Microplastic analysis using chemical extraction followed by LC-UV analysis: a straightforward approach to determine PET content in environmental samples

Axel Müller1,2, Caroline Goedecke1, Paul Eisentraut1, Christian Piechotta1 and Ulrike Braun1*

Abstract

Background: The ubiquitous occurrence of microplastic particles in marine and aquatic ecosystems was intensively investigated in the past decade. However, we know less about the presence, fate, and input paths of microplastic in terrestrial ecosystems. A possible entry path for microplastic into terrestrial ecosystems is the agricultural application of sewage sludge and solid bio-waste as fertilizers. Microplastic contained in sewage sludge also includes polyethylene terephthalate (PET), which could originate as fiber from textile products or as a fragment from packaging products (foils, bottles, etc.). Information about microplastic content in such environmental samples is limited yet, as most of the used analytical methods are very time-consuming, regarding sample preparation and detection, require sophisticated analytical tools and eventually need high user knowledge.

Results: Here, we present a simple, specific tool for the analysis of PET microplastic particles based on alkaline extraction of PET from the environmental matrix and subsequent determination of the monomers, terephthalic acid, using liquid chromatography with UV detection (LC-UV). The applicability of the method is shown for different types of PET in several soil-related, terrestrial environmental samples, e.g., soil, sediment, compost, fermentation residues, but also sewage sludge, suspended particles from urban water management systems, and indoor dust. Recoveries for model samples are between 94.5 and 107.1%. Limit of determination and limit of quantification are absolute masses of 0.031 and 0.121 mg PET, respectively. In order to verify the measured mass contents of the environmental samples, a method comparison with thermal extraction-desorption-gas chromatography–mass spectrometry (TED-GC/MS) was conducted. Both methods deliver similar results and corroborated each other. PET mass contents in environmental samples range from values below LOQ in agriculture soil up to 57,000 mg kg$^{-1}$ in dust samples.

Conclusions: We demonstrate the potential of an integral method based on chemical extraction for the determination of PET mass contents in solid environmental samples. The method was successfully applied to various matrices and may serve as an analytical tool for further investigations of PET-based microplastic in terrestrial ecosystems.

Keywords: Microplastic, Fibers, PET, Analysis, LC-UV, Soil, Sewage sludge, Depolymerization

*Correspondence: ulrike.braun@bam.de
1 Bundesanstalt für Materialforschung und -prüfung (BAM), Unter den Eichen 87, 12205 Berlin, Germany
Full list of author information is available at the end of the article

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likely, however, that a further significant increase in the number of microplastic particles will have environmental impacts, whether it is a change in the accumulation behavior of natural particles [2] or the sorption and permeation of persistent pollutants of larger quantities [3, 4]. Marine ecosystems were intensively studied in the last decade, while the status of terrestrial ecosystems remains mainly unknown [5]. This knowledge gap is particularly alarming as plastic contamination in terrestrial ecosystems might be 4–23 times higher than in aquatic ecosystems [6].

Besides littering, a possible input path of plastic in soils is the application of sewage sludge as fertilizer [7]. Fibers and primary microplastic particles from laundry, cosmetic, and hygienic products are mostly retained in water treatment plants and end up in the sewage sludge [8]. Sewage sludge and municipal solid waste compost [9] are globally used as fertilizers in agriculture. A microplastic input of 63,000–430,000 t a⁻¹ was estimated for European farmlands through the use of sludge as fertilizer [10]. Studies of agricultural soil detected concentrations between 78 particles kg⁻¹ [11] and 18,760 particles kg⁻¹ [12] and 0.88–0.95 mg g⁻¹ [17]. Studies investigating the occurrence of microplastic in the soil after the applications of sewage sludge resulted in concentrations between 1100 and 3500 particles kg⁻¹ [13]. Depending on the sewage sludge treatment process, sewage sludge shows concentration up to 34,000 particles kg⁻¹ [14, 15], or mass content up to 12,000 mg kg⁻¹ [16] or 3300 mg kg⁻¹ [17]. 92% of the detected microplastic in soil aggregates from croplands in China consisted of synthetic fibers [12]. Synthetic fibers comprise 70% of polyethylene terephthalate (PET) regarding the production volume [18], and could remain up to 15 years in soils, after the application of sludge [19]. These fibers might negatively influence soil biota. A recent study indicate that the intake of PET fibers by snails leads to reduced food intake and induced damage in the villi of the gastrointestinal walls [20]. Furthermore, fibers might negatively affect the germination of ryegrass [21].

The few existing conducted studies about microplastic in soils use different analytical methods with non-comparable results. Therefore, relevant sources, occurrences, transport pathways, and the fate of microplastic are difficult to assess. Most studies use optical or microscopic–spectroscopic methods as Fourier-transform infrared spectroscopy (FTIR) or Raman spectroscopy. These methods deliver information about particle numbers, size, and shape. Those spectroscopic methods are very time-consuming for environmental samples, especially because often, complex sample pretreatment is needed. It is mandatory to reduce the inorganic matrix and enrich the microplastic concentration before analysis [22, 23]. Usually, density separation is used for this purpose. To reduce the organic matrix before FTIR or microscopy analysis, an additional oxidation step with peroxide is needed [11, 12, 22, 24]. A faster, spectroscopic method is Fourier-transform near-infrared (FT-NIR) spectroscopy by the use of a fiber-optic reflection probe. It was successfully used to determine the presence of PET in soil samples without pretreatment, but requires minimum mass contents of 1%, which is very high and practically not observed in reality [25].

An alternative is the use of integral methods, which are faster and deliver mass fractions, an important value for monitoring. Thermoanalytical methods determine polymer mass fractions by the detection of specific decomposition products formed under thermal treatment. Common are pyrolysis–gas chromatography—mass spectrometry (Py–GC/MS) [26, 27] or thermal extraction–desorption–gas chromatography—mass spectrometry (TED–GC/MS) [28, 29].

Furthermore, chemical methods are possible with two principal strategies. Either polymer chains can be extracted by solvents and detected afterward [17, 30, 31]. Regarding the poor solubility of polymers, including PET, these methods often require poisonous or expensive solvents. This is especially disadvantageous when large sample volumes must be analyzed, which is mandatory to account for the heterogeneity of soil samples [1]. The second principle is based on the decomposition of the polymeric chain and subsequent analysis of specific decomposition products. The latter is generally only possible when the polymer structure offers specific functionality for bond cut, which, for example, is present in polyesters or polyamides.

Regarding the analysis of suspected high contents of PET in soil from fertilization, such extraction techniques developed by Wang et al. [32] are of high interest for routine application. Wang et al. described a way to depolymerize PET in its monomers, among others, terephthalic acid, and to subsequently detect the terephthalic acid by LC–MS/MS (liquid chromatography coupled with tandem mass spectrometry). LC-based methods are widespread in analytical service laboratories, could be used in routine analysis, and have particularly low detection limits.

The objective of the present study is to adapt the sample extraction procedure described by Wang et al. to soil samples and to transfer the analysis to a LC-UV system (liquid chromatography with ultraviolet detection) [32]. LC–MS/MS is a complex method for the structure elucidation of trace substances in environmental samples, especially for chemical compounds of unknown composition (non-target analysis). The application of LC-UV, however, can only provide an assignable signal if the
structure of the components is known. Since the terephthalic acid is already known as a target for the detection of PET, we have therefore adapted the LC–MS/MS method by Wang et al. to LC-UV. LC-UV is a more robust and cost-effective method because the prerequisites for MS measurement (ultra-high vacuum) are not necessary, and the maintenance procedures for the MS, such as sticky char contaminations cannot occur. Here we present for the first time the use of LC-UV for the determination of PET mass content in soil. Such LC techniques require moderate conditions of sample preparation without critical chemicals in terms of costs and toxicity [17, 30, 31], and therefore allow investigations of high sample masses. This method is not limited to PET analysis in soil but suitable for the detection of PET particles in any solid environmental matrix, such as sewage sludge or bio-waste. It is also applicable for analysis of PET in dust or airborne particles and could even be used in the future for the monitoring analysis of PET fibers from washing machines or PET particles in bottled water.

**Materials and methods**

**Materials and environmental samples**

As a reference material bottle-grade PET pellets (PET Lighter C 93, provided by Equipolymers, Schkopau, Germany) were cryo-milled under constant cooling with liquid nitrogen in a Retsch CryoMill (Retsch GmbH, Haan, Germany). A pre-cooled 50-ml milling beaker equipped with a 25-mm steel ball was used for milling. After milling at 5 Hz for 5 min, the frequency was increased to 25 Hz for 5 min and subsequently reduced again to 5 Hz for 0.5 min. Furthermore, additional real polymeric samples of PET were used. They came from everyday applications, such as fibers from textiles and fragments cut from a commercial single-use soda bottle.

| Environmental sample | Origin | Presumed source of PET entry | Pyrolysable organic content (180–600 °C) |
|----------------------|--------|-----------------------------|---------------------------------------|
| Beach sediment       | Warnemuende 54.194910N 12.140777E | Marine littering                | <1%                                   |
| Agriculture soil     | Baden-Wuerttemberg 48.787672N 8.179543E | Performed sewage sludge fertilization | 3.6%                                |
| Compost              | Lower Saxony 48.179543E           | Littering from urban waste       | 18–24%                               |
| Fermentation residue | Berlin                              | Littering from urban waste       | 34–39%                               |
| Suspended particles  | Kaiserslautern                      | No content suspected due to WWTP process | 59%                                |
| Filter residue of greywater | Rhineland-Palatinate                | Fibers of textile washing        | 62%                                  |
| Sewage sludge        | Berlin                              | Separation and enrichment due to WWTP process | 47%                                |
| Dust samples         | Berlin                              | Textile fibers in indoor air      | 72–78%                               |
homogenized by milling in a liquid nitrogen-cooled centrifugal mill (ZM 200, Retsch GmbH, Haan, Germany) with 1-mm mesh size.

Although the paper focuses on soils, we investigated also samples from urban water management. Here we expected PET, especially from fragmented littering or from textile products (fibers). Suspended particles were collected using a sedimentation trap in the effluent of a wastewater treatment plant (WWTP) in Kaiserslautern. Here we did not expect a PET signal from the purification line of the plant. Particles from greywater were collected by filtration (>100 µm), from a farmstead in Rhineland-Palatinate. Sewage sludge samples were taken from a WWTP in Berlin.

Finally, we chose two indoor dust samples with an expected high loading of textile fibers. Dust samples were collected in two households in Berlin.

For sample preparation butanol (>99.5%), methanol (>99.9%), hydrochloric acid (analytical grade), and potassium hydroxide (analytical grade) were obtained from Merck-Schuchardt (Hohenbrunn, Germany). SPE cartridges, type OASIS VAC HLB 6 cc with 200 µg solid phase and 30 µm particle size, were from Waters (Eschborn, Germany). Glass fiber filters, type 9, with 90 mm diameter and syringe filters based on cellulose acetate with a pore size of 0.45 µm were both purchased from TH Geyer (Berlin, Germany).

**LC-UV**

LC-UV analysis was carried out on an HP-1100 system (Agilent Technologies, Santa Clara, California, USA), equipped with a binary pump (G1312A), autosampler (G1313A), thermostatted column compartment and diode array detector (1315A). The system was equipped with an *acquicore reversed phase column* (100 mm × 2.1 mm; particle size 2.6 µm; Thermo Fisher Scientific, Waltham, Massachusetts, USA). The column oven was set to 35 °C. The injection volume was 50 µl. The eluent flow was set to 200 µL min⁻¹. Mobile phases were ultrapure water with 1% acetic acid (Solvent A) and 1/1 v/v methanol/acetonitrile (Solvent B). The starting gradient was 95% A at 0 min, 80% A at 7 min, 80% A at 15 min, 95% A at 20 min and 95% A at 30 min. The LC system was operated with a UV-detector at a wavelength of 240 nm. At this wavelength, the UV/Vis spectra of terephthalic acid own a broad absorption peak [34]. Chemstation (A.09.03 version, Agilent Technologies, Santa Clara, California, USA) was used for data acquisition, processing, and peak integration.

**TED-GC/MS**

To check whether the method provides reliable and comparable results, environmental samples were additionally measured with a thermal analytical method. Thermal extraction-desorption-gas chromatography–mass spectrometry (TED-GC/MS) was used as a reference method. Measurements were conducted as described by Eisenbrandt et al. [28, 29]. In routine analysis, sample masses of 20 mg (composts, fermentation residue) were analyzed and variable masses between 6 and 15 mg were analyzed for samples from water and air (samples from WWTP, dust). In TED-GC/MS, ethyl benzoate was used as a specific marker for PET (m/z value of 105).

**Results and discussion**

**Optimization of the extraction process for terephthalic acid**

The sample extraction was performed in principle on the procedure described by Wang et al. [32]. However, according to the different detection systems and different samples, several steps were simplified and adapted to higher sample masses. In general, 20 g of sample was used, but sample mass varied for other matrices depending on expected PET content and available sample mass. 20 ml of 1-butanol was added to the solid sample in a 100 ml round flask. 1 g of potassium hydroxide pellets were added. The mixture was heated to 115 °C in an oil bath under constant stirring for 1 h. A tubular cooler was used for reflux. 50 ml of ultra-pure water was added, and the 2-phase system was mixed for 1 h on a stirring plate at 300 rpm. The extract was then vacuum filtrated with a glass fiber filter to remove the sediment/solids and collected. The buoyant organic phase was removed with a pipet. 10 ml of the aqueous phase was transferred to a 100 ml flask using a graduated pipette. The solution was diluted 1:10 with ultra-pure water, and the pH was adjusted to 2.5 by adding hydrochloric acid (conc. = 10%).

Before the LC-UV measurements, the aqueous extracts were cleaned up with a solid-phase extraction to avoid interferences of the organic matrix during analysis. A constant vacuum of 0.6 bar was applied using a vacuum chamber connected to a membrane pump. Cartridges were subsequently equilibrated and conditioned with 10 ml methanol and 10 ml ultra-pure water, respectively. Afterward, 10 ml of the aqueous extracts were passed over the cartridge. 5 ml of a 90:10 ultra-pure water/methanol mixture was used for washing the cartridges, before elution of the analyte with 10 ml methanol. The eluate was collected in 20 ml screw cap vials, and the solvent got evaporated. Precipitated terephthalic acid was subsequently reconstituted in 10 ml ultra-pure water, of which 1 ml was filtrated with a 0.45 µm syringe filter and transferred to LC-vials with PTFE-lined screw caps. The sample was acidified to a pH of 1–1.5 by adding 25 µl HCl (conc. = 1%). Acidification ensures complete protonation of the terephthalic acid. Pre-experiments showed a
sharpened peak form in the chromatogram and increased analytical sensitivity after acidification.

**Determination of the quality parameters**

A 6-level calibration with concentrations from 0 to 269 mg PET kg\(^{-1}\) in the reference soil was prepared, following the described sample preparation. Therefore, reference soil samples of 20 g were spiked with PET particles, which originated from cryo-milled PET pellets. All concentration levels were prepared in duplicates. All values were based on triplicate measurements. By using reference measurements, the peak in the chromatogram with a retention time of 7.7 min at a wavelength of 240 nm was identified as terephthalic acid (Fig. 1).

Figure 2 shows the linear relationship between the initially weighed PET and the signal for terephthalic acid. The coefficient of determination was found to be \(R^2 > 0.998\). Linearity was given up to 300 mg PET kg\(^{-1}\) reference soil and defined the working range. This maximum mass content of 300 mg PET kg\(^{-1}\) corresponded to a concentration of 12 mg l\(^{-1}\) terephthalic acid in the analyzed aqueous phase. The maximum solubility of terephthalic acid in water was 15 mg l\(^{-1}\) at pH 7 and 25 °C [35]. The measurement uncertainty based on a triplicate measurement was < 0.5%. According to DIN 32645, with a significance level of 0.9, the calculated limit of determination (LOD) and limit of quantification (LOQ) were 0.031 mg and 0.121 mg absolute PET, respectively. Based on the conducted calibration with 20 g reference soil matrix, this corresponds to a LOD of 1.55 mg kg\(^{-1}\) and LOQ of 6.05 mg kg\(^{-1}\). The determined LOQ and LOD were specific for the used reference soil matrix and can be varying for other matrices. The calculated LOQ of 6.05 mg kg\(^{-1}\) PET of our developed method was significantly higher, compared to Wang et al., they calculated a LOQ of 53.0 µg kg\(^{-1}\) PET. This LOQ was two magnitudes lower than our LOQ, mainly due to the use of different detectors. However, it is noteworthy that the calculated LOQ in our study was not based on a calibration using water and purified terephthalic acid, but on a calibration based on the initially spiked PET in a reference soil matrix.

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**Fig. 1** Exemplary chosen chromatograms (left) and UV-spectra (right) of terephthalic acid (PTA) standard solution and extracts of PET, PET-spiked reference soil, and PET-spiked sand with peat

**Fig. 2** Calibration slope for PET-spiked reference soil samples measured by LC-UV. PET mass refers to the initial spiked PET mass. All measurements were performed as triplicates
To achieve a high level of quality control, control experiments were conducted before each series of measurements to test the measurement quality of the LC-UV system. These were measurements of a terephthalic acid solution as well as repeated measurements of calibration standards of reference soil with different PET contents. Additionally, blanks of purified water were measured after environmental samples and showed no carry-overs of terephthalic acid or organic matrix compounds.

Table 2 summarizes the results of different model matrices containing different spiked PET types. We selected two different mass contents for all sample sorts, which covered the linear calibration area of the method as well as a compromise of microplastic contents in existing works [12, 16, 17]. Recoveries were based on reference soil calibration, as shown in Fig. 2. Sample masses of 20 g were analyzed in all cases.

Calculated recoveries for the sand and peat matrix were 94.5 and 107.1% and proofed the robustness of the method and the applicability for different environmental matrices. Calculated recoveries for different PET types, were 97.3 and 94.8% for bottle fragments, and 101.9 and 98.5% for fabric fibers. Generally, lower recoveries for the samples spiked with higher PET masses was explained by the limited solubility of terephthalic acid in water. Interference of the solid organic matter of the matrix on the detection of the terephthalic acid was not evident, as recoveries in the sand–peat matrix were not affected by the higher solid organic matter content compared to the reference soil. Hence, the method applied to different PET types and different soil matrices.

### Analysis of environmental samples

In the next step, real environmental samples were analyzed. The values of PET mass contents were based on extraction as described above and subsequent duplicate or triplicate measurement. Table 3 shows the detected mass contents and the extracted sample mass, which differed from 150 to 300 mg for dust samples up to 10,000–50,000 mg for sewage sludge, agriculture soil, and sediment samples. Figure 3 presents exemplary chromatograms and UV-spectra of selected samples.

PET was found in all investigated environmental samples, except for the samples of beach sediment.

### Table 2. Recovery rates of different PET types in model samples, with two different spiking levels

| Model sample                        | Spiked mass contents/mg kg⁻¹ | Recovery rate%/ |  |
|-------------------------------------|-----------------------------|----------------|---|
| Reference soil + bottle fragments   | 213                         | 97.3           |   |
|                                     | 431                         | 94.8           |   |
| Reference soil + fabric fibers      | 139                         | 101.9          |   |
|                                     | 191                         | 98.5           |   |
| Sand + 5% peat + PET particles      | 210                         | 107.1          |   |
|                                     | 475                         | 94.5           |   |

Measurements were performed as duplicate

### Table 3. Detected PET mass contents in environmental samples using PET extraction and TED-GC/MS

| Environmental sample             | PET extraction | TED-GC/MS       |
|---------------------------------|----------------|----------------|
|                                 | Sample weight/mg | Mass content/mg kg⁻¹ | Sample weight/mg | Mass content/ mg kg⁻¹ |
| Beach sediment, 1. extraction   | 50,000          | Below LOD       | –              | –                     |
| Beach sediment, 2. extraction   | 50,000          | Below LOD       | –              | –                     |
| Agriculture soil, 1. extraction | 20,000          | Below LOQ       | –              | –                     |
| Agriculture soil, 2. extraction | 20,000          | Below LOQ       | –              | –                     |
| Compost A                       | 5000            | 81.5            | 21.3           | Below LOD             |
| Compost B                       | 5000            | Below LOQ       | 23.9           | Below LOD             |
| Compost C                       | 5000            | 102             | 22.1           | Below LOD             |
| Fermentation residue A          | 3000            | 777             | 20.4           | Below LOD             |
| Fermentation residue B          | 3000            | 210             | 20.7           | Below LOD             |
| Fermentation residue C          | 3000            | 475             | 20.5           | Below LOD             |
| Particles of WWTP effluent      | 5000            | 38.6            | 14.9           | Below LOD             |
| Filter residue of greywater     | 500             | 1430            | 9.87           | Below LOQ             |
| Sewage sludge, 1. extraction    | 10,000          | 98.7            | –              | –                     |
| Sewage sludge, 2. extraction    | 10,000          | 142             | –              | –                     |
| Dust sample A                   | 276             | 12,500          | 6.11           | 7100                  |
| Dust sample B                   | 149             | 43,600          | 8.23           | 57,000                |
Lowest PET mass contents of 3.85 and 3.99 mg kg\(^{-1}\) were detected in the agriculture soil samples, however, this is below LOQ.

PET mass contents of compost samples were between 10.2 (below LOQ) and 102 mg kg\(^{-1}\). Compared to the compost, fermentation residues showed all over higher mass contents with values up to 776 mg kg\(^{-1}\) in the sample from the aerobization container. Mass contents in the liquid and solid fermentation residues were 209 and 475 mg kg\(^{-1}\), respectively. A correlation between the organic content and the observed PET findings could not be found for individual samples.

In the urban water management samples, an even higher PET content of 1430 mg kg\(^{-1}\) was found in the filter residue of greywater. The investigated sewage sludge showed mass contents of 98.7 and 141 mg kg\(^{-1}\), while only 38.6 mg kg\(^{-1}\) PET was found in the suspended particles of the WWTP effluent.

The highest PET mass contents of 12,500 and 57,000 mg kg\(^{-1}\) were found in the indoor dust samples. The determined polymer type was congruent with the findings of Lui [36] and Vianello [37] and suggests a future application in the field of dust analysis.

Some of the most promising samples were analyzed in parallel using TED-GC/MS. It should be noticed, that all experiments run with lower samples masses. The results were also summarized in Table 3. The findings of biowaste samples were all below LOD. Additional investigation using TED-GC/MS of samples with high PET findings from urban waste management did also not result is significant signals. However, PET was clearly found in the dust samples with amounts of 7100 mg kg\(^{-1}\) and 57,000 mg kg\(^{-1}\). The results of LC-UV and TED-GC/MS corroborated each other.

Using TED-GC/MS the LOD for PET was 0.88 µg absolute (fivefold signal-to-noise ratio). In environmental samples the variability of this value is influenced by inherent matrix components and LOD values of up to 1.37 µg were detected (Kittner et al. 2020, unpublished data). The determination of LOQ for TED-GC/MS according to DIN 32645 was not possible, because the standard deviation was found to change with relative PET content in the environmental matrix.

If the values of the PET extraction were extrapolated to the masses used for the TED-GC/MS measurement (clearly less, namely 20.4 mg and 9.87 mg) a signal can only be expected for the fermentation residues and the greywater sample. Whereas in TED-GC/MS measurement the result of greywater indicates a minor content of PET, no evidence of any PET presence was found in the fermentation residue samples.

Different reasons might be responsible for missing PET signal in the TED-GC/MS measurements of the fermentation residue samples. First, the organic environmental matrix influenced the formation of PET marker for LC-UV measurement stronger than expected from previous measurements. However, such a correlation to the background value related to the organic content in those samples (see Table 1) could not be determined. Second, the PET extraction procedure did not discriminate different kinds of polyesters, such as polyethylene terephthalate, polybutylene terephthalate or polybutylene adipate
teredephthalate. The latter is a biodegradable polymer that might be present in larger quantities in fertilizer samples from urban waste and could, therefore, be responsible for the high terephthalic acid findings in LC-UV. In contrast to that, TED-GC/MS detected ethyl benzoate as a specific marker, which could only be formed by the decomposition of PET. However, such a reason was not plausible for the greywater sample. In fact, there could also be a third, and this could be the main reason for explaining these differences. Terrestrial samples own a high heterogeneity, and 20 mg of sample in TED-GC/MS measurement might be too low for a representative measurement. For the chemical extraction method, the used extracted samples masses were always higher and will, therefore, catch a higher number of representative particles.

Even if there were still uncertainties about background signals, contributions of possible false-positive signals from other polymers, or the representatives of the analyzed samples, it became clear that a relative classification of the signal intensities of both methods was possible.

Conclusions
The goal of this study was to adapt and further simplify the chemical extraction of PET, proposed by Wang et al. [32]. It was optimized for higher soil sample masses and transferred to an LC-UV system. LC-UV is commonly available in routine labs, no poisonous chemicals were needed when larger sample volumes must be analyzed (regarding the representativeness of the sample), it was easy to automatize, a measurement took only 30 min and evaluation of the data did not need any high user knowledge.

Within this study, we showed the application of this new LC-UV analysis, including optimized sample preparation for the determination of PET in environmental samples. The recovery of spiked samples was between 95 and 107%, using different PET sorts and soil types. The sample size was up to 50 g and can be adjusted easily to even higher masses. This is of high interest for routine analysis of soils in future, because this would avoid additional sample preparation steps for analysis (e.g., density separation). A LOD and LOQ was determined of 0.031 mg and 0.121 mg absolute PET, respectively.

Using LC-UV, PET signals was found in all environmental samples: terrestrial samples, bio-waste residues, urban wastewater, and indoor dust. Mass contents varied strongly dependent on the sample between values below LOQ in agricultural soil and 57,000 mg kg\(^{-1}\) in indoor dust. Values for sewage sludge, compost, and fermentation residue were up to three magnitudes higher than in agricultural soil. Therefore, fertilization with the latter could be a possible source for microplastic in agroecosystems.

The proposed extraction can be used to quantify the mass input of PET into agroecosystems, thus providing a better understanding of the microplastic input by fertilization with sewage sludge and bio-waste. Advantageous of the method is the applicability for complex matrices. The method is a cost-effective and simple integral method and applicable for diverse environmental matrices.

The present method is a useful addition in the field of microplastic analysis and could be used for the analysis of PET in samples, but also where high findings expected. This could be PET particles in bottled water or samples with a high load of especially PET fibers, e.g., indoor dust samples or washing machine effluent. However, it should be noticed, that for environmental samples in general (e.g., like surface water, air samples) with more complex mixtures of different MP particles other method are more suitable.

Finally, this work demonstrates that also simple methods can be applied for specific questions of microplastic analytics. The use of such alternatives depends on the scientific issue of the analysis, i.e., the aim of the investigation. The present method, but also TED-GC/MS determine mass contents and are not comparable with methods for determining particle numbers, which can provide an even more in-depth description of microplastic (including particle shape and size). However, the described methods aim to be faster applicable (including sample preparation and detection) and less complicated in use. This is the only way to perform comprehensive measurements, e.g., for the screening of hot-spot entries.

However, in the end it will always be the individual goal of an investigation campaign, which investigation result should be achieved with which time horizon and analytical depth.

Abbreviations
FTIR: Fourier-transform Infrared spectroscopy; FT-NIR: Fourier-transform near-infrared spectroscopy; LC–MS/MS: Liquid chromatography–tandem mass spectrometry; LC-UV: Liquid chromatography with UV-detection; LOD: Limit of detection; LOQ: Limit of quantification; PET: Polyethylene terephthalate; PTFE: Polytetrafluoroethylene; Py-GC–MS: Pyrolysis-gas chromatography–mass spectrometry; rpm: Revolutions per minute; TED-GC/MS: Thermal extraction-desorption-gas chromatography–mass spectrometry.

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Authors’ contributions
AM, CG, and UB designed the experimental set-up. Sample extractions were carried out by AM. AM, CG, and CP were responsible for development of the LC method. AM performed LC measurements and data acquisition and processing. PE conducted TED-GC/MS measurements. AM and UB drafted the manuscript. All authors edited and reviewed the manuscript. All authors read and approved the final manuscript.

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Competing interests
The authors declare that they have no competing interests.

Author details
1 Bundesanstalt für Materialforschung und -prüfung (BAM), Unter den Eichen 87, 12205 Berlin, Germany. 2 RWTH Aachen University, Worringer Weg 1, 52062 Aachen, Germany.

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