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Article

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Prediction of the formation of biogenic non-extractable residues during degradation of environmental chemicals from biomass yields

Stefan Trapp^a, Andreas Libonati Brock^a, Karolina Nowak^b, Matthias Kästner^{b,*}

^a Department of Environmental Engineering, Technical University of Denmark, Bygningstorvet
bd. 115, DK-2800 Kgs. Lyngby, Denmark.

^b Helmholtz-Centre for Environmental Research – UFZ, Department of Environmental
Biotechnology, Permoserstr. 15, 04318 Leipzig, Germany.

** Corresponding author: Matthias Kästner; phone: 0049-341-235-1235; fax: 0049-341-235-451235; e-mail: matthias.kaestner@ufz.de*

Abstract

Degradation tests with radio or stable isotope labeled compounds enable the detection of the formation of non-extractable residues (NER). In PBT and vPvB assessment, remobilisable NER are considered as a potential risk while biogenic NER from incorporation of labeled carbon into microbial biomass are treated as degradation products. Relationships between yield, released CO₂ (as indicator of microbial activity and mineralization) and microbial growth can be used to estimate the formation of biogenic NER. We provide a new approach for calculation of potential substrate transformation to microbial biomass (theoretical yield) based on Gibbs free energy and microbially available electrons. We compare estimated theoretical yields of biotechnological substrates and of chemicals of environmental concern with experimentally determined yields for validation of the presented approach. A five-compartment dynamic model is applied to simulate experiments of ¹³C-labeled 2,4-D and ibuprofen turnover. The results show that bioNER increases with time, and that most bioNER originates from microbial proteins. Simulations with pre-calculated input data demonstrate that pre-calculation of yields reduces the number of fit parameters considerably, increases confidence in fitted kinetic data and reduces the uncertainty of the simulation results.

Key words: bound residues, modeling, Gibbs free energy, pesticides, carbon conversion, carbon turnover, microbial biomass, Nernst, NER assessment, OECD tests.

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Introduction

Degradation is a key parameter in risk assessment and registration of industrial chemicals, veterinary medicinal products and pesticides.¹⁻⁵ Microbial degradability tests are often performed with radio-labeled tracer compounds. Guidelines have been developed for fate assessment in water, sediments and soil, e.g., OECD 304, 307, 308 and 309.⁶⁻⁹ For the interpretation of results, concepts for modeling the turnover kinetics have been developed.^{10,11} Unfortunately, there is still no robust and reliable way to predict the fate of organic molecules in environmental matrices in terms of biotic transformation, mineralization, conversion to microbial biomass and the formation of so-called non-extractable residues (NER).¹²

Chemicals may persist in the environment due to several reasons. Relatively well studied is the persistence due to limited bioavailability. Chemicals being strongly adsorbed or sequestered in soil and sediments are often not available for biodegradation.¹²⁻¹⁵ Examples are the five- or six-ring polycyclic aromatic hydrocarbons. Chemicals newly introduced to the biosphere may persist due to the absence of enzymes capable of transforming such compounds. However, after some time for adaptation microbes can “learn” to degrade recalcitrant compounds.¹⁶ A third reason for persistence is that chemicals are poor growth substrates because they do not provide energy, carbon or nutrients to microbes under the specific environmental conditions. For example, alkanes have persisted over millions of years in reservoirs where no suitable electron acceptor (oxygen) was available. Under aerobic conditions, alkanes are excellent substrates with higher microbial biomass yields than glucose.^{17,18} For chlorinated solvents, e.g. trichloroethylene (TCE), the opposite was observed: the chemical provides no energy to microbes under aerobic conditions and is therefore quite persistent, while it can be reductively dehalogenated as electron acceptor in anaerobic groundwater.¹⁹ Another reason for persistence of chemicals can be toxic or inhibitory effects on the microorganisms.^{20,21}

In the PBT assessment of industrial chemicals and of veterinary pharmaceuticals, NER are differentiated into remobilisable and irreversibly bound fractions. The irreversibly bound fraction is assessed as a potential removal pathway, while the remobilisable fraction is considered a potential risk for the environment.⁵ Remobilisable NER are sequestered compounds (type I NER) and covalently bound parent compounds or metabolites (NER type II), which may be slowly released. The third fraction is labeled carbon, or other essential elements like nitrogen, transferred to living or dead biomass and eventually

fixed in soil organic matter (SOM) derived from decaying microbial biomass (type III NER). These biogenic NER (bioNER) do not constitute any risk.¹² There is thus a need to distinguish harmless, irreversibly bound bioNER from potentially toxic and remobilisable NER (i.e., type I and type II NER) in the risk assessment of chemicals.

Determining the microbial biomass yield derived from degradation of a chemical sheds light into the 'black box' of NER. The microbial yield is defined as mass of microbes that can grow on a given amount of substrate (unit g microbial biomass dry weight per g substrate, g g⁻¹).¹⁸ However, most studies with labeled carbon compounds typically express results as g C per g C, and we report these values with their original unit. The unit conversion is shown in the SI. The yield multiplied by the enzymatic substrate removal determines the growth rate of a microbe. High yield can therefore be an indication for the biodegradability of a substrate. Several methods to estimate theoretical microbial yields of a substrate from its energy of formation (Gibbs free energy) have been developed.²²⁻²⁶

The yield can be used to predict the likely range of bioNER formed during degradation of environmental chemicals. We i) provide a relationship between formation of bioNER, CO₂ release and yield; ii) present a new and pathway-independent method to estimate yields from thermodynamics combined with an approach to account for the electrons usable by degrading microbes; iii) confirm the yield estimates by comparison to results derived with existing methods²³ and to measured yields of easily degradable carbohydrates, pesticides and other chemicals of environmental concern. iv) Finally, we use the estimated yields as input to the simulation of 2,4-D and ibuprofen biodegradation under formation of microbial biomass, study the performance of the simulation, and compare pre-calculated kinetic parameters with data derived by pure model fit. Data were provided from experimental degradation studies with multi-labeled compounds (¹⁴C, ¹³C) in soil.²⁷⁻²⁹

Methods

Theoretical background

Enzymatic reactions are typically described by the *Michaelis-Menten* equation:^{30,31}

$$\frac{dm}{dt} = v_{\max} \times \frac{a}{K_M + a} \times X \quad (\text{eq. 1})$$

where m is the mass of chemical substrate metabolized (g), X is the bacterial mass (g bacteria), t is time (d), $v_{\max}$ (g substrate g bacteria⁻¹ d⁻¹) is the maximal substrate consumption rate, a is the chemical activity of the substrate (equivalent to the truly dissolved concentration) (g m⁻³);³² K_M (g m⁻³) is the chemical activity at which the substrate consumption rate is half of its maximum (half saturation or *Michaelis-Menten* constant).

The yield Y (g g⁻¹) connects metabolism and growth:

$$\mu_{\max} = v_{\max} \times Y \quad (\text{eq. 2})$$

where $\mu_{\max}$ is the maximum growth rate (d⁻¹). Microbes use part of the energy gained from the substrate for growth, and part for maintenance purposes. Experimentally observed net yields equal the true yield minus cell decay. Introducing a term for cell decay or maintenance into the *Monod* equation for microbial growth leads to eq. 3.³³

$$\frac{dX}{dt} = \frac{\mu_{\max} a}{K_M + a} \times X - b \times X \quad (\text{eq. 3})$$

where dX/dt is the change of microbial biomass with time (g microbial biomass dw per day), and b is the decay rate of microbes (death rate, d⁻¹). For the calculation of the fate of chemicals in soils or sediments, a two-compartment-sorption model^{34,35} calculating rapid (adsorption) and slow (sequestration) kinetics was combined with the equations for microbial metabolism and growth (eqs. 1 and 3).^{12,36} The complete model is described in the Supporting Information (SI).

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108 *Carbon budget and calculation of bioNER*

109 Few experimental studies deliver compound concentrations and biomass formation in a resolution that
110 allows fitting of dynamic models to the data. In degradation studies according to OECD guidelines, only
111 the fractions of NER, CO₂ and metabolites at the end of the experiment are reported.³⁷ Some general

rules and patterns can be derived concerning the distribution of the initially applied labeled carbon and the formation of biomass (here all units are g C).

We define S as the total mineralized substrate, S is initial amount of labeled carbon minus non-metabolized parent compound minus intermediate metabolites and minus $NER^{I,II}$. $NER^{I,II}$ denotes non-extractable residues due to sequestration (I) and co-valent binding (II).¹² The biomass produced from mineralization of the substrate is per definition the yield, hence, as long as growth alone is considered, $X = Y S$. The remaining labeled carbon is oxidized to carbon dioxide, thus $CO_2 = (1-Y) S$. Under these assumptions, the ratio of X to CO_2 is

$$\frac{[X]}{[CO_2]} = \frac{Y}{(1-Y)} \quad (\text{eq. 4})$$

The labeled carbon fixed in biomass due to substrate mineralization is part of the bioNER. Eq. 4 does not take decay of biomass into consideration. Earlier long-term studies³⁸⁻⁴⁰ over 224 days showed that microbial necromass is a significant source of non-living soil organic matter. In these experiments, approximately 40% of the labeled carbon initially fixed in biomass X (mainly the protein fraction) turned into SOM (which also is part of bioNER) and 10% remained within living biomass X . It follows that for $t \rightarrow \infty$ the fraction f (approximately 0.5) of the decaying X turns into bioNER, and $1-f$ forms CO_2 .⁴⁰ The resulting ratio of bioNER to CO_2 in long-term experiments with decomposition of dead biomass (and neglecting slow decomposition of SOM to CO_2) is

$$\frac{[bioNER]}{[CO_2]} = \frac{f \times Y}{(1-Y) + (1-f) \times Y} \quad (\text{eq. 5})$$

Yield estimates

The microbial yield of a substrate can be estimated from thermodynamics or from empirical equations. Approaches for yield estimation have been presented and tested by a number of researchers.^{22-24, 26,41} The approaches of McCarty²³ and Xiao and vanBriesen²⁶ require information on the metabolic pathway of the compound, which is often not given for environmental chemicals' degradation. Therefore, we based our estimates of yields on a modified approach of Diekert²², which uses the Gibbs free energy of formation

and the structural formula. We modified the method by specifying how much of the formation energy can be used by microbes.

Microbial Turnover to Biomass (MTB) - a pathway-independent thermodynamic yield estimation method

The approach considers that a substrate can be utilized for anabolism and catabolism:

$$N = N_{\text{anabolic}} + N_{\text{catabolic}} = 1/Y \quad (\text{eq. 6})$$

where N is the nutritional value (g substrate needed per g microorganism formed), the inverse of yield Y :

$$\frac{1}{Y} = \frac{1}{Y_{\text{anabolic}}} + \frac{1}{Y_{\text{catabolic}}} \quad (\text{eq. 7})$$

Anabolism

The yield associated to anabolism is

$$Y_{\text{anabolic}} = \frac{n_c \times M_c}{f_c \times M_s} \quad [\text{g microbial biomass dw g}^{-1} \text{ substrate}] \quad (\text{eq. 8})$$

where n_c is mol C per mol substrate, M_c and M_s are the molar masses of carbon (index C) and of substrate (index S), respectively, and f_c is fraction of C in bacterial dry weight (default 0.53 g carbon g⁻¹ microbial biomass dw).²³

Catabolism

The yield due to catabolic energy gain can be calculated in five steps.

Step i) Free energy of the reaction: The free energy of the reaction (change of Gibbs free energy, kJ mol⁻¹) is the sum of the Gibbs free energy of formation of products minus educts:

$$\Delta G_r^{0'} = \sum \Delta G_f^{0'} \text{ products} - \sum \Delta G_f^{0'} \text{ educts} \quad (\text{eq. 9})$$

where $G^{0'}$ is the Gibbs free energy (subscript f for formation, r for reaction) at standard-state conditions (1 mol L⁻¹, indicated by superscript 0) and at a pH of 7 (indicated by superscript '). At activities differing from 1 mol L⁻¹, the change of Gibbs free energy of the reaction $\Delta G_r'$ is

$$\Delta G_r' = \Delta G_r^{0'} + RT \ln \left(\frac{[products]}{[educts]} \right) \quad (\text{eq. 10})$$

where R (8.314 J mol⁻¹ K⁻¹) is the universal gas constant and T (K) is the absolute temperature.

Step ii) Electron transfer during the reaction (*Nernst* equation): The *Nernst* equation states that the change of Gibbs free energy ΔG_r is related to the number of electrons n transferred during the reaction, and the redox potential E (V) of the reaction:

$$\Delta G_r = n \times F \times E \quad (\text{eq. 11})$$

where F is the Faraday constant. The number of electrons n transferred in the reaction can be calculated from the change of the oxidation state (OS) of carbon during the reaction,

$$n = (\text{OS}_{\text{Product}} - \text{OS}_{\text{Substrate}}) \times n_C \quad (\text{eq. 12})$$

where n_C is again the carbon atoms in the substrate (mol C per mol substrate) which is the same as the moles of CO₂-molecules formed during complete mineralization. The oxidation state of carbon in the substrate is:

$$\text{OS}_{\text{Substrate}} = \frac{-1 \times H + 2 \times O + 3 \times N - 3 \times P + 2 \times S + 1 \times Cl}{n_C} \quad (\text{eq. 13})$$

where the letters stand for the number of the respective atoms in the molecule. After complete oxidation to CO₂, the carbon in the product has the oxidation state 4, hence OS_{Product} = +4.

Step iii) Energy available for the microbe: During biological oxidation, the organisms can use only some types of electron transfers. The free energy of the reaction is thus the upper limit ("maximum") for the energy that can be provided by the chemical. The maximum energy gained by the organism during catabolism may be considerably lower than that. As a general rule, when compounds containing hydrogen atoms connected to carbon atoms are oxidized to CO₂ and H₂O, the electrons transferred in this reaction

are available for microbes to gain energy, i.e. 2 electrons per C–H bond. Thus the number of electron transfers that can at least be used by microorganisms in a redox reaction is $n_{bio} \geq 2 \times H$ (only H bound to C atoms are counted). Subsequently, the minimum energy available for ATP synthesis by an organism ($\Delta G'_{bio}$) is:

$$\Delta G'_{bio} = \frac{n_{bio}}{n} \Delta G'_r \quad (\text{eq. 14})$$

Step iv) ATP production: With an efficiency of 40% of the microbial catabolism,²² the synthesis of 1 mol ATP from 1 mol ADP requires 80 kJ (the Gibbs free energy of the reaction is -32 kJ mol^{-1}).⁴¹ Thus, the microbes can generate β mol ATP per mol substrate:

$$\beta = \frac{\Delta G'_{bio}}{-80 \text{ kJ/mol}} \quad (\text{eq. 15})$$

Step v) Catabolic energy is used for the formation of new cell material: The produced ATP provides the energy to form new cell material.⁴¹ Y_{ATP} is the microbial biomass dw that can be formed per mol ATP.⁴¹ Diekert²² provided a range from 2 (CO_2) to 12 (glucose) g microbial biomass dw per mol ATP. Hence, we use the value of 5 g microbial biomass dw mol^{-1} ATP as default for xenobiotic chemicals, but higher values for compounds similar to glucose (for details, see SI). The yield due to catabolic energy gain can finally be calculated by

$$Y_{catabolic} = \frac{\beta \times Y_{ATP}}{M_s} \quad [\text{g microbial biomass dw g}^{-1} \text{ substrate}] \quad (\text{eq. 16})$$

The five steps can be summarized in one equation:

$$Y_{catabolic} = \frac{\beta \times Y_{ATP}}{M_s} = \frac{n_{bio}}{n} \times \frac{\Delta G'_r}{-80 \text{ kJ/mol}} \times \frac{Y_{ATP}}{M_s} \quad (\text{eq. 17})$$

The more detailed approaches of McCarty²³ and Xiao and vanBriesen²⁶ estimate β dependent on the biochemical pathway. Knowledge of the pathway is not required in the method presented because all substrate used for catabolic yield is completely oxidized to CO₂.

Choice of compounds for yield estimation

The estimation of yields is commonly applied in biotechnology or wastewater treatment. In environmental chemistry, it has been used by Helbling *et al.*⁴² to estimate the yields of two pesticides and by Yuan and vanBriesen⁴³ to estimate the yield of two chelating agents. First, we investigated the performance and the variance of results of the estimation methods with common substrates in biotechnological applications, for which measured yields are widely available. Second, we applied the method to a set of chemicals of environmental concern. The selection of chemicals for this study was based on: i) availability of measured data on bioNER (2,4-D, ibuprofen), ii) availability of biomass yield data (nitrilotriacetic acid [NTA], linuron, carbofuran, toluene), iii) knowledge on specific degradation pathways, electron acceptors or persistence (pentachlorophenol PCP, carbon tetrachloride, trichloroethene, DDT), and iv) availability of Gibbs free energies of formation (Table S2).

Brief description of experiments

Nowak *et al.*^{27,28} thoroughly balanced the formation of bioNER in a fate study with the ¹³C-labeled pesticide 2,4-D and the medical drug ibuprofen in soil. The authors also analysed the amount of ¹³C converted to total amino acids (tAA), total fatty and phospholipid fatty acids (PLFA). The tAA increased over time although the PLFA as marker for living biomass declined already after one (2,4-D) to three (ibuprofen) weeks. The details of the turnover experiments are provided in Nowak *et al.*^{27,28}; the results of the experiments are shown in Tables S9 and S10.

Description of simulations

2,4-D and ibuprofen experiments were simulated to confirm the relation between yield and bioNER formation. For a detailed description of model and input data see SI S2. The model is composed of five compartments describing the five possible states of labeled carbon: dissolved (D), adsorbed (A), sequestered (S) state, or (following metabolism) carbon dioxide (CO₂) and living and dead biomass (X and X_{dead}). The model was implemented as a set of ordinary differential equations (ODEs) in MATLAB. The model was also successfully implemented in Microsoft Excel and produced equal results. The calculated sum of living and dead biomass was considered to be bioNER, and the sum of sequestered fraction and bioNER was compared to measured total NER. No kinetic data were available to separately simulate the formation of type II NER. Hence, any type II NER formed in the experiments were considered to be included in the sequestered compartment of the model. The calculated labeled carbon in the dissolved and adsorbed compartment was compared to the measured extractable labeled carbon.

Calculation and fitting of input parameters

Input parameters for the simulations were derived as follows. The initial amount of ¹³C was assumed to be distributed between the dissolved and solid phase according to the soil-water distribution coefficient (K_d). The sequestered fraction was assumed to be equal to the NER measured at $t = 0$ and corrected by the reported recovery. $NER(t = 0)$ was subtracted from the calculated ¹³C present in the solid phase to yield the adsorbed fraction.

We adjusted the input parameters step-wise, similar to Rein *et al.*³⁶:

Step i) Yield: Calculation with the MTB method.

Step ii) Death rate b : Towards the end of the experiments, the substrate is used up and the microbes decline. Then

$$\ln \left(\frac{X(t)}{X(0)} \right)_{final\ phase} \approx -b t \quad (\text{eq. 18})$$

where X here is the measured concentration of microbial biomass; it is calculated from the measured PLFA times a factor of 20 (5% content of PLFA in native biomass).

Step iii) Growth: During the initial growth phase the microbial growth can be described as

$$\ln \left(\frac{X(t)}{X(0)} \right)_{initial\ phase} = (\mu - b)t \quad (\text{eq. 19})$$

The resulting growth rate μ at given time t is used to estimate μ_{max} (SI 3.2).

Step iv) Half-saturation constant: For 2,4-D, a literature value for the ratio μ_{max}/K_M is given in Tuxen *et al.*⁴⁴. For ibuprofen, K_M was fitted using the CO₂ development as criterion.

Step v) Initial degrader biomass: $X(0)$ was adjusted to fit the peak biomass concentration and the lag phase. The sum of root mean square errors (RMSE) was used to describe the “goodness-of-fit” (SI S3.3).

During the model calibration against the 2,4-D data we found that the sequestration (slow adsorption, leading to NER) of the labeled carbon of 2,4-D was better described by using the K_{OC} of 2,4-dichlorophenol (2,4-DCP) instead of the K_{OC} of the parent compound 2,4-D. 2,4-DCP is the transformation product of 2,4-D and has a K_{OC} much higher than 2,4-D. For the rapid adsorption (part of the extractable ¹³C), the K_{OC} of 2,4-D was kept. It is well known that chlorinated phenols tend to form abiotic NER,¹² thus the better fit of the 2,4-DCP K_{OC} may provide an indication for NER type II bonding via covalent bonds triggered by oxidative coupling. The most appropriate way to accommodate this change would be the inclusion of step-wise degradation (e.g., 2,4-D to 2,4-DCP to CO₂), but this increase in model complexity would not be justified by the available data.

Uncertainty analysis and parameter identification

Aside pre-calculation, two optimization routines were used for calibration of v_{max} , K_{OC} , $X(0)$, Y , and K_M (only for ibuprofen). The Pattern Search optimization function is an algorithm that finds local minima from a mesh around the initial values and stops when the optimization function cannot be further minimized.⁴⁵ The Bayesian optimization method DiffeRential Evolution Adaptive Metropolis algorithm (DREAM_(ZS))^{46,47} uses the Bayesian framework and also allows for the assessment of uncertainties related to the parameter estimates and the model predictions. The Bayesian optimization was done with and without Y as a pre-calculated parameter in order to assess the effect of its inclusion on the parameter estimation and on the uncertainty of the model predictions. For further details on the parameter settings see the SI (S4).

Results

Comparison of yield estimates

Table 1 and Table 2 list the observed and estimated yields of substrates relevant to biotechnology (unit g C g⁻¹ C) and of chemicals of environmental concern (unit g microbial biomass dw g⁻¹ substrate). The biotechnological substrates are easily degradable compounds for which experimental yield data are available.⁴⁸ Both the TEEM2²³ and the presented MTB yield estimation methods give relatively close estimates with a mean absolute error (MAE) of less than 0.1 g C g⁻¹ C. Few experimental yield data are available for chemicals of environmental concern. The estimates are less accurate, with the highest deviation for linuron, which had a very low measured yield.⁴² Despite its simplicity, the MTB method overall gave results with lower deviation compared to TEEM2 for the chemicals of environmental concern.

<Table 1>

<Table 2>

Dynamic model simulations

Figure 1 shows the experimental and the simulation results for ¹³CO₂, extractable ¹³C (dissolved and adsorbed) and non-extractable ¹³C (which is the sum of ¹³C-label sequestered and in living or dead biomass). For both compounds, the model with pre-calculated input data is able to reasonably describe the fate of ¹³C in the different compartments. However, CO₂ and NER are predicted to increase at an earlier time point than observed. For 2,4-D, this can be seen already at the first data points, whereas for ibuprofen it is evident after 28 days. Based on the *Michaelis-Menten* equation, it was assumed that the formation of CO₂ and new biomass occurs as soon as the labeled compound is transformed. In reality, internal storage of metabolites and HCO₃⁻ delays the release of CO₂. This may be overcome by the introduction of new parameters; only, this would considerably increase model complexity which was not desired. In the beginning of the simulation, most NER are sequestered, but towards the end of the simulation, the NER originate mainly from living and dead biomass.

The experimentally determined extractable ¹³C-ibuprofen was declining within four weeks (Fig. 1b). The extractable ¹³C-label had initially similar values but remained relatively high throughout the experiment. After 90 days, 13.4% of the ¹³C was detected in the solvent-extractable portion, but only 0.5% was ¹³C-

ibuprofen (Girardi *et al.*²⁹, Table S10). This indicates rapid formation of transformation products and incomplete mineralization with only little 2-hydroxy-ibuprofen (Table S10).

Figure 2 depicts the simulated living biomass (X), dead biomass (X_{dead}) and the sum of both. This is compared to the measured ^{13}C in PLFA multiplied with a factor 20 as a marker for living biomass (5% PLFA content), and to the measured ^{13}C -label in tAA. Living biomass contains about 50% proteins (amino acids), hence also tAA multiplied with a factor of two was plotted. It can be seen that PLFA/0.05 and tAA/0.5 as well as the simulated sum of X and X_{dead} are close until day 4 (2,4-D, Fig. 2a) or day 14 (ibuprofen, Fig. 2b), as long as the living biomass predominates. Later PLFA declines, which indicates a decline of living biomass X . The dotted line in Figure 2 is the decay halftime ($\ln 2 / b$) after maximum measured PLFA. The line indicates where >50% of microbes have died. From this point, the simulated sum of X and X_{dead} is much closer to tAA than to tAA/0.5. In decaying microbes, labile constituents like sugars and fatty acids are turned over first and the more stable amino acids (tAA) in proteins persist (also see SI S2.13).^{39,40} Thus, towards the end of the simulation, the sum of X and X_{dead} is dominated by proteins and should be compared to tAA and not to tAA/0.5.

Calculated bioNER versus measured tAA

The measured $^{13}\text{CO}_2$ release in the 2,4-D experiment was 57.6% of the initially applied ^{13}C (SI Table S9), and the calculated yield of 2,4-D was $0.28 \text{ g } ^{13}\text{C g}^{-1} ^{13}\text{C}$. Using these values in the equation for the ratio of biomass growth to CO_2 production (eq. 4) we calculated that 22% of the applied ^{13}C -label was fixed in the biomass. The measured tAA was at 23.3% (SI Table S9). For ibuprofen, with a measured $^{13}\text{CO}_2$ -release of 45.2% (Table S10) and a calculated yield of $0.43 \text{ g } ^{13}\text{C g}^{-1} ^{13}\text{C}$, the calculated ^{13}C -label in biomass was 34% (measured tAA: 28.4%). In the case of these two experiments, the measured ^{13}C -label within amino acids (tAA) was remarkably constant towards the end of the experimental period, and there was no need to consider turnover of dead biomass. The calculated ^{13}C -label fixed in bioNER with eq. 5 was 9.4% (2,4-D) and 12.4% (ibuprofen). Once the fraction of bioNER is known from Y and CO_2 , the potentially remobilisable NER type I and type II can be quantified from the total NER. In the PBT/vPvB assessment of

chemicals, the bioNER fraction can be subtracted from the total NER and counted as degraded. In a follow-up study, we used this method to estimate the bioNER for 40 chemicals of environmental concern.⁴⁹

<Figure 1>

<Figure 2>

Discussion

Yield estimates

We presented the new MTB approach for estimation of microbial biomass yields. Considering the variability of the experimental data, this approach showed fairly similar deviations from experimental yield data in comparison to the more advanced and widely applied TEEM2 approach²³, without the need for specific information about the catabolic pathway, primary oxidation processes or N sources. For environmentally relevant chemicals and pesticides the deviation of the experimental yields is even lower than estimated with the MTB. The MTB approach can be applied for many tasks, e.g. yield assessment in biological wastewater treatment or maximum transfer of labeled carbon into microbial biomass and bioNER assessment, as shown with the simulations. Yield estimates can thus contribute to an improved risk assessment of environmentally relevant chemicals. The method may be added as a module in biodegradation databases (like EAWAG-BBD/PPS <http://eawag-bbd.ethz.ch/>, KEGG <http://www.genome.jp/>) and QSAR approaches (like ChemProp www.ufz.de/ecocem/chemprop or EPI suite <https://www.epa.gov/tsca-screening-tools/epi-suitetm-estimation-program-interface>). Combined with the unified model for sorption and biodegradation (Kästner *et al.*¹², Rein *et al.*³⁶, and this study) the MTB yield estimation method can be used for modeling the entire turnover process of a chemical in the environment.

Comparison to other findings

Yield estimations are rarely applied to chemicals of environmental concern. One exception is the study of Helbling *et al.*⁴² with linuron and carbofuran. The estimated theoretical yields for carbofuran are 0.51 g g⁻¹ (MTB method) or 0.59 g g⁻¹ (TEEM2 method) and 0.41 g C g⁻¹ C = 0.50 g g⁻¹ with the related adapted TEEM1 method.⁴² The experimentally determined yield of carbofuran was 0.52 g g⁻¹ (0.42 g C g⁻¹ C). The experimental yields obtained for linuron in the Helbling *et al.*⁴² study were very low (0.06 g C g⁻¹ C = 0.05 g g⁻¹) despite a theoretical yield similar to carbofuran (0.40 g C g⁻¹ C = 0.33 g g⁻¹). Maximum growth rates μ_{max} were determined to be 7.8 d⁻¹ (carbofuran) and 1.3 d⁻¹ (linuron), corresponding to a v_{max} = 15.1 and 26.4 g (g d)⁻¹, respectively.

Kinetic parameters and yields of polycyclic aromatic hydrocarbons PAH have been determined in several studies. Wick *et al.*⁵⁰ grew *Mycobacterium* sp. LB501T on solid anthracene and obtained yields between 0.158 and 0.196 g g⁻¹ and v_{max} of 18.4 g (g d)⁻¹. Adam *et al.*⁵¹ found for the growth of three degrader strains on phenanthrene the same yield of 0.21 g g⁻¹, with v_{max} from 12 to 18 g (g d)⁻¹. Rein *et al.*³⁶ tested growth of *Mycobacterium* sp. on phenanthrene and pyrene and found yields from 0.20 to 0.32 g g⁻¹ and v_{max} from 8 to 10 g (g d)⁻¹. Toräng *et al.*⁵² estimated a yield of about 0.3 g C g⁻¹ C for the degradation of U-ring-labeled phenoxy-acetic acids (MCPD and 2,4-D) using the ¹⁴C-MPN (most probable number).⁵³

Most of the experimental yields (Table 2) are lower while v_{max} -values and growth rates are higher than those obtained here (Table 3), and there could be several reasons for this: In these studies, known pure degrader strains were tested under optimal nutrient conditions, which explains the faster growth and the lower K_M -values compared to the studies simulated here, in which natural microbial communities were used according to the OECD guidelines. Compound turnover and the related yields in experiments with natural inoculum and multiple substrates may be lower than single-strain/single-substrate experiments due to the enrichment of metabolites (incomplete mineralization) or to the use of multiple carbon sources derived from dissolved organic carbon or SOM.^{31,42}

Uncertainty analysis and parameter identification

The pre-calculated model input parameters were compared to those fitted by the DREAM_(ZS) and the Pattern Search algorithms (Table 3). For 2,4-D, the fitted yields are higher than the pre-calculated one.

Conversely, fitted yields for ibuprofen are substantially lower than the pre-calculated theoretical yield, which may be again an indication of incomplete mineralization of ibuprofen. Both $v_{\max}$ and K_M derived by the DREAM_(ZS) algorithm are clearly higher than the pre-calculated values and those derived by the Pattern Search algorithm, and this affects also the $\mu_{\max}$ -values. However, the ratio between $v_{\max}$ and K_M , which is effectively determining metabolism (eq. 1), is for 2,4-D comparable amongst all four methods. For ibuprofen this ratio is higher for the DREAM_(ZS) algorithm but within a factor of two of the values derived by the other methods. The DREAM_(ZS) algorithm returned K_{OC} -values for the 2,4-D simulation that are very close to the K_{OC} of 2,4-DCP. The value found with the Pattern Search algorithm is in between the K_{OC} -values of 2,4-D and 2,4-DCP. A large disagreement between fitted and pre-calculated values is observed for the K_{OC} of ibuprofen, where the pre-calculated value was obtained by a regression equation⁵⁴. Without exception, the pre-calculated parameters are within the 95% credibility interval given by the DREAM_(ZS) method. This gives additional confidence to the identified system kinetics.

A simultaneous fit of all parameters, as it is often done (for example, Brimo *et al.*⁵⁵), can produce a better fit to experimental data. This was also the case in our simulations, where the RMSE of the simulated results was lower when the input parameters were fitted (Table 3). Still, estimating the yield with an independent method showed some advantages for the simulation. The parameter identifiability improved, as can be seen from a decrease of the correlation between the fit parameters (Table S7). Using the criteria of Frutiger *et al.*⁵⁶ ($r < 0.7$, $\sigma/\mu < 0.5$), all parameters were identifiable via model calibration to the 2,4-D data. For ibuprofen, only Y , K_{OC} , and K_M but not $v_{\max}$ and $X(0)$ were identifiable (Table S8) (details in SI section S4). The largest effect was seen on the uncertainty of the prediction: omitting Y from the fit procedure greatly reduced the uncertainty in the model predictions, as shown by the width of the 95th-percentile credibility interval (Figures S5-S8), in particular for NER and X . Importantly, as we showed in this study, the knowledge of the yield gives insight into the degradation processes. It is now possible to elucidate the nature of non-extractable residues by a combination of novel analytics, basic principles, and dynamic simulation.

<Table 3>

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420

421 The dynamic degradation model with description is available in a public version at

422 <http://www.magicpah.org/links/> or <http://homepage.env.dtu.dk/stt/>.

423 Supporting Information available comprises: more detailed equations and parameter of the modeling
424 approach. This information is available free of charge via the Internet at <http://pubs.acs.org/>

425

426 The **MTB theoretical yield tool** is available as excel or Python code on request to the first author.

427

428

429 **References**

430 (1) EU. Regulation (EC) No 1907/2006 of the European Parliament and the Council of 18 December 2006
431 concerning Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH). *Official*
432 *Journal of the European Union*; 2006, L 136.

433 (2) EU. Regulation (EC) No 1107/2009 of the European Parliament and the council of 21 October 2009
434 concerning the placing of plant protection products on the market and repealing Council Directives
435 79/117/EEC and 91/414/EEC. *Official Journal of the European Union*; 2009, L 309/1.

436 (3) EU. Commission regulation (EU) No 283/2013 setting out the data requirements for active substances,
437 in accordance with Regulation (EC) No 1107/2009 of the European Parliament and the Council
438 concerning the placing of plant protection products on the market. *Official Journal of the European*
439 *Union*; 2013, L 93/1.

- (4) EMA European Medicines Agency. Guideline on the assessment of persistent, bioaccumulative and toxic (PBT) or very persistent and very bioaccumulative (vPvB) substances in veterinary medicinal products. September 2015.
- (5) ECHA European Chemical Agency. Guidance on Information Requirements and Chemical Safety Assessment, Chapter R.11: Endpoint specific guidance (PBT/vPvB assessment). Draft version 3.0, March 2017.
- (6) OECD. *Test No. 304A: Inherent Biodegradability in Soil*. OECD Publishing Paris; 1981. DOI 10.1787/9789264070448-en
- (7) OECD. *Test No. 307: Aerobic and Anaerobic Transformation in Soil*. OECD Publishing: Paris 2002a. DOI 10.1787/9789264070509-en
- (8) OECD. *Test No. 308: Aerobic and Anaerobic Transformation in Aquatic Sediment Systems*. OECD Publishing: Paris 2002b. DOI 10.1787/9789264070523-en
- (9) OECD. *OECD Test No. 309: Aerobic Mineralisation in Surface Water – Simulation Biodegradation Test*. OECD Publishing: Paris 2004. DOI 10.1787/9789264070547-en
- (10) Matthies M.; Witt J.; Klasmeier J. Determination of soil biodegradation half-lives from simulation testing under aerobic laboratory conditions: A kinetic model approach. *Environ. Pollut.* **2008**, *156*, 99-105.
- (11) Honti, M.; Hahn, S.; Hennecke, D.; Junker, T.; Shrestha, P.; Fenner, K. Bridging across OECD 308 and 309 data in search of a robust biotransformation indicator. *Environ. Sci. Technol.* **2016**, *50* (13), 6865-6872; DOI 10.1021/acs.est.6b01097
- (12) Kästner, M.; Nowak, K. M.; Miltner, A.; Trapp, S.; Schäffer, A. Classification and modelling of non-extractable residue (NER) formation of xenobiotics in soil – a synthesis. *Crit. Rev. Env. Sci. Tec.* **2014**, *44* (19), 1-65.
- (13) Alexander M. Aging, bioavailability, and overestimation of risk from environmental pollutants. *Environ. Sci Technol.* **2000**, *34*, 4259-4264.
- (14) Bosma, T. N.; Middeldorp, P. J. M.; Schraa, G.; Zehnder, A. J. B. Mass transfer limitation of biotransformation: Quantifying bioavailability. *Environ. Sci. Technol.* **1997**, *31*, 248-252.
- (15) Katayama, A.; Bhula, R.; Burns, G. R.; Carazo, E.; Felsot, A.; Hamilton, D.; Harris, C.; Kim, Y.H.; Kleter, G.; Koerdel, W.; Linders, J.; Peijnenburg, J.G.M.W.; Sabljic, A.; Stephenson, R.G.; Racke,

- 469 D.K.; Rubin, B.; Tanaka, K.; Unsworth, J.; Wauchope, R. D. Bioavailability of Xenobiotics in the
470 Soil Environment. In: *Reviews of Environmental Contamination and Toxicology*; D. M. Whitacre
471 (Ed.); Springer, New York, 2010; pp 1–86.
- 472 (16) Ingerslev, F; Baun, A; Nyholm, N. Aquatic biodegradation behavior of pentachlorophenol assessed
473 through a battery of shake flask die-away tests. *Environ. Toxicol. Chem.* **1998**, 17 (9), 1712-1719.
- 474 (17) Schlegel, H.G. *Allgemeine Mikrobiologie*, 5th, ed.; Verlag Georg Thieme: Stuttgart, Germany, 1981.
- 475 (18) Madigan, M.T.; Martinko, J.; Parker, J. *Biology of Microorganisms*. International Student Edition -
476 Pearson Inc.; 2011.
- 477 (19) Heimann, A.C.; Friis, A.K.; Scheutz, C.; Jakobsen, R. Dynamics of reductive TCE dechlorination in
478 two distinct H₂ supply scenarios and at various temperatures. *Biodegradation* **2007**, 18 (2), 167-
479 179.
- 480 (20) Won, W.D.; DiSalvo, L.H.; James, N.G. Toxicity and mutagenicity of 2,4,6-trinitrotoluene and its
481 microbial metabolites. *Appl. Environ. Microbiol.* **1975**, 31, 576–580.
- 482 (21) Honeycutt, M.E.; Jarvis, A.S.; McFarland, V.A. Cytotoxicity and mutagenicity of 2,4,6-trinitrotoluene
483 and its metabolites. *Ecotoxicol. Environ. Saf.* **1996**, 35, 282–287.
- 484 (22) Diekert G. Grundmechanismen des Stoffwechsels und der Energiegewinnung. In:
485 *Umweltbiotechnologie*; Ottow, J. C. G.; Bidlingmaier, W. (Eds.); Fischer Verlag, Stuttgart,
486 Germany, 1997; pp 1-38.
- 487 (23) McCarty, P. L. Thermodynamic electron equivalents model for bacterial yield prediction: Modifications
488 and comparative evaluations. *Biotechnol. Bioeng.* **2007**, 97 (2), 377–388.
- 489 (24) Heijnen, J. J. A new thermodynamically based correlation of chemotrophic biomass yields. *Anton.*
490 *Leeuw. Int. J. G.* **1991**, 60, 235-256.
- 491 (25) vanBriesen, J. M. Evaluation of methods to predict bacterial yield using thermodynamics.
492 *Biodegradation* **2002**, 13 (3), 171–90.
- 493 (26) Xiao, J.; VanBriesen, J. M. Expanded thermodynamic true yield prediction model: adjustments and
494 limitations. *Biodegradation* **2008**, 19 (1), 99–127.
- 495 (27) Nowak, K.M.; Miltner, A.; Gehre, M.; Schäffer, A.; Kästner, M. Formation and fate of bound residues
496 from microbial biomass during 2,4-D degradation in soil. *Environ. Sci. Technol.* **2011**, 45, 999-
497 1006.

- 498 (28) Nowak, K.M.; Girardi, C.; Miltner, A.; Gehre, M.; Schäffer, A.; Kästner, M. Contribution of
499 microorganisms to non-extractable residue formation during biodegradation of ibuprofen in soil.
500 *Sci. Tot. Environ.* **2013**, *445*, 377-384.
- 501 (29) Girardi, C.; Nowak, K. M.; Carranza-Diaz, O.; Lewkow, B.; Miltner, A.; Gehre, M., Schäffer, A;
502 Kästner, M. Microbial degradation of the pharmaceutical ibuprofen and the herbicide 2,4-D in
503 water and soil - Use and limits of data obtained from aqueous systems for predicting their fate in
504 soil. *Sci. Total Environ.*, **2013**, *444*, 32–42.
- 505 (30) Cornish-Bowden, A. *Fundamentals of enzyme kinetics*; Portland Press, London, U.K., 1995.
- 506 (31) Kovárová-Kovar K.; Egli T. Growth kinetics of suspended microbial cells: From single-substrate-
507 controlled growth to mixed-substrate kinetics. *Microbiol. Mol. Biol. Rev.* **1998**, *62* (3), 646–666.
- 508 (32) Trapp, S.; Franco, A.; MacKay, D. Activity-based concept for transport and partitioning of ionizing
509 organics. *Environ. Sci. Technol.* **2010**, *44* (16), 6123–6129
- 510 (33) van Uden N. Transport-limited growth in the chemostat and its competitive inhibition; a theoretical
511 treatment. *Archiv für Mikrobiologie* **1967**, *58*, 145-154.
- 512 (34) Brusseau, M.L.; Larsen, T.; Christensen, T.H. Rate-limited sorption and nonequilibrium transport of
513 organic chemicals in low organic carbon aquifer materials. *Wat. Resour. Res.* **1991**, *27*(6), 1137-
514 1145.
- 515 (35) Johnson, M.D.; Keinath II, T.M.; Weber Jr., W.J. A distributed reactivity model for sorption by soils
516 and sediments. 14. Characterization and modeling of phenanthrene desorption rates. *Environ.*
517 *Sci. Technol.* **2001**, *35*, 1688-1695.
- 518 (36) Rein, A.; Adam, I. K. U.; Miltner, A.; Brummer, K.; Kästner, M.; Trapp, S. Impact of bacterial activity
519 on turnover of insoluble hydrophobic substrates (phenanthrene and pyrene) - Model simulations
520 for prediction of bioremediation success. *J. Hazard. Mater.* **2016**, *306*, 105–114.
- 521 (37) Barriuso, E.; Benoit, P.; Dubus, I. G. Formation of pesticide nonextractable (bound) residues in soil:
522 Magnitude, controlling factors and reversibility. *Environ. Sci. Technol.* **2008**, *42*, 1845-1854.
- 523 (38) Kindler, R; Miltner, A; Richnow, H.-H.; Kästner, M. Fate of gram-negative bacterial biomass in soil—
524 mineralization and contribution to SOM. *Soil Biol. Biochem.* **2006**, *38*, 2860–2870.
- 525 (39) Kindler, R.; Miltner, A.; Thullner, M.; Richnow, H.-H.; Kästner M. Fate of bacterial biomass derived
526 fatty acids in soil and their contribution to soil organic matter. *Org. Geochem.* **2009**, *40*, 29–37.

- 527 (40) Miltner, A.; Bombach, P.; Schmidt-Brücken, B.; Kästner, M. SOM genesis: Microbial biomass a
528 significant source. *Biogeochemistry* **2012**, *111*, 41-55.
- 529 (41) Thauer, R. K.; Jungermann, K.; Decker, K. Energy Conservation in Chemotrophic Anaerobic Bacteria.
530 *Bacteriol. Rev.* **1977**, *41* (1), 100-180.
- 531 (42) Helbling, D.E.; Hammes, F.; Egli, T.; Kohler, H.-P. E. Kinetics and yields of pesticide biodegradation
532 at low substrate concentrations and under conditions restricting assimilable organic carbon. *Appl.*
533 *Environ. Microb.* **2014**, *80* (4), 1306–1313.
- 534 (43) Yuan, Z.; vanBriesen, J. M. Bacterial growth yields on EDTA, NTA, and their biodegradation
535 intermediates. *Biodegradation* **2008**, *19* (1), 41–52; DOI <http://doi.org/10.1007/s10532-007-9113-y>
- 536 (44) Tuxen, N.; De Liptay, J. R.; Albrechtsen, H.-J.; Aamand, J.; Bjerg, P.L. Effect of exposure history on
537 microbial herbicide degradation in an aerobic aquifer affected by a point source. *Environ. Sci.*
538 *Technol.* **2002**, *36*, 2205-2212.
- 539 (45) MathWorks 2016. Help to “patternsearch”. <https://se.mathworks.com/help/gads/patternsearch.html>
- 540 (46) ter Braak, C.J.F.; Vrugt, J.A. Differential evolution Markov chain with snooker updater and fewer
541 chains. *Stat. Comput.* **2008**, *18* (4), 435-446.
- 542 (47) Vrugt, J. A. Markov chain Monte Carlo simulation using the DREAM software package: Theory,
543 concepts, and MATLAB implementation, *Environ. Model. Softw.* **2016**, *75*, 273-316.
- 544 (48) Heijnen, J. J.; Dijken, J. P. In search of a thermodynamic description of biomass yields for the
545 chemotrophic growth of microorganism. *Biotechnol. Bioeng.* **1992**, *39*, 833–852.
- 546 (49) Brock, A.; Kästner, M.; Trapp, S. Microbial growth yield estimates from thermodynamics and its
547 importance for degradation of pesticides and formation of biogenic non-extractable residues, *SAR*
548 *QSAR Environ. Res.*, 28:8, 629-650, DOI: 10.1080/1062936X.2017.1365762.
- 549 (50) Wick, L.Y.; Colangelo, T.; Harms, H. Kinetics of mass transfer-limited bacterial growth on solid PAHs,
550 *Environ. Sci. Technol.* **2001**, *35*, 354–361.
- 551 (51) Adam, I. K. U.; Rein, A.; Miltner, A.; Fulgêncio, A. C. D.; Trapp, S.; Kästner, M. Experimental Results
552 and Integrated Modeling of Bacterial Growth on an Insoluble Hydrophobic Substrate
553 (Phenanthrene). *Environ. Sci. Technol.* **2014**, *48* (15), 8717–8726.

- 554 (52) Toräng, L.; Nyholm, N.; Albrechtsen, H.-J. Shifts in biodegradation kinetics of the herbicides MCPP
555 and 2,4-D at low concentrations in aerobic aquifer materials. *Environ. Sci. Technol.* **2003**, *37* (14),
556 3095-3103.
- 557 (53) Lehmicke, L. G.; Williams, R. T.; Crawford, R. L.. 14C-most-probable-number method for
558 enumeration of active heterotrophic microorganisms in natural waters. *Appl. Environ. Microbiol.*
559 **1979**, *38* (4), 644–649.
- 560 (54) Franco, A.; Trapp, S. Estimation of the soil-water partition coefficient normalized to organic carbon for
561 ionizable organic chemicals. *Environ. Toxicol. Chem.* **2008**, *27* (10), 1995–2004.
- 562 (55) Brimo, K.; Garnier, P.; Sun S.; Bertrand-Krajewski, J.-L.; Cebon, A.; Ouvrard, S. Using a Bayesian
563 approach to improve and calibrate a dynamic model of polycyclic aromatic hydrocarbons
564 degradation in an industrial contaminated soil. *Environ. Pollut.*, **2016**, *215*, 27-37; DOI
565 [dx.doi.org/10.1016/j.envpol.2016.04.094](https://doi.org/10.1016/j.envpol.2016.04.094)
- 566 (56) Frutiger, J.; Marcarie, C.; Abildskov, J.; Sin, G. A Comprehensive Methodology for Development,
567 Parameter Estimation, and Uncertainty Analysis of Group Contribution Based Property Models-An
568 Application to the Heat of Combustion. *J. Chem. Eng. Data* **2016**, *61* (1), 602–613.
- 569 (57) Chong, N. M.; Tsai, S. C.; Le, T. N. The biomass yielding process of xenobiotic degradation.
570 *Bioresource Technol.* **2010**, *101* (12), 4337–4342.
- 571 (58) Oh, Y.-S.; Shareefedeen, Z.; Baltzis, B. C.; Bartha, R. Interactions between benzene, toluene, and p-
572 xylene (BTX) during their biodegradation. *Biotechnol. Bioeng.* **1994**, *44*, 533–538.
- 573

Tables

Table 1. Comparison of yield estimates (g biomass carbon g⁻¹ substrate carbon, g C g⁻¹ C) using the TEEM2²³ and the MTB methods for biotechnological substrates. AE is absolute error and MAE is mean absolute error. Measured yields are taken from ref. 48.

Biotech. substrates	Measured	TEEM2	AE	MTB	AE
Acetate [−]	0.41	0.40	0.01	0.47	0.05
Citrate ^{3−}	0.365	0.34	0.025	0.29	0.075
Ethanol	0.53	0.67	0.14	0.60	0.07
Formaldehyde	0.47	0.51	0.04	0.58	0.11
Glucose	0.61	0.48	0.13	0.61	0.0
Glycerol	0.67	0.55	0.12	0.62	0.05
Glyoxylate	0.22	0.27	0.05	0.27	0.05
Methanol	0.54	0.56	0.02	0.66	0.12
Propionate [−]	0.48	0.47	0.01	0.50	0.02
Pyruvate [−]	0.32	0.39	0.07	0.39	0.07
MAE			0.0615		0.0615

Table 2. Comparison of yield estimates (g microbial biomass dw g⁻¹ substrate, g g⁻¹) using the TEEM2²³ and the MTB methods for chemicals of environmental concern. AE is absolute error and MAE is mean absolute error.

Environmental chemicals	Observed	TEEM2	AE	MTB	AE	Reference for observed Y
2,4-D (¹² C)	0.25	0.39	0.14	0.23	0.02	⁵⁷
2,4-D (¹³ C ring-labeled)	0.18; 0.25					This study; ⁵²
2,4-DCP	0.30	0.41	0.11	0.21	0.09	⁵⁷
Benzene	0.71	0.84	0.13	0.65	0.06	⁵⁸
Carbofuran	0.52	0.59	0.07	0.51	0.01	⁴²
Carbon tetrachloride		0		0		Persistent
DDT		0.42		0.25		
Ibuprofen (¹² C)	0.43	0.61	0.18	0.62	0.19	This study
Ibuprofen (¹³ C ring-labeled)	0.39					²⁸
Linuron	0.05	0.33	0.28	0.32	0.27	⁴²
Nitrilotriacetic acid	0.23	0.23	0.00	0.27	0.04	⁴³
Pentachlorophenol		0.19		0		Persistent ¹⁶
Phenanthrene	0.21	0.82	0.61	0.53	0.32	⁵¹
Pyrene	0.32	0.54	0.22	0.44	0.12	³⁶
Trichloroethene		0.16		0.11		Persistent
Toluene	0.71	0.86	0.15	0.69	0.02	⁵⁸
MAE			0.189		0.114	

Table 3. Input and fit parameters used for the simulation of degradation experiments of 2,4-D and ibuprofen described in Nowak *et al.*^{27,28} and Girardi *et al.*²⁹. Parameter values highlighted in bold were not fitted but pre-calculated.

Parameter	unit	Manual w/ pre-estimated yield	Pattern Search w/o pre- estimated yield	DREAM w/ pre- estimated yield (95% credibility interval)	DREAM w/o pre-estimated yield (95% credibility interval)
2,4-D					
Y	$\text{g } ^{13}\text{C}_{\text{biomass}} (\text{g } ^{13}\text{C}_{\text{substrate}})^{-1}$	0.28^a	0.36	0.28^a	0.31 (0.21; 0.52)
b	d^{-1}	0.05^b	0.05^b	0.05^b	0.05^b
μ_{max}	$\text{g } ^{13}\text{C}_{\text{biomass}} (\text{g } ^{13}\text{C}_{\text{substrate}} \text{ d})^{-1}$	1.1 ^c	1.43	1.61 (0.38; 2.72)	1.73 (0.40; 3.85)
$v_{\text{max}}/K_{\text{M}}$	$\text{m}^{-3} (\text{g } ^{13}\text{C d})^{-1}$	2.72 ^d	2.08 ^d	2.72 ^d	2.47 ^d (1.45; 3.61)
$X(0)$	g m^{-3}	0.172	0.28	0.87 (0.16; 1.34)	0.88 (0.16; 1.4)
K_{OC}	L kg^{-1}	2,4-D: 71.4^e ; 2,4-DCP: 689^f	300	655 (218; 977)	668 (506; 836)
sum	$\text{g } ^{13}\text{C m}^{-3}$	5.57	1.56	2.17	2.13
RMSE ^g					
Ibuprofen					
Y	$\text{g } ^{13}\text{C}_{\text{biomass}} (\text{g } ^{13}\text{C}_{\text{substrate}})^{-1}$	0.43^a	0.28	0.43^a	0.32 (0.11; 0.53)
b	d^{-1}	0.03^b	0.03^b	0.03^b	0.03^b
μ_{max}	$\text{g } ^{13}\text{C}_{\text{biomass}} (\text{g } ^{13}\text{C}_{\text{substrate}} \text{ d})^{-1}$	0.50 ^c	0.28	1.41 (0.18; 4.1)	1.34 (0.12;

	$^{13}\text{C}_{\text{substrate}} \text{ d}^{-1}$				3.7)
$X(0)$	g m^{-3}	0.069	1.2	0.69 (0.05; 1.3)	0.62 (0.05, 1.3)
$v_{\text{max}}/K_{\text{M}}$	$\text{m}^{-3} (\text{g}^{13}\text{C d})^{-1}$	0.39	0.35	0.51 (0.10; 2.4)	0.67 (0.12; 2.7)
K_{OC}	L kg^{-1}	108^{h}	552	558 (83.8; 788)	546 (99.3; 789)
sum	$\text{g}^{13}\text{C m}^{-3}$	4.62	1.89	2.86	2.97
RMSE ^g					

588 a: estimated with MTB method; conversion factor 2,4-D = 0.822 and ibuprofen = 1.41 for conversion to g
589 microbial biomass dw per g substrate; b: from slope of $\ln X$ at the end of the experiment; c: from slope of
590 $\ln X$ in the initial growth phase; d: K_{M} of 2,4-D was calculated⁵²; e: K_{OC} of 2,4-D (estimated⁵⁴) was used for
591 rapid adsorption; f: K_{OC} of 2,4-DCP (estimated⁵⁴) was used for slow adsorption (sequestration); g
592 description of sum RMSE see SI S3.3; h: estimated⁵⁴
593

594 **Figure legends:**

595 **Figure 1.** Measured and simulated ^{13}C -label distribution. A) Top: $^{13}\text{C}_6$ -2,4-D, and B) bottom: $^{13}\text{C}_6$ -
 596 ibuprofen. Symbols show measured data, curves show the simulated turnover. Symbols: CO_2 (black
 597 square), NER (grey circle), extractable ^{13}C -label (dark grey triangle), and added compound, i.e., ibuprofen
 598 or 2,4-D (black circle). Curves: Sequestered (black dashed (—)), extractable compound (dark grey (—)),
 599 CO_2 (black (—)), and living biomass + dead biomass + sequestered compound (=NER) (light grey). Error
 600 bars show the standard deviation of the measurements as reported by Girardi *et al.*²⁹ and Nowak *et*
 601 *al.*^{27,28}.

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 603
 604 **Figure 2.** Simulations results for the growth of biomass. A) Top: $^{13}\text{C}_6$ -2,4-D and B) bottom: $^{13}\text{C}_6$ -ibuprofen.
 605 Symbols show measured data, curves show simulation of the formation of living and dead biomass.
 606 Symbols: Phospholipid fatty acids (PLFA; black circles), total amino acids (tAA; dark grey squares), and
 607 total amino acids multiplied with a factor of two to yield total dead and alive biomass (empty squares).
 608 Curves: Concentration of living biomass X (black), concentration of dead biomass X_{dead} (light grey), and
 609 concentration of living and dead biomass $X + X_{\text{dead}}$ (dark grey). Error bars show the standard deviation of
 610 the measurements as reported by Nowak *et al.*^{27,28}. The dotted vertical line is the halftime of decay ($\ln 2 /$
 611 *b*) after maximum measured PLFA and indicates were > 50% of tAA is necromass.

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Figure 1

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Figure 2