubhydric, [8] N/A=information not available). Concentrations refer to mg kg\(^{-1}\) dry soil, if not specially marked.

| organism | experimental environment | plastic type | aging | coating | additives | shape | size span [µm] | concentrations | exposure time | passive transport | active uptake | bioaccum. dynamics | measured adverse effects | reference |
|----------|--------------------------|--------------|-------|---------|-----------|-------|---------------|----------------|---------------|-------------------|--------------|-------------------|-------------------------|-----------|
| Acari    | diverse mites            | microcosm    | PE    | no      | N/A       | N/A   | fr <4800       | 0.90% v/v (manure) | 16 d          | N/A              | N/A           | N/A              | ≥5% v/v: abundance ↓ | Stamatiadis and Dindal (1990) |
|          | Hypoaspis aculeifer      | Petri dish   | PVC   | N/A     | no        | N/A   | p 80.250      | 5000 items per dish | N/A          | yes              | N/A           | N/A              | N/A         | Zhu et al. (2018a) |
| T. Onisc.| Porcellio scaber         | mesocosm     | PHB   | no      | N/A       | N/A   | fr 4 cm\(^2\)  | 1 item per cosm       | 14 d         | N/A              | yes           | N/A              | feeding ↓    | Wood and Zimmer (2014) |
| T. Onisc.| Porcellio scaber         | Petri dish   | PE    | N/A     | N/A       | N/A   | fr 183±93     | 0.4% w/w (food)     | 14 d         | N/A              | yes           | N/A              | no          | Kokalj et al. (2018) |
| T. Tardigrades | diverse tardigrades \^a | in situ      | N/A   | N/A     | N/A       | N/A   | mf 137±51     | N/A            | N/A          | no               | N/A           | N/A              | N/A         | Guzmão et al. (2016) |
| T. Tardigrades | Folsomia candida      | microcosm    | PVC   | N/A     | no        | N/A   | p 80.250      | 1000           | 56 d         | N/A              | N/A           | N/A              | microbiome, growth ↓, reproduction ↓ | Zhu et al. (2018a) |
| T. Tardigrades | Folsomia candida      | microcosm    | PVC   | N/A     | no        | N/A   | p 80.250      | 5000 items per dish | N/A          | yes              | N/A           | N/A              | N/A         | Zhu et al. (2018a) |
| Collembola| Folsomia candida         | microcosm    | PE    | N/A     | no        | N/A   | mb <500       | 0..10000       | 7 d          | N/A              | N/A           | N/A              | ≥5000: avoidance ≤1000: reproduction ≤1000: microbiome | Ju et al. (2019) |
|          | Lobella sokamensis      | soil sample  | PS    | no      | N/A       | N/A   | carboxyl fluorescence mb 0.5 | 4.8        | yes          | yes          | N/A              | N/A         | avoidance, motivity ↓ | Kim and An (2019) |
3.3 Annelida

Land-based Annelida comprise another large group of invertebrates. The Lumbricidae (earthworms) are a well-studied family (Darwin, 1881; Lavelle et al., 2006), represented in high abundance and diversity in many ecosystems all around the world (Phillips et al., 2019). Earthworms are often used as indicators for soil health (Fründ et al., 2011; Pulleman et al., 2012), as they are ecosystem engineers which through their burrowing activity influence various soil physical, chemical and biological processes (Jouquet et al., 2006; Lavelle et al., 2006).

By far the most of the studies on the influence of MP on earthworms are performed with PE and the species Lumbricus terrestris or Eisenia fetida, but there are also single studies with Aporrectodea rosea (Boots et al., 2019) and Eisenia andrei (Rodriguez-Seijo et al., 2017) and with the less common species Metaphire californica (Wang et al., 2019b). We found one field study of earthworms and MPs (Huerta Lwanga et al., 2017a) among many laboratory experiments with MPs mixed into soil volumes (concentrations ranging up to 20000 mg kg$^{-1}$ dry soil) or applied with litter on top of the soil surface (≤60% w/w). The particles sizes were usually <1 mm in diameter, but some were even up to 2x2 cm$^2$, and the duration of experiments was generally 14 to 28 days, few lasted up to 60 days.

The uptake of MPs of a broad size range by earthworms was shown in studies based on particles in earthworm casts of Lumbricus terrestris (Huerta Lwanga et al., 2016; Cao et al., 2017; Hodson et al., 2017; Rillig et al., 2017; Prendergast-Miller et al., 2019; Yu et al., 2019; Huerta Lwanga et al., 2017a), Eisenia fetida (Rodriguez-Seijo et al., 2018; Chen et al., 2020; Wang et al., 2019c), Eisenia andrei (Rodriguez-Seijo et al., 2017) and Metaphire californica (Wang et al., 2019b). Zhang et al. (2018) showed that relatively large PE particles of 1.5 x 1.5 cm$^2$ are not ingested by Lumbricus terrestris, but partial ingestion of such large particles of biodegradable MPs does take place after initial weathering in soil or in compost has occurred. In some laboratory experiments, MPs were found in the gut of dissected earthworms (Huerta Lwanga et al., 2016; Hodson et al., 2017; Rodriguez-Seijo et al., 2017), but the concentration of MPs in the gut was not significantly different between treatments nor significantly different from the bulk soil concentration, so there was no evidence of accumulation of MPs in the earthworm bodies (Hodson et al., 2017). Chen et al. (2020) assume an accumulation of MP takes place in Eisenia fetida, based on an observed increase of MP concentrations in the casts in the course of 4 weeks. Huerta Lwanga et al. (2017a) supposed an accumulation of MPs in the food chain as the concentration of MPs in chicken gizzards is strongly increased compared to that in the earthworm casts in the same experiments. However, mainly the amount of large particles, i.e. macroplastics, in the gizzards was very large, thus it seems likely that the chicken directly fed on plastics and an accumulation through the food chain cannot be proven with the current knowledge and should be further investigated.
Several studies did not find significant negative effects of MPs on earthworms’ avoidance behaviour (Judy et al., 2019), nor on growth (Hodson et al., 2017; Rodriguez-Seijo et al., 2017; Judy et al., 2019; Wang et al., 2019c), mortality (Hodson et al., 2017; Rillig et al. (2017); Rodriguez-Seijo et al. (2017); Judy et al. (2019); Prendergast-Miller et al. (2019) or reproduction (Huerta Lwanga et al., 2016; Rodriguez-Seijo et al., 2017). However, other studies do show adverse effects of the uptake of MP in different degrees and on different aspects of earthworms’ fitness: A reduced growth was shown by Cao et al. (2017) for *Eisenia Fetida* and the mortality increased at an exposure of concentrations ≥10000 mg kg\(^{-1}\) dry soil. At lower concentrations no significant effects were found. The growth of *Aporrectodea rosea* was also inhibited when exposed to biodegradable polylactic acid, conventional high-density polyethylene (at 1000 mg kg\(^{-1}\) dry soil), and MP clothing fibers (at 10 mg kg\(^{-1}\) dry soil) (Boots et al., 2019). Huerta Lwanga et al. (2016) showed a decrease in growth and increased mortality at concentrations ≥28\% w/w in litter and after 60 days, though after just 14 days no mortality occurred in these experiments.

In some studies, additional effects such as histopathological changes or stress biomarkers were measured. For *Eisenia fetida* Chen et al. (2020) observed skin damage at 1500 mg MP kg\(^{-1}\) in soil, measured an increase in catalase activity and malondialdehyde content at 1000 mg kg\(^{-1}\) and at ≥1000 mg kg\(^{-1}\) acetylcholine esterase was significantly stimulated. Wang et al. (2019c) tested *Eisenia fetida* and found that MPs only increased the catalase and peroxidase levels as well as the level of lipid peroxidation and decreased the activity of superoxide dismutase and glutathione S-transferase at an exposure of 200000 mg kg\(^{-1}\) dry soil for 14 days. No discernible influence was found at 100000 mg kg\(^{-1}\). However, Rodríguez-Seijo et al. (2018) also found for *Eisenia fetida* a significant positive correlation of MP concentration with different biomarker responses: catalase, glutathione S-transferase, lactate dehydrogenase and thiobarbituric acid reactive substances. In addition, Rodríguez-Seijo et al. (2017) observed histological damage of the gut and occurrence of inflammatory processes as well as an increase of stress response indicators associated with MP exposure of *Eisenia andrei*. For *Lumbricus terrestris* Prendergast-Miller et al. (2019) showed an increase in metallothionein expression at an exposure with ≥1000 mg kg\(^{-1}\) dry soil and a decrease in heat shock protein 70 at a concentration of ≥10000 mg kg\(^{-1}\).

Due to the large differences in experimental conditions – e.g. size of the MPs, addition of MPs to soil or to litter, duration of experiments, earthworm species – the current knowledge is not sufficient to detect whether there is a threshold in MP size and concentration at which the MP become harmful for earthworms and how this threshold differs for different earthworms species and MP shapes. The results of Huerta Lwanga et al. (2016), who found no effects of MPs on earthworms at 14 days, but significant influence on growth and mortality after 60 days, indicate the importance of longer measurements. This is consistent with Pelosi et al.
(2015), who concluded that the influence of pesticides on earthworm communities should be tested in long term field experiments.

Earthworms activity also increased the transport of MP in soil columns to deeper soil layers (Rillig et al., 2017; Yu et al., 2019; Huerta Lwanga et al., 2017b). The smaller the MP the stronger the transport. Particles are transported both actively – ingested and later cast out – and passively after attachment to the earthworm's body or by water flow through the biopores. As Huerta Lwanga et al. (2018) showed that the bacteria in the gut of Lumbricus terrestris can decompose MPs, it seems likely that particles taken up at the surface are egested as smaller particles in deeper layers.

Microplastics might well serve as a vector for contaminant transport to soil organisms. Though adsorption on plastics was seen to be lower than on the soil matrix, the desorption of Zn was seen to be higher in synthetic earthworm guts. However, there was no measurable negative effect of Zn or the PE on Lumbricus terrestris (Hodson et al., 2017). Wang et al. (2019b) studied the influence of MP on arsenic uptake and negative effects on Metaphire californica and concluded that MPs decreased the uptake of arsenic and that MPs reduced the influence of arsenic on the gut bacterial communities. Rodriguez-Seijo et al. (2019) showed altered enzyme activities and enhanced avoidance behavior in face of LD-PE pellets spiked with the insecticide chlorpyriphos. Yang et al. (2019a) studied the influence of MPs on the transport of glyphosate, however they mainly showed that the glyphosate transport was increased by earthworm activity, the role of MPs in this transport could not be determined with this study. These studies show that MP might have very different influences on the uptake and the adverse effects of different pollutants on earthworms and further investigation is needed in order to understand the influence of MPs on pollutant transport.

In contrast to the recently well-researched Lumbricidae, a near relative, the family of Megascolecidae (giant earthworms), is not yet mentioned in literature. Another branch within the Annelida, the small Enchytraeidae (potworms), were shown to suffer adverse effects on body weight and microbiome with PS microspheres (0.05 to 0.1 µm) at concentrations of ≥10 % w/w within their food source, but an unexpected increase of reproduction at 0.5 % w/w (Zhu et al., 2018b). The reproduction was reduced at abnormal concentrations of 90 g kg⁻¹ dry soil of polyamid particles (13 to 150 µm), but not with PVC (Lahive et al., 2019).

The edaphon of semisubhydric soils is often treated as a marginal group between the area of interest of soil and aquatic scientists. As a highly diverse soil biocenosis outside the focus of this paper, the benthos along seashores and fresh waters is also affected by MPs and should therefore be shortly mentioned by reviewing the lugworm Arenicola marina, a well examined deposit-feeder of the tidal flats. In situ, MP accumulates within its tissue and feces (Van Cauwenbergh et al., 2015). In laboratory experiments, PS particles ≥500 µm were avoided as food-source and passively translocated within the sediment at concentrations of ~2 g kg⁻¹.
(Gebhardt and Forster, 2018), but were measured within the feces at ~74 g kg\(^{-1}\) causing effects on feeding activity and body weight with no influence on the survival rate (Besseling et al., 2012). PS microspheres ≤30 µm remained within the animal without any adverse effects regardless of particle size (Van Cauwenberghe et al., 2015). Other studies found adverse effects on respiration, energy reserves, feeding, egestion and casting after uptake of PVC particles ≤478 µm at different sediment concentrations of >2 g kg\(^{-1}\), but neither on biomass and survival nor due to HD-PE (Wright et al., 2013; Green et al., 2016). There is further a difficulty in distinguishing between the adverse effects of MPs and substances adsorbed on or leached from MPs (Besseling et al., 2012). When adding PCB-spiked PE to mud flat sediment with concentrations up to 5000 mg kg\(^{-1}\) dry mass, there was no significant change of survival rate or body weight. The decreased feeding activity and heap mass could be attributed to increasing plastic concentrations, but not to enhanced PCB bioaccumulation via PE uptake (Besseling et al., 2017). However, all these studies found adverse effects at MP concentrations orders of magnitude above natural values.
Table 4: Microplastic studies on Lumbricidae (p=particles, ms=microspheres, b=beads, f= fibers, m=micromicelles, [b] N/A=information not available). Concentrations refer to mg kg⁻¹ dry soil, if not specially marked.

| Organism               | Experimental environment | Plastic type | Aging | Coating | Additives | Shape | Size span [µm] | Concentrations | Exposure time | Passive transport | Active uptake | Bioaccumulation dynamics | Measured adverse effects | Reference                                |
|------------------------|--------------------------|--------------|-------|----------|-----------|-------|----------------|----------------|---------------|-------------------|---------------|------------------------|-------------------------------|-----------------------------------------|
| Lumbricus terrestris   | mesocosm                 | PE           | washed | (C,H₃,C,H₃) | N/A       | N/A   | p              | <150           | 0.60% w/w (litter) | 14 d / 60 d | yes                | yes            | N/A                     | at 60 d, ≥29% w/w survival k, growth f | Huerta Lwanga et al. (2016)               |
| Eisenia fetida         | glass beaker             | PS           | N/A   | N/A       | N/A       | ms    | 50.80          | 0.20000        | 30 d          | N/A                | yes            | N/A                    | ≥5000: survival f                  | Cao et al. (2017)                          |
| Lumbricus terrestris   | bag                      | PE           | N/A   | N/A       | N/A       | p     | 0.92±1.09 mm²  | 3500           | 28 d          | N/A                | yes            | N/A                    | no                            | Hodson et al. (2017)                         |
| Lumbricus terrestris   | home yard                | diverse      | yes   | N/A       | N/A       | N/A   | N/A            | 0.87±1.9 items g⁻¹ | N/A           | yes                | conc. in chickens  > 48 h / 56 d | N/A                      | Huerta Lwanga et al. (2017a)                    |
| Lumbricus terrestris   | mesocosm                 | PE           | washed | (C,H₃,C,H₃) | N/A       | N/A   | p              | <150           | 0.60% w/w (litter) | 14 d          | yes                | N/A            | N/A                    | ≥1000: : weight f                  | Huerta Lwanga et al. (2017b)                   |
| Eisenia fetida         | mesocosm                 | PE           | N/A   | N/A       | N/A       | p     | 710.2800       | 750 µg on 2.5 kg soil | 21 d          | yes                | yes            | N/A                    | no                            | Rillig et al. (2017)                          |
| Eisenia andrei         | glass beaker             | PE           | N/A   | N/A       | N/A       | p     | 150            | 0.1000         | 28 d          | N/A                | yes            | N/A                    | ≥62.5: intestinal damage             | Rodriguez-Seijo et al (2017)                  |
| Eisenia fetida         | glass bottle             | PE           | washed | (C,H₃,C,H₃) | N/A       | N/A   | p              | 150            | 7% w/w (litter) | 10000          | N/A                | yes            | N/A                    | N/A                                          | Huerta Lwanga et al. (2018)                     |
| Aporrectodea rosea     | mesocosm                 | PLA, PE      | washed | (EIOH)    | N/A       | N/A   | N/A            | 1000           | 28 d          | N/A                | yes            | N/A                    | ≥125: altered enzyme activity         | Rodriguez-Seijo et al (2018)                   |
| Eisenia fetida         | mesocosm                 | PE           | washed | (EIOH)    | N/A       | N/A   | p              | 1000           | 28 d          | N/A                | yes            | N/A                    | growth f                          | Boots et al. (2019)                          |
| Eisenia fetida         | mesocosm                 | PE           | washed | (EIOH)    | N/A       | N/A   | f              | ≤2000          | 48 h / 56 d   | N/A                | N/A            | N/A                    | no                            | Judy et al. (2019)                           |
| Lumbricus terrestris   | bag                      | PE           | N/A   | N/A       | N/A       | mf    | Φ40.7±8.8 x 361.6±367.0 | 0.1000          | 35 d          | N/A                | yes            | N/A                    | ≥1000: metallothionein expression f              | Prendergast-Miller et al. (2019)              |
| Eisenia fetida         | mesocosm                 | LD-PE        | washed | (EIOH)    | N/A       | N/A   | p              | 250.1000       | 40 items on 0.5 kg soil | 14 d          | N/A                | N/A            | N/A                    | ≥1000: heat shock protein 70 f              | Rodriguez-Seijo et al (2019)                  |
| Eisenia fetida         | mesocosm                 | LD-PE        | washed | (EIOH)    | N/A       | N/A   | p              | 250.1000       | 40 items on 0.5 kg soil | 14 d          | N/A                | N/A            | N/A                    | ≥1000: altered enzyme activity with CPF: altered enzyme activity, avoidance of MPs | Rodriguez-Seijo et al (2019)                  |
| Lumbricus terrestris   | bag                      | PE           | N/A   | N/A       | N/A       | mf    | Φ40.7±8.8 x 361.6±367.0 | 0.1000          | 35 d          | N/A                | yes            | N/A                    | ≥1000: heat shock protein 70 f              | Prendergast-Miller et al. (2019)              |
| Lumbricus terrestris   | bag                      | PE           | N/A   | N/A       | N/A       | p     | 300             | 0.20000        | 14 d          | N/A                | yes            | N/A                    | ≥200000: altered enzyme activity         | Wang et al. (2019c)                          |
| Lumbricus terrestris   | mesocosm                 | PE           | washed | (EIOH)    | N/A       | N/A   | 0.7% w/w (litter) | 14 d          | N/A          | N/A                | yes            | N/A                    | ≥1000: heat shock protein 70 f              | Yang et al. (2019a)                          |
| Lumbricus terrestris   | bag                      | PE           | washed | (EIOH)    | N/A       | N/A   | p              | 150            | 7% w/w (litter) | 10000          | N/A                | yes            | N/A                    | ≥1000: heat shock protein 70 f              | Yu et al. (2019)                             |
| Lumbricus terrestris   | PE and div. - grade       | PE           | washed | (EIOH)    | N/A       | p     | 1.5x1.5 cm²  | 0.1500         | 28 d          | N/A                | yes            | yes                    | ≥1000: heat shock protein 70 f              | Zhang et al. (2018)                          |
| Eisenia fetida         | PE and div. - grade       | PE           | washed | (EIOH)    | N/A       | p     | 1.5x1.5 cm²  | 0.1500         | 28 d          | N/A                | yes            | yes                    | ≥1000: heat shock protein 70 f              | Chen et al. (2020)                           |

**Lumbricidae**
| organism               | experimental environment | plastic type | aging | coating | additives | size span [µm] | concentrations | exposure time | passive transport | active uptake | bioaccum. dynamics | measured adverse effects | reference          |
|------------------------|--------------------------|---------------|-------|---------|------------|----------------|----------------|---------------|-------------------|---------------|---------------------|------------------------|-----------------|
| Enchytraeus crypticus  | Petri dish               | PS            | N/A   | N/A     | mb         | 0.05..0.1     | 0..10% w/w (food) | 7             | N/A               | yes           | N/A                | at 0.5% w/w: reproduction ‡  
|                        |                          |               |       |         |            | 0.1..10       | 20000..120000 | 50000         | 20 h / 21 d       | N/A           | ≥10% w/w: microbiome, weight ‡   
|                        |                          |               |       |         |            | 10..150       | 106..150      | 90000         | N/A               | yes           | ≥90000: reproduction ‡         | Zhu et al. (2018c) |
| Enchytraeus crypticus  | microcosm                | PA            | N/A   | N/A     | fluorescence | 13.150..150   | 100..200       | 135000        | 200 h / 21 d     | N/A           | yes                | N/A                    | Lahive et al. (2018) |
| Arenicola marina †     | in situ                  | N/A           | N/A   | N/A     | ms         | N/A           | 10..90         | 10000..50000   | 300       | yes               | 1.2±2.8 items g⁻¹        | Cauwenberghe et al. (2015) |
|                        | liquid culture           | PS            | no    | N/A     |            | N/A           | 10..90         | 10000..50000   | 300       | yes               | N/A                    | Cauwenberghe et al. (2015) |
| Arenicola marina †     | mesocosm                 | PE            | N/A   | PCBs    | fluorescence mb | 10..180       | 0.50000        | 28 d           | N/A               | yes           | N/A                | ≥10000: feeding ‡, weight ‡      | Besseling et al., (2012) |
|                        |                          |               |       |         |            |               |               | 28 d           | N/A               | yes           | N/A                | >2000: respiration ‡, casting ‡, no | Green et al. (2016) |
| Arenicola marina †     | mesocosm                 | HD-PE         | N/A   | N/A     |            | 9..478        | 0.20000 mg kg⁻¹ | 31 d          | N/A               | N/A           | N/A                | ≥74000: feeding ‡, weight ‡      | Besseling et al., (2012) |
|                        |                          |               |       |         |            | 3..16         | 0..100         | 31 d          | N/A               | N/A           | N/A                | >2000: respiration ‡, casting ‡, no | Besseling et al., (2017) |
| Arenicola marina †     | mesocosm                 | PVC           | N/A   | N/A     | not leaching | −130          | 0..50000       | 28 d           | N/A               | N/A           | N/A                | ≥10000: energy reserves ‡, ≥50000: feeding ‡, egestion ‡, casting ‡ | Wright et al. (2013) |
3.4 Further invertebrates

As part of the microfauna, the phylum Nematoda (nematodes or roundworms) is an ecologically important branch containing >25000 species (Zhang, 2013) in freshwater, marine, endobiotic and soil habitats. Due to their diverse trophic interactions nematodes hold a central position in both bottom-up and top-down controlled food webs (Yeates, 2001; Ferris, 2010) and thus most likely the uptake and transfer of MP.

Active feeding of adults and larvae of different species on 0.5 to 6 µm PS/latex microspheres (the size of their bacterial prey) was proven by Nika et al. (2016) and Fueser et al. (2019). However, most MP experiments on Nematodes are based on the bacterial-feeding model organism Caenorhabditis elegans. Kiyama et al. (2012) showed the favored uptake of PS microspheres with sizes of 0.5 to 3 µm by adult and 0.5 µm by larval C. elegans. The ingestion of MP decreased in the presence of bacteria as the natural food source.

When larval stages and adults ingested PS between 0.05 and 5 µm within an aqueous suspension or on agar plates, adverse effects such as oxidative stress, neurodegeneration, intestinal and DNA damage or dysfunction in motility, growth, life span, defecation, reproduction or energy metabolism appeared from a wide spectrum of concentrations from ≥1 µg l⁻¹ up to ≥86.3 mg l⁻¹ (Zhao et al., 2017; Dong et al., 2018; Kim et al., 2019; Lei et al., 2018a; Lei et al., 2018b; Qu et al., 2019a). These effects are not seen below 1 µg l⁻¹ (Qu et al., 2019b), and are enhanced due to amino modifications on micropshere surfaces (Qu et al., 2019c). The incubation on agar plates with PE, PP and PVC particles <70 µm caused similar influences on survival, fertility, brood size and intestinal function (Lei et al., 2018b). Leachates from soils amended with 5 mg kg⁻¹ dry soil of HD-PE and PVC decreased reproduction in laboratory cultures, but there was no effect shown on survival and after application of PET (Judy et al., 2019). Furthermore, silica nanoparticles (0.05 µm) are not only taken up orally but also via the vulva and spermathecae and migrate into gonad cells (Scharf et al., 2013), This process was confirmed for PS nanoparticles with the potential of a transfer to the progeny (Zhao et al., 2017).

The clear adverse effects of these studies are limited in their representativity by a narrow restriction to liquid cultures and a single model organism. Broader studies like on prominent soil-born nematodes such as Acrobeoides buetschlii (Frey, 1971) are still lacking. When assuming in first proximity mg l⁻¹ solution = mg kg⁻¹ dry soil, the applied concentrations between 0.001 and 86.8 mg l⁻¹ match lower levels of soil contamination.

Feeding studies on the phylum Rotifera with MPs are fully based on PS microbeads and model organisms of the planktonic genus Brachionus. However, this data can carefully be transferred to soil environments as also soil rotifers are aquatic organisms living in water-filled pores and waterfilms. Different Brachionus sp. ingest microbeads <10 µm with strong preference for particles the size of their natural food source, namely bacteria and algae with
2 to 5 µm in diameter (Vadstein et al., 1993; Heerkloß and Hlawa, 1995; Baer et al., 2008; Jeong et al., 2016). The uptake appears to be selective as microbeads are fewer incorporated than bacteria and algae (Vadstein et al., 1993). The egestion of particles ≤0.5 µm is hindered compared to 6 µm (Jeong et al., 2016). In suspension, microbeads ≤0.5 µm cause adverse effects on fertility and life span at ≥0.1 mg l\(^{-1}\) as well as oxidative stress and less growth at ≥10 mg l\(^{-1}\) (Jeong et al., 2016; Sun et al., 2019).

Terrestrial mollusks comprise snails and slugs within the class of **Gastropoda**. These grazers feed on bacterial biofilms, fungi and plant tissue (Parkyn and Newell, 2013). Studies on terrestrial species are sparse, but data on the benthic *Littorina* sp. imply passive transport and non-selective MP uptake by feeding on surfaces with contaminated feces and mucus trails of other snails (Gutow et al., 2019). With focus on *benthic* snails, Imhof and Laforsch (2016) found no significant influence on growth parameters and fertility of juveniles and adult *Potamoeryrgus antipodarum* even when a food source with 70 % w/w of 5 to 600 µm sized fragments was given (a mixture of PA, PC, PET, PS, PVC). In contrast, adverse effects were found in recent work on the terrestrial snail *Achatina fulica*, that showed uptake and complete gastrointestinal passage within 48 h with partial degradation of PET fibers (appr. 1258x76 µm), but reduced excretion and food intake as well as increased oxidative stress at concentrations of ≥0.01 g kg\(^{-1}\), ≥0.14 g kg\(^{-1}\) and ≥0.71 g kg\(^{-1}\) dry soil, respectively (Song et al., 2019).
Table 6: Microplastic studies on nematods (ms=microspheres, fr=fragments, np=nanoparticles, mb=microbeads, ms=microspheres, ox.=oxidative, [8] N/A=information not available). Concentrations refer to mg kg⁻¹ dry soil, if not specially marked.

| organism                  | experimental environment | plastic type       | aging | coating | additives | shape | size span [µm] | concentrations | exposure time | passive transport | active uptake | bioaccum. dynamics | measured adverse effects | reference            |
|----------------------------|--------------------------|---------------------|-------|---------|-----------|-------|---------------|----------------|---------------|-------------------|---------------|-------------------|--------------------------|-----------------------|
| Caenorhabditis elegans     | agar plate               | PS                  | N/A   | carboxyl| sulfate amino fluorescence | ms     | 0.1..6.6      | N/A            | 0.5..2 h      | N/A               | yes           | 0.5..3 µm          | N/A                      | Kiyama et al. (2012)   |
| Caenorhabditis elegans     | liquid culture           | PS                  | N/A   | N/A     | carboxyl fluorescence | ms     | 0.1           | 0.001..10 mg l⁻¹ | 4.5 d         | N/A               | Yes           | N/A               | 0.01 mg l⁻¹: motivity ↓, growth ↓, defecation ↓, within gonads | Zhao et al. (2017)     |
| Caenorhabditis elegans     | liquid culture           | PS                  | N/A   | N/A     | N/A 10 mV fluorescence | ms     | 0.1           | 0.0001..0.001 mg l⁻¹ | N/A           | N/A               | Yes           | N/A               | 0.001 mg l⁻¹: motivity ↓, ox. stress ↑ | Dong et al. (2018)    |
| Caenorhabditis elegans     | liquid culture           | PS                  | N/A   | N/A     | preservatives, fluorescence | ms     | 0.05..0.2     | 0.01..8.68 mg l⁻¹ | 24 h         | N/A               | Yes           | N/A               | 17.3 mg l⁻¹: motivity ↓, reproduction ↓, metabolic dysf. | Kim et al. (2019)     |
| Caenorhabditis elegans     | liquid culture           | PS                  | N/A   | N/A     | carboxyl fluorescence | ms     | 0.1           | 0.001..1 mg l⁻¹  | N/A           | N/A               | Yes           | N/A               | 0.01 mg l⁻¹: neurodegeneration | Qu et al. (2019a)     |
| Caenorhabditis elegans     | liquid culture           | PS                  | N/A   | N/A     | N/A no N/A | ms | 0.1..5       | 1 mg l⁻¹       | 3 d           | N/A               | Yes           | N/A               | 0.01 mg l⁻¹: motivity ↓, survival ↓, growth ↑, ox. stress ↑, neurotoxicity | Lei et al. (2018a)    |
| Caenorhabditis elegans     | agar plate               | PE, PP, PVC, PS     | no    | N/A     | N/A fluorescence | ms     | 0.1..200      | 0.5..10.0 mg m⁻³ | 2 d           | N/A               | Yes           | N/A               | 0.5 mg m⁻³: survival ↓, growth ↓, reproduction ↓, ox. stress ↑, intestinal damage, mainly 1µm: intestinal damage | Lei et al. (2018b)    |
| Caenorhabditis elegans     | agar plate               | PS                  | N/A   | N/A     | silica gel fluorescence | ms     | 0.1           | 2500 mg l⁻¹     | 7 d            | N/A               | Yes           | N/A               | within tissue and gonads | Scharf et al. (2013)  |
| Caenorhabditis elegans     | liquid culture           | HDPE, PET, PVC      | no    | N/A     | N/A np | no fr <2000 soil extract | 72 h | N/A           | N/A            | N/A               | N/A           | N/A               | reproduction ↓ | Judy et al. (2019)    |
| Caenorhabditis elegans     | agar plates              | latex               | N/A   | N/A     | fluorescent mb | 0.5 | N/A           | 30 min         | N/A           | N/A               | yes           | N/A               | N/A                      | N/A                   |
| Caenorhabditis elegans     | liquid culture           | PE, PP, PVC         | no    | N/A     | fluorescent | ms     | 0.5..6       | 3·10⁻⁴...10⁸ items l⁻¹ (~0.2..1200 mg l⁻¹) | 4.73 h        | N/A               | N/A           | N/A               | N/A                      | N/A                   |
| Caenorhabditis elegans     | liquid culture           | PS                  | N/A   | N/A     | N/A no amino | ms | 0.1          | 0.0001..0.001 mg l⁻¹ | N/A           | N/A               | N/A           | N/A               | 0.01 mg l⁻¹: reproduction ↓, DNA damage | Qu et al. (2019b)     |
| Caenorhabditis elegans     | liquid culture           | PS                  | N/A   | N/A     | N/A no amino | ms | 0.1          | 0.001..1 mg l⁻¹   | N/A           | N/A               | yes           | N/A               | 0.01 mg l⁻¹: reproduction ↓, DNA damage | Qu et al. (2019c)     |
| *Pristionchus pacificus*   | liquid culture           | PS                  | N/A   | N/A     | N/A no amino | ms | 0.1          | 0.001..1 mg l⁻¹   | N/A           | N/A               | yes           | N/A               | 0.01 mg l⁻¹: reproduction ↓, DNA damage | Qu et al. (2019c)     |
| *Acrobolosida nanus*       | liquid culture           | PS                  | N/A   | N/A     | N/A no amino | ms | 0.1          | 0.001..1 mg l⁻¹   | N/A           | N/A               | yes           | N/A               | 0.01 mg l⁻¹: reproduction ↓, DNA damage | Qu et al. (2019c)     |

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Table 7: Microplastic studies on Rotifera and Gastropoda (ms=microspheres, mb=microbeads, fr=fragments, f=fibers, ox.=oxidative, pref.=preferential, p=planctic, b=benthic, N/A=information not available). Concentrations refer to mg kg⁻¹ dry soil, if not specially marked.

| organism              | experimental environment | plastic type | aging | coating | additives | shape | size span [µm] | concentrations | exposure time | passive transport | active uptake | bioaccum. dynamics | measured adverse effects | reference                  |
|-----------------------|--------------------------|--------------|-------|---------|-----------|-------|----------------|----------------|---------------|-------------------|----------------|---------------------|----------------------------|---------------------------|
| Brachionus plicatilis ² | liquid culture           | PS           | N/A   | carboxyl| fluorescence| ms   | 1.6..20        | 5·10⁷ µm⁻¹ l⁻¹  | 35 min        | N/A               | ≤10 µm | pref. 4.5 µm        | N/A                        | Bear et al. (2008)          |
| Brachionus plicatilis ² | liquid culture           | latex        | N/A   | N/A     | fluorescence| mb   | 0.3..3.1       | 3·10⁻⁷..7·10⁸ items⁻¹ | 20 min | N/A               | yes          | pref. 22 µm        | N/A                        | Vadstein et al. (1993)     |
| Brachionus koreanus ² | liquid culture           | PS           | no    | N/A     | fluorescence| mb   | 0.05..6        | 0..20 mg⁻¹      | 1 d           | N/A               | yes          | egestion rate 0.05 µm < 0.5 µm < 6 µm | ≤0.5 µm, >0.1 mg⁻¹; reproduction ↓, survival ↓ | Jeong et al. (2016)        |
| Brachionus plicatilis ² | liquid culture           | PS           | N/A   | N/A     | N/A       | mb   | 0.07..7        | 0..20 mg⁻¹      | N/A           | N/A               | yes          | N/A                | ≤0.07 µm, >10 mg⁻¹; reproduction ↓, growth ↓ | Sun et al. (2019)          |
| Brachionus quadridentatus ² | liquid culture         | PS           | N/A   | N/A     | N/A       | ms   | 2..10          | 2..10 µm⁻¹      | 8..10 d       | N/A               | pref. 3.5 µm pref. 2 µm | N/A        | N/A                | N/A                        | Heerkloß and Hlawa (1993)  |
| Littorina littorea ²   | microcosm               | PMMA         | N/A   | N/A     | fluorescence| ms   | 2..10          | N/A            | 18 h          | N/A               | yes          | N/A                | N/A                        | Gutow et al. (2019)        |
| Potamopyrgus antipodarum ² | aquarium               | PET, PVC, PA, PC | N/A | N/A     | no        | fr   | 0.70% w/w (food) | s141 d       | N/A           | yes               | N/A        | N/A                | N/A                        | Imhof and Laforsch (2016)  |
| Achatina fulica ²      | mesocosm                | PET           | N/A   | no / stained| f       | approx. 1258x76 µm | 10..710       | 28 d          | N/A               | yes          | excretion after 48 hours | ≥140: food intake ↓, 210: excretion ↓, ≥170: ox. stress ↓, gastrointestinal damage | Song et al. (2019)         |
3.5 Vertebrates

Different taxa of the class of Amphibia have a predator function within the edaphic food web (e.g. preying on invertebrates) (Hebrard et al., 1992). While no data on the reaction to soil MPs are available neither for the legless Gymnophiona nor for adults of the order Anura, sparse data on tadpoles of aquatic frogs suggest uptake followed by regular excretion of PS microspheres as shown with *Xenopus tropicalis* (Hu et al., 2016). Further, there exist no data on the families Serpentes (snakes) and Anguidae within the class of Reptilia, residing at the outer rim of the food web.

Within the broad field of Mammalia, studies on MP ingestion are sparse and focus on mice as a rodent model organism. Feeding of mice with PS microspheres of 1 to 14 µm in concentrations of $1.49 \times 10^6$ to $4.55 \times 10^7$ particles at a volume of $10 \text{ ml kg}^{-1}$ body weight for 4 weeks showed no adverse effects (Stock et al., 2019). In contrast, longer exposition (6 weeks) with lower concentrations of particles with the same shape and size range changed the mouse microbiome and caused metabolic and intestinal dysfunction (Lu et al., 2018; Jin et al., 2019), which comes along with bioaccumulation within organs (Yang et al., 2019b). These studies are regularly conducted with passive feeding and exclude active foraging on perceptible plastic particles. However, the uptake via prey or feeding on contaminated roots and litter is highly probable. Further Rodentia – Cricetidae (hamsters, lemmings, voles), Bathyergidae (blesmols, mole-rats), Octodontidae as well as Spermophilus (ground squirrels) and Marmota (marmots) within the family of Sciuridae – were not yet studied, just as other mammalian (sub)orders like Chrysochloridae (golden moles), Cingulata (armadillos), Macroscelidea (elephant shrews), Notoryctemorphia and Peramelemorphia.
Table 8: Microplastic studies on Anura (An.) and Rodentia (ms=microspheres, =aquatic, N/A=information not available).

| organism     | experimental environment | plastic type | aging | coating | additives | shape | size span [µm] | concentrations | exposure time | passive transport | active uptake | bioaccum. dynamics | measured adverse effects          | reference                      |
|--------------|--------------------------|--------------|-------|---------|-----------|-------|----------------|----------------|---------------|-------------------|---------------|-------------------|-------------------------------|-------------------------------|
| Xenopus tropicalis a | Petri dish             | PS           | N/A   | N/A     | fluorescence | ms    | 1..10          | 100..10⁰ items l⁻¹ (55·10⁻⁴..55 mg l⁻¹) | 48 h          | N/A                      | yes            | egestion within days | N/A                          | Hu et al. (2016)               |
| transgenic mice | in vivo                | PS           | N/A   | N/A     | sulfate    | fluorescence | ms    | 4             | 4.55·10⁰ items per mouse (0.025 mg per mouse) | 28 d          | N/A                      | yes            | N/A                          | no                           | Stock et al. (2019)              |
| mice         | in vivo                | PS           | N/A   | N/A     | sulfate    | fluorescence | ms    | 5             | 0.1..1 mg l⁻¹ (food) | 42 d          | N/A                      | yes            | N/A                          | ≥0.1 mg l⁻¹: microbiome, metabolic dysfunction | Jin et al. (2019)                     |
| mice         | in vivo                | PS           | N/A   | N/A     | ms         | 0.5..50       | 0.1..1 mg l⁻¹ (food) | 35 d          | N/A                      | N/A            | N/A                          | N/A                          | Lu et al. (2018)                     |
| Mus musculus | in vivo                | PS           | N/A   | N/A     | fluorescence | ms    | 5..20         | 200 mg l⁻¹ (food) | 28 d          | N/A                      | yes            | 8x, 8±5 and 0.71±0.14 mg kg⁻¹ body weight | N/A                          | Yang et al. (2019b)                |
4 Synthesis

4.1 Summarized observations

Our systematic search comprised recent research on the interaction of soil organisms with MP, but also studies with focus on feeding experiments, that are published much earlier than the awareness on plastic in the environment appeared. The numerous studies found with focus on the ingestion of MPs consistently showed the active uptake by diverse soil organisms with few exceptions spread over the whole branch of invertebrates. In addition, also studies on adverse effects caused by the intake of MP contaminated food (e.g. of food pallets by dung beetles) imply the ingestion into the test organism. Distinct size preferences are observed in dung beetles, nematodes, rotifers and ants showing that mainly particles are ingested, that are small enough to enter the gastrointestinal tract. In contrast, active comminution by gnawing on larger particles was tested only for a few taxa and confirmed for woodlice, termites and mealworms, and in the case of earthworms only after initial weathering.

After the ingestion, MP is translocated actively until excretion or death of the transporting organism, which was only directly shown in experiments with earthworms. The passive transport by attachment, dragging and pushing was investigated in a few experiments with earthworms, mites and springtails that partly worked without soil substrate and consistently showed positive results.

After exposition to MP, a pattern of adverse effects can be seen: Across various taxa, altered microbiomes, reduced motility, body mass, fertility and life span as well as increased oxidative stress and metabolic malfunctioning occur in different combinations mainly due to µm-sized MP in and above the whole known natural range of concentrations. For some taxa such as Nematodes, Gastropoda and Rotifera these effects appear at natural and increased MP concentrations (<100 mg kg\(^{-1}\) dry soil), for Collombola and Lumbricidae at concentrations like in highly contaminated sites (≥1000 mg kg\(^{-1}\) dry soil) and for Enchytraeidae, Arenicola marina and in further experiments with earthworms at implausibly high values. The data show a tendency, that the effects occur at lower concentrations, when the added particles are smaller. Small sized particles also provide the highest surface/volume ratio and thus the highest reactive surface per weight.

Most studies work with defined increasing MP concentrations and particle sizes in soil substrates and food sources, which can be used to determine relationships between environmental concentrations and adverse effects. However, the lack of information about intake rates, grades of accumulation and effective prey-predator transfer leads to a gap within the chain of explanation for toxic effects on the soil organisms. In some experiments, the intestinal passage of MP and sizes preferably retained within the gut were shown, but there are no experiments that could demonstrate quantitative bioaccumulation. In contrast,
quantification of the retained and egested MP particle size fractions might be biased due to
gnawing and intestinal comminution as shown for woodlice, termites, mealworms, snails and
earthworms.

In order to improve our understanding of processes underlying adverse effects of MP on soil
organisms, data on ingestion rates, dwell times, biodegradation and egestion rates are
important bricks e.g. to reveal bioaccumulation dynamics. However, there are only a few data
on biodegradation (mealworms, snails, earthworms), egestion (rotifers, frogs, snails,
earthworms) and remaining concentrations in the body (lugworm, mice, earthworms).
4.2 Limitations of previous studies

The available studies worked with items within the full size span of micro- and nanoplastics (≤5000 µm). Approximately 72 % of the experiments used microplastic (0.1 to 5000 µm), only 6 % nanoplastic (<0.1 µm), 10 % included macroplastic (>5000 µm) and 12 % used microplastic of undefined size. When MP ≥50 µm was applied, mainly particles and fragments made of PE and PVC were used, whereas PS/latex microspheres were mainly applied for sizes ≤10 µm (Table 1). The latter are readily available, highly standardized and are mostly used with fluorescent dyes and either without additional functional groups, carboxylated or, more rarely, with amino or sulfate groups. However, there are indications that the spectrum of particle type and shape used in experiments does not correspond to the properties of particles in soils. In different natural as well as agriculturally and industrially contaminated terrestrial and semisubhydric sites, fibers and fragments of PE and PP, mostly ≤100 µm, were much more abundant than PVC, PET and PS items (Claessens et al., 2011; Vianello et al., 2013; Nor and Obbard, 2014; Naji et al., 2017; Zhang and Liu, 2018; Li et al., 2018a). This is probably caused by high loads of MP fibers in discharged waste water and sewage sludge, which is used in agricultural sites worldwide (Mahon et al., 2016; Li et al., 2018b). It is likely that shape plays an important role for the ingestion of MP items. Unfortunately, we did not find studies that have carried out a complete classification of sampling sites according to plastic origin, size and type, that could help to evaluate differences between former experimental and natural plastic composition to achieve the most realistic experimental conditions. Little knowledge about the size distribution of MP in soils furthermore complicates the determination of realistic concentrations for the addition of a certain particle size spectrum. All reviewed studies either arbitrarily set their applied concentrations or had to base them on measurements of total specific MP masses, regardless of how much of this mass is in the tested size range. This may lead to a false estimation of total adverse MP concentrations.

In contrast to particle type and shape, the documentation of chemical properties of MP samples in most of these studies is fragmentary. Some experiments explicitly mentioned that the added plastic was unweathered, whereas most studies lack information about the degree of aging implying that unweathered items were used. Only a few experiments involved aging of MP, but without comparison to results of natural weathering (Tsunoda et al., 2010; Gebhardt and Forster, 2018). That is in conflict with natural conditions, as plastic that remains within the soil after littering, sewage sludge application or plastic mulching shows signs of weathering, e.g. modified carbonyl indices (Andrady, 2017), while unweathered soil MP might be rare. In addition, Zhang et al. (2018) showed that earthworms actively comminute only weathered bioplastics. In experiments using PS microspheres, carboxylation is often used to imitate a reduced hydrophobicity due to weathering. However, according to manufacturer information microplastics only have little influence on hydrophobicity.
Weathering of MP surfaces within soils comes along with biofilm growth and adsorption of organic molecules, which could potentially affect the attractiveness or toxicity for grazers and other organisms. Such coatings were applied only in a few cases (Besseling et al., 2017; Angotti et al., 2018; Gebhardt and Forster, 2018), but were not documented in most studies. Similarly, the type and concentration of additives such as flame retardants, anti-oxidants or stabilizers often remained undocumented, with exception of fluorescent dyes, that are well mentioned. The release of additives can have a harmful effect on the test organism, as shown for aquatic environments (e Silva et al., 2016). Some studies on the ingestion of MP by the soil mesofauna indicate that the diameter of the gastrointestinal tract is a useful upper size limit for added particles, as far as the organism is unable to crush them (Heerkloß and Hlawa, 1995; Holter, 2000; Holter et al., 2002; Holter and Scholtz, 2005; Baer et al., 2008; Fueser et al., 2019). However, using only ingestible particle sizes in their natural concentrations neglect the adverse effects of plastic leachates, which can also get into the soil solution and onto the mineral phase from larger particles and affect soil life.

The conditions of incubation differ considerably in terms of habitats and duration of exposure. In most studies, the exposure ranges from a few minutes to a few days in experiments with micro- and small mesofauna and hours to several weeks in experiments with large meso- and macrofauna and is mainly based on excretion or reproductive cycles. Long-term studies, which are indeed difficult to carry out in mesocosms, practically do not exist. However, certain adverse effects might only establish themselves after long term trials, as was shown for the influence of pesticides (Pelosi et al., 2015).

Some experiments were carried out in soil-free test environments such as liquid cultures or Petri dishes with nutrient solutions or a specific food source (nematods, rotifers, mice). Therefore, motivity is less restricted and feeding behavior can be altered compared to cultivation within soil environments. For example, the ingestion of MP by nematodes decreases in the presence of an alternative and more natural food source like bacteria, which can significantly reduce the bioaccumulation and thus the effective toxicity (Kiyama et al., 2012). This can lead to less consumption of MP in soil environments and an overestimation of the toxicity in liquid culture experiments. Also, all laboratory feeding experiments were carried out by use of only one species. The complexity of the food web in soils is thereby excluded and the potential accumulation from prey to predators still unexplored.
4.3 Pinpoints for future research

Most studies reviewed in this work have a pioneering role in MP research and, thus, are subject to some experimental limitations caused by an early state of knowledge. The adverse effects recently found are alarming, but must be considered under the restrictions named above. We propose the following points as part of a *modus operandi* for future MP research.

In past studies, particular adverse effects of MP were measured only for certain sizes, shapes, coatings, leachates or adsorbed substances (*Tables 2 to 8*). Experimental concentrations were assumed randomly or derived from cumulative concentrations of one or more MP types measured in natural soils (approx. 1 to some 1000 mg kg\(^{-1}\) dry soil), regardless of size. For those specific experiments coming, the spectrum of concentrations used should be adapted to the quantities of the size spectrum, that occurs within the soil. For future studies on mixed contaminations, we recommend to evaluate the overall adverse effects of PE, PP, PVC, PET, PU and PS to certain test organisms by use of typical MP-specific spans of concentration, size and shape distribution in natural soils or food samples. This previously requires well-structured data of appropriate MP type, shape and size for different soils in differently contaminated areas.

Experiments on adverse effects should be applied within soil matrices to allow the interplay of plastic, natural organic and mineral matter. The MP should be weathered, as plastic in soils underlie broad environmental aging. Pre-weathering of MP should therefor not only be performed in climate chambers (e.g. following DIN EN ISO 4892-2/3), but also include subsequent leaching and equilibration of additives or coatings within the soil matrix before the main experiment. Furthermore, the experimental design may consider coatings with biofilms or attractants and even particle color to regulate the preference of the test organisms.

Most detailed information about ingestion are available for dung beetles, nematods and earthworms, data on adverse effects on nematods, earthworms, lugworms and collembola. Future experiments should focus on a larger variety of ecologically relevant taxa like *Coleoptera, Formicidae, Acari, Oniscidea, Collembola, Lumbricidae, Enchytraeidae, Nematoda* and *Gastropoda*. The studies are recommended to conduct with emphasis on uptake, accumulation and key adverse effects like on survival rate, motility, growth and fertility as well as on the stability of the intestinal microbiome. Further studies with more than one test organism are important to foster our understanding of MP within certain food chains. Also long-term experiments might reveal adverse effects, which evolve slowly within populations. This may enable the assessment of the distribution and effects of MP within the food web and the resulting long-term impact on soil ecosystems.
5 Conclusion

Our review of 77 studies on the impact of microplastic on the soil fauna shows a considerable diversity and distribution of adverse effects within the soil tree of life. However, these effects have to be considered carefully, as many experiments did not use plastic matching properties within natural soils and found adverse effects only at concentrations like in highly contaminated soils or above. To elucidate effective concentrations and properties for short and long-term effects on soil faunal health, the most exact reproduction of plastic properties within the soil matrix and natural living conditions of the test organisms is necessary together with a better knowledge on common concentrations and size distributions of soil microplastic. For future experiments we therefore recommend to choose compositions of type, shape, size, concentration, grade of weathering, leachability and coating with biofilms and other organic matter as expected in the habitat to be examined. Furthermore, coming studies should include long-term exposure and food chain experiments to get a better look at the effect of even smaller MP concentrations and their enrichment within the food web. This may give us a better way of assessing the impact of global microplastic contamination on e.g. soil biodiversity, soil carbon cycles and soil quality.
Author contribution
Frederick Büks developed the review concept, collected data and prepared the manuscript except for earthworms. Nicolette Loes van Schaik did all the work on earthworms. Martin Kaupenjohann supervised the study by participating in structural discussions on the idea and concept of the paper as well as the final corrections.

The authors declare that they have no conflict of interest.

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Competing interests
The authors declare that they have no conflict of interest.
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What do we know about how the terrestrial multicellular soil fauna reacts to microplastic?

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Final responses to all referees plus marked-up version
Dear Referee #1

First I would like to express my sincere thanks to you for thoroughly reviewing our manuscript and for your very helpful and precise suggestions. In the following I will answer your points. Our corrections are marked-up with yellow numbers within the corrected manuscript at the end of this document.

Best regards,
Frederick Büks

Abstract

[1] Lines 20-21: “Most of the studies applied MP concentrations similar to amounts in slightly to very heavily polluted soils.” This sentence makes the reader expect that generally, the concentrations in the experimental environments are mostly the same as expected in the environment, but is this really the case? I would suggest showing the percentage of experiments with high microplastic exposure that is not representative of most soils.

Thanks a lot for this point. We now write: “About 58% of the studies thereby use inappropriate concentrations or units, but 42% applied MP concentrations similar to amounts in slightly to very heavily polluted soils.”

Introduction

[2] Line 53: Instead of “microbial decay”, I’d suggest “processing by soil organisms”, since this includes any process relevant for the generation of smaller plastic particles.

Done.

[3] Line 61: I’d suggest changing the sentence to “intensive use of plastic mulching and sewage sludge”, for the former, Huang et al. (2020) show an increase in microplastic by approx. 1 order of magnitude between fields with 5 and 24 continuous years of plastic mulching.

Done.

[4] Line 95: Suggest changing “feed on” to “inadvertently ingest”, otherwise it sounds like the organisms are actually able to metabolize the microplastics.

Done and reference added.

Search pattern

[5] The cut-off dates (time period that was considered) of the search should be mentioned somewhere.

Information added to this chapter (see the answer to referee #3)

[6] Figure 1: This figure shows the phylogenetic tree of edaphic fauna, rather than “edaphic tree of faunal life”.

Thank you. And done.

Data collection

[7] Line 113-122: I’ve been having some difficulties understanding the search methodology and table 8 (table 8 should be moved at the appropriate place to become table 1).

We moved the table to line 124 and mentioned that it contains the number of found studies. All table numbers were adjusted within the text.

It would be great if the authors could re-word this, specifying:

What does it mean that some combinations would have caused too much search effort?

It means e.g. that searching for a taxon only in combination with “PET” gives results for PET bottles for cultivation and experiments and also the “use” as pets, if the search is not case sensitive. We now tried to clarify this in our text.
“Organism-plastic” is not a type-shape combination.
Oh, yes, that’s right. Corrected.

What exactly does the number of studies in table 8 mean? The number of articles or single experiments (sometimes more than one taxon or plastic type is used in one article)?
The number counts for how often type-shape combinations were used in all reviewed experimental setups independently of organism.

Some articles are included that studied the uptake of macroplastics by organisms, mainly termites and ant species. It is reasonable to include these studies, but it should be mentioned more prominently, in the abstract and aims of the review, that macroplastics are included. Where macroplastics were used in the reviewed studies, the size was explicitly mentioned in the article text, so we do not see a necessity for elaborating the text. We did add a mention of macroplastics to the abstract.

Maybe also in the synthesis, a sentence about the proportions of experiments using macro-, micro-, and nanoplastic would be a helpful piece of information.
Now mentioned in “4.2 Limitations of previous studies”

[8] Tables 1-7: What does N/A mean in the tables? In some cases I assume “not analysed” (e.g., passive transport), but in other cases it should mean “not mentioned” (e.g., aging, coating, etc.) or “not observed” (e.g., measured adverse effects). I think this needs to be specified. Usually, N/A refers to “not applicable”, but this doesn’t fit in the tables.
In this work it means “(data) not available”. We marked it at the tables.

Synthesis

Lines 549-550: Could you cite the studies that imitated weathering in the described way?
We did so. Tsunoda et al. (2010) artificially aged their plastic by soaking in hot water at 90°C for 21 days, and then it was sanded/scratched with medium-grade paper prior to the test. Gebhard and Forster (2018) incubated particles in seawater for 4 weeks to stimulate the formation of biofilms.

[9] Lines 555-557: This is true, but it should be acknowledged that these additives are mainly present in commercial plastics, and therefore, mentioning of additives is not expected for “clean” microbeads specifically synthesized for the experiments. Nevertheless, the disadvantages of using these microbeads has been clearly discussed earlier in this section.
Done.

Conclusions

[10] Line 620-621: I am a little concerned about describing the results as “alarming”. Is it really? The following sentences actually refute this rather strong statement.
Replaced with “considerable”.

[11] Lines 624-629: I would suggest changing the sentence to: “To elucidate […], the most exact reproduction of plastic concentrations and properties […].” However, the difficulty here is that very scarce data of limited quality is available on concentrations of microplastic in soils, so a range of concentrations need to be used for future experiments in order to match the “real world” concentrations in soil, while expecting a decrease in uncertainty in analytic results in the future. Especially in the lower size ranges (<100μm) quantification is currently challenging. Therefore, little is known about size distributions occurring in soils. It might be worth mentioning this dilemma in a sentence.
Done.

[12] Technical corrections:
All done.
Dear PD Dr. Werner Kratz (referee #2) 

Thank you very much for your review. In the following I will try to answer your comments at my best. Our corrections are marked-up with **green numbers** within the corrected manuscript at the end of this document.

Best regards,
Frederick Büks

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1. **Line 53**: Is that only “microbial” decay?
   We agree, we will change this to “processing by soil organisms” as it is actually micro- as well as macroorganisms.

2. **Figure 1**: The taxonomic group “further Panarthropods” is placed centrally, the other groups are not.
   Done.

3. **Table 7**: The last three experiments within this table were conducted by feeding the mice with a MP suspension. You might write “(food)” behind the concentration data as in the other tables.
   Thanks a lot. Done.

4. **Line 507**: “Preferably” instead of “p preferrably”.
   Done.

5. **Table 8**: Could you explain the meaning of the numbers within the table. Are these the numbers of experiments with the named type-shape combinations?
   Yes. Please see the answer to referee #1 (Table 8 is now Table 1).

6. **Lines 549-507**: Is that proved that carboxylation of microspheres decreases hydrophobicity in an appreciable extent?
   We ask the manufacturers of Polyciences Europe GmbH, a leading producer of PS microspheres, and they said no. We added this important information to the review.
Dear Referee #3

Thank you very much for your critical review of our manuscript. It has helped us to see some points which still need clarification. In the following, we want to explain how we propose to adjust our article based on the reviewer’s comments and also explain why in some cases we do not agree with the reviewer’s proposed changes. Our corrections are marked-up with purple numbers within the corrected manuscript at the end of this document.

(1) First and foremost, please have the manuscript edited by a professional (!) native (!) biologist (!). The English of your text is largely understandable, but rough. Apart from annoying typos, I found sentences the meaning of which I only understood when trying to translate them to German (my native language). So, your text will heavily benefit from thorough native editing.

Rereading our article we did indeed see that some typos had escaped our notice. We are slightly surprised by the request of the reviewer to have the manuscript edited by a “professional (!) native (!) biologist (!)”. We rephrased some stiff sentences and corrected grammatical errors. If a proofreading is indeed wished, we will have a scientific translator (English native speaker) correct the article.

(2) Then, the text lacks conciseness, it is overly long. For example, I suggest to omit all biological/ecological details you provide when introducing a taxon. This is per se interesting, but not to the point here (except when the reader needs background to understand microplastic effects). Then, figure 1 does not contribute to the understanding of your presentation, omit it. And I do not think it necessary to present taxa for which there is no information available, especially if the taxa are of minor or no importance in soil (e.g. line 183ff, 205, 220, 227, 450ff) or if the literature is not on edaphic species (435ff). As a reviewer, you are of course required to address blind spots of research (thus pointing out important taxa that are missing in literature), but you need to better balance completeness with a concise presentation.

Your suggestion to omit the ecological presentation of some key taxa is understandable. If we would expect all readers to be well acquainted with the soil fauna, we would definitely go along with this. However, SOIL is a multi-disciplinary journal connecting a broad spectrum of soil scientists. Therefore, we think it is helpful to provide a short overview of information on the soil fauna, such as ecological functionalities (marker function, transport, degradation, habitat and food selection), which might influence how they cope with microplastics. We have critically gone through the article and here we summarize which parts we will shorten.

- **Proposal**: We shortened the introduction of the springtail section, as it is indeed oversized.
- **Proposal**: We would agree with moving it to the supplements in order to save space, in case this is wished.
- **Proposal**: We shortened the chapter about Onychophora. 
  - **Proposal**: Unstudied taxa are still presented, but their importance for future research is now additionally mentioned in section 4.3 to better “balance completeness”.
  - **Proposal**: We shortened the chapter about Onychophora.
  - **Proposal**: We use this benthic species to show more clearly how inconsistent the few results for benthic and terrestrial snails are.

(3) I miss a convincing argumentation why you focus on multicellular animals (but then, you provide many details about bacteria, fungi, algae, plant roots in 72ff... omit this). A good line of reasoning could be that you follow up on the Rillig and Bonkowski (2018) paper. The aim of this review is to depict the influence of microplastic contamination in soils to the soil fauna. But, to present a holistic view on the food web, we refer to microorganisms, plant roots and biofilms within the introduction section. Being large fields of knowledge on their own, these organisms are not part of the focus in this review, however they are food sources for meso- and macroorganisms and, thus, worthy of mention. Given that we only use 22 lines to describe these other parts of the phylogenetic tree of soil life, we think this is merited and wish to leave this part in the review.

Unfortunately, we do not understand how Rillig and Bonkowski (2018), a paper on soil protozoa, matches your point. We have read this paper and do mention it elsewhere in the review.

(4) Please provide details of your literature search (123ff). When did you search? Which time span did you cover? Which search strings? Please consider the literature on meta-analyses how to properly specify these technical aspects.
The search was applied between June 2019 and January 2020, repeated in the first week of January 2020 and covers publications until January 2020. The search strings result from combinations of taxon, plastic type and particle shape shown in Table 1 (formerly Table 8).

- [Proposal]: Information added to section 2.

(5) Thank you very much for the positive note.

(6) line 636f: Please reconsider including your supervisor as a co-author. What “supervision” means is nowhere clearly defined, however, co-authorship is only justified for significant contributions to the manuscript. Honorary authorship violates the principle of scientific honesty. We understand this point completely and agree that it is not good practice to include scientists who have not contributed significantly to a paper. We also acknowledge that supervision is a very broad term and would like to specify the contribution of Martin Kaupenjohann to the paper. [6] Martin Kaupenjohann was involved in the development of the idea and concept for this paper. During the literature reading and writing phase he has supported the work with frequent discussions of the contents of the article. And finally he has critically revised the manuscript.

Best regards,

Dr. Frederick Büks
Dr. Loes van Schaik
Prof. Dr. Martin Kaupenjohann
Dear Dr. Maha Deeb (referee #4)

Thank you very much for the repeated check and your friendly report.

Best regards,

Dr. Frederick Büks
What do we know about how the terrestrial multicellular soil fauna reacts to microplastic?

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Abstract. The ubiquitous accumulation of microplastic (MP) particles across all global ecosystems comes along with the uptake into soil food webs. In this review, we analyzed studies on passive translocation, active ingestion, bioaccumulation and adverse effects within the phylogenetic tree of multicellular soil faunal life. The representativity of these studies for natural soil ecosystems was assessed using data on the type of plastic, shape, composition, concentration and time of exposure.

Available studies cover a wide range of soil organisms, with emphasis on earthworms, nematodes, springtails, beetles and lugworms, each focused on well-known model organisms. [1] About 58% of the studies thereby used inappropriate concentrations or units, but 42% applied MP concentrations similar to amounts in slightly to very heavily polluted soils. In many cases, however, polystyrene microspheres have been used, a combination of plastic type and shape, that is easily available, but does not represent the main plastic input into soil ecosystems. In turn, MP fibers are strongly underrepresented compared to their high abundance within contaminated soils. [7] A few studies also examined the comminution of macroplastic by the soil fauna. Further properties of plastic such as aging, coating and additives were insufficiently documented. Despite these limitations, there is a recurring pattern of active intake followed by a population shift within the gut microbiome and adverse effects on motility, growth, metabolism, reproduction and mortality in various combinations, especially at high concentrations and small particle sizes.

For the improvement of future studies, we identified problems of past experiments and recommend that coming studies take into account the type, shape, grade of aging, specific concentrations of MP fractions and long-term incubation, in natural and contaminated soils.
1 Introduction

Imagine a compact plastic cube of nearly 2 km side length and a weight of 7300000000 tons, with major percentages by weight of 36% polyethylene (PE), 21% polypropylene (PP), 12% polyvinyl chloride (PVC) and 10% of each polyethylene terephthalate (PET), polyurethane (PU) and polystyrene (PS). That is the accumulated global non-fiber production of the six main plastic types until 2015. It accounts to 87% of the all-time plastic production, which evolved exponentially, since the early 1950s, from some megatons (Mt) to 8300 Mt in 2015, with only 260 Mt annual output in 2009 increased to 380 Mt in 2015 (Thompson et al., 2009; Geyer et al., 2017). Of this ever produced plastic, 6300 Mt became waste until 2015, of which only 21% were recycled or incinerated, whereas 5000 Mt ended up in landfills and nature (Geyer et al., 2017). As a corollary of production, use and disposal, a certain part of plastic waste is constantly released into the environment through various pathways, but our knowledge about rates of mass flow into global ecosystems is very limited. Based on waste generation in coastal countries, Jambeck et al. (2015) calculated the global plastic input to marine ecosystems to be roughly 4.8 to 12.7 Mt in 2010. Such data on soils are lacking, but Nizzetto et al. (2016) estimated that the load of microplastic (MP) to agricultural sites in Europe is in the same order of magnitude as that in marine environments.

By littering, plastic mulching, the application of sewage sludge, digestates and composts as well as windblown dispersal (Bertling et al., 2018; Weithmann et al., 2018; Zhang et al., 2019; Wang et al., 2019a), plastic from our technosphere arrives in soil ecosystems in various forms as large and small fragments, fibers and particles. Exposed to UV radiation, mechanical stress and processing by soil organisms, plastic items become weathered and prone to a successive comminution towards the size range of MP with increased surface, charge and biofilm cover (Kale et al., 2015; Andrady, 2017). However, the resistance of plastic to metabolization causes a constant accumulation in soils as long as the release rate from human processes is above the very slow rate of degradation.

Due to a lack of monitoring programs, data on MP concentrations in terrestrial soils are rare, and those using w/w concentrations represent only a small part compared to item concentrations. In soils with only slightly contaminated conditions, amounts seem to average about 1 mg kg\(^{-1}\) soil dry weight (and approx. 200 items kg\(^{-1}\) dry soil) (Rezaei et al., 2019). On sites with industrial activity or intensive use of plastic mulching and sewage sludge in agriculture, concentrations can be increased by 2 to 4 orders of magnitude (Fuller and Gautam, 2016; Zhang and Liu, 2018; Huang et al., 2020). Semisubhydric soils such as beaches, mudflats, mangroves or lagoons, that are additionally contaminated from the aquatic side, contain MP of the order of 10 to 100 items kg\(^{-1}\) dry soil and single extreme samplings contained several thousand items (Nor and Obbard, 2014; Naji et al., 2017; Garcés-Ordóñez et al., 2019; Li et al., 2018a). More informative data using mg kg\(^{-1}\) are only available for beaches and coastal deconstruction yards in municipal neighborhoods and amount to 0.5 and
70 mg kg\(^{-1}\) dry soil, 0.00005 and 0.007 % w/w, respectively (Reddy et al., 2006; Claessens et al., 2011). All these concentration data represent a wide range of particle sizes between 0 and 5000 \(\mu\)m with different materials, shapes and degrees of aging.

Plastic particles can possibly enter and accumulate in the food web by either direct uptake from soil or by consumption of other soil biota contaminated by adhesion or ingestion (Huerta Lwanga et al., 2017a). There is evidence, that MP is incorporated even by plants and unicellular organisms at the base of the food web. Bacteria, for example, that are reasonably assumed to avoid MP uptake due to their minor size and the prevalent lack of phagocytosis, were shown to take up inorganic nanoparticles of a few nanometers (Kumar et al., 2011). Although the physiochemical properties of weathered nanoparticulate plastics might differ from these, also their uptake seems likely.

A similar argument can be made for fungi and soil algae, but studies on incorporation are lacking, whereas the transfer into a freshwater food web by adhesion of nanoplastic on algae has been shown by Chae et al. (2018). The uptake of MP into plant roots is also inhibited (Rillig et al., 2019), but occurred for nanoplastics that permeate into the plant tissue (Li et al., 2019). Also the integration into root tissue after adsorption to the rhizodermis has yet to be studied.

In contrast, protozoa feature phagocytosis for the active ingestion of particles. Diverse soil, freshwater and marine ciliates ingest PS/latex beads of 0.1 to 14.4 \(\mu\)m in laboratory experiments, with preferences to their natural prey size (Fenchel, 1980; Jonsson, 1986; Lavin et al., 1990). Soil amoebas act similarly, but additionally select according to food quality (Weisman and Korn, 1967; Vogel et al., 1980; Bowers and Olszewski, 1983; Avery et al., 1995; Elloway et al., 2006).

Finally, many soil microbiota live protected within biofilms. Plastic particles were shown to be a potential surface for the formation of those biofilms (Lobelle and Cunliffe, 2011), which are a food source for grazing primary consumers. Inadvertent ingestion might also transfer occluded or abrased MP to higher trophic levels.

But what about the larger organisms that feed on all these, free plastic particles, contaminated microorganisms, biofilms and one another? Recent work discussed the effects of MP on soil biota (Chae and An, 2018) or called for intensified research on certain taxonomic groups (Rillig and Bonkowski, 2018). Thus, we were motivated to give on our part a review with focus on the most-produced plastics and their passive translocation, ingestion, bioaccumulation and adverse effects on the multicellular soil fauna. The types, sizes and shapes of plastic used in former laboratory studies were compared with the available knowledge on plastic in the environment, and recommendations are given for future research. This analysis aims to support the assessment of the influence of MP on the ecosystem services provided by diverse soil organisms.
2 Search pattern

Within the tree of life, edaphic branches were identified comprising taxa that permanently inhabit the soil, are both-sided part of the soil food web and/or the burrowing macro- and megafauna or have active subterranean larval stages. The resulting tree of soil life based on the NCBI taxonomy database (Fig. 1) was drawn using the software phyloT and shows the leading taxonomic rank, which is mainly the family, but in exceptions – e.g. if one species represents the only soil-born between many aquatic – a lower rank.

Figure 1: [2] Tree of [6] edaphic fauna. Taxonomic ranks, that were examined in this qualitative study, are placed at the outer rim of the diagram. The length of the connecting line between two taxa represents the grade of phylogenetic relationship.
A pattern of search terms was established ([7] Table 1), consisting of „taxon“ (Linné's binominal nomenclature, common name, plural-sensitive search), „plastic type“ (plastic, microplastic, nanoplastic, PE or polyethylene, PP or polypropylene, PVC or polyvinyl chloride, PS or polystyrene, PU or polyurethane, PET or polyethylene terephthalate and latex) and „common shapes“ (fragments, particles, fibers, microfibers, beads, microbeads, microspheres). [7] Some type-shape combinations caused problems, as they led to a very large amount of unuseful, off-topic papers – e.g. using any taxon combined with PET, papers with the use of PET bottles in experimental set-ups were selected or also studies on pets. Those combinations of search terms were excluded from this pattern. Further plastic types and shapes occurring within the found studies were also included in the review. Data on microspheres and microbeads were pooled, as both names describe one and the same.

Table 1: [5] Types and shapes of microplastic particles in edaphon studies within this review. (X) symbolizes combinations excluded from the search pattern. [7] The number counts for how often type-shape combinations were used in all reviewed experimental setups independently of organism. Empty fields stand for zero results. Microbeads and microspheres are often mixed up terms and, thus, counted together.

| Organism: | Linné's systematic names | OR common name | fragments | particles | microfibers | beads | microbeads | microspheres | other, diverse | N/A |
|-----------|--------------------------|----------------|-----------|-----------|------------|-------|------------|--------------|---------------|-----|
| plastic   | X                        |                |           |           |            |       |            |              |               |     |
| microplastic |                         |                |           |           |            |       |            |              |               |     |
| nanoplastic |                          |                |           |           |            |       |            |              |               |     |
| PE OR polyethylene |                      |                | X         | 4         | 10         | 1     | 1          | 4             | 7              |     |
| PP OR polypropylene |                       |                | X         | 1         |            |       |            |              |               |     |
| PVC OR polyvinyl chloride |                  |                | X         | 4         | 6          | 1     |            |              |               |     |
| PS OR polystyrene |                          |                | X         | 6         | 3          | 24    | 4          |              |               |     |
| PU OR polyurethane |                         |                | X         |            |            |       |            |              |               |     |
| PET OR polyethylene terephthalate |              |                | X         | 3         | 2          |       | X          |              |               |     |
| latex     |                          |                | X         |            |            |       |            |              | 6              |     |
| other     | 6 3 1 1                  |                |           |           |            |       |            |              |               |     |
| N/A       | 1 1 2 3                  |                |           |           |            |       |            |              |               |     |

The search was applied [5][5] between June 2019 and January 2020 within the Web of Science Core Collection Database, repeated in the first week of January 2020 and covers publications until January 2020. The search strings result from combinations of taxon, plastic type and particle shape shown in Table 1. Based on the search pattern, data on passive transport,
ingestion, bioaccumulation and adverse effects were collected for each edaphic group. Studies that only use uncommon, local, outdated, weird or nicknames are excluded by the search pattern. Studies testing injection to tissues, lymph or blood were excluded, as they do not represent natural ways to incorporate MPs. Data on inhalation by the megafauna in fact represent a natural way of uptake, but were also excluded as they are exclusively related to above-ground organisms, that only occur on the outer edge of the food-web. Also running debates on phylogenetic classifications are not part of this work and the taxonomists will be able to adjust the branches accordingly to their purpose.

The data of related taxonomic groups were pooled and evaluated for their environmental representativity based on exposure time, plastic concentrations and properties used. From this synthesis recommendations for a structured experimental design were derived for application in future studies.
3 Data collection

3.1 Insects

Within the Panarthropoda, the insects comprise the highest taxonomic diversity. And, regarding MPs, they represent an unevenly studied taxonomic group. Within the Insecta, the **Coleoptera** (beetles) are an extraordinarily diverse and abundant taxon. Studies on plastic uptake into adult individuals mainly focus on the subfamily of Scarabaeinae (dung beetles). Comprehensive experiments with latex microbeads showed, that many species only ingest smaller particles with maximum diameters of about 10 to 83 µm and retain them within the gut – with a slightly positive dependency on body size. Larger particles were rejected by a filtering mechanism within the mouth region and not ground with the mandibles (Holter, 2000; Holter et al., 2002; Holter and Scholtz, 2005). Beside those on Nematods, these data comprise by far the most detailed information about size-dependent uptake of MP particles compared to other edaphic taxa. This gives a good foundation for future studies on adverse concentrations. In addition, several studies with plastic as predominant food source could show chewing, ingestion and intestinal degradation of different PS and PE foams in feeding experiments with *Tenebrio sp.* larvae (mealworms). These experiments also pointed out an alteration of the gut microbiome, but no adverse effects on reproduction and survival, with only one case of a non-significant tendency to higher mortality after 1 month of exposure (Yang et al., 2015; Brandon et al., 2018; Yang et al., 2018; Peng et al., 2019).

The **Isoptera** (termites), recently categorized as part of the order Blattodea, are the oldest social insects with a tribal history of about 130 million years (Korb, 2008). Especially in arid ecosystems with a lack of earthworms they play an important role in homogenization of soils, but also in sorting of soil mineral particles for building mounds as well as decomposition and distribution of organic matter (De Bruyn and Conacher, 1990). Tsunoda et al. (2010) and Lenz et al. (2012) could show, that different termite species are picky feeders and erode PE, but avoid other plastic cable sheathings. This suggests the excretion of ground MP particles by termites, but metabolic impacts are unknown. In contrast to termites, data on other Blattodea (e.g. cockroaches) were not found.

The suborder **Apocrita** comprises some flying insects, that inhabit burrows within the soil, such as ground-dwelling wasps within the Vespidae superfamily, mining bees within the Apoidea superfamily and the Spheciformes. They mostly do not prey and feed on subterrestrial organisms, but may move MP particles into the ground, as implied by a report of Allasino et al. (2019) on solitary bees, which built nests fully made of plastic fragments. The Apocrita also contain the Formicidae (ants). Some ant species are considered an important factor for seed dispersal, a behavior, that could also be shown for artificial plastic seeds with ~2 mm diameter (Hughes and Westoby, 1992; Angotti et al., 2018). Robins and
Robins (2011) found that this also includes differently shaped cultural objects: *Rhytidoponera metallica*, a representative of ground-nesting, omnivore ants, is capable not only of a remarkable bioturbation, but also of an active, apparently random burying of anthropogenic plastic artifacts >1 mm. Seeds are used as a food source, thus, the ingestion of plastic bites is conceivable, but not documented. The uptake of latex microspheres ≥0.88 µm with liquids by larvae of *Solenopsis invicta* seems to be prevented by filtration within the mouth and the particles are released as larger aggregates, whereas other species ingest by far larger particles up to 150 µm (Glancey et al., 1981). However, also here data on adverse effects are missing.

Further insects with edaphic adult stages, e.g. *Dermoptera* (earwigs), *Heteroptera* (true bugs) and *Zygentoma* (silverfish, fishmoth, firebrat) or soil- or litter-dwelling larvae such as *Embioptera* (webspinners, footspinners), *Thysanoptera* (thrips), *Psocoptera* (booklice, barklice, barkflies), *Neuroptera* (lacewings), *Raphidioptera* (snakeflies) or *Zoraptera* (angel insects) were not yet researched with focus on soil MP.

Regarding insects, mainly studies on translocation and uptake of MP were carried out. In contrast, work on bioaccumulation is completely lacking and adverse effects are sparsely tested using *Tenebrio sp.* larvae. Such studies could provide information whether or not the input of MP in soil ecosystems is one of many factors causing the global decline of the entomofauna (Oliveira et al., 2019; Sánchez-Bayo and Wyckhuys, 2019).
Table 2: Microplastic studies on Coleoptera, Blattodea (Blattod.), Apoidea (A.) and Formicidae (mb=microbeads, fr=fragments, ms=microspheres, b=beads, [a] N/A=information not available). Concentrations refer to mg kg⁻¹ dry soil, if not specially marked.

| organism | experimental environment | plastic type | aging | coating | additives | shape | size span [µm] | concentrations | exposure time | passive transport | active uptake | bioaccum. dynamics | measured adverse effects | reference |
|----------|--------------------------|--------------|-------|---------|-----------|-------|---------------|---------------|---------------|-------------------|--------------|-------------------|-------------------------|-----------|
| Aphodius erraticus | Petri dish | latex | N/A | N/A | mb | 5 | 2..39 | N/A | 45 min | N/A | no | ≤14 µm | N/A | Holter (2000) |
| Aphodius rufipes | Petri dish | latex | N/A | N/A | mb | 2..39 | N/A | 45 min | N/A | ≤10..≤50 µm | N/A | no | ≤14 µm | N/A | Holter et al. (2002) |
| Aphodius ater | vial | latex | N/A | N/A | mb | 2..83 | N/A | 45 min | N/A | ≤18 µm | N/A | no | ≤18 µm | N/A | Holter and Scholtz (2005) |
| Aphodius fimetarius | Petri dish | latex | N/A | N/A | mb | 2..83 | N/A | 45 min | N/A | ≤18 µm | N/A | no | ≤50 µm | N/A | Holter et al. (2006) |
| Aphodius contaminatus | Petri dish | latex | N/A | N/A | mb | 2..83 | N/A | 45 min | N/A | ≤10..≤50 µm | N/A | no | ≤18 µm | N/A | Holter et al. (2006) |
| Tenebrio molitor larvae | container | LD-PE | N/A | no foam retardant | foam | 8..27 cm² | 100% w/w (food) | 31 d | N/A | yes | biodegrad. | N/A | Yang et al. (2015) |
| Tenebrio molitor larvae | container | LD-PE | N/A | no flame retardant | foam | 8..27 cm² | 50..100% w/w (food) | 32 d | N/A | yes | biodegrad. | microbe | Brandon et al. (2018) |
| Tenebrio molitor larvae | container | PS | N/A | no foam | foam | N/A | 4..100% w/w (food) | 32 d | N/A | yes | biodegrad. | no | Yang et al. (2018) |
| Tenebrio obscurus larvae | container | PS | N/A | no foam | foam | N/A | 86..100% w/w (food) | 31 d | N/A | yes | biodegrad. | microbe | Peng et al. (2019) |
| Coptotermes formosanus | mesocosm | LD-PE | others | yes/no | N/A | cable sheets | 4 cm, ∅ 0.8 cm | N/A | 42 d | N/A | yes | no | N/A | Tsunoda et al. (2010) |
| diverse termites | in situ | MD-PE | no | N/A | anti-oxidant stabilizer | cable sheets, cable sheets | 30 cm, ∅ 1.4 cm | N/A | 6 yr. | N/A | yes | no | N/A | Lenz et al. (2012) |
| Formicidae | Petri dish | latex | N/A | N/A | fluorescence | ms | 0.9..4.5 | 2.5% w/w (food) | direct | N/A | filtration | N/A | Glancey et al. (1982) |
| Aphaenogaster longiceps | in situ | N/A | N/A | N/A | b | 50 items per nest | 3 d | yes | N/A | N/A | Hughes and Westoby (1992) |
| Pheidole sp. | in situ | N/A | N/A | N/A | b | <75.5 cm | N/A | 26 mos. | yes | N/A | N/A | Robins and Robins (2011) |
| Pheidole sp. | in situ | N/A | N/A | N/A | b | 1.8 cm | N/A | 1 d | yes | N/A | N/A | Angotti et al. (2018) |
### 3.2 Other panarthropods

Apart from the insects, **Acari** (mites) comprise many abundant soil-living taxa, that feed on litter, fungi and fauna as predators and parasites and are bioindicators, as they are sensitive to changes in the soil physiochemical environment (Gulvik, 2007). Experiments indicated, that mites passively transport MP due to pushing and dragging after attachment to their cuticle, as shown with 80 to 250 µm sized PVC particles in a Petri dish experiment without soil (Zhu et al., 2018a). The population within manure pats slightly declines when exposed to mm-sized unweathered PE and PS particles at concentrations of 5 % v/v and declines strongly at ≥60 % v/v (Stamatiadis and Dindal, 1990). This could probably be an effect of moisture deficiency due to a reduced water holding capacity in an unnaturally enriched substrate, but not necessarily through plastic intake. In contrast, no data was found on their arachnoid, preying relatives, the order of **Pseudoscorpiones** (false scorpions).

Just as many other highly abundant and diverse representatives of the soil mesofauna, the **Oniscidea** (woodlice) contribute to the decomposition of litter by chewing and passage through their digestive system (Warburg, 1987) and react strongly to environmental pollution, thus, they are potentially used as bioindicators (van Gestel et al., 2018). They practice a strict selection of natural food sources (Hassall and Rushton, 1984). This is also demonstrated for starch and cellulose based plastic films (4 cm²), which were consumed and digested in experiments with the model organism *Porcellio scaber*, in contrast to PHB (polyhydroxybutyrate) films, that reduces the feeding rate (Wood and Zimmer, 2014). Smaller PE particles (137±51 µm and 183±93 µm) embedded into food pellets (0.4 % w/w) were taken up easily by *Porcellio scaber*, and the smaller fraction caused a slight and non-significant reduction of body mass after 14 days of exposure, but not of feeding, defecation or energy reserves (Kokalj et al., 2018).

Other panarthropodean groups are even less studied in terms of MP. We did not find literature on the subphylum of **Myriapoda** containing the classes of **Diplopoda** (millipedes), **Chilopoda** (centipedes), **Pauropoda** and **Symphyla** (pseudocentipedes or symphilids), which are important litter-feeders and predators within various soil ecosystems.

The situation is nearly similar with the phylum of **Tardigrada** (water-bears or tardigrades), that has many ecologically relevant and well studied species feeding on microorganisms and detritus particles. Sparse field research in semisubhydric environments showed no uptake of MP fibers by tardigrada (Gusmão et al., 2016), but comprehensive data on terrestrial soils are lacking. Similarly, the related phylum of **Onychophora** (velvet worms), primordial invertebrates that are mainly native in litter and soils with high water holding capacity under pleistocene-like forest vegetation within tropical and moderate regions (Monge-Nájera, 1994). Another branch within the panarthropoda, the phylum of **Onychophora** (velvet worms), comprises primordial invertebrates that are mainly native in litter and soils with high water...
holding capacity under Pleistocene-like forest vegetation within tropical and moderate regions (Monge-Nájera, 1994). As predators, they most likely take up plastic debris appearing within or on their prey, but no studies on MP are available, most likely due to their remote habitats, low-abundance and little scientific focus.

The phylum of Collembola (springtails) \textsuperscript{11}, together with the Diplura and Protura (Westheide and Rieger, 1996; Pass et al., 2011), an abundant, diverse and ubiquitous soil-borne phylum with a broad spectrum of food sources (Hopkin, 1997), also represent an intensively studied group within the Arthropoda. Together with the Diplura (which mainly live in tropic and subtropic regions in litter and humid topsoil and feed on fungal hyphae, POM and prey) (Westheide and Rieger, 1996) and the Protura (Pass et al., 2011), the Collembola build an intensively studied morphological group, that \textsuperscript{12} exhibits similar ecological functions, such as distribution and decomposition of organic matter as well as the control of fungal abundance (Hopkin, 1997). Springtails provide up to 27% of the soil biomass and up to 33% of the total soil respiration (with higher shares in colder ecosystems) (Petersen, 1994) with up to 100 000 individuals per square meter (Hopkin, 1997). Thus, their well-being plays an important role for ecosystem functioning.

In a Petri dish experiment without soil, Maaß et al. (2017) showed the passive transport of urea-formaldehyde particles <400 µm and undefined PET fragments by two Collembola species (Folsomia candida and Proisotoma minuta) due to attachment, but found no ingestion. Within a soil matrix, trials of Kim and An (2019) indicated hindrance of collembolan migration by larger PS particles (44±39, 282±131 and 676±479 µm) at concentrations of 1000 mg kg\(^{-1}\) corresponding to highly contaminated soils. In addition, they found suppressed mobility due to the attachment of even smaller PS microbeads (0.47 to 0.53 µm) at concentrations of 8 mg kg\(^{-1}\) dry soil, which is equivalent to values found in nature. Small particles <50 µm were moved, while larger particles were most likely \textsuperscript{12} cast off. When F. candida encounters two of its predators, the mites Damaeus exspinosus and Hypoaspis aculeifer, the dispersal of 80 to 250 µm PVC particles is enhanced as shown by Zhu et al. (2018a) in a Petri dish experiment. Without proving the ingestion or the minimal effective MP concentration, Zhu et al. (2018b) published an alteration of the gut microbiome and adverse effects on growth and reproduction of F. candida by 80 to 250 µm PVC particles mixed in soil at concentrations of 1000 mg kg\(^{-1}\) dry soil. These data were not considered robust (van Gestel and Selonen, 2018), but match with a later study that found inhibited reproduction at ≥1000 mg kg\(^{-1}\) and avoidance behavior as well as microbiome alteration at ≥5000 mg kg\(^{-1}\) (Ju et al., 2019). Such concentrations can occur in highly contaminated soils (Fuller and Gautam, 2016). However, documentations on the active uptake, gnawing and grinding of MP by springtails proposed by Rillig (2012) is still lacking and also studies on Diplura and Protura were not found.
Table 3: Microplastic studies on Acari, Oniscidea (Onisc.), Tardigrada (T.) and Collembola (fr=fragments, p=particles, mf=microfibers, mb=microbeads, ms=microspheres, s=semisubhydric, [8] N/A=information not available). Concentrations refer to mg kg⁻¹ dry soil, if not specially marked.

| Organism | Experimental environment | Plastic type | Aging | Coating | Additives | Shape | Size Span [μm] | Concentrations | Exposure Time | Passive Transport | Active Uptake | Bioaccumulation Dynamics | Measured Adverse Effects |
|----------|--------------------------|--------------|-------|---------|-----------|-------|---------------|---------------|--------------|-------------------|--------------|--------------------------|------------------------|
| Acari    | Microcosm                | PE           | no    | N/A     | N/A       | fr    | <4800-2000    | 0.90% v/v (manure) | 16 d         | N/A               | N/A          | ≥5% v/v: abundance ‡  | Stamatiadis and Dindal (1990) |
| Hypoaspis aculeifer | Petri dish          | PVC         | N/A   | no      | N/A       | p     | 80.250        | 5000 items per dish | N/A          | yes               | N/A          | N/A                | Zhu et al. (2018a)          |
| Porcellio scaber | Mesocosm           | PHB          | no    | N/A     | N/A       | fr    | 4 cm²         | 1 item per cosm    | 14 d         | N/A               | yes          | N/A                | Wood and Zimmer (2014)      |
| Porcellio scaber | Petri dish          | PE           | N/A   | N/A     | N/A       | fr    | 183±53        | 0.4% w/w (food)    | 14 d         | N/A               | yes          | N/A                | Kokalj et al. (2018)        |
| T. in situ | In situ                | N/A          | N/A   | N/A     | N/A       | mf    | N/A           | N/A            | N/A          | no                | N/A          | N/A                | Gusmão et al. (2016)        |
| Folsomia candida | Cup               | UF, PET      | N/A   | no      | N/A       | p,fr  | <400          | 2.5-5 mg per cup   | N/A          | yes               | N/A          | N/A                | Maaß et al. (2017)          |
| Folsomia candida | Petri dish          | PVC          | N/A   | no      | N/A       | p     | 80.250        | 1000           | 56 d         | N/A               | N/A          | N/A                | Zhu et al. (2016a)          |
| Folsomia candida | Microcosm          | PVC          | N/A   | no      | N/A       | p     | 80.250        | 1000           | 56 d         | N/A               | N/A          | N/A                | Zhu et al. (2018b)          |
| Folsomia candida | Microcosm          | PE           | N/A   | no      | N/A       | mb    | <500          | 0.1-10000      | 7 d          | N/A              | N/A          | ≥5000: avoidance       | Ju et al. (2019)            |
| Lobella sokamensis | Soil sample       | PS           | N/A   | Carboxyl fluorescence | MB        | 0.5    | 4.8           | 1000          | ≤3 min       | N/A               | yes          | N/A                | Kim and An (2019)          |

‡ ≥50% v/v: abundance

| Experimental Environment | Plastic Type | Aging | Coating | Additives | Shape | Size Span [μm] | Concentrations | Exposure Time | Passive Transport | Active Uptake | Bioaccumulation Dynamics | Measured Adverse Effects |
|--------------------------|--------------|-------|---------|-----------|-------|---------------|---------------|--------------|-------------------|--------------|--------------------------|------------------------|
| Acari                    | Microcosm    | PE    | no      | N/A       | N/A   | fr            | 0.90% v/v (manure) | 16 d         | N/A               | N/A          | ≥5% v/v: abundance ‡  | Stamatiadis and Dindal (1990) |
| Hypoaspis aculeifer     | Petri dish   | PVC   | N/A    | no        | N/A   | p             | 80.250        | 5000 items per dish | N/A          | yes               | N/A          | N/A                | Zhu et al. (2018a)          |
| Porcellio scaber        | Mesocosm     | PHB   | no     | N/A       | N/A   | fr            | 4 cm²         | 1 item per cosm    | 14 d         | N/A               | yes          | N/A                | Wood and Zimmer (2014)      |
| Porcellio scaber        | Petri dish   | PE    | N/A    | N/A       | N/A   | fr            | 183±53        | 0.4% w/w (food)    | 14 d         | N/A               | yes          | N/A                | Kokalj et al. (2018)        |
| T. in situ              | In situ      | N/A   | N/A    | N/A       | N/A   | mf            | N/A           | N/A          | no                | N/A          | N/A                | Gusmão et al. (2016)        |
| Folsomia candida        | Cup          | UF, PET| N/A   | no        | N/A   | p,fr          | <400          | 2.5-5 mg per cup   | N/A          | yes               | N/A          | N/A                | Maaß et al. (2017)          |
| Folsomia candida        | Petri dish   | PVC   | N/A    | no        | N/A   | p             | 80.250        | 1000           | 56 d         | N/A               | N/A          | N/A                | Zhu et al. (2016a)          |
| Folsomia candida        | Microcosm    | PVC   | N/A    | no        | N/A   | p             | 80.250        | 1000           | 56 d         | N/A               | N/A          | N/A                | Zhu et al. (2018b)          |
| Folsomia candida        | Microcosm    | PE    | N/A    | no        | N/A   | mb            | <500          | 0.1-10000      | 7 d          | N/A              | N/A          | ≥5000: avoidance       | Ju et al. (2019)            |
| Lobella sokamensis      | Soil sample  | PS    | N/A    | Carboxyl fluorescence | MB        | 0.5    | 4.8           | 1000          | ≤3 min       | N/A               | yes          | N/A                | Kim and An (2019)          |

‡ ≥50% v/v: abundance

≥5000: avoidance

≥1000: reproduction

≥5000: microbiome

Avoidance, motivity

| Experimental Environment | Plastic Type | Aging | Coating | Additives | Shape | Size Span [μm] | Concentrations | Exposure Time | Passive Transport | Active Uptake | Bioaccumulation Dynamics | Measured Adverse Effects |
|--------------------------|--------------|-------|---------|-----------|-------|---------------|---------------|--------------|-------------------|--------------|--------------------------|------------------------|
| Acari                    | Microcosm    | PE    | no      | N/A       | N/A   | fr            | 0.90% v/v (manure) | 16 d         | N/A               | N/A          | ≥5% v/v: abundance ‡  | Stamatiadis and Dindal (1990) |
| Hypoaspis aculeifer     | Petri dish   | PVC   | N/A    | no        | N/A   | p             | 80.250        | 5000 items per dish | N/A          | yes               | N/A          | N/A                | Zhu et al. (2018a)          |
| Porcellio scaber        | Mesocosm     | PHB   | no     | N/A       | N/A   | fr            | 4 cm²         | 1 item per cosm    | 14 d         | N/A               | yes          | N/A                | Wood and Zimmer (2014)      |
| Porcellio scaber        | Petri dish   | PE    | N/A    | N/A       | N/A   | fr            | 183±53        | 0.4% w/w (food)    | 14 d         | N/A               | yes          | N/A                | Kokalj et al. (2018)        |
| T. in situ              | In situ      | N/A   | N/A    | N/A       | N/A   | mf            | N/A           | N/A          | no                | N/A          | N/A                | Gusmão et al. (2016)        |
| Folsomia candida        | Cup          | UF, PET| N/A   | no        | N/A   | p,fr          | <400          | 2.5-5 mg per cup   | N/A          | yes               | N/A          | N/A                | Maaß et al. (2017)          |
| Folsomia candida        | Petri dish   | PVC   | N/A    | no        | N/A   | p             | 80.250        | 1000           | 56 d         | N/A               | N/A          | N/A                | Zhu et al. (2016a)          |
| Folsomia candida        | Microcosm    | PVC   | N/A    | no        | N/A   | p             | 80.250        | 1000           | 56 d         | N/A               | N/A          | N/A                | Zhu et al. (2018b)          |
| Folsomia candida        | Microcosm    | PE    | N/A    | no        | N/A   | mb            | <500          | 0.1-10000      | 7 d          | N/A              | N/A          | ≥5000: avoidance       | Ju et al. (2019)            |
| Lobella sokamensis      | Soil sample  | PS    | N/A    | Carboxyl fluorescence | MB        | 0.5    | 4.8           | 1000          | ≤3 min       | N/A               | yes          | N/A                | Kim and An (2019)          |

‡ ≥50% v/v: abundance

≥5000: avoidance

≥1000: reproduction

≥5000: microbiome

Avoidance, motivity
3.3 Annelida

Land-based Annelida comprise another large group of invertebrates. The Lumbricidae (earthworms) are a well-studied family (Darwin, 1881; Lavelle et al., 2006), represented in high abundance and diversity in many ecosystems all around the world (Phillips et al., 2019). Earthworms are often used as indicators for soil health (Fründ et al., 2011; Pulleman et al., 2012), as they are ecosystem engineers which through their burrowing activity influence various soil physical, chemical and biological processes (Jouquet et al., 2006; Lavelle et al., 2006).

By far the most of the studies on the influence of MP on earthworms are performed with PE and the species Lumbricus terrestris or Eisenia fetida, but there are also single studies with Aporrectodea rosea (Boots et al., 2019) and Eisenia andrei (Rodriguez-Seijo et al., 2017) and with the less common species Metaphire californica (Wang et al., 2019b). We found one field study of earthworms and MPs (Huerta Lwanga et al., 2017a) among many laboratory experiments with MPs mixed into soil volumes (concentrations ranging up to 20000 mg kg\(^{-1}\) dry soil) or applied with litter on top of the soil surface (≤60% w/w). The particles sizes were usually <1 mm in diameter, but some were even up to 2x2 cm\(^2\), and the duration of experiments was generally 14 to 28 days, few lasted up to 60 days.

The uptake of MPs of a broad size range by earthworms was shown in studies based on particles in earthworm casts of Lumbricus terrestris (Huerta Lwanga et al., 2016; Cao et al., 2017; Hodson et al., 2017; Rillig et al., 2017; Prendergast-Miller et al., 2019; Yu et al., 2019; Huerta Lwanga et al., 2017a), Eisenia fetida (Rodriguez-Seijo et al., 2018; Chen et al., 2020; Wang et al., 2019c), Eisenia andrei (Rodriguez-Seijo et al., 2017) and Metaphire californica (Wang et al., 2019b). Zhang et al. (2018) showed that relatively large PE particles of 1.5 x1.5 cm\(^2\) are not ingested by Lumbricus terrestris, but partial ingestion of such large particles of biodegradable MPs does take place after initial weathering in soil or in compost has occurred. In some laboratory experiments, MPs were found in the gut of dissected earthworms (Huerta Lwanga et al., 2016; Hodson et al., 2017; Rodriguez-Seijo et al., 2017), but the concentration of MPs in the gut was not significantly different between treatments nor significantly different from the bulk soil concentration, so there was no evidence of accumulation of MPs in the earthworm bodies (Hodson et al., 2017). Chen et al. (2020) assume an accumulation of MP takes place in Eisenia fetida, based on an observed increase of MP concentrations in the casts in the course of 4 weeks. Huerta Lwanga et al. (2017a) supposed an accumulation of MPs in the food chain as the concentration of MPs in chicken gizzards is strongly increased compared to that in the earthworm casts in the same experiments. However, mainly the amount of large particles, i.e. macroplastics, in the gizzards was very large, thus it seems likely that the chicken directly fed on plastics and an accumulation through the food chain cannot be proven with the current knowledge and should be further investigated.
Several studies did not find significant negative effects of MPs on earthworms’ avoidance behaviour (Judy et al., 2019), nor on growth (Hodson et al., 2017; Rodriguez-Seijo et al., 2017; Judy et al., 2019; Wang et al., 2019c), mortality Hodson et al. (2017); Rillig et al. (2017); Rodriguez-Seijo et al. (2017); Judy et al. (2019); Prendergast-Miller et al. (2019) or reproduction (Huerta Lwanga et al., 2016; Rodriguez-Seijo et al., 2017). However, other studies do show adverse effects of the uptake of MP in different degrees and on different aspects of earthworms’ fitness: A reduced growth was shown by Cao et al. (2017) for Eisenia Fetida and the mortality increased at an exposure of concentrations ≥10000 mg kg\(^{-1}\) dry soil. At lower concentrations no significant effects were found. The growth of Aporrectodea rosea was also inhibited when exposed to biodegradable polylactic acid, conventional high-density polyethylene (at 1000 mg kg\(^{-1}\) dry soil), and MP clothing fibers (at 10 mg kg\(^{-1}\) dry soil) (Boots et al., 2019). Huerta Lwanga et al. (2016) showed a decrease in growth and increased mortality at concentrations ≥28% w/w in litter and after 60 days, though after just 14 days no mortality occurred in these experiments.

In some studies, additional effects such as histopathological changes or stress biomarkers were measured. For Eisenia fetida Chen et al. (2020) observed skin damage at 1500 mg MP kg\(^{-1}\) in soil, measured an increase in catalase activity and malondialdehyde content at 1000 mg kg\(^{-1}\) and at ≥1000 mg kg\(^{-1}\) acetylcholine esterase was significantly stimulated. Wang et al. (2019c) tested Eisenia fetida and found that MPs only increased the catalase and peroxidase levels as well as the level of lipid peroxidation and decreased the activity of superoxide dismutase and glutathione S-transferase at an exposure of 200000 mg kg\(^{-1}\) dry soil for 14 days. No discernible influence was found at 100000 mg kg\(^{-1}\).

However, Rodríguez-Seijo et al. (2018) also found for Eisenia fetida a significant positive correlation of MP concentration with different biomarker responses: catalase, glutathione S-transferase, lactate dehydrogenase and thiobarbituric acid reactive substances. In addition, Rodriguez-Seijo et al. (2017) observed histological damage of the gut and occurrence of inflammatory processes as well as an increase of stress response indicators associated with MP exposure of Eisenia andrei. For Lumbricus terrestris Prendergast-Miller et al. (2019) showed an increase in metallothionein expression at an exposure with ≥1000 mg kg\(^{-1}\) dry soil and a decrease in heat shock protein 70 at a concentration of ≥10000 mg kg\(^{-1}\).

Due to the large differences in experimental conditions – e.g. size of the MPs, addition of MPs to soil or to litter, duration of experiments, earthworm species – the current knowledge is not sufficient to detect whether there is a threshold in MP size and concentration at which the MP become harmful for earthworms and how this threshold differs for different earthworms species and MP shapes. The results of Huerta Lwanga et al. (2016), who found no effects of MPs on earthworms at 14 days, but significant influence on growth and mortality after 60 days, indicate the importance of longer measurements. This is consistent with Pelosi et al.
(2015), who concluded that the influence of pesticides on earthworm communities should be tested in long term field experiments.

Earthworms activity also increased the transport of MP in soil columns to deeper soil layers (Rillig et al., 2017; Yu et al., 2019; Huerta Lwanga et al., 2017b). The smaller the MP the stronger the transport. Particles are transported both actively – ingested and later cast out – and passively after attachment to the earthworm's body or by water flow through the biopores. As Huerta Lwanga et al. (2018) showed that the bacteria in the gut of *Lumbricus terrestris* can decompose MPs, it seems likely that particles taken up at the surface are egested as smaller particles in deeper layers.

Microplastics might well serve as a vector for contaminant transport to soil organisms. Though adsorption on plastics was seen to be lower than on the soil matrix, the desorption of Zn was seen to be higher in synthetic earthworm guts. However, there was no measurable negative effect of Zn or the PE on *Lumbricus terrestris* (Hodson et al., 2017). Wang et al. (2019b) studied the influence of MP on arsenic uptake and negative effects on *Metaphire californica* and concluded that MPs decreased the uptake of arsenic and that MPs reduced the influence of arsenic on the gut bacterial communities. Rodríguez-Seijo et al. (2019) showed altered enzyme activities and enhanced avoidance behavior in face of LD-PE pellets spiked with the insecticide chlorpyriphos. Yang et al. (2019a) studied the influence of MPs on the transport of glyphosate, however they mainly showed that the glyphosate transport was increased by earthworm activity, the role of MPs in this transport could not be determined with this study. These studies show that MP might have very different influences on the uptake and the adverse effects of different pollutants on earthworms and further investigation is needed in order to understand the influence of MPs on pollutant transport.

In contrast to the recently well-researched Lumbricidae, a near relative, the family of *Megascoleidae* (giant earthworms), is not yet mentioned in literature. Another branch within the Annelida, the small *Enchytraeidae* (potworms), were shown to suffer adverse effects on body weight and microbiome with PS microspheres (0.05 to 0.1 \(\mu m\)) at concentrations of \(\geq 10\%\) w/w within their food source, but an unexpected increase of reproduction at 0.5 % w/w (Zhu et al., 2018b). The reproduction was reduced at abnormal concentrations of 90 g kg\(^{-1}\) dry soil of polyamid particles (13 to 150 \(\mu m\)), but not with PVC (Lahive et al., 2019).

The edaphon of semisubhydric soils is often treated as a marginal group between the area of interest of soil and aquatic scientists. As a highly diverse soil biocenosis outside the focus of this paper, the benthos along seashores and fresh waters is also affected by MPs and should therefore be shortly mentioned by reviewing the lugworm *Arenicola marina*, a well examined deposit-feeder of the tidal flats. In situ, MP accumulates within its tissue and feces (Van Cauwenbergh et al., 2015). In laboratory experiments, PS particles \(\geq 500\mu m\) were avoided as food-source and passively translocated within the sediment at concentrations of \(\sim 2\ g\ kg^{-1}\).
(Gebhardt and Forster, 2018), but were measured within the feces at ~74 g kg\(^{-1}\) causing effects on feeding activity and body weight with no influence on the survival rate (Besseling et al., 2012). PS microspheres ≤30 µm remained within the animal without any adverse effects regardless of particle size (Van Cauwenberghe et al., 2015). Other studies found adverse effects on respiration, energy reserves, feeding, egestion and casting after uptake of PVC particles ≤478 µm at different sediment concentrations of >2 g kg\(^{-1}\), but neither on biomass and survival nor due to HD-PE (Wright et al., 2013; Green et al., 2016). There is further a difficulty in distinguishing between the adverse effects of MPs and substances adsorbed on or leached from MPs (Besseling et al., 2012). When adding PCB-spiked PE to mud flat sediment with concentrations up to \([\text{12}]5000\ \text{mg kg}^{-1}\) dry mass, there was no significant change of survival rate or body weight. The decreased feeding activity and heap mass could be attributed to increasing plastic concentrations, but not to enhanced PCB bioaccumulation via PE uptake (Besseling et al., 2017). However, all these studies found adverse effects at MP concentrations orders of magnitude above natural values.
Table 4: Microplastic studies on Lumbricidae (p=particles, ms=microspheres, b=beads, f=fibers, ms=micofibers, [8] N/A=information not available). Concentrations refer to mg kg⁻¹ dry soil, if not specially marked.

| organism                  | experimental environment | plastic type | aging       | coating | additives | shape | size span [µm] | concentrations | exposure time  | passive transport | active uptake | bioaccum. dynamics | measured adverse effects                                      | reference                                      |
|---------------------------|--------------------------|---------------|-------------|---------|-----------|-------|----------------|----------------|----------------|------------------|---------------|------------------|-------------------------------------------------------------|-----------------------------------------------|
| *Lumbricus terrestris*    | mesocosm                 | PE            | washed      | N/A     | N/A       | p     | <150           | 0.50% w/w (litter) | 14 d / 60 d    | yes             | yes             | N/A             | ≥60 d, ≥29% w/w, survival, k, growth †                      | Huerta Lwanga et al. (2016)                      |
| *Eisenia fetida*          | glass beaker             | PS            | N/A         | N/A     | ms        | 50–80 | 0.02–0.09 mm²   | 3500           | 28 d           | N/A             | yes             | N/A             | ≥5000: survival †                                      | Cao et al. (2017)                               |
| *Lumbricus terrestris*    | bag                      | PE            | N/A         | N/A     | p         | 0.92±1.9 items g⁻¹ | N/A           | 28 d           | N/A             | yes             | no              | no                                               | Hodson et al. (2017)                             |
| *Lumbricus terrestris*    | home yard                | diverse       | yes         | N/A     | N/A       | N/A   | 0.87±1.9 items g⁻¹ | N/A           | 14 d           | yes             | N/A             | N/A             | Hospers et al. (2017)                                   | Huerta Lwanga et al. (2017a)                     |
| *Lumbricus terrestris*    | mesocosm                 | PE            | washed      | N/A     | N/A       | p     | <150           | 0.50% w/w (litter) | 14 d           | yes             | N/A             | N/A             | N/A             | Hospers et al. (2017)                                   | Huerta Lwanga et al. (2017b)                     |
| *Eisenia andrei*          | mesocosm                 | PE            | N/A         | no      | no        | b     | 710–2800       | 750 µg on 2.5 kg soil | 21 d           | yes             | N/A             | N/A             | no                                               | Rillig et al. (2017)                              |
| *Lumbricus terrestris*    | glass bottle             | PE            | washed      | N/A     | N/A       | p     | 150            | 7% w/w (litter)  | 60 d (earthworms) | yes             | yes             | N/A             | ≥62.5: intestinal damage                                 | Huerta Lwanga et al. (2018)                      |
| *Eisenia fetida*          | mesocosm                 | PE            | washed      | N/A     | N/A       | p     | 0.001          | 1000           | 28 d           | N/A             | yes             | N/A             | ≥125: altered enzyme activity                             | Rodriguez-Seijo et al. (2018)                   |
| *Aporrectodea rosea*      | mesocosm                 | PLA, PE       | washed      | N/A     | N/A       | p     | 0.100          | 1000           | 30 d           | N/A             | yes             | N/A             | growth †                                                | Boots et al. (2019)                              |
| *Eisenia fetida*          | mesocosm                 | PE            | washed      | N/A     | N/A       | p     | <2000          | soil extract    | 48 h / 56 d    | N/A             | N/A             | no                                               | Judy et al. (2019)                                |
| *Lumbricus terrestris*    | bag                      | PE            | N/A         | N/A     | p         | 1.7±0.8 x 361.6±367.0 | 0.1000       | 35 d           | N/A             | yes             | N/A             | ≥1000: metallothionein expression †                       | Prendergast-Miller et al. (2019)                 |
| *Eisenia fetida*          | mesocosm                 | PVC           | washed      | N/A     | N/A       | mf    | 250–1000       | 40 items on 0.5 kg soil | 14 d           | N/A             | N/A             | N/A             | ≥1000: heat shock protein 70 †                             | Rodriguez-Seijo et al. (2019)                   |
| *Eisenia fetida*          | mesocosm                 | PE            | washed      | N/A     | N/A       | p     | 300            | 0.20000        | 14 d           | N/A             | N/A             | ≥20000: altered enzyme activity                            | Wang et al. (2016c)                              |
| *Lumbricus terrestris*    | bag                      | PE            | washed      | N/A     | N/A       | p     | <300           | 0.100          | 28 d           | N/A             | yes             | N/A             | ≥1000: altered enzyme activity                            | Wang et al. (2016c)                              |
| *Lumbricus terrestris*    | glass beaker             | PE            | washed      | N/A     | N/A       | p     | <150           | 0.7% w/w (litter) | 14 d           | N/A             | N/A             | N/A             | ≥1000: altered enzyme activity                            | Yang et al. (2019a)                              |
| *Lumbricus terrestris*    | mesocosm                 | PE            | washed      | N/A     | N/A       | p     | <1500          | 7% w/w (litter)  | 14 d           | yes             | yes             | N/A             | ≥1000: altered enzyme activity                            | Yu et al. (2019)                                 |
| *Lumbricus terrestris*    | mesocosm                 | PE            | unwashed,   | N/A     | N/A       | p     | 1.5x1.5 cm²   | 4 items per dish | 14 d           | yes             | no              | N/A             | ≥1000: altered enzyme activity                            | Zhang et al. (2018)                              |
| *Eisenia fetida*          | petri dish               | PE            | washed      | N/A     | N/A       | p     | <400           | 0.1500         | 28 d           | N/A             | yes             | yes             | ≥1000: oxidative stress; ≥1000 mg/kg: neurotoxicity †    | Chen et al. (2020)                                |
Table 5: Microplastic studies on Enchytraeidae and Arenicola marina (mb=microbeads, p=particles, ms=microspheres, sed.=sediment, s=semisubhydric, [a] N/A=information not available). Concentrations refer to mg kg\(^{-1}\) dry soil in terrestrial soils and mg kg\(^{-1}\) dry sediment in semisubhydric soils, if not specially marked.

| Organism | Experimental environment | Plastic type | Aging | Coating | Additives | Shape | Size span [µm] | Concentrations | Exposure time | Passive transport | Active uptake | Bioaccum. dynamics | Measured adverse effects | Reference |
|----------|--------------------------|--------------|-------|---------|-----------|-------|----------------|----------------|---------------|------------------|--------------|-------------------|------------------------|-----------|
| Enchytraeus crypticus | Petri dish | PS | N/A | N/A | N/A | mb | 0.05..0.1 | 0..10% w/w (food) | 7 | N/A | yes | N/A | 1.2±0.8 items g\(^{-1}\) | Zhu et al. (2018c) |
| Enchytraeus crypticus | microcosm | PA, PVC | N/A | N/A | fluorescence | N/A | 13.150..106.150 | 20000..120000 | 20 h / 21 d | N/A | yes | N/A | ≥10% w/w: microbiome, weight | Lahive et al. (2018) |
| Arenicola marina | in situ | N/A | N/A | N/A | N/A | N/A | 10..90 | N/A | 10000..500000 items kg\(^{-1}\) | 14 d | N/A | yes | 10 µm: 9600±1800 items kg\(^{-1}\) 30 µm: 800±700 items kg\(^{-1}\) | Cauwenbergh et al. (2015) |
| Arenicola marina | liquid culture | PS | no | N/A | N/A | ms | 10..90 | N/A | 10000..500000 items kg\(^{-1}\) | 14 d | N/A | yes | 1.2±0.8 items g\(^{-1}\) | no |
| Arenicola | mesocosm | PS, PA | yes | biofilm | N/A | p | 500..1000 | 10..200 | 106..240 d | yes | no | N/A | N/A | N/A | Gebhardt and Forster (2018) |
| Arenicola marina | mesocosm | PS, PVC | N/A | N/A | N/A | p | 400..1300 | 0..74000 | 28 d | N/A | ≥400 µm | no | ≥74000: feeding, weight | Besseling et al. (2012) |
| Arenicola marina | mesocosm | HD-PE | N/A | N/A | N/A | p | 9.478..3.316 | 0..200000 mg kg\(^{-1}\) wet sed. | 31 d | N/A | N/A | N/A | ≥2000: respiration, casting | no | Green et al. (2016) |
| Arenicola marina | mesocosm | PE, PCBs | fluorescence | mb | 10..180 | 0..5000 | 28 d | N/A | yes | no | feeding activity, heap mass | Besseling et al. (2017) |
| Arenicola marina | mesocosm | PVC | N/A | N/A | not leaching | p | −130 | 0..500000 | 28 d | N/A | N/A | N/A | ≥10000: energy reserves, ≥50000: feeding, egestion, casting | Wright et al. (2013) |
3.4 Further invertebrates

As part of the microfauna, the phylum Nematoda (nematodes or roundworms) is an ecologically important branch containing >25000 species (Zhang, 2013) in freshwater, marine, endobiotic and soil habitats. Due to their diverse trophic interactions nematodes hold a central position in both bottom-up and top-down controlled food webs (Yeates, 2001; Ferris, 2010) and thus most likely the uptake and transfer of MP.

Active feeding of adults and larvae of different species on 0.5 to 6 µm PS/latex microspheres (the size of their bacterial prey) was proven by Nika et al. (2016) and Fueser et al. (2019). However, most MP experiments on Nematodes are based on the bacterial-feeding model organism Caenorhabditis elegans. Kiyama et al. (2012) showed the favored uptake of PS microspheres with sizes of 0.5 to 3 µm by adult and 0.5 µm by larval C. elegans. The ingestion of MP decreased in the presence of bacteria as the natural food source.

When larval stages and adults ingested PS between 0.05 and 5 µm within an aqueous suspension or on agar plates, adverse effects such as oxidative stress, neurodegeneration, intestinal and DNA damage or dysfunction in motility, growth, life span, defecation, reproduction or energy metabolism appeared from a wide spectrum of concentrations from ≥1 µg l⁻¹ up to ≥86.3 mg l⁻¹ (Zhao et al., 2017; Dong et al., 2018; Kim et al., 2019; Lei et al., 2018a; Lei et al., 2018b; Qu et al., 2019a). These effects are not seen below 1 µg l⁻¹ (Qu et al., 2019b), and are enhanced due to amino modifications on micropshere surfaces (Qu et al., 2019c). The incubation on agar plates with PE, PP and PVC particles <70 µm caused similar influences on survival, fertility, brood size and intestinal function (Lei et al., 2018b). Leachates from soils amended with 5 mg kg⁻¹ dry soil of HD-PE and PVC decreased reproduction in laboratory cultures, but there was no effect shown on survival and after application of PET (Judy et al., 2019). Furthermore, silica nanoparticles (0.05 µm) are not only taken up orally but also via the vulva and spermathecae and migrate into gonad cells (Scharf et al., 2013). This process was confirmed for PS nanoparticles with the potential of a transfer to the progeny (Zhao et al., 2017).

The clear adverse effects of these studies are limited in their representativity by a narrow restriction to liquid cultures and a single model organism. Broader studies like on prominent soil-born nematodes such as Acrobeloides buetschlii (Frey, 1971) are still lacking. When assuming in first proximity mg l⁻¹ solution = mg kg⁻¹ dry soil, the applied concentrations between 0.001 and 86.8 mg l⁻¹ match lower levels of soil contamination.

Feeding studies on the phylum Rotifera with MPs are fully based on PS microbeads and model organisms of the planktonic genus Brachionus. However, this data can carefully be transferred to soil environments as also soil rotifers are aquatic organisms living in water-filled pores and waterfilms. Different Brachionus sp. ingest microbeads <10 µm with strong preference for particles the size of their natural food source, namely bacteria and algae with
2 to 5 µm in diameter (Vadstein et al., 1993; Heerkloß and Hlawa, 1995; Baer et al., 2008; Jeong et al., 2016). The uptake appears to be selective as microbeads are fewer incorporated than bacteria and algae (Vadstein et al., 1993). The egestion of particles ≤0.5 µm is hindered compared to 6 µm (Jeong et al., 2016). In suspension, microbeads ≤0.5 µm cause adverse effects on fertility and life span at ≥0.1 mg l⁻¹ as well as oxidative stress and less growth at ≥10 mg l⁻¹ (Jeong et al., 2016; Sun et al., 2019).

Terrestrial mollusks comprise snails and slugs within the class of **Gastropoda**. These grazers feed on bacterial biofilms, fungi and plant tissue (Parkyn and Newell, 2013). Studies on terrestrial species are sparse, but data on the benthic *Littorina sp.* imply passive transport and non-selective MP uptake by feeding on surfaces with contaminated feces and mucus trails of other snails (Gutow et al., 2019). With focus on *benthic* snails, Imhof and Laforsch (2016) found no significant influence on growth parameters and fertility of juveniles and adult *Potamopyrgus antipodarum* even when a food source with 70 % w/w of 5 to 600 µm sized fragments was given (a mixture of PA, PC, PET, PS, PVC). In contrast, adverse effects were found in recent work on the terrestrial snail *Achatina fulica*, that showed uptake and complete gastrointestinal passage within 48 h with partial degradation of PET fibers (appr. 1258x76 µm), but reduced excretion and food intake as well as increased oxidative stress at concentrations of ≥0.01 g kg⁻¹, ≥0.14 g kg⁻¹ and ≥0.71 g kg⁻¹ dry soil, respectively (Song et al., 2019).
Table 6: Microplastic studies on nematods (ms=microspheres, fr=fragments, np=nanoparticles, mb=microbeads, ms=microspheres, ox.=oxidative, [8]). Concentrations refer to mg kg⁻¹ dry soil, if not specially marked.

| organism | experimental environment | plastic type | aging | coating | additives | shape | size span [µm] | concentrations | exposure time | passive transport | active uptake | bioaccum. dynamics | measured adverse effects | reference |
|----------|--------------------------|--------------|-------|---------|-----------|-------|---------------|----------------|---------------|-------------------|-------------|-------------------|--------------------------|-----------|
| Caenorhabditis elegans | agar plate | PS | N/A | carboxyl | sulfate amino | fluorescence | ms | 0.1...6.0 | N/A | 0.5...2 h | N/A | yes | 0.5...3 µm | N/A | Kiyama et al. (2012) |
| Caenorhabditis elegans | liquid culture | PS | N/A | carboxyl | fluorescence | ms | 0.1 | 0.001...10 mg l⁻¹ | 4.5 d | N/A | Yes | N/A | ≥0.01 mg l⁻¹: motility ↑, growth ↓, defecation ↑, within gonads | Zhao et al. (2017) |
| Caenorhabditis elegans | liquid culture | PS | N/A | (ζ=10mV) | fluorescence | ms | 0.1 | 0.00001...0.001 mg l⁻¹ | N/A | N/A | Yes | N/A | ≥0.01 mg l⁻¹: motivity ↑, ox. stress ↓ | Dong et al. (2018) |
| Caenorhabditis elegans | liquid culture | PS | N/A | N/A | preservatives, fluorescence | ms | 0.05...0.2 | 17.3...86.8 mg l⁻¹ | 24 h | N/A | Yes | N/A | ≥17.3 mg l⁻¹: motivity ↑, reproduction ↓, growth ↓, metabolic dysf. | Kim et al. (2019) |
| Caenorhabditis elegans | liquid culture | PS | N/A | (ζ=10mV) | fluorescence | ms | 0.1 | 0.001...1 mg l⁻¹ | N/A | N/A | Yes | N/A | ≥1 mg l⁻¹: neurodegeneration | Qu et al. (2019a) |
| Caenorhabditis elegans | liquid culture | PS | N/A | N/A | N/A | ms | 0.1...5 | 1 mg l⁻¹ | 3 d | N/A | Yes | N/A | motivity ↑, survival ↓, growth ↑, ox. stress ↓, neurotoxicity | Lei et al. (2018a) |
| Caenorhabditis elegans | agar plate | PE, PP, PVC, PS | no | N/A | N/A | fr, ms | 0.1...0.2 | 0.5...10.0 mg m⁻³ | 2 d | N/A | Yes | N/A | ≥0.5 mg m⁻³: survival ↑, at 5 mg m⁻³: growth ↓, reproduction ↓, ox. stress ↑, intestinal damage mainly ≤1μm: intestinal damage | Lei et al. (2018b) |
| Caenorhabditis elegans | agar plate | PS | N/A | N/A | N/A | np | 0.05 | 2500 mg l⁻¹ | 7 d | N/A | Yes | N/A | within tissue and gonads | Scharf et al. (2013) |
| Caenorhabditis elegans | liquid culture | HDPE, PET, PVC | no | N/A | N/A | np | <2000 | soil extract | 72 h | N/A | N/A | N/A | reproduction ↓ | Judy et al. (2019) |
| Caenorhabditis elegans | agar plates | latex | N/A | N/A | fluorescence | mb | 0.5 | N/A | 30 min | N/A | yes | N/A | motility ↓, growth ↑, motility ↓, reproduction ↓, motility ↓, ox. stress ↓ | Nika et al. (2016) |
| Panagrolaimus thienemann | agar plate | PE, PP, PVC | no | N/A | N/A | fluorescence | ms | 0.1...2 | 3.10⁶...10⁶ items l⁻¹ | 4.73 h | N/A | ≤3µm | ≤0.5µm | ≤1µm | ≤1µm | ≤1µm | ≤1µm | ≤1µm | ≤1µm | ≤1µm | ≤1µm | ≤1µm | ≤1µm | N/A | Fueser et al. (2019) |
| Plectus acuminatus | liquid culture | PS | N/A | N/A | N/A | fluorescence | ms | 0.5...6 | 0.5...10¹⁰ | N/A | N/A | N/A | N/A | N/A | N/A | N/A | N/A | N/A | N/A | N/A | N/A | N/A | Fueser et al. (2019) |
| Poikilolaimus regenfussi | agar plate | PE, PP, PVC | no | N/A | N/A | fluorescence | ms | 0.1 | 0.0001...0.001 mg l⁻¹ | N/A | N/A | N/A | N/A | ≥0.01 mg l⁻¹: reproduction ↓, DNA damage | Qu et al. (2019b) |
| Acrobelesus nanus | liquid culture | PS | N/A | N/A | N/A | ms | 0.1 | 0.001...1 mg l⁻¹ | N/A | N/A | yes | N/A | ≥0.01 mg l⁻¹: reproduction ↓, DNA damage | Qu et al. (2019c) |
| Pristionchus pacificus | agar plate | PE, PP, PVC | no | N/A | N/A | fluorescence | ms | 0.1 | 0.001...1 mg l⁻¹ | N/A | N/A | yes | N/A | ≥0.01 mg l⁻¹: reproduction ↓, DNA damage | Qu et al. (2019c) |
| Aphelenchoides parthenin | liquid culture | PS | N/A | N/A | N/A | ms | 0.1 | 0.001...1 mg l⁻¹ | N/A | N/A | yes | N/A | ≥0.01 mg l⁻¹: reproduction ↓, DNA damage | Qu et al. (2019c) |

N/A=information not available.
Table 7: Microplastic studies on Rotifera and Gastropoda (ms=microspheres, mb=microbeads, fr=fragments, f=fibers, ox.=oxidative, pref.=preferential, p=planctic , b=benthic, [a] N/A=information not available). Concentrations refer to mg kg⁻¹ dry soil, if not specially marked.

| Organism                        | Experimental Environment | Plastic Type | Aging | Coating | Additives | Shape | Size Span [µm] | Concentrations | Exposure Time | Passive Transport | Active Uptake | Bioaccum. Dynamics | Measured Adverse Effects                             | Reference                      |
|---------------------------------|--------------------------|--------------|-------|---------|-----------|-------|----------------|----------------|---------------|------------------|--------------|-------------------|-----------------------------------------------------|-------------------------------|
| Brachionus plicatilis           | liquid culture           | PS           | N/A   | N/A     | carboxyl fluorescence | ms    | 1.6..20        | 5·10⁻⁶ [µm]³ l⁻¹ (~5.25 mg l⁻¹) | 35 min       | N/A             | ≤10 µm          | pref. 4.5 µm       | N/A                                                              | Bear et al. (2006)             |
| Brachionus plicatilis           | liquid culture           | latex        | N/A   | N/A     | fluorescence | mb    | 0.3..3.1       | 3·10⁻⁷..7·10⁸ items l⁻¹ (~0.0004..11 mg l⁻¹) | 20 min       | N/A             | yes             | pref. 2 µm         | N/A                                                              | Vadstein et al. (1993)          |
| Brachionus koreanus             | liquid culture           | PS           | no    | N/A     | N/A       | fluorescence | mb    | 0.05..6        | 0...20 mg l⁻¹    | 1 d           | N/A             | yes             | ≥0.5 µm, >0.1 mg l⁻¹; reproduction ↓, survival ↓; oxidative stress ↑ | Jeong et al. (2016)            |
| Brachionus plicatilis           | liquid culture           | PS           | N/A   | N/A     | N/A       | mb    | 0.07..7        | 0...20 mg l⁻¹    | N/A          | N/A             | yes             | N/A                                                              | Sun et al. (2019)              |
| Brachionus quadridentatus ⚫     | mesocosm                 | PS           | N/A   | N/A     | N/A       | ms    | 2..10          | 5..100          | 8..10 d       | N/A             | yes             | N/A                                                              | Heerkloß and Hlawa (1993)      |
| Littorina littorea              | microcosm                | PMMA         | N/A   | N/A     | N/A       | fluorescence | fr    | 0.0..20        | increasing      | 18 h          | N/A             | ≥140: food intake ↓ | N/A                                                              | Gutow et al. (2019)            |
| Potamopyrgus antipodarum ⚫     | aquarium                 | PET, PS, PVC, PA, PC | N/A | N/A     | no / stained | fr    | 5..600         | 0..70% w/w (food) | ≤141 d  | N/A             | yes             | N/A                                                              | Imhof and Laforsch (2016)      |
| Achatina fulica                | mesocosm                 | PET           | N/A   | N/A     | no / stained | f     | approx. 125.8x76 µm | 10..710        | 28 d          | N/A             | yes             | ≥140: food intake ↓; excretion ↓; excretion ↓; gastrointestinal damage | Song et al. (2019)             |

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3.5 Vertebrates

Different taxa of the class of Amphibia have a predator function within the edaphic food web (e.g. preying on invertebrates) (Hebrard et al., 1992). While no data on the reaction to soil MPs are available neither for the legless Gymnophiona nor for adults of the order Anura, sparse data on tadpoles of aquatic frogs suggest uptake followed by regular excretion of PS microspheres as shown with Xenopus tropicalis (Hu et al., 2016). Further, there exist no data on the families Serpentes (snakes) and Anguidae within the class of Reptilia, residing at the outer rim of the food web.

Within the broad field of Mammalia, studies on MP ingestion are sparse and focus on mice as a rodent model organism. Feeding of mice with PS microspheres of 1 to 14 µm in concentrations of \(1.49 \times 10^6\) to \(4.55 \times 10^7\) particles at a volume of 10 ml kg\(^{-1}\) body weight for 4 weeks showed no adverse effects (Stock et al., 2019). In contrast, longer exposition (6 weeks) with lower concentrations of particles with the same shape and size range changed the mouse microbiome and caused metabolic and intestinal dysfunction (Lu et al., 2018; Jin et al., 2019), which comes along with bioaccumulation within organs (Yang et al., 2019b). These studies are regularly conducted with passive feeding and exclude active foraging on perceptible plastic particles. However, the uptake via prey or feeding on contaminated roots and litter is highly probable. Further Rodentia – Cricetidae (hamsters, lemmings, voles), Bathyergidae (blesmols, mole-rats), Octodontidae as well as Spermophilus (ground squirrels) and Marmota (marmots) within the family of Sciuridae – were not yet studied, just as other mammalian (sub)orders like Chrysochloridae (golden moles), Cingulata (armadillos), Macroscelidea (elephant shrews), Notoryctemorphia and Peramelemorphia.
Table 8: Microplastic studies on Anura (An.) and Rodentia (ms=microspheres, a=aquatic, N/A=information not available).

| Organism     | Experimental environment | Plastic type | Aging | Coating | Additives | Shape | Size Span [µm] | Concentrations | Exposure Time | Passive Transport | Active Uptake | Bioaccum. Dynamics | Measured Adverse Effects | Reference          |
|--------------|--------------------------|--------------|-------|---------|-----------|-------|---------------|----------------|---------------|-------------------|---------------|-------------------|------------------------|------------------------|
| Xenopus tropicalis a | Petri dish                | PS           | N/A   | N/A     | fluorescence | ms    | 1.10          | 100..10^7 items l^-1 (55·10^-9..55 mg l^-1) | 48 h          | N/A              | yes         | egestion within days | N/A             | Hu et al. (2016) |
| transgenic mice | in vivo                   | PS           | N/A   | N/A     | sulfate    | fluorescence | ms    | 1             | 4.55·10^7 items per mouse (0.025 mg per mouse) | 28 d          | N/A              | yes         | N/A              | no             | Stock et al. (2019) |
| mice         | in vivo                   | PS           | N/A   | N/A     | sulfate    | fluorescence | ms    | 5             | 0.1..1 mg l^-1 (food) | 42 d          | N/A              | yes         | N/A              | ≥0.1 mg l^-1: microbiome, metabolic dysfunction | Jin et al. (2019) |
| mice         | in vivo                   | PS           | N/A   | N/A     | N/A        | ms    | 0.5..50       | 0.1..1 mg l^-1 (food) | 35 d          | N/A              | N/A         | N/A              | ≥0.1 mg l^-1: metabolic dysfunction | Lu et al. (2018) |
| Mus musculus | in vivo                   | PS           | N/A   | N/A     | N/A        | fluorescence | ms    | 5..20         | 200 mg l^-1 (food) | 28 d          | N/A              | yes         | 8x, 8±5 and 0.71±0.14 mg kg^-1 body weight | N/A             | Yang et al. (2019b) |
4 Synthesis

4.1 Summarized observations

Our systematic search comprised recent research on the interaction of soil organisms with MP, but also studies with focus on feeding experiments, that are published much earlier than the awareness on plastic in the environment appeared. The numerous studies found with focus on the ingestion of MPs consistently showed the active uptake by diverse soil organisms with few exceptions spread over the whole branch of invertebrates. In addition, also studies on adverse effects caused by the intake of MP contaminated food (e.g. of food pallets by dung beetles) imply the ingestion into the test organism. Distinct size preferences are observed in dung beetles, nematodes, rotifers and ants showing that mainly particles are ingested, that are small enough to enter the gastrointestinal tract. In contrast, active comminution by gnawing on larger particles was tested only for a few taxa and confirmed for woodlice, termites and mealworms, and in the case of earthworms only after initial weathering.

After the ingestion, MP is translocated actively until excretion or death of the transporting organism, which was only directly shown in experiments with earthworms. The passive transport by attachment, dragging and pushing was investigated in a few experiments with earthworms, mites and springtails that partly worked without soil substrate and consistently showed positive results.

After exposition to MP, a pattern of adverse effects can be seen: Across various taxa, altered microbiomes, reduced motility, body mass, fertility and life span as well as increased oxidative stress and metabolic malfunctioning occur in different combinations mainly due to µm-sized MP in and above the whole known natural range of concentrations. For some taxa such as Nematodes, Gastropoda and Rotifera these effects appear at natural and increased MP concentrations (<100 mg kg⁻¹ dry soil), for Collembola and Lumbricidae at concentrations like in highly contaminated sites (≥1000 mg kg⁻¹ dry soil) and for Enchytraeidae, Arenicola marina and in further experiments with earthworms at implausibly high values. The data show a tendency, that the effects occur at lower concentrations, when the added particles are smaller. Small sized particles also provide the highest surface/volume ratio and thus the highest reactive surface per weight.

Most studies work with defined increasing MP concentrations and particle sizes in soil substrates and food sources, which can be used to determine relationships between environmental concentrations and adverse effects. However, the lack of information about intake rates, grades of accumulation and effective prey-predator transfer leads to a gap within the chain of explanation for toxic effects on the soil organisms. In some experiments, the intestinal passage of MP and sizes preferably retained within the gut were shown, but there are no experiments that could demonstrate quantitative bioaccumulation. In contrast,
quantification of the retained and egested MP particle size fractions might be biased due to
gnawing and intestinal comminution as shown for woodlice, termites, mealworms, snails and
earthworms.

In order to improve our understanding of processes underlying adverse effects of MP on soil
organisms, data on ingestion rates, dwell times, biodegradation and egestion rates are
important bricks e.g. to reveal bioaccumulation dynamics. However, there are only a few data
on biodegradation (mealworms, snails, earthworms), egestion (rotifers, frogs, snails,
earthworms) and remaining concentrations in the body (lugworm, mice, earthworms).
4.2 Limitations of previous studies

The available studies worked with items within the full size span of micro- and nanoplastics (≤5000 µm). Approximately 72% of the experiments used microplastic (0.1 to 5000 µm), only 6% nanoplastic (<0.1 µm), 10% included macroplastic (>5000 µm) and 12% used microplastic of undefined size. When MP ≥50 µm was applied, mainly particles and fragments made of PE and PVC were used, whereas PS/latex microspheres were mainly applied for sizes ≤10 µm (Table 1). The latter are readily available, highly standardized and are mostly used with fluorescent dyes and either without additional functional groups, carboxylated or, more rarely, with amino or sulfate groups. However, there are indications that the spectrum of particle type and shape used in experiments does not correspond to the properties of particles in soils. In different natural as well as agriculturally and industrially contaminated terrestrial and semisubhydric sites, fibers and fragments of PE and PP, mostly ≤100 µm, were much more abundant than PVC, PET and PS items (Claessens et al., 2011; Vianello et al., 2013; Nor and Obbard, 2014; Naji et al., 2017; Zhang and Liu, 2018; Li et al., 2018a). This is probably caused by high loads of MP fibers in discharged waste water and sewage sludge, which is used in agricultural sites worldwide (Mahon et al., 2016; Li et al., 2018b). It is likely that shape plays an important role for the ingestion of MP items. Unfortunately, we did not find studies that have carried out a complete classification of sampling sites according to plastic origin, size and type, that could help to evaluate differences between former experimental and natural plastic composition to achieve the most realistic experimental conditions. Little knowledge about the size distribution of MP in soils furthermore complicates the determination of realistic concentrations for the addition of a certain particle size spectrum. All reviewed studies either arbitrarily set their applied concentrations or had to base them on measurements of total specific MP masses, regardless of how much of this mass is in the tested size range. This may lead to a false estimation of total adverse MP concentrations.

In contrast to particle type and shape, the documentation of chemical properties of MP samples in most of these studies is fragmentary. Some experiments explicitly mentioned that the added plastic was unweathered, whereas most studies lack information about the degree of aging implying that unweathered items were used. Only a few experiments involved aging of MP, but without comparison to results of natural weathering (Tsunoda et al., 2010; Gebhardt and Forster, 2018). That is in conflict with natural conditions, as plastic that remains within the soil after littering, sewage sludge application or plastic mulching shows signs of weathering, e.g. modified carbonyl indices (Andrady, 2017), while unweathered soil MP might be rare. In addition, Zhang et al. (2018) showed that earthworms actively comminute only weathered bioplastics. In experiments using PS microspheres, carboxylation is often used to imitate a reduced hydrophobicity due to weathering. However, according to manufacturer information microplastics only have little influence on hydrophobicity.
Weathering of MP surfaces within soils comes along with biofilm growth and adsorption of organic molecules, which could potentially affect the attractiveness or toxicity for grazers and other organisms. Such coatings were applied only in a few cases (Besseling et al., 2017; Angotti et al., 2018; Gebhardt and Forster, 2018), but were not documented in most studies. Similarly, the type and concentration of additives such as flame retardants, anti-oxidants or stabilizers often remained undocumented, with exception of fluorescent dyes, that are well mentioned. The release of additives can have a harmful effect on the test organism, as shown for aquatic environments (e Silva et al., 2016). Some studies on the ingestion of MP by the soil mesofauna indicate that the diameter of the gastrointestinal tract is a useful upper size limit for added particles, as far as the organism is unable to crush them (Heerkloß and Hlawa, 1995; Holter, 2000; Holter et al., 2002; Holter and Scholtz, 2005; Baer et al., 2008; Fueser et al., 2019). However, using only ingestible particle sizes in their natural concentrations neglect the adverse effects of plastic leachates, which can also get into the soil solution and onto the mineral phase from larger particles and affect soil life.

The conditions of incubation differ considerably in terms of habitats and duration of exposure. In most studies, the exposure ranges from a few minutes to a few days in experiments with micro- and small mesofauna and hours to several weeks in experiments with large meso- and macrofauna and is mainly based on excretion or reproductive cycles. Long-term studies, which are indeed difficult to carry out in mesocosms, practically do not exist. However, certain adverse effects might only establish themselves after long term trials, as was shown for the influence of pesticides (Pelosi et al., 2015).

Some experiments were carried out in soil-free test environments such as liquid cultures or Petri dishes with nutrient solutions or a specific food source (nematods, rotifers, mice). Therefore, motivity is less restricted and feeding behavior can be altered compared to cultivation within soil environments. For example, the ingestion of MP by nematodes decreases in the presence of an alternative and more natural food source like bacteria, which can significantly reduce the bioaccumulation and thus the effective toxicity (Kiyama et al., 2012). This can lead to less consumption of MP in soil environments and an overestimation of the toxicity in liquid culture experiments. Also, all laboratory feeding experiments were carried out by use of only one species. The complexity of the food web in soils is thereby excluded and the potential accumulation from prey to predators still unexplored.
4.3 Pinpoints for future research

Most studies reviewed in this work have a pioneering role in MP research and, thus, are subject to some experimental limitations caused by an early state of knowledge. The adverse effects recently found are alarming, but must be considered under the restrictions named above. We propose the following points as part of a modus operandi for future MP research.

In past studies, particular adverse effects of MP were measured only for certain sizes, shapes, coatings, leachates or adsorbed substances (Tables 2 to 8). Experimental concentrations were assumed randomly or derived from cumulative concentrations of one or more MP types measured in natural soils (approx. 1 to some 1000 mg kg\(^{-1}\) dry soil), regardless of size. For those specific experiments coming, the spectrum of concentrations used should be adapted to the quantities of the size spectrum, that occurs within the soil. For future studies on mixed contaminations, we recommend to evaluate the overall adverse effects of PE, PP, PVC, PET, PU and PS to certain test organisms by use of typical MP-specific spans of concentration, size and shape distribution in natural soils or food samples. This previously requires well-structured data of appropriate MP type, shape and size for different soils in differently contaminated areas.

Experiments on adverse effects should be applied within soil matrices to allow the interplay of plastic, natural organic and mineral matter. The MP should be weathered, as plastic in soils underlie broad environmental aging. Pre-weathering of MP should therefor not only be performed in climate chambers (e.g. following DIN EN ISO 4892-2/3), but also include subsequent leaching and equilibration of additives or coatings within the soil matrix before the main experiment. Furthermore, the experimental design may consider coatings with biofilms or attractants and even particle color to regulate the preference of the test organisms.

Most detailed information about ingestion are available for dung beetles, nematods and earthworms, data on adverse effects on nematods, earthworms, lugworms and collembola. Future experiments should focus on a larger variety of ecologically relevant taxa like Coleoptera, Formicidae, Acari, Oniscidea, Collembola, Lumbricidae, Enchytraeidae, Nematoda and Gastropoda. The studies are recommended to conduct with emphasis on uptake, accumulation and key adverse effects like on survival rate, motility, growth and fertility as well as on the stability of the intestinal microbiome. Further studies with more than one test organism are important to foster our understanding of MP within certain food chains. Also long-term experiments might reveal adverse effects, which evolve slowly within populations. This may enable the assessment of the distribution and effects of MP within the food web and the resulting long-term impact on soil ecosystems.
5 Conclusion

Our review of 77 studies on the impact of microplastic on the soil fauna shows a considerable diversity and distribution of adverse effects within the soil tree of life. However, these effects have to be considered carefully, as many experiments did not use plastic matching properties within natural soils and found adverse effects only at concentrations like in highly contaminated soils or above. To elucidate effective concentrations and properties for short and long-term effects on soil faunal health, the most exact reproduction of plastic properties within the soil matrix and natural living conditions of the test organisms is necessary together with a better knowledge on common concentrations and size distributions of soil microplastic. For future experiments we therefore recommend to choose compositions of type, shape, size, concentration, grade of weathering, leachability and coating with biofilms and other organic matter as expected in the habitat to be examined. Furthermore, coming studies should include long-term exposure and food chain experiments to get a better look at the effect of even smaller MP concentrations and their enrichment within the food web. This may give us a better way of assessing the impact of global microplastic contamination on e.g. soil biodiversity, soil carbon cycles and soil quality.
Author contribution
Frederick Büks developed the review concept, collected data and prepared the manuscript except for earthworms. Nicolette Loes van Schaik did all the work on earthworms. Martin Kaupenjohann supervised the study by participating in structural discussions on the idea and concept of the paper as well as the final corrections.

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Competing interests
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