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2 Article

3 Metabolic Pathway Analysis for Nutrient Removal of 4 the Consortium between *C. vulgaris* and *P.* 5 *aeruginosa*

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18 **Abstract:** Anthropogenic activities have increased the amount of urban wastewater discharged into
19 natural aquatic reservoirs confining in them a high amount of nutrients and organics contaminants.
20 Several studies have reported that an alternative to reduce those contaminants is using consortiums
21 of microalgae and endogenous bacteria. In this research, a genome-scale biochemical reaction
22 network is reconstructed for the co-culture between the microalga *Chlorella vulgaris* and the bacterium
23 *Pseudomonas aeruginosa*. Metabolic Pathway Analysis (MPA), is applied to understand the metabolic
24 capabilities of the co-culture and to elucidate the best conditions in removing nutrients such
25 as Phosphorus (inorganic phosphorous and phosphate) and Nitrogen (nitrates and ammonia).
26 Theoretical yields for Phosphorus removal under photoheterotrophic conditions are calculated,
27 determining their values as 0.042 mmol of PO₄/ g DW of *C. vulgaris*, 19.53 mmol of inorganic
28 Phosphorus /g DW of *C. vulgaris* and 4.90 mmol of inorganic Phosphorus/ g DW of *P. aeruginosa*.
29 Similarly, according to the genome-scale biochemical reaction network the theoretical yields for
30 Nitrogen removal are 10.3 mmol of NH₃/g DW of *P. aeruginosa* and 7.19 mmol of NO₃ /g DW of
31 *C. vulgaris*. Thus, this research proves the metabolic capacity of these microorganisms in removing
32 nutrients and their theoretical yields are calculated.

33 **Keywords:** Extreme Pathways; Nutrients Removal; *C. vulgaris*; *P. aeruginosa*

34 1. Introduction

35 Diverse human activities have increased the amount of urban wastewater effluents discharged
36 into natural aquatic reservoirs, confining in them a high amount of nutrients and organics contaminants
37 such as NH₄⁺, NO₃⁻ and PO₄³⁻. These compounds have been identified as the main cause leading
38 to eutrophication in natural aquatic reservoirs. Therefore, finding new strategies for secondary
39 wastewater treatments have received an important attention to decrease the amount of these
40 compounds before being discharged into the water bodies [1].

41 Microalga offers a promising approach to remove and to re-use nutrients such as Nitrogen (N) and
42 Phosphorus (P), because they can be assimilated into its biomass [2]. It has been reported that the
43 microalga *Chlorella* accumulates concentrations ranged between 5.0 to 10.1 % for N and between

44 0.5 to 1.3 % for P [2]. The advantages of using microalgae for this purpose include the low cost
45 for the growing process by using solar energy; the metabolic capability of microalgae which can
46 use endogenous Carbon sources and, the possibility of recycling those assimilated nutrients as a
47 fertilizer, avoiding a sludge handling problem [3]. In addition to the wastewater effluent treatments,
48 microalgae can be also used for biodiesel production [4] and even as a food source [5]. Thus, making
49 the secondary wastewater treatment more affordable and sustainable [6,7].
50 Nevertheless, a pure culture of microalgae is not always maintained. Microalgae always coexist with
51 endogenous bacteria which are able to thriving in natural aquatic systems [8]. Hence, it is natural that
52 some simultaneous interactions must exist between these microorganisms; on one hand, bacteria are
53 benefited from the exudates of microalgae, like oxygen and starch and, on the other hand, the growth
54 of microalgae is promoted by bacterial products such as Carbon dioxide (CO₂), inorganic substances
55 and some growth factors [9,10]. Therefore, natural interactions between microalgae and bacteria are
56 considered as an innovative technology to improve the wastewater nutrient removal.
57 There are some experimental studies which have been working with different species of microalgae
58 and bacteria for urban and industrial wastewater treatments [3,11]. The consortium of *C. vulgaris* and
59 *P. putida*, has demonstrated a good simultaneous nutrients removal (ammonium and phosphate) and
60 organic contaminants in synthetic municipal wastewater, compared with the axenic cultures [12,13].
61 Lananan, et al. 2014 [14], reported a removal up to 99.15 % of the total Phosphorus concentration in
62 domestic wastewater treatment, using the co-culture of *Chlorella* with an effective microorganism
63 (EM-1). Moreover, it also has been reported the co-culture of *C. vulgaris* with *Azospirillum brasilense* in
64 cellular immobilization increases the ammonia and the Phosphorus removal [9]. Again, the co-culture
65 of *Chlorella* with other bacteria removed up to 80 % of total N presents in animal feed wastewater
66 production. However, studies with pure cultures have not effect on Nitrogen or Phosphorus removal
67 in industrial wastewater [15]. The above studies prove that the consortium of microalgae-bacteria is a
68 better biological system to remove nutrients than pure cultures of these microorganisms. *Pseudomonas*
69 is a common bacteria present in wastewater and mentioned in many studies [16,17]. However the
70 metabolic activity and capability of these microorganisms can be altered by varying the culture
71 conditions in the wastewater processes, including those associated with the microflora, particularly
72 the α -Proteobacteria group, as *Pseudomonas*. To the knowledge of the authors, there is a lack of
73 studies regarding the interaction between these microorganisms - [*C. Vulgaris* - *P. aeruginosa*]-, their
74 metabolisms, the upper and lower bounds for nutrients removal according to their biochemical
75 network and the possible metabolic phenotypes.

76
77 Currently, most of the genomic information from one specific microorganism is available from
78 biological databases which is collect from high-throughput technologies, describing the metabolisms
79 and components such as genes, proteins and metabolites. From those databases, it is possible to
80 reconstruct genome-scale biochemical reaction networks for microorganisms and then analyze them
81 using metabolic engineering tools [18]. Varma *et al.* (1993) [19], were the first authors in modeling a
82 metabolic network from an entire organism (*E. coli*), obtaining the optimal carbon flux distribution
83 using Flux Balance Analysis (FBA). Metabolic Pathway Analysis (MPA), is another technique used to
84 analyze genome-scale metabolic networks, to find their phenotypic capabilities calculating a set of
85 systematically independent and unique Extreme Pathways (ExPas), [20]. Extreme pathways (ExPas)
86 are mathematically derived vectors that can be used to characterize the phenotypic potential of a
87 defined metabolic network [20,21]. ExPas describe the conversion of substrates into products, while
88 creating all byproducts needed to maintain the systemic elemental balance and the cofactor pools at
89 steady state [18,20]. By calculating the ExPas from a metabolic network, it is possible to explain the
90 active metabolisms in a particular pathway and the theoretical yields of products with respect to the
91 sources of carbon or nutrients. Thus, calculating and analyzing ExPas from the metabolic network
92 of the consortium between *C. vulgaris* and *P. aeruginosa*, it is possible to estimate their phenotypic
93 potential under different schemes. Our research group has been working in the evaluation of nutrients

94 removal by using different microorganisms and the consortium between them [13]. Consequently, this
 95 project provides a fundamental approach to enhance our understanding of biological system where
 96 microalgae and bacteria coexist as it occurs in most wastewater treatment.

97

98 2. Results

99 2.1. Stoichiometric matrix S

100 The stoichiometric matrix S has a dimension of 286×293 , representing the metabolites and the
 101 set of internal fluxes and exchange fluxes, such as photons (Pho), external glucose (GLUext), sulfate
 102 (SO_4ext), Magnesium (Mg), Potassium (K), iron (Fe), Calcium (Ca), Zinc (Zn), Copper (Cu), Manganese
 103 (Mn) and more important the nutrients which are studied in this article (PO_4ext , Piext , NO_3ext and
 104 NH_3ext). The outputs fluxes were biomass from each microorganism, polyhydroxyalkanoates, maltose,
 105 Carbon dioxide (COext) and oxygen production (O_2ext).

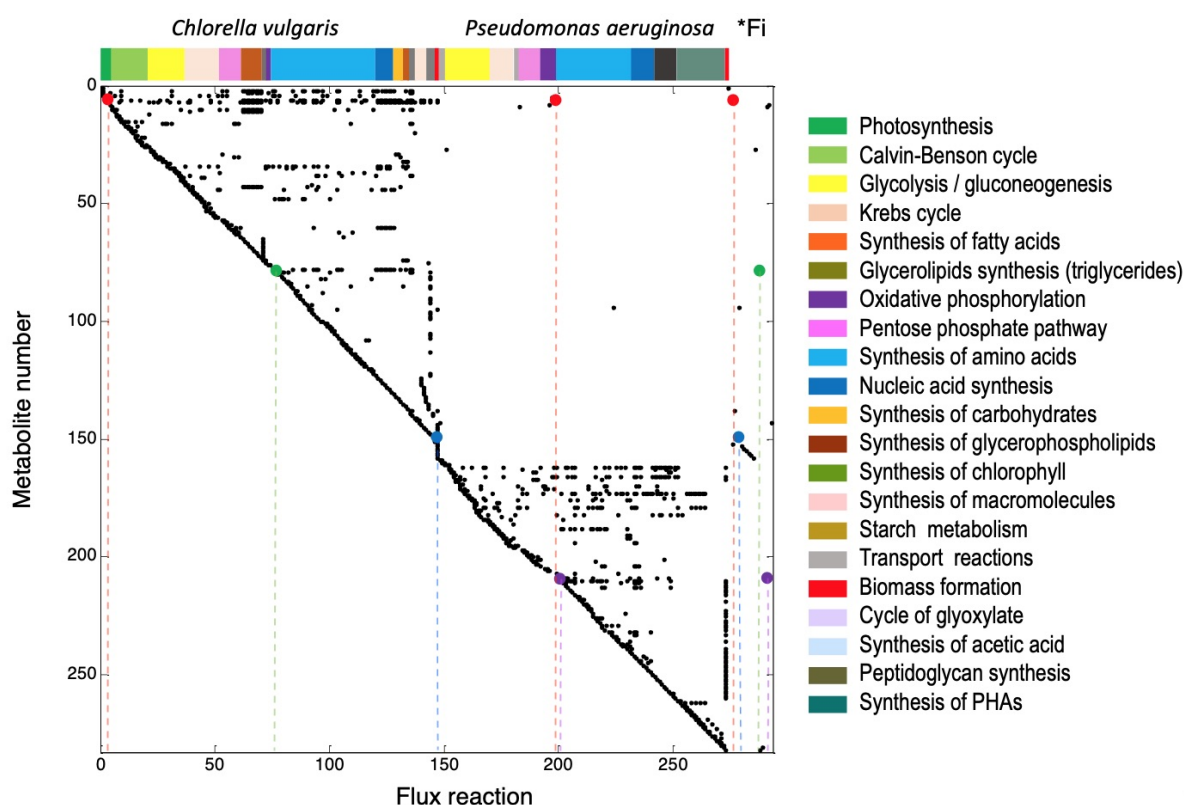


Figure 1. Estequiometric matrix S , of co-culture of *C. vulgaris* and *P. aeruginosa*. Main metabolisms of microorganisms are show at the top of the figure; *Fi represents the exchange fluxes considered in the metabolic network -biochemical reactions from 273-293, see Appendix A- Piext is the inorganic phosphorous (●), phosphate, PO_4ext (●); nitrate, NO_3ext (●) and ammonia, NH_3ext (●).

106 The Figure 1 represents a novel way to describe graphically the obtained matrix S from the
 107 reconstructed metabolic model at genomic scale of this particular consortium of microalgae-bacteria.
 108 The abscissas axis represents the internal and exchange fluxes and the ordinates axis denotes the
 109 metabolites in order of appearance in the stoichiometric model. It is also represented in Figure 1 the
 110 different metabolism for each microorganisms. Hence, in Figure 1 can be noticed in which biochemical
 111 fluxes, the external nutrients are incorporated and, also it is possible to relate those fluxes with a
 112 metabolism belonging to a particular microorganism.

113 2.2. Extreme pathways analysis for phosphorous species removal

114 Phosphorous nutrient is presented in two forms; inorganic phosphorous (Pi_{ext}) and phosphate
115 (PO_{4ext}). The inorganic phosphorous (red circle in Figure 1, (●)) is related with both microorganisms.

116 First, it could enter as an external flux into endogenous inorganic Phosphorus (Pi_{Cv}) as part of
117 the requirements for photosynthesis metabolism in the microalgae; subsequently, Pi_{Cv} takes part
118 of other fifty-two biochemical reactions which are mainly related with glycolysis, Krebs cycle and
119 oxidative phosphorylation. Therefore, Pi_{Cv} is one of the metabolites that shows a greater connectivity
120 between the fluxes in the matrix, because it is mostly required by microalga as part of its anabolism as it
121 can observed in the Figure 2. For instance, the flux number 2, in red circle (Pi_{ext}→Pi_{Cv}), is activated
122 in 2844 (100 %) of the ExPas obtained. However, only 2572 ExPas correspond to an assimilation by
123 microalga towards biomass generation. The rest, 273 ExPas are related with maltose production which
124 is an endogenous organic compound destined for the growth of the *P. aeruginosa*. These last ExPas could
125 suggest a commensalism interaction where microalgae metabolic machinery, works for bacteria supply
126 Carbon in a photoautotrophic scheme. Until now, the active fluxes and their belonged metabolisms
127 are elucidated. However, it is also possible to find theoretical yields and having a quantitative result
128 like inorganic phosphorous removal with respect to biomass of each microorganism. For instance, the
129 highest yield was 180.23 mmol Pi_{Cv}/ g DW of *C. vulgaris*, and it was discerned from 2572 calculated
130 yields. The major ExPa removal corresponds with a pure culture of *C. vulgaris* in a photoheterotrophic
131 scheme and it needs a requirement of 50.79 mmol of glucose/ g DW of *C. vulgaris* and 1252.06 mmol of
132 photons/ g DW of *C. vulgaris* to be held.

133 Second, in the metabolism of *P. aeruginosa* (from reaction 148 to 273), the Pi_{ext} (●) enter as part
134 of the oxidative phosphorylation for ATP synthesis and after it is incorporated in twenty-eight
135 biochemical reactions belonging to the metabolisms of glycolysis, Krebs cycle and synthesis of acetic
136 acid (Figure 1). In 33 % of the total calculated ExPas, it was found the maximum theoretical inorganic
137 phosphorous removal by the bacteria which was 4.90 mmol Pi_{Pa}/ g DW of *P. aeruginosa*. Even so,
138 this maximum yield by the bacteria can occur either in a photoautotrophic or photoheterotrophic
139 scheme of the consortium, its value is less than the one obtained by the microalgae.

140
141 The phosphate specie (PO_{4ext}) which is in green circle in Figure 1 (●), is assimilated by *C. vulgaris*
142 (PO_{4ext}→PO_{4Cv}) as a part of the substrate for biomass synthesis (flux number 147), meaning a
143 removal of this nutrient in 90.43% of the ExPas, when there is microalgae biomass production 59
144 PO_{4ext} m mol / g DW of *C. vulgaris* (Figure 2).

145 2.3. Extreme pathways analysis for Nitrogen species removal

146 Nitrogen nutrient is represented in two nitrogenous species in the metabolic network, as nitrate
147 (NO_{3ext}) and ammonia (NH_{3ext}). The Nitrate must be first converted into endogenous ammonia
148 (NH_{3Cv}), in order to be used by the microalgae. The anterior condition is contemplated in flux
149 number 75 (●) and it is active in the 90.43 % of the total ExPas (Figure 2) having a maximum removal
150 of 7.19 mmol of NO₃/g DW of *C. vulgaris*. NH_{3Cv} can be assimilated and incorporated into
151 other twenty-two fluxes inside of the metabolisms of *C. vulgaris*. These fluxes go to the amino acids
152 synthesis (glutamate, glutamine, glycine, proline, arginine, histidine, isoleucine, leucine, methionine,
153 phenylalanine, chorismate and valine), which are part of the protein synthesis (number flux reaction
154 148) and nucleic acids (number flux reactions 140 and 141).

155 On the other hand, the removal of external ammonia (NH_{3ext}, (●)) by bacteria in Figure 2, can be seen
156 in purple circle (●) in the number flux reaction 200, which represents the 33.33 % of the ExPas. The
157 anterior flux is active in the proportion as biomass formation (number flux reaction 273), and it has a
158 maximum removal of 10.30 mmol of NH₃/g DW of *P. aeruginosa* either under a photoautotrophic or
159 photoheterotrophic scheme.

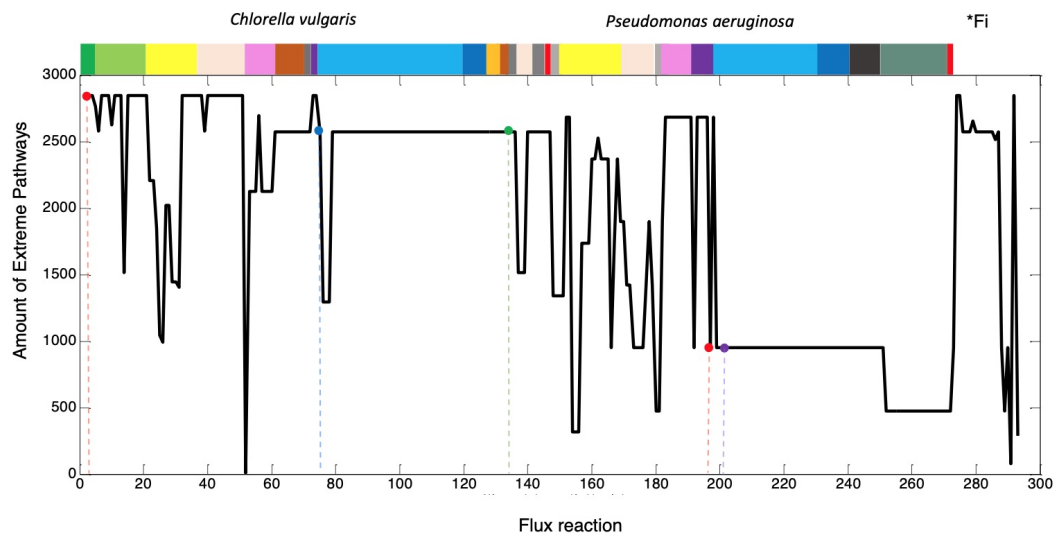


Figure 2. Participation of flux reaction in the extreme pathways. * Fi: exchange fluxes. Piext (●), PO₄ext (●), NO₃ext (●) and NH₃ext (●).

160 2.4. Analysis of the best Extreme pathways analysis for nutrients removal

161 Considering only the ExPas that showed removal for the four nutrients species, there were
 162 obtained 864 feasible ExPa, 96 were for a photoautotrophic scheme and 768 for photoheterotrophic
 163 scheme. Nevertheless, with the announced ExPas it was necessary to reduce them even more.
 164 Therefore, another parameter to consider in reducing the amount of ExPas, was the degradation of
 165 organic carbon presented in wastewater, in this case was represented as an external glucose (GLUext)
 166 in the model. This last criterion reduced to 336 feasible extremes pathways, considered as the best
 167 ExPas for nutrient removal. Thus, one of the best ExPa for nutrient removal by the co-culture is
 168 schematized in Figure 3; and it accounts for 246 biochemical flux reactions, 84 % of the total metabolic
 169 network. The above means that for the nutrient removal purpose, is not needed the entire metabolism
 170 machinery of this co-culture.

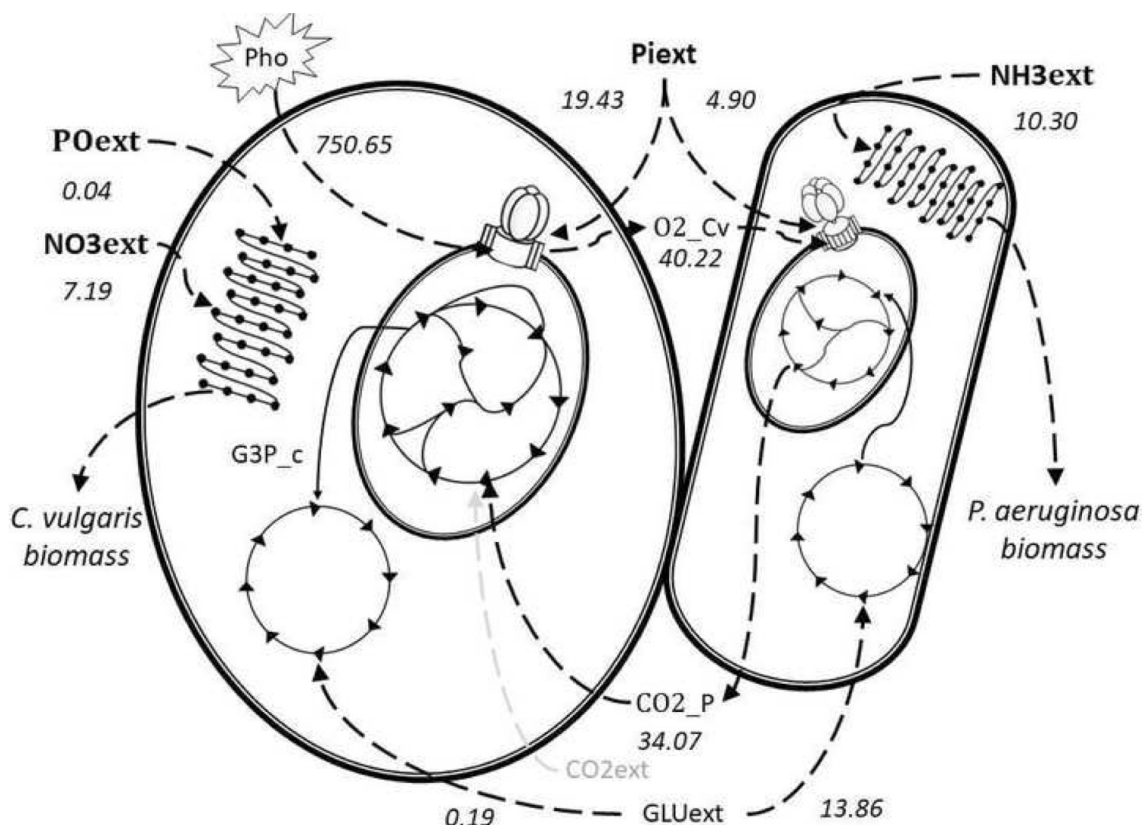


Figure 3. Schematic representation of the best ExPa for nutrients removal under a photoheterotrophic condition for the consortium of *C. vulgaris* and *P. aeruginosa*. The numbers represent the yields for consuming or producing the indicated compounds with units of mmol of nutrient / g DW of microorganism, respectively.

171 The mentioned ExPa corresponds to a photoheterotrophic scheme, where there is organic carbon
 172 source as glucose (GLUext), and the inorganic carbon source is the endogenous product of bacterial
 173 respiration (CO₂_P); this behavior could represent a decrease in operation costs due to aeration or
 174 external Carbon dioxide supply in common wastewater treatment.

175

176 The glucose entry goes for both microorganisms, having a maximum removal of 0.93 mmol of
 177 glucose/ g DW of *C. vulgaris* and 13.85 mmol of glucose/ g DW of *P. aeruginosa*. Even so, most of
 178 the organic source is directed towards the bacteria at the glycolysis metabolism. This agrees with
 179 experimental reports where the growth of a bacterium is related with glucose uptake at the expense of
 180 microalgae development [22]. While the inorganic Carbon source comes from the internal respiration
 181 of bacteria, the total yield 34.07 mmol of CO₂ goes to the Calvin cycle in the microalga metabolism to
 182 be fixed into the triose glyceraldehyde 3 phosphate (G3P_c). This last metabolite goes for chlorophyll
 183 synthesis and it is also incorporated into the five step of glycolysis. In this particular ExPa, G3P_c
 184 is not needed as substrate to produce starch or maltose. The mentioned metabolites are only used
 185 as an important energy compound when there is no carbon dioxide or nutrients. The last idea
 186 can be reinforced because there is simultaneous *C. vulgaris* and *P. aeruginosa* biomass synthesis, so
 187 either microorganism is not in competition with each other. Therefore, the best obtained yields for
 188 phosphorous and Nitrogen removals with microalgae were 19.43 mmol of inorganic phosphorous/
 189 g DW of *C. vulgaris*, 0.04 mmol of phosphate/ g DW of *C. vulgaris* and 7.19 mmol of nitrate/ g of
 190 DW of *C. vulgaris*. For bacteria nutrient removals, yields of 4.90 mmol of inorganic phosphorous/
 191 g DW of *P. aeruginosa* and 10.30 mmol of ammonia / g DW of *P. aeruginosa* were obtained. The last
 192 results indicated a more efficient removal of inorganic phosphorous from microalgae than bacteria,

193 due to its requirement during photosynthesis metabolism according to Figure 1, in fact it is important
194 to mention that all the ExPas (2844) no matter the scheme, they presented inorganic phosphorous
195 removal. On the other hand, bacteria only exhibited a phosphorous removal in 33.3 % of the total
196 ExPas.

197 3. Methods

198 3.1. Reconstruction of a genome-scale biochemical reaction networks

199 The first step to rebuild a genome-scale metabolic network for the consortium of *C. vulgaris*
200 and *P. aeruginosa* was to assemble the stoichiometric reactions base on their genome annotation.
201 Different metabolic databases exist to match the biochemical reactions with the specific genes for each
202 microorganism. In this research, the reconstruction of the metabolic networks was made manually
203 using the databases of BRENDA (BRaunschweig ENzyme DAtabase), NCBI (National Center of
204 Biotechnology Information), MetaCyc, KEGG (Kyoto Encyclopedia of referenced literature as [23,24]
205 and [5]. These references contain genomic, genetic, enzymatic, taxonomic and biochemical information,
206 available for a large number of microorganisms including *Chlorella* and *Pseudomonas* genres.

207 The considered metabolisms for microalgae were those related with the autotrophy and
208 photoheterotrophy schemes such as photosynthesis, chlorophyll synthesis (*Chla* and *Chlb*),
209 Calvin-Benson cycle, starch metabolism, glycolysis/gluconeogenesis and finally, the basic metabolism
210 for biomass formation such as TCA cycle, fatty acids synthesis, triglycerides synthesis, oxidative
211 phosphorylation, pentose phosphate pathway, protein synthesis (18 amino acids), nucleic acids
212 synthesis, carbohydrate synthesis, glycerophospholipids and maintenance. These metabolisms were
213 represented in the first 147 biochemical reactions (Appendix A).

214 Otherwise, biochemical reactions from 148 to 273 denoted the metabolisms for *P. aeruginosa*. This
215 bacterium has a huge metabolic capacity, therefore for the purpose of this project, it was considered
216 the metabolisms related with central metabolism in a prokaryotic cell such as: starch metabolism,
217 glycolysis, TCA cycle, glyoxylate cycle, pentose phosphate pathway, oxidative phosphorylation,
218 amino acid synthesis, nucleic acid synthesis, peptidoglycan synthesis, synthesis of fatty acids
219 and biomass formation. In the same way metabolisms related with synthesis of acetic acid and
220 polyhydroxyalkanoates. Moreover, it was included the transport and exchange fluxes at the end of
221 these metabolic networks.

222
223 Additionally, to the information from the databases, our research group obtained experimentally
224 the elementary composition of *P. aeruginosa* using an elemental analyzer (Fisons model 1108) [25]. The
225 results were employed to establish the biochemical reaction for the production of biomass (biochemical
226 reaction 273, Appendix A). Metabolites such as CoA, NAD, NADP, FAD, ADP, and H₂O, were omitted
227 because of they are present in the same concentration as their analogous pairs such as AcCoA, NADH,
228 NADPH, FADH, and ATP [26]. All the stoichiometric coefficients have the units of mmol unless they
229 were specified as grams. The nomenclature of compounds are given in Appendix B.

231 3.2. Extreme Pathway Analysis (ExPas)

After rebuilding the genome-scale metabolic network for the consortium, the biochemical reactions
were ordered in a matrix S with dimension $m \times n$, whose rows (m) represent the mass balance for each
metabolite (X), and n is the number of internal and exchange fluxes (v) participating in each mass
balance [21,27,28], respectively.

$$\frac{dX}{dt} = S \cdot v, \quad (1)$$

The ExPas were calculated base on the explained and analyzed principles elsewhere [20,21,27], satisfying the constraints of steady state

$$S \cdot v = 0, \quad (2)$$

the restrictions of non-negative internal fluxes -satisfying the thermodynamics of the biochemical reactions-

$$v \geq 0, \quad (3)$$

and the appropriate low and up boundaries α and β for the exchange fluxes b , such as the nutrient fluxes

$$\alpha_j \leq b \leq \beta_j. \quad (4)$$

232 An algorithm was developed in the MATLAB platform (The Mathworks, Inc., USA). From every single
233 ExPa, the theoretical yield of nutrient removal was calculated with respect to the produced biomass of
234 each microorganism. Then, a rigorous analysis was carried out to select the ExPas with the highest
235 theoretical yield for nutrient removal.

236

237 4. Conclusions

238 For our knowledge, there is not reported a metabolic genomic reconstruction of the co-culture
239 of microalgae and bacteria working together to study their interaction for nutrient removal. Thereby,
240 the contributions of this work were the reconstruction of a genome-scale metabolic network for the
241 consortium of the co-culture of the microalgae *C. vulgaris* and the symbiotic bacteria *P. aeruginosa*, and
242 also the obtaining results of theoretical yields for the maximum metabolic capacity of nutrients such as
243 Nitrogen and Phosphorus removal under a photoheterotrophic scheme, which is considered as the
244 most related with practical wastewater treatment. The last contributions were important because of
245 they could mean an improvement of biologic systems designs such as photon intensity, oxygen, and
246 dioxide Carbon requirements among others to ensure the demands of microalgae and bacteria for
247 achieving the maximal nutrients removal.

248 The in silico removal of Nitrogen and Phosphorous by this consortium were carried out only when
249 there was biomass formation for both microorganisms. Therefore, that means that the elimination of
250 these nutrients was only by assimilating them into biomass and hence, an interaction of symbiosis is
251 critical to obtain the best removal yields. The above results indicated that there was no formation of
252 others byproducts containing Nitrogen or phosphorous compounds with this particular co-culture
253 and all are directed to microalgae and bacteria growth.

254 Even with a lack of detailed experimental information, like kinetics constants and flux limitations for
255 most reactions, we have been able to show a closed behavior that could happen in a biologic system as
256 a quaternary wastewater treatment. Finally, it should be noted that the developed genomic model of
257 microalgae and bacteria is not limited to only one purpose.

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260 **Conflicts of Interest:** The authors declare NO conflict of interest with the publication of this paper

261 Abbreviations

262 The following abbreviations are used in this manuscript:

CO2-Cv	Carbon dioxide in <i>C. vulgaris</i>
CO2-P	Carbon dioxide in <i>P. aeruginosa</i>
Glu-Cv	L-Glutamate in <i>C. vulgaris</i>
Glu-P	L-Glutamate in <i>P. aeruginosa</i>
NH3ext	External ammonia
NO3ext	External nitrate
263 O2-Cv	Oxygen in <i>C. vulgaris</i>
O2-P	Oxygen in <i>P. aeruginosa</i>
Piext	External phosphorus
Pho	Photon
PO4ext	External phosphate
ExPas	Extreme Pathways

264 References

- 265 1. Ruiz-Marin, A.; G., L.; Mendoza-Espinosa.; Stephenson, T. Growth and nutrient removal in free and
266 immobilized green algae in batch and semi-continuous cultures treating real wastewater. *Bioresource*
267 *Technology* **2010**, *101*, 58–64.
- 268 2. Beuckels, A.; Smolders, E.; Muylaert, K. Nitrogen availability influences phosphorus removal in
269 microalgae-based wastewater treatment. *Water Research* **2014**, *77*, 98–106.
- 270 3. Alsan, S.; Kapdan, I.K. Batch kinetics of nitrogen and phosphorus removal from synthetic wastewater by
271 algae. *Ecological Eng.* **2006**, *28*, 64–70.
- 272 4. Mohan, S.V.; Rohit, M.; Chiranjeevi, P.; Chandra, R.; Navaneeth, B. Heterotrophic microalgae cultivation to
273 synergize biodiesel production with waste remediation: Progress and perspectives. *Bioresource Technology*
274 **2015**, *184*, 169–178.
- 275 5. Oberhardt, M.A.; Puchatka, J.; Fryer, K.E.; dos Santos, V.A.P.M.; Papin, J.A. Genome-Scale Metabolic
276 Network Analysis of the Opportunistic Pathogen *Pseudomonas aeruginosa* PAO1. *Journal of Bacteriology*
277 **2008**, *190*, 2790–2803.
- 278 6. Renuka, N.; Sood, A.; Prasanna, R.; Ahluwalia, A.S. Phycoremediation of wastewaters: a synergistic
279 approach using microalgae for bioremediation and biomass generation. *International Journal of*
280 *Environmental Science and Technology* **2015**, *12*, 1443–1460.
- 281 7. Matamoros, V.; Gutierrez, R.; Ferrer, I.; Garcia, J.; Bayona, J.M. Capability of microalgae-based wastewater
282 treatment systems to remove emerging organic contaminants: A pilot-scale study. *J. Hazardous Mat.* **2015**,
283 *288*, 34–42.
- 284 8. Kouzuma, A.; Watanabe, K. Exploring the potential of algae/bacteria interactions. *Current Opinion in*
285 *Biotechnology* **2015**, *33*, 125–129.
- 286 9. Choix, F.; de Bashan, L.; Bashan, Y. Enhanced accumulation of starch and total carbohydrates in
287 algininate-immobilized *Chlorella* spp. induced by *Azospirillum brasilense*: I. Autotrophic conditions.
288 *Enzyme and Microbial Technology* **2012**, *51*, 294–299.
- 289 10. Qu, L.; Wang, R.; Zhao, P.; Chen, R.; Zhou, W.; Tang, L.; Tang, X. Interaction between *Chlorella vulgaris*
290 and bacteria: interference and resource competition. *Acta Oceanologica Sinica* **2014**, *33*, 135–140.
- 291 11. Pires, J.C.M.; Alvim-Ferraz, M.C.M.; Martins, F.G.; Simoes, M. Wastewater treatment to enhance the
292 economic viability of microalgae culture. *Environmental Science and Pollution Research* **2013**, *20*, 5096–5105.
- 293 12. Mujtaba, G.; Rizwan, M.; Lee, K. Simultaneous removal of inorganic nutrients and organic carbon
294 by symbiotic co-culture of *Chlorella vulgaris* and *Pseudomonas putida*. *Biotechnology and Bioprocess*
295 *Engineering* **2015**, *20*, 114–122.
- 296 13. Gómez-Guzmán, A.; Jiménez-Magaña, S.; Guerra-Rentería, A.S.; Gómez-Hermosillo, C.; Parra-Rodríguez,
297 F.J.; Velázquez, S.; Aguilar-Uscanga, B.R.; Solis-Pacheco, J.; González-Reynoso., O. Evaluation of nutrients
298 removal (NO₃-N, NH₃-N and PO₄-P) with *Chlorella vulgaris*, *Pseudomonas putida*, *Bacillus cereus* and a
299 consortium of these microorganisms in the treatment of wastewater effluents. *Water Science Technology*
300 **2017**, *76*, 49–56.

- 301 14. Lananan, F.; Hamid, S.H.A.; Din, W.N.S.; Ali, N.; Khatoon, H.; Jusoh, A.; Endut, A. Symbiotic
302 bioremediation of aquaculture wastewater in reducing ammonia and phosphorus utilizing Effective
303 Microorganism (EM-1) and microalgae (Chlorella sp.). *International Biodeterioration Biodegradation* **2014**,
304 *26*, 127–134.
- 305 15. Posadas, E.; Bochon, S.; Coca, M.; Garcia-González, M.; Muñoz, P.G.E.R. Microalgae-based agro-industrial
306 wastewater treatment: a preliminary screening of biodegradability. *Journal of Applied Phycology* **2014**,
307 *26*, 2335–2345.
- 308 16. Bergero, M.F.; Lucchesi, G.I. Immobilization of *Pseudomonas putida* A (ATCC 12633) cells: A promising
309 tool for effective degradation of quaternary ammonium compounds in industrial effluents. *International*
310 *Biodeterioration Biodegradation* **2015**, pp. 38–43.
- 311 17. Cooper, M.B.; Smith, A.G. Exploring mutualistic interactions between microalgae and bacteria in the omics
312 age. *Current Opinion in Plant Biology* **2015**, *26*, 147–153.
- 313 18. Price, N.D.; Reed, J.L.; Palsson, B.O. Genome-scale models of microbial cells: evaluating the consequences
314 of constraints. *Nature Reviews Microbiology* **2004**, *2*, 886–897.
- 315 19. Varma, A.; Bosech, B.W.; Palsson, B.O. Stoichiometric Interpretation of *Escherichia coli* Glucose Catabolism
316 under Various Oxygenation Rates. *Applied and Environmental Microbiology* **1993**, pp. 2465–2473.
- 317 20. Papin, J.A.; Price, N.D.; Palsson, B.O. Extreme Pathway Lengths and Reaction Participation in
318 Genome-Scale Metabolic Networks. *Genome Research* **2002**, *12*, 1889–1900.
- 319 21. Schilling, C.H.; Letscher, D.; Palsson, B.O. Theory for the Systemic Definition of Metabolic Pathways and
320 their use in Interpreting Metabolic Function from a Pathway-Oriented Perspective. *Journal of Theoretical*
321 *Biology* **2000**, *203*, 229–248.
- 322 22. Guo, Z.; Tong, Y.W. The interactions between *Chlorella vulgaris* and algal symbiotic bacteria under
323 photoautotrophic and photoheterotrophic conditions. *Journal of Applied Phycology* **2013**, *26*, 1483–1492.
- 324 23. Allen, J.F. Photosynthesis of ATP: Electrons, Proton Pumps, Rotors, and Poise. *Cell* **2002**, *110*, 273–276.
- 325 24. Orth, J.D.; Thiele, I.; Palsson, B.O. *KEGG: Kyoto Encyclopedia of Genes and Genomes*;
326 <https://www.genome.jp/kegg/>, 2016.
- 327 25. Rojas Rosas O, González Reynoso O, N.A.J. Producción de Polihidroxicarbohidratos y análisis de la capacidad
328 metabólica de *Pseudomonas aeruginosa* ATCC 9027, mediante el estudio de la distribución de flujos de
329 carbono. PhD thesis, Universidad de Guadalajara, 2006.
- 330 26. Stephanopoulos, G.; Aristidou, A.A.; Nielsen, J. *Metabolic Engineering: Principles and Methodologies*; Elsevier,
331 1998.
- 332 27. Bell, S.L.; Palsson, B.O. Expa: a program for calculating extreme pathways in biochemical reaction networks.
333 *Bioinformatics* **2005**, *21*, 1739–1740.
- 334 28. Orth, J.D.; Thiele, I.; Palsson, B.O. What is flux balance analysis? *Nature Biotechnology* **2013**, *28*, 245–248.