Probabilistic indicators for soil and groundwater contamination risk assessment

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Abstract

Deterministic assessments of whether, when, and where environmental safety thresholds are exceeded by pollutants are often unreliable due to uncertainty stemming from incomplete knowledge 2 of the properties of environmental systems and limited sampling. We present a global sensitivity 3 analysis to rank the contribution of uncertain parameters to the probability, P, of a target quantity to 4 exceed user-defined environmental safety thresholds. To this end, we propose a new index (AMAP) 5 which quantifies the impact of a parameter on P and can be readily employed in probabilistic risk 6 assessment. We apply AMAP, along with existing moment-based sensitivity indices, to quantify 7 the sensitivity of soil and aquifer contamination following herbicide glyphosate (GLP) dispersal to 8 soil hydraulic parameters. Target quantities are GLP and its toxic metabolite aminomethylphosc phonic acid (AMPA) concentrations in the top soil as well as their leaching below the root zone. 10 The global sensitivity analysis encompasses six scenarios of managed water amendments and rain-11 fall events. The biodegradation of GLP and AMPA varies slightly across scenarios, while leaching 12 below the root zone is greatly affected by the assumed hydrologic boundary conditions. AMAP 13 shows that, among the tested uncertain parameters, absolute permeability, air-entry suction, and 14 porosity have the greatest impact on GLP and AMPA probability to pollute the aquifer by exceed-15 ing the aqueous concentration thresholds. Our results show that AMAP is effective to thoroughly 16 explore time histories arising from model-based predictions of environmental pollution hazards. 17

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¹⁸ The proposed methodology may support informed decision making in risk assessments and help

¹⁹ assessing ecological indicators through threshold-based analyses.

Keywords: global sensitivity analysis; uncertainty quantification; modeling; pollution; soil; groundwater; glyphosate; AMPA; environmental risk assessment

20 1. Introduction

Surface waters and aquifers are the recipients of contaminants resulting from anthropogenic 21 activities such as agriculture, industry and waste treatment. According to the Lancet Commission 22 on Pollution and Health, more than 140,000 synthetic molecules have been developed since 1950; 23 of those, the 5,000 most produced molecules can be found in the environment worldwide (Landri-24 gan et al., 2018). These molecules and their metabolites can be persistent in the environment and, 25 therefore, their detrimental effects can dramatically extend over space and time, thus harming not 26 only humans and other living organisms but also their descendants (Kubsad et al., 2019). In order 27 to minimize and control harmful impacts, the use of these molecules has to be properly planned, 28 managed, and regulated, thus requiring a good understanding of their dynamics in the environment. 29 Mathematical models are often used as decision support tools to evaluate contaminant degra-30 dation and transport (e.g., EPA, 2008; Porta et al., 2018; Manheim et al., 2019). Predictive models 31 are also employed to carry out assessments on future scenarios such as climate change, land-use 32 change, and global environmental change (e.g., Hiscock et al., 2007; Armitage et al., 2011; Brack 33 et al., 2015). Processes and environmental factors controlling contaminant dynamics (e.g., soil-34 water dynamics, contaminant sorption to soil minerals and organic matter, biochemical degra-35 dation, microbial-nutrient interactions, and soil-plant interactions) are described and coupled in 36 models through mathematical equations with parameters often sourced from literature or estimated 37 against laboratory or field experiments (Jackson et al., 2000; Barrios et al., 2019). Since envi-38 ronmental systems are complex and open to energy and mass flows, it is difficult to constrain and 39

⁴⁰ model all controlling processes (Oreskes *et al.*, 1994). Deterministic models neglect the inherent
⁴¹ uncertainty associated with model structure and parameters (Uusitalo *et al.*, 2015).

To overcome the shortcoming of deterministic models, uncertainty quantification and sensi-42 tivity analysis methods are used within a stochastic framework to provide decision makers with 43 estimates of the potential outcomes of tested scenarios (e.g., Bates et al., 2003; Walker et al., 2003; 44 Uusitalo et al., 2015). There exist many different approaches for sensitivity analysis (e.g., Razavi 45 & Gupta, 2015; Pianosi et al., 2016; Ceriotti et al., 2018). Among these, global sensitivity analysis 46 (GSA) is one of the most comprehensive approaches because it allows (a) quantifying sensitivity 47 across the entire parameter space and (b) accounting for the joint effects resulting from the uncer-48 tainty in diverse parameters even in nonlinear models. Output sensitivity to model parameters is 49 commonly quantified using variance-based techniques such as the Sobol's indices (Sobol', 1993); 50 more recently, the AMA family of indices was introduced (Dell'Oca et al., 2017) to quantify sen-51 sitivity in terms of any statistical moment of the probability density function (pdf) of the model 52 outputs. Available sensitivity indices do not provide a straightforward assessment of how a pa-53 rameter influences the probability for a model output (e.g., the concentration of a contaminant) 54 to exceed a user-defined threshold value. In environmental risk assessment and management, the 55 policies and protection strategies often rely on regulatory guidelines that specify a safety limit for 56 a certain contaminant. For example, the European Commission has set two severe safety limits to 57 protect water resources quality from pesticide contamination: (a) the concentration of single pesti-58 cides and their relevant metabolites must not exceed 0.1 μ g l⁻¹; and (b) the sum of pesticides and 59 their metabolites concentration must not exceed 0.5 μ g l⁻¹ (2006/118/EC, 2006). 60

In this study we introduce a new global sensitivity index (*AMAP*) to rank parameters based on their impact on the probability to exceed a defined safety limit for single contaminants and mixtures. *AMAP* complements the available *AMA* moment-based indices by targeting sensitivity with respect to the exceedance probability rather than the statistical moments of the output

pdf. We demonstrate the use of AMAP to the case study of dispersal of the herbicide glyphosate 65 (GLP) throughout a soil profile in an irrigated winter wheat field, using the results of la Cecilia 66 et al. (2018) as a reference. The case study selection is motivated by the observation that GLP is 67 currently the most widely used herbicide worldwide (Maggi et al., 2019). Modeling GLP biodegra-68 dation pathways requires a complex network of bioreactive processes coupled to water flow and 69 solute transport. Hence, quantitative indicators that can identify relevant parameters and processes 70 are important to reduce the uncertainty involved in risk assessment and help constraining the whole 71 decision-making process. Societal implications of the uncertainties underlying environmental risk 72 assessment of GLP have been widely discussed in the recent literature (e.g., Van Straalen & Legler, 73 2018). The relevance of hydraulic parameters on the fate of GLP and AMPA in soil has been doc-74 umented in field experiments (e.g. Soracco et al., 2018; Lupi et al., 2019), as well as in numerical 75 studies (Heuvelink et al., 2010). In this work we analyze the impact of uncertainty in soil hydraulic 76 parameters on risk assessment of GLP and AMPA accumulation and leaching. We apply a number 77 of different boundary conditions affecting transport and biodegradation processes to gain a wider 78 understanding of how (i) AMAP informs on the effects of parameter uncertainty within pollution 79 risk assessment and (ii) GLP and AMPA biodegradation predicted by the model is affected by 80 the assumed boundary water fluxes. We designed such specific scenarios to represent managed 81 (irrigated) and unmanaged (not irrigated) cropping. Analyses are also accompanied by specific 82 robustness tests of the proposed AMAP index to show limits and advantages of its generalized 83 application beyond the test case presented here. While we focus on soil and water contamination 84 risk assessment, we emphasize that the proposed sensitivity index is readily applicable in other 85 contexts, such as to assess the response of ecological and environmental systems, and particularly 86 within threshold-based analysis of ecological indicators reported in recent literature (e.g., Libralato 87 et al., 2019; Fu et al., 2019). 88

89 2. Materials and Methods

We introduce here the definition of the sensitivity indices (or metrics) as well as the approach 90 we employ for their application within contamination risk assessments. In Section 2.1 we intro-91 duce the proposed sensitivity metrics and provide an operational framework for their application in 92 a generic environmental problem. Next, we illustrate the application of our method to soil contam-93 ination as a result of glyphosate (GLP) dispersal. We start by presenting an overview of the kinetic 94 model used to describe GLP contamination in a winter wheat field (Section 2.2). Target outputs for 95 risk analysis and their corresponding safety thresholds are identified in Section 2.3. In Section 2.4, 96 six scenarios with different ecohydrological boundary conditions are designed to be used in GSA. 97 Finally, we select the uncertain parameters and describe the sampling methodology in Section 2.5. 98

99 2.1. Sensitivity indices and application to environmental problems

The *AMA* sensitivity indices (Dell'Oca *et al.*, 2017) quantify the impact of each uncertain parameter on the statistical moments of the pdf of the target model outputs. Let $g(\mathbf{p})$ be an output of interest, and $\mathbf{p} = (p_1, \dots, p_i, \dots, p_N)$ a vector gathering *N* uncertain parameters. The *AMAM_i* indices quantify the impact of variability in parameter p_i on the statistical moment M_i of $g(\mathbf{p})$ (e.g., *AMAE* for the expected value *E* and *AMAV* for the variance *V*). *AMAM_i* is defined as

$$AMAM_{i} = \begin{cases} \frac{\int_{\Gamma_{i}} |\mathbf{M}[g(\mathbf{p}|p_{i})] - \mathbf{M}[g(\mathbf{p})]|\rho(p_{i})dp_{i}}{|\mathbf{M}[g(\mathbf{p})]|} & \text{if } \mathbf{M}[g(\mathbf{p})] \neq 0\\ \int_{\Gamma_{i}} |\mathbf{M}[g(\mathbf{p}|p_{i})]|\rho(p_{i})dp_{i} & \text{if } \mathbf{M}[g(\mathbf{p})] = 0 \end{cases}$$
(1)

where $\rho(p_i)$ is the pdf of p_i defined in the parameter space Γ_i . Along with Eq. (1), and to cast our work within a risk assessment framework, we introduce the new index *AMAP*, which allows quantifying the expected variation of the probability of exceedance of a threshold value *thr* as

$$AMAP_{i} = \int_{\Gamma_{i}} \left| P_{thr} - P\left[g(\mathbf{p}|p_{i}) > thr \right] \right| \rho(p_{i}) dp_{i},$$
(2)

where $P_{thr} = P[g(\mathbf{p}) > thr]$ is the unconditional probability that the quantity $g(\mathbf{p})$ exceeds the threshold *thr* and $P[g(\mathbf{p})|p_i]$ indicates the same probability conditional to parameter p_i . Note that *AMAP* provides the probability-weighted average distance between conditional and unconditional exceedance probability within Γ_i and is limited between 0 and 1. The output $g(\mathbf{p})$ and its related threshold *thr* can be any quantity of interest, including a contaminant concentration, and can be used in a generalized way for the purpose of, but not limited to, risk analysis as shown later in this work.

A flowchart is presented in Figure 1 to illustrate the workflow for the AMAP application within 115 model-based environmental pollution assessment. First, prior information needs to be collected 116 to define (i) a model structure and reference values of model parameters, and (ii) target outputs 117 of interest and the related safety thresholds, where the latter can be user-defined or taken from 118 guidelines. Additionally, various scenarios can be selected to explore the system response in di-119 verse conditions (e.g., diverse hydrologic or climatic regimes, socio-economic and/or legislative 120 constraints). As environmental models typically embed a large number of parameters, a subset of 121 these is selected to conduct sensitivity analysis. The latter may be then used to (a) rank parameter 122 importance (b) design and prioritize experimental campaigns aimed at constraining the uncertainty 123 of the selected output. A pdf $\rho(p_i)$ must be defined for each uncertain input to compute AMAP_i 124 through Eq. (2). This can be determined from available prior information through empirically 125 defined frequency distributions or according to general pdf models (e.g., Gaussian or uniform dis-126 tributions), thus defining a probability space for the selected parameters set. Different choices for 127 the input pdfs $\rho(p_i)$ can be performed and the results of the analyses may depend on the chosen 128 input distribution. Therefore, the chosen $\rho(p_i)$ should reflect available information as closely as 129 possible. Stochastic sampling of the parameters within this set is then performed N times, ren-130 dering N values of the output $g(\mathbf{p})$. These latter are employed to evaluate the conditional and 131 unconditional probability to exceed a given threshold needed in (2) to compute AMAP. The work-132

- flow is replicated for each of the selected scenarios, thus providing a scenario-dependent sensitivity
- ranking that can guide in implementing strategies to reduce uncertainty.



Figure 1: Flowchart of the steps used in this study for the application of AMAP index.

135 2.2. Reference site and modeling description

¹³⁶ We apply the framework introduced in Sec. 2.1 to the case of GLP biodegradation considered ¹³⁷ in la Cecilia *et al.* (2018). The reference winter wheat field is located in the Modena Municipality, ¹³⁸ Italy (44°40′57″N; 10°57′48″E). The soil is a typical alluvial deposit of the Po Valley region ¹³⁹ characterized by a silt loam and loam layers (SGSS, 2016). Two regions of interest were identified ¹⁴⁰ along the soil profile: the root zone (RZ) with thickness $h_{RZ} = 1$ m and the soil below RZ (BRZ) ¹⁴¹ with thickness $h_{BRZ} = 4$ m.

Rainfall data in the period 2006-2016 were collected (Arpae-Simc, 2016) and post-processed
 to compute the water infiltration after assuming a 20% rainfall interception by the crop. The actual

crop evapotranspiration was calculated from data in Arpae-Simc (2016) with the time-varying crop 144 coefficient $K_{\rm C}$ in Allen *et al.* (1998). Irrigation was estimated to match groundwater table depth ob-145 servations in Chiari et al. (2016) as described in la Cecilia et al. (2018). The 11 year precipitation, 146 actual evapotranspiration, and irrigation time series were repeated periodically to build 50 years of 147 daily boundary conditions. GLP was applied annually at rate 1.2×10^{-3} mol m⁻² (i.e., 2 kg ha⁻¹ 148 year⁻¹) in a single application event. An interception fraction of 0.2 and a drift fraction of 0.2 for 149 post-emergence pesticide application on winter wheat (Trevisan et al., 2009) were accounted for 150 as losses of the applied GLP rate, thus resulting in a net application rate $A = 7.2 \times 10^{-4}$ mol m⁻² 151 (i.e., $1.2 \text{ kg ha}^{-1} \text{ year}^{-1}$). 152

The GLP reaction network includes GLP and AMPA biodegradation and neglects chemical 153 degradation because it has been shown to only occur in soils rich in Mn oxides (e.g., birnessite 154 mineral, la Cecilia & Maggi, 2018). Inhibition of the reaction by heavy metals (e.g. Cu²⁺)(Barrett 155 & McBride, 2005; la Cecilia & Maggi, 2018; la Cecilia et al., 2018; Li et al., 2015) is also not 156 considered. GLP and AMPA degradation is tightly coupled with the nitrogen (N) cycle and a pool 157 of soil organic matter, which releases ammonium (NH_4^+) , ortophosphate (PO_4^{-3}) , and monomeric 158 organic carbon (represented by CH₂O) for microbial metabolic purposes (Maggi et al., 2008). Six 159 microbial functional groups describe the soil microbial ecology and include: GLP and AMPA 160 hydrolizing and oxidizing bacteria (B_{HyO}) , and aerobic and anaerobic bacteria $(B_{AER}$ and $B_{ANAER})$ 161 that consume organic carbon (la Cecilia & Maggi, 2018; la Cecilia et al., 2018), and NH₄⁺ and 162 NO₂⁻ oxidizing bacteria (B_{AOB} and B_{NOB}) that mediate a two-steps nitrification, and denitrifying 163 bacteria (B_{DEN}) that perform a three-step NO_3^- denitrification reduction to N_2 (Maggi *et al.*, 2008). 164 Dynamic stability of soil microbial ecology resorts to group-specific biomass background recovery 165 rates after Porta et al. (2018) showed that some functional groups can be outcompeted for some 166 parameter combinations. Inhibition on various reactions include O2 effects to anaerobic reactions 167 and pH below 6 and above 8 for microbial activity (Boon & Laudelout, 1962). Protection of 168

aqueous species, including GLP and AMPA, to the mineral phase is modeled as a linear equilibrium process. Biodegradation neglects the protected phase because it is assumed to not be accessible to exoenzymes as suggested in Riley *et al.* (2014). Benchmarking of the reaction network has been performed in Maggi *et al.* (2020) against field measurements of GLP and AMPA concentrations reported in the literature. All the details of the biodegradation reaction network used in this work are available in la Cecilia *et al.* (2018).

Deterministic simulations of GLP biodegradation were conducted using the BRTSim-v3.1a 175 general-purpose solver for reaction-advection-diffusion processes in variably saturated soils (Maggi, 176 2019). BRTSim numerically resolves the mass, momentum and energy conservation laws, bio-177 chemical kinetics, and equilibrium reactions using hybrid explicit-implicit finite volumes solvers, 178 which are described in detail in the User Guide and Technical Manual (Maggi, 2018). Under the 179 assumption that the gas phase undergoes negligible pressure gradients, advection was neglected 180 here for gaseous species while diffusion of gaseous species was explicitly included. A steady tem-181 perature profile was assigned linearly changing from 20 °C at the top soil to 14 °C at 5 m depth. 182

In the following we describe the key equations used to describe flow and transport processes, i.e. where the investigated uncertain parameters are directly involved (see Section 2.5). We indicate variables dimensions using the notation $[M,L,T,\Theta]$ for mass, length, time and temperature, respectively. The mass and momentum conservation laws for water in a one dimensional variably saturated soil with constant porosity ϕ can be written as (Richards, 1931)

$$\phi \frac{\partial S_l}{\partial t} = -\frac{\partial q}{\partial z} - ET(z,t) - \Delta B(z,t), \quad \text{with} \quad q = -\frac{\rho_l g k}{\mu} k_r(S_l) \left(\frac{\partial \psi(S_l)}{\partial z} - 1 \right), \quad (3)$$

where *t* [T] is time, S_l [-] is water saturation, *z* [L] is the vertical coordinate, *k* [L²] and k_r are the absolute and relative permeabilities, *g* [LT⁻²] is the gravitational acceleration, *q* [LT⁻¹] is the Darcy's velocity, ψ [L] is the water potential, and ρ_l [ML⁻³] and μ [ML⁻¹T⁻¹] are the water density and viscosity, respectively. Terms ET(z, t) and $\Delta B(z, t)$ [T⁻¹] describe contributions to the soil water saturation due to actual evapotranspiration and immobilization in the microbial biomass, respectively. Precipitations P(t) and irrigation I(t) [LT⁻¹] per unit planar surface are assigned at the soil surface. Eq. (3) was solved assuming the following empirical formulations for k_r and ψ proposed by Brooks & Corey (1964)

$$k_r = S_{le}^{2b+3},$$
 (4a)

$$\psi = \psi_s S_{le}^{-b},\tag{4b}$$

where *b* [-] is the pore volume distribution index, ψ_s [L] is the air-entry suction at water saturation, and *S*_{*le*} is the effective water saturation defined as

$$S_{le} = \frac{S_l - S_{lr}}{1 - S_{lr} - S_{gr}},$$
(5)

¹⁹⁸ S_{lr} and S_{gr} indicate the water and gas residual saturations, respectively.

¹⁹⁹ Mass balance of chemical species is expressed by

$$\phi \frac{\partial X^{i}}{\partial t} = \frac{\partial J_{X^{i}}}{\partial z} + \hat{r}^{i} \qquad J_{X^{i}} = -qX^{i} + \phi D_{m} \frac{\partial X^{i}}{\partial z}$$
(6)

where X^i [ML⁻³] is aqueous concentration, D_m [L²T⁻¹] is molecular diffusion, J_{X^i} [ML⁻²T⁻¹] is the advective diffusive solute flux, \hat{r}^i [ML⁻³T⁻¹] lumps the contribution from all chemical and biochemical reactions.

²⁰³ Chemical and biochemical kinetic reactions are assumed to occur only in the aqueous phase.
 ²⁰⁴ The modelling approach employed to represent the feedbacks between water stress, temperature
 ²⁰⁵ and bioreactive processes is detailed in Appendix A.

²⁰⁶ The microbial biomass was initialized with a simulation of 100 years to allow the water flow

and microbial processes related to GLP and the N cycle reaching a stationary state. The initialized 207 system was then used repeatedly for the purpose of GSA to simulate a period of $T_s = 50$ years, 208 setting GLP and AMPA initial concentrations to zero in each realization. As a consequence, this ini-209 tialization presumed the ecological capability to degrade GLP, i.e., the microbial functional groups 210 have already adapted to degrade GLP when they receive the first application. Recent experiments 211 show that adaptation may imply a time lag in the order of months before soil microorganisms can 212 effectively degrade GLP (Tang et al., 2019a), or in the order of hours due to catabolite repression 213 mechanisms triggered by substrate preference and memory of previous growth conditions (la Ce-214 cilia et al., 2019). As these observed adaptation times are significantly smaller than the considered 215 time window of 50 years we neglect these effects in our analysis. This assumption may generally 216 be reasonable because agricultural soils have typically been exposed to a wide suite of xenobiotics 217 before the application of GLP. 218

219 2.3. Target quantities and thresholds for pollution assessment

²²⁰ The target quantities selected for risk analysis are (see also Figure 2):

• the depth-averaged aqueous concentration of GLP and the mixture MXT = GLP + AMPA in BRZ (1 to 5 m depth), labeled as $C_{GLP}|_{BRZ}$ and $C_{MXT}|_{BRZ}$. The threshold values used are 0.1 $\mu g l^{-1}$ for GLP and 0.5 $\mu g l^{-1}$ for MXT, as prescribed by the Directive 2006/118/EC (2006) for the tolerable contamination of groundwater.

• GLP and AMPA mass in the top 30 cm of soil. These variables are indicated as $M_{\text{GLP}|TOP}$ and $M_{\text{AMPA}|TOP}$. We used the ecotoxicological concentration (i.e., LC₅₀) of GLP and AMPA mass fractions to earthworms as thresholds, which are set to $M_{thr,\text{GLP}} = 5,600 \text{ mg kg-soil}^{-1}$ and $M_{thr,\text{AMPA}} = 1,000 \text{ mg kg-soil}^{-1}$ for GLP and AMPA, respectively (Lewis *et al.*, 2006).

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• the yearly cumulative leaching rate of GLP and MXT between RZ and BRZ soil, F_{GLP} and F_{MXT} , corresponding to advective-diffusive fluxes between the two soil regions, defined pos-

itive for downward fluxes (i.e., from RZ to BRZ). As threshold fluxes we set 0.02 mg m⁻² y^{-1} and 0.1 mg m⁻² y^{-1} for GLP and MXT, respectively. These values correspond to the 0.01% and 0.05% of the gross GLP application rate $A_{GLP} = 2$ kg ha⁻¹ y^{-1} , following the rationale employed in 2006/118/EC (2006). We explore here the impact of using a threshold mass rate to reflect the risk of aquifer contamination and pollution. This definition is motivated by the possibility of direct comparison with application and biodegradation rates, and detachment from water saturation-dependent assessment indicators.



Figure 2: Graphical representation of the outputs considered for pollution and contamination risk analysis

In the following, we employ the term 'contamination' for a non-negligible concentration of a molecule in an environmental compartment where it should not exist, and 'pollution' when a given quantity exceeds a prescribed safety threshold following Rodriguez Eugenio *et al.* (2018). Therefore, contamination refers here to any positive level of concentration or flux detected below the root zone.

AMAP_i indices were calculated considering all the selected outputs within the simulation time window $T_s = 50$ y. Results obtained at year 15 are emphasized in the discussion because 15 y is the maximum approval period for pesticides in the EU. Note that while concentrations and top soil masses of contaminants are considered as continuous functions of time, fluxes are annual averages ²⁴⁷ used to detect persistent contaminant leaching.

²⁴⁸ Finally, we calculated the annual biodegradation efficiency in RZ of GLP

$$E_{\text{GLP}}(n) = \frac{1}{n} \sum_{j=1}^{n} \frac{1}{A\Delta t} \int_{h_{RZ}} \int_{(n-1)\Delta t}^{n\Delta t} \left[R_{\text{GLP}}^{\text{Oxi}}(z,t) + R_{\text{GLP}}^{\text{Hyd}}(z,t) \right] dz dt \tag{7}$$

as the ratio of biodegraded GLP through oxidation R_{GLP}^{Oxi} and hydrolysis R_{GLP}^{Hyd} calculated as in Eq. A.1 relative to the GLP application rate *A* in year *n*. Similarly, the biodegradation efficiency of AMPA is defined as

$$E_{\text{AMPA}}(n) = \frac{1}{n} \sum_{j=1}^{n} \int_{h_{RZ}} \int_{(n-1)\Delta t}^{n\Delta t} \frac{R_{\text{AMPA}}(z,t)}{R_{\text{GLP}}^{\text{Oxi}}(z,t)} dz dt$$
(8)

where R_{AMPA} is biodegraded AMPA. In both (7)-(8) we set $\Delta t = 1$ year. Note that Eq. (8) measures the ratio between biodegraded and produced AMPA, where the latter is a metabolite of GLP biodegradation through oxidation, see la Cecilia & Maggi (2018) for details. The biodegradation efficiencies in Eqs. (7)-(8) are not used for *AMAM* or *AMAP* analyses because there is no legislation that recommends threshold biodegradation efficiencies in field applications. However, Eq. (7)-(8) were used to interpret how parameter uncertainty affected GLP and AMPA biodegradation across different scenarios.

259 2.4. Selection of scenarios

We consider six hydrometeorological scenarios (Table 1). The first scenario corresponds to the reference (REF) case study described in la Cecilia *et al.* (2018), where the precipitation $P = P_{\text{REF}}$ and actual evapotranspiration $ET = ET_{\text{C, REF}}$ are used to estimate irrigation rates $I = I_{\text{REF}}$ necessary to match locally-measured water table depths. This case study is representative of managed agricultural crops. In the second and third scenarios, P is decreased and increased by 20% of P_{REF} to simulate drier (DRY) and wetter (WET) conditions, respectively, while irrigation is maintained as in the REF scenario. In the fourth scenario, steady state (SS) boundary conditions are set equal to time-averaged fluxes $P = \overline{P_{\text{REF}}}$, $ET = \overline{ET_{\text{C, REF}}}$, $I = \overline{I_{\text{REF}}}$ calculated over the whole simulation time and applied as constant boundary flows. In this case, the GLP application rate is also timeaveraged and applied throughout the simulated time period as $A = \overline{A_{\text{REF}}}$. DRY and SS unmanaged scenarios without irrigation are also considered.

The managed and unmanaged SS scenarios are tested to investigate the extent to which accounting of time-resolved as compared to constant hydrological fluxes can influence the prediction of GLP and AMPA biodegradation.

Boundary	Managed		Unmanaged			
conditions	(REF)	(DRY)	(WET)	(SS)	(DRY)	(SS)
Р	P _{REF}	$P_{\rm REF} imes 0.8$	$P_{\text{REF}} \times 1.2$	$\overline{P_{\text{REF}}}$	P _{REF}	$\overline{P_{\text{REF}}}$
ET	$ET_{C, REF}$	$ET_{C, REF}$	$ET_{C, REF}$	$\overline{ET_{C, REF}}$	$ET_{C, REF}$	$\overline{ET_{C, REF}}$
Ι	I _{REF}	I_{REF}	$I_{\rm REF}$	$\overline{I_{\text{REF}}}$	0	0
A	$A_{\rm REF}$	A_{REF}	A_{REF}	$\overline{A_{\text{REF}}}$	$A_{\rm REF}$	$\overline{A_{\text{REF}}}$

Table 1: Boundary conditions applied in the tested scenarios. REF, DRY, WET and SS indicate the reference scenario, dry and wet scenarios, and steady state scenarios, respectively.

274 2.5. Selection of uncertain parameters

To illustrate the use of the sensitivity metrics defined in Section 2.1, we analyze here the im-275 pact of uncertainty related to soil hydraulic properties on the target quantities defined in Section 276 2.3. We consider a total of six uncertain parameters indicated in Eqs (3)-(4), i.e. $k, b, \psi_s, \phi, S_{lr}$ and 277 S_{gr} . In principle, permeability k can be expressed as a function of the other parameters by means 278 of empirical or semi-empirical correlations (e.g. Brutsaert, 2000). However, we have included k279 within the set of uncertain parameters because of the reported non-exact dependence between ϕ 280 and k (e.g., Maggi & Porporato, 2007). These parameters are assumed to be mutually independent 281 and uniformly distributed within given ranges $\delta(\mathbf{p})$. Our choice of using a uniform pdf to charac-282 terize the uncertain model inputs rests on the idea of assigning equal weight to each value of the 283 distribution (i.e., an equal prior probability). The range $\delta(p_i)$ associated with k, b, ψ_s and ϕ (Table 284 2) is retrieved from naturally occurring soils, i.e., from hydrothermal properties database by Dai 285

et al. (2013) and the SoilGrids database in Hengl et al. (2017). Measured residual liquid satura-286 tion S_{lr} range from 0.046 to 0.31 (Ghanbarian-Alavijeh et al., 2010), while S_{gr} observations vary 287 between 0.092 and 0.22 (Smith & Browning, 1943; Peck, 1969). Based on these observations, S_{1r} 288 and S_{gr} are here considered as uniformly distributed parameters with values comprised between 289 0.05 and 0.2 (Table 2). Our analysis neglects vertical heterogeneity of soil properties. We solve 290 one-dimensional flow and transport along a vertical soil column. This choice is consistent with 291 models used for assessing pesticide leaching in regulatory frameworks (Jene, 1998; Carsel et al., 292 1985; Van den Berg et al., 2012; Carsel et al., 1985; 2006/118/EC, 2006). The assumed uncertainty 293 in the soil hydraulic parameters can be used to assess the impact of their spatial variability, such as 294 rendered by geo-referenced databases discussed in Heuvelink et al. (2010). 295

Sampling of the parameters space was conducted using a quasi Monte Carlo (QMC) technique
(Sobol', 1998). In total, 5,000 parameter realizations were generated and applied to each scenario.
Upon performing a forward modeling run for each of the selected sampling points, we obtain a
QMC ensemble of our target outputs of interest that is next used to conduct the sensitivity analysis.
We verified the convergence of the QMC samples in terms of the outputs sample pdfs, see Appendix
B. The CPU time for each QMC flow and transport simulation is 600 s (Intel Xeon Platinum 8160
@ 2.10 GHz).

k	b	ψ_s	ϕ	S_{lr}	S_{gr}
$\times 10^{-13} [m^2]$	[-]	[m] water	[-]	[-]	[-]
(0.50, 10)	(3, 7)	(-0.6, -0.1)	(0.4,0.5)	(0.05, 0.2)	(0.05, 0.2)

Table 2: Parameter value ranges.

303 3. Results and discussions

In the following we apply the global sensitivity indices *AMAM* and *AMAP* introduced in Section 2 to the reference scenario and we then analyze the impact of the considered hydrological and management regime on the system response. Finally we consider the impact of the selected scenarios on the biodegradation efficiency, to assess the relevance of the hydrologic boundary conditions
 on the GLP biodegradation reaction network.

309 3.1. Global sensitivity indices in the reference (REF) scenario

Figure 3 reports $AMAE_i$, $AMAV_i$ and $AMAP_i$ computed for all uncertain parameters p_i and for all target quantities evaluated at the time corresponding to the maximum approval period for pesticides in the EU, i.e. at t = 15 y.

AMAE_i (Figure 3a-d) suggests that the soil permeability k has the greatest influence on the sample average of all target quantities analyzed followed by (a) ψ_s and ϕ for the concentration targets and (b) by the gas residual saturation S_{gr} and ψ_s for the flux targets. Other parameters display moderate to minor effects. Similar results have been obtained for AMAV_i for the two concentrations (Figure 3e-f), while the fluxes variances are also greatly influenced by the pore volume distribution index b (Figure 3g-h).

AMAP_i (Figure 3i to 1) shows that the probability of C_{GLP} and (to a lesser extent) F_{MXT} to 319 exceed their thresholds are impacted by the variability of the uncertain parameters at t = 15 y. Oth-320 erwise, AMAP_i obtained for $C_{MXT}|_{BRZ}$ and F_{GLP} are negligible. This result is explained observing 321 that the threshold MXT concentration/GLP flux is never or always exceeded in the investigated 322 sample regardless of the parameters' values (see also Sections 3.4 and 3.3). The investigated pa-323 rameters have then a negligible influence on the probability of exceeding the threshold, while they 324 still influence the outputs mean and variance as shown by the corresponding $AMAE_i$ and $AMAV_i$ 325 values. 326

Overall Figure 3 suggests that an accurate characterization of k, ψ_s , ϕ should be prioritized to predict agrochemicals' concentrations in the aquifer. In addition, estimating agrochemical fluxes from the root zone to the aquifer would benefit from an accurate knowledge of S_{gr} and b.



Figure 3: Global sensitivity indices (a)-(d) $AMAE_i$; (e)-(h) $AMAV_i$; and (i)-(l) $AMAP_i$ evaluated for GLP and MXT aqueous concentrations and fluxes from RZ to BRZ. Analyses are relative to REF scenario at time t = 15 years.

330 3.2. Range Impact Analysis - RIA

In this section, we quantify the variability of $AMAP_i$ with respect to the level of uncertainty assumed for each parameter p_i upon considering GLP concentration $C_{\text{GLP|BRZ}}(t = 15y)$. We keep the average value for each parameter probability distribution constant and we increase/decrease the ranges of variability in $\delta(p_i)$ by a prescribed factor comprised between 0.7 and 1.1. This allows testing the robustness of parameter ranking upon maintaining uniformly distributed parameters and without violating physical constraints (i.e., positive permeability and porosity comprised between zero and one). Figure 4 shows that the *AMAP_i* indices smoothly vary. The ranking of parameters importance is also consistent for all the investigated ranges with k, ψ_s and ϕ chiefly influencing the system response, while the effect of b and S_{lr} and S_{gr} appears negligible. Increasing values of *AMAP_i* are obtained for increasing parameters ranges, which reflects the increase of assumed uncertainty in the parameter values.



Figure 4: AMAP - Range Impact Analysis for $C_{GLP|BRZ}(t = 15y)$.

343 3.3. Analysis of contamination and pollution

Figure 5a-b displays the temporal evolution of the expected values (or sample-averages) $E[M_{GLP|_{TOP}}]$ and $E[M_{AMPA|_{TOP}}]$, respectively. Both quantities are significantly smaller (by three orders of magnitude) than their ecotoxicological threshold (i.e., LC₅₀ for earthworms) and the probability to exceed the thresholds is negligible across the whole sample. Managed and unmanaged steady state scenarios result in the largest $E[M_{GLP|_{TOP}}]$ and $E[M_{AMPA|_{TOP}}]$ values, which are likely caused by reduced flushing as compared to scenarios where intense precipitations caused fast GLP and AMPA
 transport to BRZ.

The sample-averaged aqueous concentrations $E[C_{GLP}|_{BRZ}]$ and $E[C_{MXT}|_{BRZ}]$ (Figure 5c and d) increase over time in the aquifer. Wet scenarios (REF and WET) lead to faster increase in $E[C_{GLP}|_{BRZ}]$ and $E[C_{MXT}|_{BRZ}]$ as compared to dry scenarios (DRY). Our results also suggest that pollution may occur at substantially longer time scales in steady state conditions, i.e., SS scenarios do not cause any significant contamination within the investigated 50-year time period.

Figure 5e and f show the relative frequency (or sample probability) of the exceedance time \hat{t} , 356 i.e., the time at which $C_{GLP}|_{BRZ}$ and $C_{MXT}|_{BRZ}$ exceed the corresponding threshold concentrations. 357 All transient scenarios showed more than 95% probability for $C_{GLP}|_{BRZ}$ and $C_{MXT}|_{BRZ}$ to exceed 358 the threshold concentrations within a time frame of 50 years. On the other hand, considering the 359 EU maximum approval period for pesticides of 15 years, the probability to exceed the threshold 360 concentrations is smaller than 20% for $C_{GLP}|_{BRZ}$ and negligible for $C_{MXT}|_{BRZ}$. This result is con-361 sistent with Figure 3j, showing negligible $AMAP_i$ values (for all parameters) for $C_{MXT}|_{BRZ}$. We 362 further note that the probability distributions of \hat{t} display heavier right tails for DRY than for WET 363 scenarios, i.e., GLP and MXT arrival times to BRZ are characterized by larger uncertainty in DRY 364 than in WET scenarios. This result quantifies a delay in the occurrence of water pollution BRZ in 365 DRY conditions. 366

The mean GLP and MXT leaching rates from RZ to BRZ (Figure 6a-b) vary significantly across all scenarios. As expected, WET scenarios lead to higher leaching rates than DRY ones. Soil BRZ can undergo pollution after 4 years since the first GLP application in both WET and DRY scenarios. Figure 6a and b also show that upward (negative) fluxes occur from BRZ to RZ in dry scenarios. These instances are driven by particularly dry periods and elevated $ET_{\rm C}$, which result in high water suction in TOP and RZ from BRZ. Upward fluxes are consistent with previous observations of herbicide transport during capillary driven groundwater rise (Arjoon *et al.*, 1998). This result may



Figure 5: Average GLP and AMPA mass in RZ (a-b) and aqueous concentrations in BRZ (c-d); (e)-(f) represent the corresponding probability distribution of exceedance time \hat{t} , i.e., the time at which contaminant concentrations exceeded threshold concentrations.

raise awareness for pollutants accumulation at soil depths that may be reached during water table
fluctuations, where there may be a lack of active biodegraders. The recontamination does not occur
in the SS scenarios because the saturation profile is constant in time. This result confirms our
interpretation that unsteady water inputs due to precipitations can cause rapid contaminant flushes
(i.e., positive fluxes) to BRZ as compared to SS scenarios.

Figures 6c and d show that the probability to exceed F_{thr} is very high ($P_{thr} \approx 1$) within 15 years for both GLP and MXT in all scenarios except SS. Comparing Figures 6c and d with Figure 5e and f, one can conclude that the time scale associated to pollution observed in leaching rate (from RZ to BRZ) is significantly smaller than the time scale linked to resident agrochemical concentrations in BRZ. Therefore, measurements and modeling predictions of leaching rates would provide a more conservative indicator than concentration data within a contaminant risk assessment framework.



Figure 6: Sample-averaged GLP (F_{GLP}) (a) and MXT (F_{MXT})(b) leaching rate from RZ to BRZ, respectively; (c) and (d) represent the relative frequency of the exceedance time \hat{t} , i.e., the first time at which F_{GLP} and F_{AMPA} exceeded the threshold values.

385 3.4. AMAP and scenario analyses

 $AMAP_i$ can be used as a time dependent sensitivity measure as illustrated in Figure 7. We 386 observe that parameter ranking is consistent across different outputs for the same scenario because 387 the selected outputs are inherently linked between each other. Significant differences in ranking 388 are conversely observed across scenarios. For example, parameter b has an important influence on 389 the probability that both $C_{GLP}|_{BRZ}$ and $C_{MXT}|_{BRZ}$ exceed the related thresholds in DRY scenarios 390 (Figure 7b,f,d,h). On the other hand, the parameter ψ_s is predominant over b in assessing $C_{GLP}|_{BRZ}$ 391 and $C_{MXT}|_{BRZ}$ in wet scenarios (WET and REF). This result is consistent with Eq. (4), showing 392 that the impact of b decreases as the water saturation increases. 393

Figure 8 displays $AMAP_i$ values evaluated for the transient scenarios and considering $C_{GLP}|_{BRZ}$ and F_{MXT} after 15 years from the first GLP application. Analogous results for C_{MXT} and F_{GLP} in the transient conditions as well as for all target quantities in the SS scenarios are not reported since negligible values of $AMAP_i$ have been obtained for all parameters.

Parameters *k* and ψ_s are the most influential parameters on $C_{\text{GLP}}|_{BRZ}$ and F_{GLP} . These results are consistent with Eq. (3); that is, (*i*) increasing *k* promotes faster water flows, and thereby solute transport; (*ii*) decreasing ψ_s promotes higher water capillary rise from BRZ to RZ contrasting leaching. Figures 7-8 allow identifying which parameters should be further constrained, such as through measurement campaigns, to reduce the uncertainty associated with probabilistic groundwater or soil contamination risk assessment.

404 3.5. Biodegradation and flow regime

Mean biodegradation efficiencies $E[E_{GLP}]$ and $E[E_{AMPA}]$ indicate that biodegradation starts as soon as GLP is applied and increases over time (see Figure 9). Sample-averaged GLP biodegradation efficiency does not change significantly among all investigated scenarios, because $E[E_{GLP}]$ varies only between 0.85 and 0.9. The variability slightly increases when AMPA is considered, values of $E[E_{AMPA}]$ ranging between 0.24 and 0.32. Therefore, the sensitivity of microbial activity



Figure 7: Time evolution of AMAP values: each row of plots is associated with a single output, columns distinguish the different scenarios.

and biodegradation to soil water availability is not particularly relevant in the selected scenarios
even though water availability is explicitly considered in the model via Eq. (A.3).

This result suggests that simplified SS models as compared to time-resolved hydrologic boundary conditions may be used to predict overall contaminant mass budgets and is in line with previous numerical results by Tang *et al.* (2019b). However, SS scenarios do not yield accurate predictions of contaminants concentrations and leaching rates, which are instead driven by hydrological fluc-



Figure 8: *AMAP* values of the uncertain soil hydraulic parameters relative to (a) GLP aqueous concentration BRZ and (b) MXT fluxes from RZ to BRZ. Analyses were carried out at time t = 15 years and results are grouped and colored according to each one of the six scenarios.

416 tuations, as discussed in Section 3.3.

417 3.6. Final remarks

Our results demonstrate that probabilistic indicators allow identifying the impact of soil hy-418 draulic properties on pesticide contamination and leaching. In particular, we show that leaching is 419 highly variable depending on soil properties, hydrological boundary conditions, and land manage-420 ment practices. In previous studies, Stenemo & Jarvis (2007) showed that the ranking of parameters 421 may change according to soil texture, while Jury & Gruber (1989) showed that the persistence of 422 pesticides with residence times longer than one year is more affected by soil rather than climatic 423 variability. A number of studies have investigated uncertainty quantification and sensitivity anal-424 ysis of agrochemical biodegradation and leaching (Dubus et al., 2003; Stenemo & Jarvis, 2007; 425 Heuvelink et al., 2010). These modeling works assumed first order decay of pesticides in soil in 426



Figure 9: Mean biodegradation efficiency of (a) GLP and (b) AMPA in RZ and in each scenario.

contrast with our approach that uses Michaelis-Menten-Monod kinetics. Some of these studies 427 suggest that soil hydraulics has a smaller impact on pesticide fate as compared to half life and 428 sorption properties, yet our results demonstrate that soil hydraulic properties have a relevant im-429 pact on threshold exceedance probability. In particular, the AMAP time series demonstrate the 430 impact of soil hydraulic parameters appears to be more persistent in dry than wet scenarios. We 431 emphasize that the above mentioned studies typically rely on a single output statistics for parameter 432 ranking in pesticide leaching (Heuvelink et al., 2010) or quantify sensitivity by changing param-433 eters one at a time (Dubus et al., 2003). Conversely, our results advocate for the use of a suite 434 of global sensitivity indicators to thoroughly test the system response, and are in line with recent 435 studies (Borgonovo et al., 2017; Ceriotti et al., 2018). Hence AMAP can complement moment-436 based sensitivity indicators that do not account for safety threshold considered in risk assessment 437 protocols. Likewise moment-based indicators, AMAP can be used with any input parameter dis-438 tribution, model formulation and dimensionality. The present work considers only scenario and 439 parametric uncertainty. Model structure and dimensionality has been identified as another possible 440 relevant source of uncertainty. Different results may be obtained upon considering diverse flow and 441 transport formulations as well as alternative biogeochemical reaction networks. Future work is en-442

visaged to provide a formal derivation of model structure sensitivity indices within a probabilistic
risk assessment framework, as recently discussed in Dell'Oca *et al.* (2020).

445 **4.** Conclusions

446 Our work leads to the following major conclusions:

• We test the use of a suite of sensitivity indicators for soil and groundwater environmental 447 risk assessment. To this end, we introduce a new sensitivity index (AMAP) and we show its 448 application to contamination and pollution following glyphosate (GLP) herbicide dispersion 449 in an agricultural soil column. AMAP provides a time dependent indication of the relevance 450 of each parameter on the probability for a given output to exceed a user defined threshold and 451 complements available moment-based sensitivity metrics. The index developed here can be 452 readily applied to rank uncertain parameters with respect to any arbitrary threshold related 453 to the quality of any environmental sphere (e.g., water, air, soil, and their combinations) and 454 to inform appropriate management and restoration strategies. 455

• We quantify the impact of uncertainty in soil hydraulic parameters on the time required for the concentration of GLP and its toxic metabolite AMPA to exceed pollution thresholds in the top soil, below the root zone and for the fluxes measured at 1 m depth. We repeated these analyses in different scenarios of managed and unmanaged water budgets. When concentration thresholds are considered, dry scenarios result in larger uncertainty in terms of exceedance time as compared to wet ones. Measurements of contaminant fluxes reduce such uncertainty.

• Parameter ranking varies with ecohydrological scenarios of precipitations and irrigation practice. Permeability (k) and air-entry suction (ψ_s) have the greatest effect on exceedance of water quality safety limits in the reference scenario as well as in the wet scenario. The

influence of the pore volume distribution index b emerges in dry conditions, probably due 466 to its relation with the relative permeability. Gas residual saturation may play an important 467 role in contaminant transport in some scenarios, however it has shown a relatively minor 468 role on pollution assessment. Porosity appears to have a minor effect on pollution risk as 469 compared to permeability, except in the driest investigated conditions. These results suggest 470 that AMAP can be used to refine uncertainty quantification in hazard assessment through 471 measurements campaigns or to design risk management strategies, which can be specialized 472 to local ecohydrological boundary conditions. 473

• Steady state scenarios do not allow assessing contamination and pollution and overpredict pollution time scales if compared with time-resolved simulations. Remarkably the sampleaveraged GLP and AMPA biodegradation efficiencies show only minor differences across the tested scenarios. This result shows that steady state simulation may be able to match biodegradation efficiency yielded by time-resolved boundary conditions, but are not effective in rendering contamination hazards. Average flow conditions neglect the impact of short range fluctuations that play a predominant role in triggering contamination and pollution.

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491 Appendix A. Modelling of feedbacks between water saturation and biodegradation rates

The rate of change of an aqueous species for a given reaction in (6) is $\hat{r}^i = dX^i/dt = x_iR^i$, where x_i is the stoichiometric number for species *i* and R^i [ML⁻³T⁻¹] is the reaction velocity. For a generic reaction with n_O *n*-order kinetic products, n_{MM} Michaelis-Menten-Monod terms with concentration $X_{n_{MM}}$, n_{COM} competitive reactants ($X_{n_{COM}}$), and n_{INB} inhibition terms ($X_{n_{INB}}$), the reaction velocity *R* is written as

$$R^{i} = rf_{B} \prod_{n_{O}} X_{n_{O}}^{x_{n_{O}}} \cdot \prod_{n_{MM}} \frac{X_{n_{MM}}}{X_{n_{MM}} + K_{n_{MM}} \left(1 + \sum_{n_{COM}} \frac{X_{n_{COM}}}{K_{n_{COM}}}\right)} \prod_{n_{INB}} \frac{X_{n_{INB}}}{X_{n_{INB}} + K_{n_{INB}}},$$
 (A.1)

⁴⁹⁷ where $r [T^{-1}]$ is the reaction rate constant, $K_{n_{MM}}$, $K_{n_{INB}}$, $K_{n_{COM}}$ are the Michaelis-Menten half ⁴⁹⁸ saturation, competition and inhibition constants, respectively, $f_B = 1$ if the reaction is chemical or ⁴⁹⁹ if the reaction is biochemical.

$$f_B = \min\{f(S_B), f(\theta), f(S_l) / \max\{f(S_l)\}\},$$
(A.2)

where $f(S_B)$ [-], $f(\theta)$ [-] and $f(S_l)$ [-] are the specific microbial response functions to space availability, temperature and water saturation, respectively. These terms are evaluated as

$$f(S_B) = \min\left\{1 - \frac{S_B - S_{lr}}{1 - S_{gr} - S_{lr}}, 1 - \frac{f_l S_B}{S_l}, 1 - \frac{(1 - f_l) S_B}{S_g}\right\},\tag{A.3a}$$

$$f(\Theta) = \left(\frac{e^{\theta}}{e^{\theta_{LB}} + e^{\theta}}\right)^n \cdot \left(\frac{e^{\theta_{UB}}}{e^{\theta_{UB}} + e^{\theta}}\right)^m,$$
(A.3b)

$$f(S_{l}) = \frac{S_{l}}{S_{l,LB} + S_{l}} \cdot \frac{S_{l,UB}}{S_{l,UB} + S_{l}},$$
(A.3c)

where f_l [-] is the biomass water fraction, S_B [-] is the biomass saturation, θ_{LB} and θ_{UB} [Θ] and $S_{l,LB}$ [-] and $S_{l,UB}$ [-] are the lower and upper temperature (in Kelvin) and liquid saturation response parameters. The three functions in (A.3) introduce a limitation to microbially driven reactions as a
 function of environmental factors that may limit the bacterial growth and/or activity.

The function $f(S_B)$ implies that microbial functional groups can grow as long as there is enough 506 free water to immobilize, or gas space available for the cell solid fraction $(1 - f_i)$ to occupy, or there 507 is enough pore volume to host the total microbial biomass volume. Following the scheme in Maggi 508 & Porporato (2007), the function $f(S_B)$ also implies that the total water saturation includes the free 509 (mobile) water saturation S_l and the immobilized water saturation $S_{lB} = f_l S_B$. As a consequence, 510 the term ΔB in Eq. (3) accounts for the rate of change in mobile water saturation S₁ when the 511 total microbial biomass increases (i.e., $\Delta B > 0$ expresses water immobilization) or decreases (i.e., 512 $\Delta B < 0$ expresses water remobilization). Hence, Eq. (3) is subject to the constraint $S_l + S_g + S_B = 1$. 513 Function f(T) limits R when temperature is below θ_{LB} and above θ_{UB} . Finally, function $f(S_l)$ limits 514 R when S_l is below $S_{l,LB}$ or above $S_{l,UB}$. 515

The response function $f(\Theta)$ appearing in (A.1)-(A.2) was calculated with m = 0.1, n = 0.5, $\theta_{LB} = 6 \,^{\circ}\text{C}$ and $\theta_{UB} = 45 \,^{\circ}\text{C}$ to return the microbial activity curve typical of mesophiles documented in (Rittmann & McCarty, 2001) (Figure A.1a), while $f(S_l)$ was implemented with $S_{l,LB} =$ $S_{l,UB} = 0.46$ to represent typical water stresses (e.g., Moyano *et al.*, 2012; Yan *et al.*, 2018) (Figure A.1b). The response function $f(S_B)$ was implemented with $f_l = 0.85$ after Rockhold *et al.* (2005) and varies with S_B and therefore with *t* and over *z*.



Figure A.1: Microbial response functions for (a) temperature stress $f(\Theta)$, and (b) water stress $f(S_1)$ as a function of temperature and mobile water saturation, respectively. Functions apply to all microbial functional groups accounted for in the GLP biogeochemical reaction network. The curve for mesophiles (left panel) is from (Rittmann & McCarty, 2001); experiments in the right panel are from Wickland & Neff (2008).

522 Appendix B. Stability of QMC sampling

Results in Figure B.2 shows that 5,000 stochastic realizations resulted in stable relative frequency in the target outputs. The results also allow appreciating that different scenarios resulted in different frequencies.



Figure B.2: Relative frequency of M_{GLP} , F_{GLP} , F_{GLP} at increasing number of QMC realizations (from 1,000 in light to 5,000 in dark, grey, red and blue, respectively).

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