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# Stochastic Ex-Ante LCA under Multidimensional Uncertainty

Anticipating the Production of Undiscovered Microalgal Compounds in Europe

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- <sup>1</sup> Stochastic Ex-ante LCA Under Multidimensional
- <sup>2</sup> Uncertainty: Anticipating the Production of
- <sup>3</sup> Undiscovered Microalgal Compounds in Europe
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### 19 ABSTRACT

Due to their biodiversity, microalgae represent a promising source of high-value compounds that bioprospecting is aiming to reveal. Performing an ex-ante Life Cycle Assessment (LCA) to anticipate and potentially minimize the environmental burden associated with the European production of a bioprospected microalgal compound is subject to substantial and multi-factorial uncertainty, as the compound remains undiscovered. Given that any microalgal strain could potentially host the compound of interest, the ex-ante LCA should consider this bioprospecting uncertainty together with the uncertainty on the technology and the production mix.

27 Using a parameterized cultivation simulation and consequential LCA model, and an extensive 28 stochastic pseudo Monte Carlo approach, we define and propagate techno-operational, 29 bioprospecting, and production mix uncertainties for a microalgal compound being currently 30 bioprospected in Europe. We perform global sensitivity analysis using different sampling 31 strategies to identify the main contributors to the total output variance. Overall, the uncertainty 32 propagation allowed us to define and analyze the probabilistic scope for the potential 33 environmental impacts in the emerging production of high-value microalgal compounds in Europe, 34 based on current knowledge. These findings can support policy-making as well as actors in the 35 microalgal sector towards technological paths with lower environmental impact.

36 SYNOPSIS

We anticipate the environmental impacts associated with the European production of a currently
bioprospected microalgal compound via a stochastic, ex-ante, and consequential LCA based on
microalgal cultivation simulations.

40

### 41 TEXT

### 42 **1. Introduction**

43 The biological diversity of microalgae makes them a promising biological group for 44 biotechnological applications such as the production of organic products of high commercial value. Among these products, while microalgae-based 3<sup>rd</sup> generation biofuels have so far failed to 45 46 compete economically with fossil fuels<sup>1</sup>, a few high-value microalgal compounds are already 47 commercialized<sup>2,3</sup>. Recent discoveries from bioprospecting, i.e. the search for compounds and 48 properties within the biodiversity that could be valuable for human activities, range from antimicrobial to antitumoral lipids, proteins, and carbohydrates in microalgal strains<sup>4-10</sup>. This 49 50 suggests that the European microalgae sector might develop substantially in the near future. Anticipating the environmental consequences of such a development is crucial as early-stage 51 emerging systems are characterized by a high design freedom<sup>11</sup> implying potential detrimental 52 53 scenarios and because sustainable development processes "are timely, anticipatory, integrative, 54 flexible and action focused"<sup>12</sup>. Life Cycle Assessment (LCA) constitutes a robust holistic tool to 55 quantitatively anticipate such impacts on a systemic level and offers flexibility and parametrization 56 to project different scenarios.

57 LCA studies on the environmental impacts of microalgae production address primarily 58 bioenergy-oriented microalgae growth in Open Pond Raceways<sup>13–18</sup>, while high-value compounds 59 production would likely require stable and contamination-free photobioreactors (PBR). When 60 reviewing eighteen existing LCA studies of microalgae<sup>19–27</sup>, we found that they consider only 61 seven well-studied strains. This is likely because the microalgal sector has historically focused on 62 bioenergy applications and therefore primary data and assumptions on yields and operating

63 conditions exist only for a restricted set of strains that are fit for bioenergy and have already been 64 already cultivated industrially (lipid-rich, robust etc.). This data alone cannot support a 65 comprehensive assessment of the future consequences of the microalgae sector's development 66 which could potentially take as many paths as there are microalgal strains and promising 67 compounds to be discovered via bioprospecting. Furthermore, while previous bioengineering studies<sup>28,29</sup> model the potential for microalgal productivities in different PBRs and locations, 68 69 previous LCA studies only assess productions localized in one or a few sites. Yet, it is reasonable 70 to assume that an increase in demand for new microalgal compounds would, in the long run, result 71 in the supply of microalgae from several different locations with distinct productivities, 72 thermoregulation needs, and electricity mixes. In fact, 447 microalgae or cyanobacteria farms are 73 already active in 23 European countries<sup>30</sup>.

74 Forecasting the environmental consequences of future developments in the European microalgae 75 sector therefore requires extending the scope of the previous assessments to include many possible 76 scenarios. This immense number of possibilities is inherent to bioprospecting, which implies that 77 the desired property of a bioprospected compound is known (for instance an anti-inflammatory 78 compound), but not the organism which will produce it. Besides the case of bioprospecting, it is common<sup>31</sup> to consider a large number of scenarios in ex-ante LCA as it aims at anticipating the 79 80 environmental impact of emerging technologies before these are actually implemented at industrial scale<sup>32</sup>. A major challenge in ex-ante LCA is thus the need to generate Life Cycle Inventories 81 82 (LCI) for future production systems which currently exist only at a low level of technological and market maturity<sup>33</sup> such as pilot scale applications. In a recent review of common practices in ex-83 ante LCA<sup>31</sup>, one third of the 18 included studies resorted to process simulation to obtain LCIs at 84 85 an industrial scale, mainly using technology-specific simulation software. In the case of microalgal

compounds which are still being bioprospected, the upscaling anticipation can be done using parameterized physical and biological models to simulate photobioreactors and strains in different conditions. In fact, it is common practice<sup>18,21,23–25,27,34</sup> to simulate inventories for LCAs in the microalgal sector and recent studies show advanced models taking various biophysical phenomena into account<sup>35</sup>. Microalgae LCA models with extensive parameterization can be found associated with some forms of uncertainty and sensitivity analysis via stochastic sampling<sup>14,34,36</sup>.

92 These parametrized models, however, cannot cover the scope of possibilities associated with a 93 microalgal compound that is not found yet, and whose production upscaling and development in a 94 European production market are indeterminate. Yet, the consequences of an increase in demand 95 for high value microalgal compounds must be assessed early on to inform both policy makers in 96 the sustainability domain and the microalgal sector. The ex-ante assessment should put these 97 stakeholders in a better position to evaluate the likelihood of microalgal high-value compounds' 98 environmental superiority over alternatives as well as to identify the optimal scenarios and try to 99 aim for them.

100 To do so, the present work aims at investigating the impacts of an increase in demand for a 101 microalgal compound with a desired property that is currently being bioprospected in Europe (a 102 bioprospected microalgal compound). The uncertainty space associated with this technological 103 development is shaped by several forms of uncertainty that can either be epistemic i.e. due to a lack of knowledge, or aleatory because stemming from inherently random processes<sup>37</sup>. We build 104 on previous work<sup>38</sup> developing a parameterized model to simulate the microalgae cultivation 105 106 technology from a life cycle perspective. The "techno operational uncertainty" addressed in this 107 previous work is due to the impossibility of accurately predicting the productivity of a new specific 108 strain in a specific photobioreactor and location. To tackle our research question, we here need to

109 extend the parameterized model and the scope of previous LCAs in the microalgal sector to include 110 other sources of uncertainty, namely the lack of knowledge about the exact nature of the strain and 111 the bioprospected compound, the associated PBR, and the geographic developments of the market. 112 We anticipate that despite these large uncertainties, the stochastic propagations will provide 113 insightful density-based representations of the environmental consequences of an increase in 114 demand for yet undiscovered microalgal compounds. This work also contributes to the 115 developments and discussions around uncertainty, global sensitivity analysis and the associated 116 terminologies in the field of ex-ante LCA.

117

#### 118 2. **Methods**

119 The approach consists in applying stochastically generated samples to our previously developed parametrized LCA model<sup>38</sup> on which minor changes were made (cf. 2.1) to assess the impacts for 120 121 an increase in demand and production of 1 kg of bioprospected compound in Europe. The whole model is coded on Python 3.8 with the LCA package Brightway $2^{39}$  and is available on GitHub<sup>40</sup>. 122 123 The background database is the consequential version of Ecoinvent 3.6. The impact assessment 124 categories are Global Warming with a 100-year time horizon (GW100), Freshwater Eutrophication 125 (FE), Water Depletion (WD) and Terrestrial Ecotoxicity (TETinf) from ReCiPe Midpoint (H) 126 V1.13.

### 127 **2.1 Deterministic model and LCA framework**

The functional unit is 1 kg of bioprospected compound. Consequential LCA modeling was chosen as, by looking at the future effects of decisions, and including only activities and technologies expected to be able to respond to future changes in demand, the consequential approach is prospective in nature and therefore well-suited to the assessment of emerging technologies.

132 Moreover, the consequential approach reduces the number of normative assumptions needed, and 133 since it is highly speculative to anticipate normative preferences in the future, this can be argued 134 to be an advantage when performing ex-ante LCA. The foreground product system includes the 135 cultivation of microalgae in an outdoor vertical tubular PBR with the associated energy 136 consumptions for water pumping, mixing, thermoregulation by a heat pump and centrifugation for 137 biomass harvesting. The product system also comprises nutrients, CO<sub>2</sub>, water, and glass 138 consumptions. The extraction of the compound is not modeled but cell disruption is accounted for 139 as it likely constitutes the first step of the post-harvest processing, regardless of the biochemical class of the compound (protein, lipid, carbohydrate)<sup>41,42</sup>. Drying is modeled for the whole biomass 140 141 so that the co-produced biomass (dependent co-product) is ready to substitute functionally 142 equivalent products already on the market. The cultivation technology and the foreground product system are described in detail in SI I.2 and in our previous work and model<sup>38</sup>, for which a few 143 144 modifications were made to the product system. First, thermoregulation of the PBR was assumed to be provided by a reversable heat pump<sup>43</sup> with a Coefficient of Performance (COP) of 3, which 145 146 we chose as an average value over locations and seasons, instead of electric heating and a fluid 147 thermal exchanger. Additionally, considering multiple potential strains with different 148 characteristics made it necessary to model two additional possible substitution routes for the co-149 produced biomass (cf. SI I.3). This biomass can now enter the animal feed energy and feed protein markets<sup>44</sup>, or be directly incorporated in fish feed after modification of the reference fish feed 150 151 composition as previously modeled<sup>38</sup>. As a third possible substitution route, the co-produced 152 biomass can be digested in an anaerobic digester for biogas production and substitution on the 153 biogas marginal market based on a functional unit of 1MJ of heating capacity. The biogas yields 154 depend on the composition of the microalgal strains (cf. SI I.3.2.2).

155 The life cycle inventory results from a simulation of microalgal cultivation assumed from April to 156 September within Europe. This simulation and the LCI result from the interaction of 27 parameters 157 characterizing the strain and its compound, 22 techno-operational parameters characterizing the 158 cultivation technology, the PBR geometry and setup, 3 geographic parameters defining the 159 location, and 6 physical parameters. The cultivation simulation uses climatic data from the Photovoltaic Geographic Information System (PVGIS)<sup>45</sup>. The simulation, its modules, the 160 parameters, and all the equations are detailed in our previous work<sup>38</sup> and in SI I for the additions 161 162 and modifications.

### 163 2.2 Types of uncertainty and assumptions

### 164 2.2.1 Bioprospecting uncertainty

165 The "bioprospecting uncertainty" is caused by the lack of knowledge about the strain and the 166 compound that will be found to feature the desired property (for instance anti-inflammatory) and successfully upscaled after bioprospecting. The bioprospecting uncertainty is epistemic<sup>37</sup> in the 167 168 first place as it stems from the lack of knowledge about which kinds of microalgal strains or 169 compounds classes (lipid, carbohydrate, protein) are more likely to possess the desired property., 170 We have assumed that all strains and types of compounds have equal probabilities to possess the 171 desired property, as we currently lack arguments and data to hypothesize potential correlations 172 between biological traits and desired properties. Therefore, the bioprospecting process was 173 modeled as a random draw within microalgal biodiversity, whose result is subject to aleatory 174 uncertainty. This is conceptually analogous to a draw in an opaque urn in which the balls would 175 be the strains and their compounds whose proportions in the box depend on our knowledge on 176 biodiversity(cf. 3.3).

177 To do so, we first use 27 parameters defining a "strain-compound pair", i.e. a specific compound 178 hosted in a specific strain. Some parameters define biological characteristics such as biomass 179 composition, nitrogen source, or photosynthetic efficiency. Other parameters define if the strain-180 compound pair requires PBR thermoregulation at night or characterize the fate of the co-produced 181 biomass (biogas, fish feed, or animal feed) that we consider strain-specific because depending on cell wall characteristics, digestibility, toxicity etc.<sup>46-48</sup> Finally, compound-specific parameters 182 183 define respectively: whether the bioprospected compound is a lipid, protein, or carbohydrate, i.e. 184 the biochemical class of the compound; and the mass fraction of compound in such class, for 185 example measured as the mass of compound per total mass of proteins in the strain if the compound 186 is a protein. The values of these compound-specific parameters are also characterized by 187 uncertainty which means that the same functional unit of 1 kg of bioprospected compound can be 188 provided by different reference flows of cultivated microalgal biomass.

We modeled the random draw within the microalgal biodiversity by sampling random values for the parameters for which variation ranges are reported in the literature for the microalgal biodiversity (23 out of 27 parameters, cf. SI II.1). For 17 parameters out of 23, we modeled a uniform distribution within the range in lack of further arguments to use other distributions.

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### 194 2.2.2 Techno-operational uncertainty

This epistemic uncertainty was specifically addressed in our previous work<sup>38</sup> and is due to the unpredictable behavior and growth of a strain defined by its biological parameters in a specific vertical tubular PBR and location. More precisely, we used the model from Williams and Laurens (2010)<sup>28</sup> which estimates a maximum areal yield ( $g_{dw} \cdot m^{-2} \cdot d^{-1}$ ) depending on the ground horizontal irradiance (kJ· m<sup>-2</sup>· d<sup>-1</sup>) and the strain-specific theoretical energetic yield ( $g_{dw} \cdot kJ^{-1}$ ). Since

Williams and Laurens (2010)<sup>28</sup> observe that real cultivations in PBR would often reach 30% of 200 201 this maximum yield, in this study the techno-operational uncertainty refers to the lack of 202 knowledge about which PBR geometry and operational set-up will enable the modeled strain to 203 reach this percentage. The techno-operational uncertainty was addressed by simulating random 204 values of geometrical (e.g. tube diameter and distance between tubes) and operational (flow rate, 205 biomass concentration) parameters defining the PBR for a strain and location. The values were 206 sampled within ranges reported in the literature for different vertical tubular PBRs with different 207 strains in various locations (cf. SI I.1). The sampled combinations were all assumed to have equal 208 chances of enabling the strain to reach 30% of its maximum yield.

In addition to the geometrical and operational parameters, the uncertain vertical distance between the water source (river or well) and the PBR, and the wind-dependent convective exchange coefficient ruling the thermal exchange between the PBR and the surrounding air were included in the techno-operational uncertainty.

213 In total, 7 techno-operational parameters are therefore considered uncertain.

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#### 215 2.2.3 Geographic locations and uncertainty of production mix

Similarly to any agricultural crop for which an increase in demand in Europe will be answered by different producers in distinct locations (a mix), an increase in demand for the bioprospected compound will be met by distinct microalgae plants in Europe belonging to a "compound production mix". To anticipate the development of this compound production mix in Europe, we first assumed that the production would take place in the 10 countries which have the highest potential for microalgal biomass production as identified by Skarka<sup>49</sup>: Spain (ES), Sweden (SE), Italy (IT), Portugal (PT), United Kingdom (UK), France (FR), Greece (EL), Cyprus (CY), Ireland

223 (IE) and Germany (DE). The author identified these countries as the best combinations of 224 temperature, solar irradiance, and available land after exclusion of urban, mountainous, and 225 protected areas. To account for indeterminacy regarding the plants' locations, a grid was first 226 generated with random locations drawn every 2° of latitude in each country (28 locations in total). 227 Mono-dimensional sampling was performed in each of this location, while multi-dimensional 228 sampling only could add production mix uncertainty by sampling production mixes within the grid 229 (cf. 2.3). To do so, random combinations of locations were selected within the grid in three 230 different scenarios regarding the spread of the mix over Europe. Thus, in these scenarios, 5, 15, 231 or 25 locations out of 28 were assumed to answer to the increase in demand.

232 In consequential LCA, the identification of the marginal suppliers and their shares in the mix 233 are based on the study of market trends, that are shaped by both political and economic factors<sup>50,51</sup>. 234 In our case, due to the lack of data on market performance we used areal productivity as a proxy 235 and assumed that the plants in locations with higher areal productivities will have a higher chance 236 to be part of the compound production mix. We then assumed equal production shares within this 237 mix, once its locations have been determined: each plant produces 200 g of compound in a 238 production mix for 1 kg if the mix contains 5 plants in different locations. A country-specific 239 electricity mix was used when simulating each plant.

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#### 241 2.2.4 Independence assumptions

The model was designed so that the sampled parameters are independent. For instance, our knowledge does not indicate if a specific strain has a higher chance to grow on nitrate instead of ammonium if it has a high optimal temperature and a short cell diameter. These three parameters were therefore sampled independently. Similarly, a cell diameter does not indicate if the strain has

246 a higher chance to grow at the expected yield in a certain tube diameter, all other things being 247 considered. Indeed, we did not model direct dependence between techno-operational parameters 248 and productivity for a strain but instead treated this complexity as a part of the uncertainty (cf. 249 2.2.2) that could be reduced with more knowledge on the dependencies observed across species 250 and locations. On the contrary, resorting to a parameter defining the content of the bioprospected 251 compound in the biomass which would be independent of the biomass composition would lead to 252 unrealistic scenarios. Thus, the biomass composition was first generated by randomly sampling a 253 lipid content of the ash-free dry biomass and an ash content, from which the rest of the composition is calculated<sup>28</sup>, and the bioprospected compound constitutes a random fraction of a random 254 255 biochemical class (lipid, protein, carbohydrate).

256

### 257 2.3 Sampling strategies for uncertainty propagation and sensitivity analysis

As illustrated in Figure 1, we implemented two random sampling strategies to propagate the uncertainty and perform sensitivity analysis.

The first strategy named "mono-dimensional sampling" mixes all types of uncertainties previously described (bioprospecting, techno-operational, cf. 2.2) by applying Sobol sampling to all parameters. With this sampling strategy, we obtained for each location of the grid 114688 random combinations of the 23 biological and 7 uncertain techno-operational parameters. The global sensitivity analysis associated with mono-dimensional sampling allows ranking all parameters based on their influence on the dispersion of the impact scores in one location.

To be able to simulate the development of European production mixes producing the same strain-compound pairs in different locations, the second sampling strategy, namely "multidimensional sampling", differentiates between uncertainty types. We first generated a Sobol

269 sample to obtain a set of 5376 strain-compound pairs that could potentially display the desired 270 property (bioprospecting uncertainty). The cultivation and production of each strain-compound 271 pair was then simulated for each of the 28 locations of the grid, in 150 random PBR geometries 272 and operational setups generated with Monte Carlo sampling on the 7 uncertain techno-operational 273 parameters (techno-operational uncertainty). As the same strain-compound pairs were simulated 274 in all locations of the grid, we could generate production mixes for all pairs by composing 400 275 random combinations of locations on the grid (Production mix uncertainty). The probabilities for 276 each location to be part of the marginal mix were weighted with the areal productivities (cf. 2.2.3). 277 Unlike mono-dimensional sampling, multi-dimensional sampling allows assessing the uncertainty 278 associated with the environmental impact caused by an unknown European production mix 279 producing the same strain-compound pair, after aggregating techno-operational uncertainty (cf. 280 Figure 1). This sampling strategy also allows studying how sensitive is the impact of the entire 281 European production mix (composed of the 28 locations of the grid) to the bioprospecting 282 uncertainty, i.e. to the parameters defining the strain-compound pair produced in this mix.

The global sensitivity analyses were performed with the Sobol methods from the python Salib package<sup>52</sup>.

285



286

287 Figure 1: Visualization of the sampling strategies and the associated calculations. Acronyms:

288 S-C= Strain-Compound. Blue and orange boxes respectively indicate a calculation step and a

```
sampling step. We indicated the figures of the article under the results that they display.
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#### **3. Results and discussion**

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## 296 3.1 Uncertainty propagation with mono-dimensional sampling

297 The first notable observation is that the propagation of the uncertainty resulted in a very wide range 298 of impacts scores, with for instance the global warming impact (GW100) per kg of bioprospected 299 compound ranges from -100 to +89000 kg CO<sub>2</sub>-eq / kg compound with mono-dimensional 300 sampling (cf. SI I.3). This very large dispersion is mainly due to the uncertain content of 301 bioprospected compound in the biomass, which is part of the bioprospecting uncertainty. The 302 bioprospected compound content is a secondary parameter as it results from the interaction of four 303 uncertain primary parameters (cf. 2.2.4). The propagation of the uncertainty for these four 304 parameters resulted in a target compound content ranging from 0.001 to 0.6 g gdriedbiomass<sup>-1</sup> (cf. SI 305 II.2). The highest impact score in the range corresponds to extremely unfavorable conditions: a very low content of bioprospected compound in the biomass (0.002 g·gdriedbiomass<sup>-1</sup>), coupled with 306 307 a northern location (Sweden), a high strain-specific thermal range and a large PBR volume 308 involving low volumetric productivity and substantial PBR heating requirements. GW, Freshwater 309 Eutrophication (FE), and Water Depletion (WD) impact scores all increase when the target 310 compound content decreases because of a need to produce more biomass to provide the same 311 functional unit, but a lower target compound content induces a higher Terrestrial Ecotoxicity 312 (TET) impact on average (Figure 2). The observed overall trend in Figure 2 however hides the 313 specific trends for the three equiprobable substitution routes. Indeed, the overall trend is only due 314 to the scenario in which the coproduced biomass substitutes fish feed. Despite constituting one 315 third of all the simulations, the strong positive slope observed for this scenario outweighs the 316 slightly negative slope for the two other scenarios, which results in the trend observed in Figure 2

- 317 (cf. Figure S13 in SI I. 4.2.4.1). Fish feed substitution almost always implies negative TET scores
- 318 (positive impact on the environment), which is partly the case for animal feed and never happened
- 319 for biogas substitution. The environmental superiority of fish feed substitution over biogas
- 320 production had also been found in a previous study<sup>24</sup>.



321

Figure 2: Boxplot of the environmental scores obtained in mono-dimensional sampling. The 3 211 264 LCA scores resulting from mono-dimensional sampling are divided into 3 quantiles applying to the dispersion of the uncertain bioprospected compound content in the biomass. Boundaries of the quantiles (gbioprospected compound gdriedbiomass<sup>-1</sup>): 0-001-0.090,0.090-0.185,0.185-0.60. The dots indicate the average scores per country. The regression lines are only displayed to highlight the bioprospected compound content influence on the scores. The equations of the lines

are Score=coeff\*Q+intercept, with Q the quantile numbers 1,2, and 3. More detailed plots can be found in SI I.4.2.4.1. WD= Water Depletion, GW100=Global Warming with a 100-year time *horizon, FE=Freshwater Eutrophication, TETinf=Terrestrial Ecotoxicity.* 

### 343 3.2 Uncertainty propagation with multi-dimensional sampling



344 *3.2.1 Bioprospecting and Techno-operational uncertainties* 

345

**Figure 3:** (A)Distinction of uncertainties with strain-compound-specific ridgelines and (B) Mapping of average impacts scores per country. (A) Each horizontal line corresponds to one strain-compound pair for which the LCA scores in different locations and stochastically generated PBRs constitute the density curves. The plots only show 200 of the 5376 randomly generated strain-compound pairs in multi-dimensional sampling. The strain-compound pairs were divided into 200 quantiles regarding the dispersion of the GW impact scores and one pair was chosen per quantile and displayed on the figures. (B) The color gradients indicate the mean impact score per

353 country, considering all simulations performed in each country. The red dots indicate the 28
354 locations of the randomly generated grid. The figures for TET and FEP are shown in SI I.4.1.

In Figure 3, for each horizontal line representing a strain-compound pair, the density curves of the impact (horizontal axis in Figure 3) result from the propagation of the techno-operational uncertainty in the different countries. Despite a visible shift of the impact density curves across strain-compound pairs (vertical axis in Figure 3) and different countries (different colors in Figure 3), the significant overlap between strain-compound pairs indicates a large influence of technooperational uncertainty.

361 It must be noted for GW, FE, and WD, that the shifts of the curves along the horizontal axis are 362 not mere linear transposition of the density curves but are associated with a higher standard 363 deviation of the results for strain-compound pairs with higher average impact scores (cf. SI I.4.3). 364 In other words, the dispersion of the impact scores due to techno-operational uncertainty varies 365 with the modeled strain-compound pair. This heteroskedastic statistical behavior stems from two 366 mechanisms particularly visible for GW. First, the same techno-operational input uncertainty is 367 assumed for the cultivation simulations of all strain-compound pairs, which tends to result in a 368 constant coefficient of variation (Standard deviation/Mean) and therefore a linear increase of the 369 standard deviation across the mean impact scores of different strain-compound pairs (cf. Figure 370 S16, SI I.4.3). Second, the coefficient of variation is tendentially higher for strains with higher 371 optimum temperatures of culture (cf. Figure S15, S16, S20, SI I.4.3), which are also associated 372 with high impact scores on average. This phenomenon was observed in our previous work<sup>38</sup> and 373 stems from the fact that techno-operational uncertainty propagates more when thermoregulation 374 can become a hotspot due to the combination of strain's thermal requirements and location.

375

## 376 *3.2.2 Variability across countries and production mix uncertainty*

377 Despite techno-operational and bioprospecting uncertainties which cause substantial overlap 378 between impact density curves (Figure 3.a.), the ten countries can be ranked according to the 379 average impact score for all simulations (Figure 3.b. and also observed in Figure 2 with mono-380 dimensional sampling). Latitude is an important determinant of the environmental impacts (cf. 381 Figure S5, SI I.4.2.1.1) as it affects the horizontal irradiance and therefore influences biomass 382 productivity. Latitude also determinates the outside temperature and therefore the energy required 383 to thermoregulate the culture, in particular heating requirements, which was highlighted as an environmental hotspot in other studies<sup>38,53,54</sup>. Thus, the impact scores for all impact categories and 384 385 countries tendentially increase with the strain-specific optimal temperatures Topt (part of the 386 bioprospecting uncertainty), and the regression slopes associated with log(Impact score) = f(Topt)387 are higher for northern countries (cf. SI I.4.2.1.2). Additionally, the impact scores per FU are 388 strongly influenced by the impact profiles of the national electricity mixes (cf. SI I.4.2.2). To make 389 an example, despite of its southern location, production in Portugal is associated with the highest 390 WD impact score among all countries due to the high WD impact of the Portuguese electricity mix 391 (Figure 2,3). However, it is important to note that current marginal mixes used in consequential 392 modeling only have a limited period of validity in the near/medium term future.

393 Differences between the impact scores of distinct countries do not necessarily imply additional 394 uncertainty for the impact scores associated with the production of 1 kg of bioprospected 395 compound produced by a European production mix. Figure 4 shows how the dispersion of the 396 impact scores for the strain-compound pairs varies across the generated production mixes. The 397 differences observed between the distributions, which illustrate the production mix uncertainty,

- 398 shrink when more locations are considered per mix. Indeed, the more locations there are in the 399 randomly generated mixes, the closer the latter get to a full European mix composed of the 28
- 400 locations of the grid.



402 Figure 4: Production mix and bioprospecting uncertainties in aligned boxplots. The red area 403 consists of a succession of narrow boxplots along the horizontal axis corresponding to 400 404 randomly generated production mixes. The range of each boxplot on the vertical axis corresponds 405 to the dispersion of impact score per strain-compound pair produced in the production mix. The 406 impact score for a strain-compound i in the production mix p is calculated as follows: SMimp<sub>p,i</sub> = 407  $\frac{\sum_{L=0}^{N_p} Median([imp_{i,L,0}, ..., imp_{i,L,150}])}{N_p}$ , with Np the number of locations in the production mix p, L 408 the identifier of a location being part of p and imp\_{i,L,a} the impact score calculated for the

- 409 production of strain-compound pair i, in location L and in PBR a. The black and blue dots
- 410 respectively indicate the means and medians of each boxplot. The horizontal black and blues lines
- 411 respectively represent the mean across boxplots and the means of the medians across boxplots.



#### 412 *3.3 Sensitivity Analysis*

413

414 Figure 5: Shares of the Sobol total-order sensitivity for the uncertain parameters in multi-

415 dimensional and mono-dimensional samplings. The parameters are detailed in SI II.1. In mono-

416 dimensional sampling the share of the total-order sensitivity for a parameter x is calculated as

417  $Share_{x} = \frac{\sum_{l=0}^{28} Indice_{x,L}}{\sum_{i=0}^{p} (\frac{\sum_{l=0}^{28} Indice_{i,L}}{28})}$ , with Indice<sub>x,L</sub> the total-order sensitivity indice calculated in location L

418 for parameter x, and p the number of uncertain parameters in mono-dimensional sampling. In

419 multi-dimensional sampling, 
$$Share_x = \frac{Index2_x}{\sum_{i=0}^{p} Index2_i}$$
 with  $Indice2_x$  the total-order sensitivity index

- 420 of the biological parameter x associated with the impact for a strain-compound in Europe (cf.
- 421 Figure 1). The error bars and dispersion of the different indexes are shown in SI I.5.3.

422 Mono-dimensional and multi-dimensional samplings allow for a multifaceted understanding of the 423 model's sensitivity to the different parameters. Regarding the impact scores associated with one 424 strain-compound pair in one location, Figure 5 shows for mono-dimensional sampling that the 425 uncertainty on the fraction of bioprospected compound in the biochemical class (lipid, 426 carbohydrate, protein) dominates in the output uncertainty for all impact categories. This was 427 expected as this parameter eventually affects the overall content of bioprospected compound in the 428 biomass and therefore substantially influences the reference flows in the product system. The 429 uncertainty on the parameter assigning the substitution route also dominates the output uncertainty 430 for TET and FE, which is to be put in relation to the significantly different LCA profiles of the 431 three substitution routes (cf. 3.1 and Figure S13 in SI I.4.2.4.1, SI I.4.2.3). Eventually, the techno-432 operational uncertainty on the geometry of the PBR and the resulting culture volume (tube 433 diameter, horizontal distance between stacks, gap between tubes) accounts for 15-20% of the output variance for all impact categories. As also described in our previous work<sup>38</sup> this is mainly 434 435 due to the influence of the PBR volume on the thermoregulation requirements, together with the 436 strain's thermal requirements (Topt, Tplateau) and location. The 22 other parameters explain 437 around 60% of the output uncertainty, which makes it difficult to decide on which parameters 438 could be fixed to a unique value without losing information on the output uncertainty.

While mono-dimensional-sampling can spot uncertainty hotpots in the details of the model at the techno-operational level, only multi-dimensional-sampling can investigate the sensitivity of the model to a strain-compound pair at the European production level.

442 The latter as well shows that the fraction of bioprospected compound in the biochemical class 443 (bioact fraction molec) the assigning the substitution and parameter route 444 (random market subst) dominate the output uncertainty (75 %). The similar ranking of the 445 common parameters between mono and multi-dimensional sampling shows that the same strain-446 compound-specific parameters strongly influence the impact scores for a production both in a 447 unique location and when the same strain-compound pair is produced all over Europe.

Interestingly and in accordance with former observations (cf. 3.1), almost 100% of the output uncertainty for TETinf comes from the substitution mechanisms, determined by the biomass composition and substitution route. Thus, one could theoretically provide an educated estimate of the future TETinf impact of the production in Europe as soon as the biomass composition and the substitution market for a newly found strain-compound are known. Nevertheless, this estimate should be done keeping in mind the sensitivity of the model at the techno-operational level revealed by the mono-dimensional sampling.

### 455 3.3 Reflecting on Variability and Uncertainty

456 In this work, we have consistently used the word "uncertainty" to qualify the need to resort to a 457 stochastically sampled set of values instead of using a static set of values for the parameters of our 458 LCA model. The distinction between variability and uncertainty is key within the LCA community 459 and more generally in modeling disciplines which cannot settle for a mere deterministic assessment to support decision-making<sup>55–57</sup>. While variability is intrinsic to real-world phenomena and 460 461 processes, uncertainty is often defined as being due to a lack of knowledge about the model and its parameters<sup>56–58</sup>. This semantic overlapping between variability and uncertainty depending on 462 the formulation of the research question is well described by Frey<sup>55</sup>. Our case confirms and 463 464 illustrates how uncertainty and variability merge in some cases, depending on the research

465 question. The geographic variability becomes production mix uncertainty in our ex-ante LCA, as 466 it stems from an irreducible lack of knowledge a priori about the future development of the mix. 467 Similarly, while there is a vast diversity of microalgal strains and compounds, this biological 468 variability translates into uncertainty associated with our research question about the impacts of 469 an increase in demand for *a* microalgal compound that is currently being bioprospected in Europe. 470 The uncertainty is here similar to the one applying to the result of a random draw (aleatory 471 uncertainty) within a diverse population expressing variability (the biodiversity). Finally, the 472 techno-operational conditions should also be understood as part of the uncertainty rather than just 473 variability, as generating random PBR geometries and setups does not aim at representing alternative routes for a same compound production<sup>57</sup>, but instead represent equiprobable scenarios 474 475 for one strain to reach a specific productivity according to our limited knowledge.

476 To go further, the multi-dimensional strategy uses two independent loops as in the twodimensional Monte Carlo simulation proposed by Michiels and Geeraerd<sup>56</sup> to distinguish 477 478 variability and uncertainty. The difference is that we use this approach to distinguish 479 bioprospecting uncertainty from techno-operational and production mix uncertainty. By doing so, 480 we neglect biological variability by assuming that, once found and cultivated in a European mix, 481 a strain does not express any phenotypical or genomic plasticity and features the same biological 482 parameters' values in all locations. This is an oversimplification as the same strain producing the 483 same compound in a European mix could for instance accumulate more or less lipids for different locations due to different light regimes<sup>59</sup>. In this sense, mono-dimensional sampling, while making 484 impossible to distinguish between the types of uncertainty, also tackles a less fixist<sup>60</sup> and therefore 485 486 more realistic concept of "microalgal strain" by simulating a continuum of biological parameters 487 in a continuum of different PBRs and locations.

### 488 3.4 Limits

489 Our forecast partly relies on our choices regarding the modeling of microalgal biodiversity. Adapting the generic model from Williams and Laurens<sup>28</sup> to represent a strain could be argued to 490 491 be simplistic and not cover the immense diversity of microalgae. For instance, sinking rates can in 492 reality vary from the Stokes' law estimates we used to estimate centrifugation energy consumption<sup>61</sup>, depending on the microalgal taxa and cell shapes<sup>62</sup>. Adapting the centrifugation 493 494 technology may be necessary for some strains<sup>63</sup>. Furthermore, using ranges and distributions for 495 the biological parameters based on results obtained within the known biodiversity to simulate undiscovered strains could be a good example of survivor bias<sup>64</sup>. In fact, we believe this bias 496 497 benefits the representation of uncertainty by taking into account the demonstrated difficulty to cultivate many strains which can be observed in their environment<sup>65,66</sup>. The discovered and 498 499 upscaled strain will therefore likely be relatively similar to the ones we already know. Finally, and 500 as previously mentioned (cf. 2.2.2), the model would benefit from further refinement of the 501 interactions between techno-operational, biological and geographic variables to limit the weight 502 of unlikely combinations in the uncertainty representation.

## 503 3.5 Use of the results for decision-making

The high dispersion of the results associated with a very *large LCA space*<sup>67</sup> and the complex overlapping of the uncertainties must lead us to question the usability of the estimates for decision making. Ideally, the results should be used for planning and providing insightful indications on whether this technology will likely be beneficial and compete with alternatives. We can summarize the results by using the median impact score per kg of bioprospected compound across production mixes and strain-compound pairs: 1.5 m<sup>3</sup> for WD; 96 kg CO<sub>2</sub>-eq for GW; 0.017 kg P-eq for FE and 0.007 kg 1.4-DC-eq for TET (cf. Figure 4 and Table S3 in SI I.3 for complete statistical

511 description). These values, however, are obtained by keeping one median score per strain-512 compound pair and location, thus aggregating techno-operational uncertainty (cf. Figure 1). A first 513 comparison of magnitudes can be made with other bioactive compounds such as drugs from industrial chemistry whose impacts can range from 30 to 3000 kg CO<sub>2</sub>-eq per kg of drug<sup>68</sup>. Overall, 514 515 if a solution based on a bioprospected microalgal compound was to be compared with an 516 alternative technology for decision-making prior to technology development, the whole 517 distribution of the results should be considered and different statistical measures could be used<sup>69</sup> 518 (cf. SI I.3).

519 It must be highlighted that the results presented in this article can be understood as a null model by analogy with its use in  $ecology^{70}$ . Thus, the patterns of the model's output densities are obtained 520 521 for a set of standards assumptions associated with our current level of knowledge. Additionally, 522 the understanding of the uncertainty propagation combined with the sensitivity analysis allow 523 anticipating the shape of the densities when other assumptions are made or more knowledge is 524 gained. A key assumption supporting the *null model* is that bioengineers will find the combination 525 of photobioreactor geometry and operational setup associated with 30% of the strain-specific 526 energetic yields (cf. 2.2.1), as observed for cultivated strains by Williams and Laurens<sup>28</sup>. Running 527 the model with more pessimistic or optimistic assumptions regarding the capacity of bioengineers 528 to optimize photobioreactors for specific strains would shift the impact density curves. Another 529 assumption is that there is no restriction on the possible biochemical class of the target compound 530 (protein, lipid, carbohydrate) and content in the biomass. Finally, we do not account for market 531 mechanisms that could trim the density curves by making the worst cases economically non-viable, 532 for instance due to very high energy consumption per functional unit. This assumption can be

qualified as realistic as the context of high-value compounds does not exclude cases with high
production costs provided that the market prices of the compounds follow.

#### 535 4 Outlook

536 Through a heavy stochastic simulation of microalgae cultivations across strains, technological 537 settings, and locations, this work demonstrates the use of computational resources to investigate 538 the uncertainty associated with the future environmental impacts of a technology at very early 539 stage. The stochastic approach, coupled with an explicit classification and separation of the 540 uncertainties, allowed isolating the most important uncertain parameters but also to understand 541 how techno-operational, bioprospecting and production mix uncertainties interact with each other. 542 It is key to note that, by propagating uncertainty regarding the LCA of one bioprospected 543 microalgal compound, our approach eventually drew the LCA profile of a whole biological group 544 (microalgae) and its sector (productions of high-value microalgal compounds), in a whole market 545 (Europe). An even more accurate LCA portrait of the microalgal high-value compounds sector 546 would benefit from including background uncertainty, but also the extraction procedures which 547 highly depend on the compound and strain. Overall, the approach can be generalized to 548 technologies at a conceptual level of development for which the modelers know enough about the 549 ruling biological and physical phenomena to determine the key model variables, draw 550 dependencies, and eventually parameterize a model in which uncertainties are singled out. The 551 value of the approach is enhanced when simultaneously applied to competing alternatives so that 552 probability estimates resulting from the same method and same understanding of the uncertainties 553 can be compared.

554 Finally, an additional step towards educated decision-making process and planning would be to 555 use the model to go beyond the presented *null scenario* and propagate uncertainties using

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556	prospective databases for market information and future marginal suppliers, but also climate									
557	projections that could substantially influence the forecasts.									
558										
559										
560	ASSOCIATED CONTENT									
561	"Supporting Information I" (docx): Product system, description of additional calculations to the									
562	model, additional figures, elementary composition of microalgal molecules' biochemical classes.									
563	" Supporting Information II" (xlsx):									
564 565 566 567 568 569 570	<ul> <li>Sheet "SI II.1": Table of parameters.</li> <li>Sheet "SI II.2": Statistical description of all model's outputs in mono-dimensional sampling.</li> <li>Sheet "SI II.3": Correspondence between foreground activities and background activities.</li> </ul>									
571	The code of the model allowing reproduction of the figures is available at									
572	https://github.com/PJGilmw/Bioprospected_LCA <sup>40</sup> .									
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