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Bacterial lipoxygenases are associated with host-microbe interactions and may provide cross-kingdom host jumps

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Abstract

In this bioinformatic research, we studied the association of bacterial lipoxygenases (LOXs) with pathogenic and symbiotic traits by text networks analysis, phylogenetic analysis, and statistical analysis of molecular structure. We found that bacterial lipoxygenases are associated with a broad host range — from coral to plants and humans. In humans, bacterial LOXs are associated with opportunistic and nosocomial infections as well as with affecting specific patient populations like cystic fibrosis patients. Moreover, bacterial LOXs are associated with plant-human (or human-plant) host jumps in emerging pathogens. We also inferred a possible mechanism of such host jumps working *via* a host's oxylipin signalling "spoofing".

Keywords: lipoxygenases, oxylipins, cross-kingdom pathogens, cross-kingdom host jumps, emerging pathogens, cystic fibrosis

Running title: Lipoxygenases and host-microbe interactions

Graphical abstract
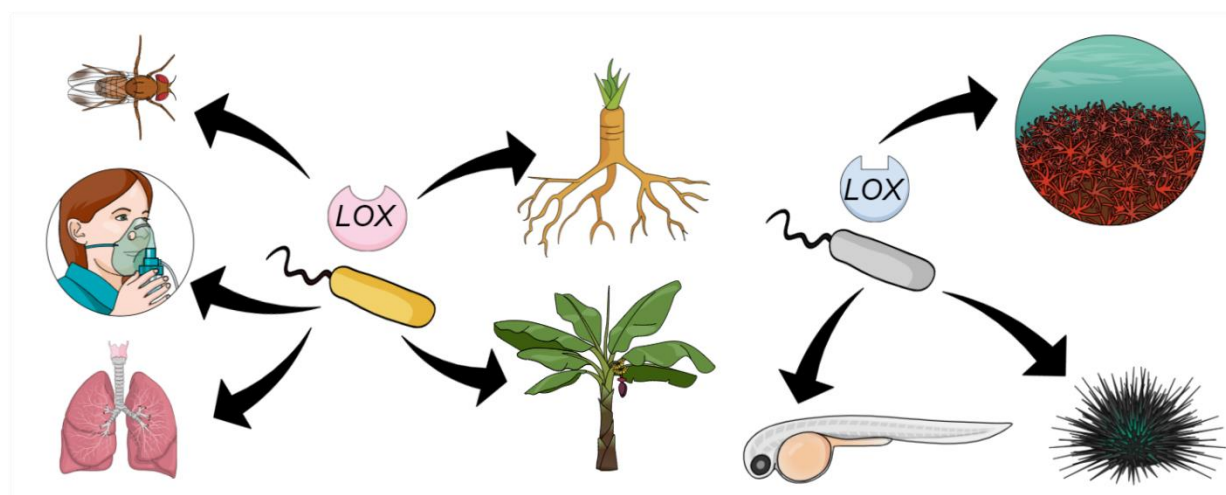
Background

Recently, the problem of host switch (also host shift) in pathogens has gained new relevance due to the growing threat posed by emerging pathogens [1]. The mechanism of host switching by the viruses, responsible for the current (coronaviruses) or possible (henipaviruses, filoviruses) pandemics, has been particularly actively studied. Host shifts typically involve two or more relatively closely related taxa with similar morphology and physiology. Coronaviruses and filoviruses pass to humans from bats, most often via intermediate hosts (palm civets, camels, primates) [2], while new human-threatening influenza viruses often originate in waterfowl [3]. In contrast, there are virtually no known examples of viruses capable of transferring from humans to plants and *vice versa*. The only possible candidate for such a virus is the hepatitis D virus [4], which is similar to plant viroids, but direct relatedness between them is controversial.

However, cross-kingdom host jump — a host switch in which the new host belongs to a different kingdom from the previous one — is common among bacteria. Most of the known cross-kingdom jumps occur exactly between plants and humans, and such bacteria have a specific clinical profile [5]. As a rule, they cause opportunistic infections (septicemia, pneumonia, surgical infections) without affecting healthy immunocompetent individuals. Less frequently, plant-animal host jumps have been recorded [6].

The mechanisms of such host jumps are intriguing. Such distant hosts have different molecular structures determining susceptibility to infection, and the biochemical basis of pathogen compatibility to different hosts is yet unclear although extensively studied. Proteases, phospholipases, toxins, and secondary metabolism regulators have been identified as virulence factors common to plant- and animal-targeted virulence. Quorum sensing systems are also discussed in this regard [5, 7].

In our recent paper [8], we had suggested the role of lipoxygenases and oxylipin signalling in cross-kingdom pathogenicity. Lipoxygenases are very conserved enzymes involved in the synthesis of oxylipins — signalling compounds regulating stress and immune responses in a wide range of multicellular eukaryotes, such as animals (including humans), plants, and different groups of algae. The main goal of our work was to trace evolutionary origins of oxylipin signalling, using lipoxygenases as a bioinformatic proxy for oxylipin biosynthetic ability. We have found that in bacteria, lipoxygenases were primarily associated with multicellularity (at least at one stage of the life cycle) which reflects the possible role of oxylipins as ancient cell-to-cell signalling compounds.

However, some bacteria from our dataset did not possess any (albeit primitive) multicellularity. For some of these bacteria, ecophysiological data was not available. Regarding the fraction with described ecophysiology, it could be divided into three groups: (1) human- or

vertebrate-associated bacteria, (2) plant-associated bacteria, (3) and bacteria associated with marine organisms. Notably, the first two groups were significantly overlapped: at least two bacteria in our dataset (*Pseudomonas aeruginosa* and *Burkholderia gladioli*) were cross-kingdom pathogens capable of infecting both plants and humans. Continued updating of our dataset resulted in adding one more species (*Pantoea ananas*) to the list of lipoxygenase-positive cross-kingdom pathogens [9]. These facts lead to an assumption that lipoxygenases may facilitate cross-kingdom host jumps and serve as versatile virulence/symbiosis factors.

In our previous paper [8], we have noticed some additional traits common for lipoxygenase-positive bacteria. They were opportunistic pathogens (these bacteria are prone to affect immunocompromised people, especially patients with cystic fibrosis). The most concerning common trait we noticed is multiple drug resistance [9].

However, these traits were noticed during the non-systematic review of literature data on bacteria in which lipoxygenases were detected. The list of these characteristic traits was not reinforced by thorough statistical analysis. The main focus of the previous papers was a link between LOXs and multicellularity, and conclusions on a link between LOXs and host-microbe interactions were preliminary.

In the current article, we continue the same research project and elaborate this association in more details. The aim of this bioinformatic research was to find additional evidence that lipoxygenases and oxylipins could contribute to host colonization and invasion, as well as to cross-kingdom host jumps.

Materials and Methods

Collecting and updating the dataset

We started with the same dataset of bacterial lipoxygenases as used in our previous research [8]. However, we are updating it in an ongoing manner because LOX-like sequences are constantly added to protein databases, and some sequences are revised and deleted.

The method of updating the dataset is the same as for collecting the initial dataset (described in our previous paper [8]). The difference was that we used pathogen and symbiont LOX sequences form the "old" dataset as queries and downloaded all hits that belong to pathogens and symbionts (they were typically sorted at the top of the hit list) along with some cyanobacterial and myxobacterial LOXs (according to our previous research, they are outgroups for all pathogen and symbiont LOXs). The methods for checking LOX-like hits were the same as described in our previous paper [8].

The resulting list of LOX-carrying bacteria was used for the estimation of a statistically reinforced ecological profile of bacterial LOXs and LOX-carrying bacteria. The criteria of inclusion into this list were:

96   1) the presence of a species in the updated LOX dataset;

97   2) availability of any ecological data for in the literature;

98   3) pathogenic/symbiotic properties or other association with any host.

99   *Ecological profile generation by network text analysis*

100   We used a combination of systematic literature review and network text analysis to

101   estimate the overall ecological profile of LOX-carrying pathogenic and symbiotic bacteria.

102   The species names from the list of LOX-carrying bacteria were used as the queries for

103   PubMed database search (https://pubmed.ncbi.nlm.nih.gov/). The first ten results for each query

104   were used for further text analysis. We used article abstracts, titles, and keywords for manual

105   extraction of terms characterizing the ecology of the respective species (referred to below as

106   "terms"). Totally, this research used an array of 130 papers for meta-analysis [10-138].

107   These terms were manually normalized to the root form and, in some cases, to the most

108   frequent form (e.g., "emergent" to "emerging", "growth promotion" to "growth-promoting"). An

109   important exclusion (that could potentially bias the statistics of the term) was any terms signing

110   antimicrobial resistance: they all were normalized as "AMR", and any terms signing multidrug

111   resistance were normalized as "MDR".

112   For each term, we counted the number of species in which it occurs to estimate the

113   abundance of each term in the collective ecological profile of the LOX-carrying bacteria. We

114   excluded the terms occurring in only one species as the least representative. Then we explored

115   associations between the terms by building a graph where each node (entity) represents a term,

116   and each edge represents a co-occurrence of two terms (represented by its nodes) in the same

117   species. The weight of each edge represents the number of species where the respective terms co-

118   occur. We classified all terms to 6 groups:

119   1) "vertebrate-related" or "human-related" terms associated with affecting humans and

120   pathogenicity to humans and/or vertebrates (i.e., "human", "lung", "abscess", cystic

121   fibrosis" etc.). Interactions with planktonic larval forms of fishes were excluded from this

122   group;

123   2) "plant-related" terms associated with plant pathogenicity or plant symbiosis (i.e., "plant",

124   "rhizosphere", "root" etc.);

125   3) "insect-related" group included only one term ("insect");

126   4) "marine-related" terms reflecting association with marine organisms;

127   5) "public health threat" group (i.e. "emerging", "AMR", "MDR", "threat" etc.);

128   6) other terms not fitting in any of the above groups.

129   This analysis was performed with the aid of Microsoft Excel 2019, Python 3, Gephi 0.9 [139],

130   and Cosmograph (https://cosmograph.app/) software.

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131 Similar ecological classification ("vertebrate-related", "plant-related" and "marine-related")

132 was applied to the bacterial species for further analysis. There was a difficulty classifying bacteria

133 which were associated with fishes. On the one hand, fishes are vertebrates, and fish-related bacteria

134 could be considered as vertebrate-related. On the other hand, they belong ecologically to aquatic

135 environment, and bacteria associated with marine fishes could be considered as marine-related.

136 Here, we used additional taxonomic data for additional clarification. If the nearest related species

137 of a bacterium affected other vertebrates (including terrestrial ones) and/or humans, this species

138 was considered as "human/vertebrate-related". If they affected marine invertebrates, this species

139 was considered as a representative of "marine-related" group.

140 *Phylogenetics and conservation analysis*

141 Phylogenetic analysis was performed with the same software and with the same protocols

142 as described in our previous paper: MAFFT online v. 7 [140], MEGA X [141], iTOL [142] for

143 phylogenetic trees and SplitsTree for phylogenetic networks [143]. Amino acid conservation

144 analysis was performed with ConSurf server [144] in ConSeq mode [145].

145 *Binding site statistical analysis*

146 For the statistical analysis of binding site structures, we used amino acid residue volumes

147 provided by Perkins (1986) [146]. Statistical analysis was performed with MS Excel 2019 and

148 PAST [147].

149 The inclusion criteria for the statistical analysis were:

150 1) a LOX is represented on any phylogenetic tree;

151 2) (preferable) the bacteria, possessing this LOX, is described in terms of their

152 ecophysiology.

153 **Results**

154 *Collecting and updating the dataset*

155 The full updated list of LOX-carrying bacteria is provided in **Table 1** with the color-coded

156 ecological functions. The list of bacterial species for the network text analysis along with

157 corresponding accession IDs of lipoxygenases and PubMed IDs (PMIDs) of articles included in

158 the analysis is provided in **Table S1**.

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Order	Species
Burkholderiales	*V. paradoxus, V. guangxiensis, V. gossypii, B. gladioli, B. singularis, B. thailandensis, B. stagnalis*
Corynebacteriales	*Nocardia seriolae, N. pseudobrasiliensis, N. brasiliensis, Mycobacteroides abscessus, Rhodococcus erythropolis, Rhodococcus sp. 66b*
Enterobacterales	*Pluralibacter gergoviae, Kosakonia sp. AG348, K. sacchari, E. hormaechei, Enterococcus faecium, Cedecea lapagei, Pantoea sp. OXWO6B1, P. ananas, Moellerella wisconsensis, Dickeya zeae*
Holosporales	*Candidatus Finniella inopinata*
Nitrospinae/ Tectomicrobia group	*Candidatus Entotheonella palauensis*
Oceanospirillales	*Gynuella sunshinyii, Endozoicomonas numazuensis*
Oligoflexales	*Pseudobacteriovorax antillogorgiicola*
Pseudomonadales	*Pseudomonas aeruginosa*
Pseudonocardiales	*Kutzneria sp. 744, Pseudonocardia acaciae, Lentzea kentuckyensis*
Vibrionales	*Vibrio penaeicida, Enterovibrio norvegicus, E. coralii, E. calviensis, E. nigricans*

Table 1. Full list of analyzed LOX-carrying species with color-coded ecological functions. Human/vertebrate-associated bacteria are depicted in **red**, plants-associated bacteria are depicted in **green**, insect-associated bacteria are depicted in **violet**, marine-associated bacteria are depicted in **blue**. If a bacterium has several host types, the corresponding colours are all represented in its names.

172   *The ecological profile reveals intriguing traits of LOX-carrying bacteria*

173   The most common terms, defined as the terms with the largest number of species in which

174   they occur (summarized in **Figure 1**, the full list provided in **Table S2**), provide some insights into

175   the most characteristic ecological traits of LOX-carrying pathogens and symbionts. The most

176   concerning fact is that the "public health threat" group is extremely prevalent among these bacteria.

177   The leader of prevalence was the "AMR" term. The analysis of the most prevalent terms of the

178   "human-related" and "plant-related" groups shed some light to some ecological details of host-

179   microbe interaction in LOX-carrying bacteria. For instance, plant-associated LOX carriers usually

180   colonize roots or are endophytic. LOX carriers pathogenic for humans are prone to affect lungs

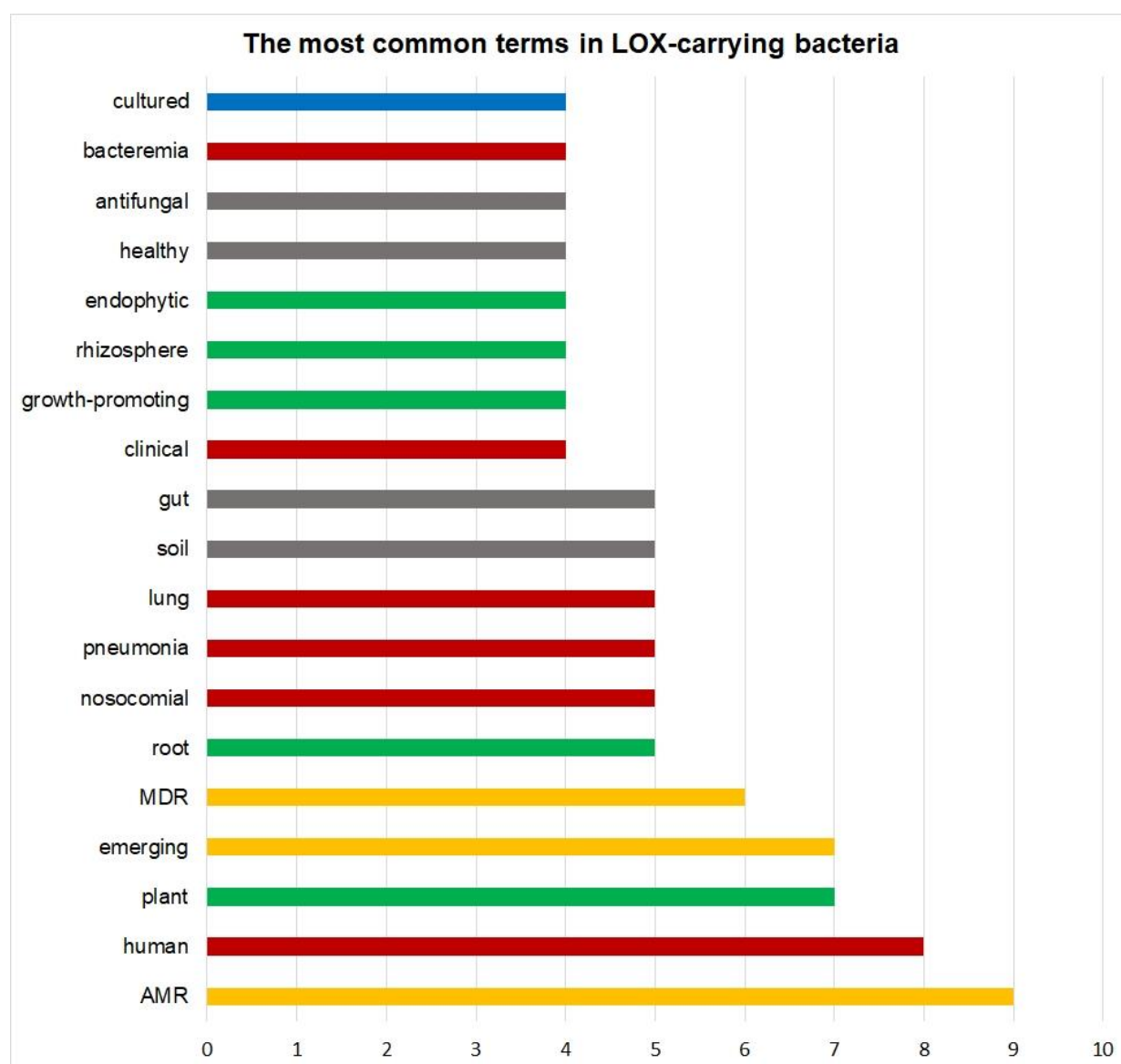181   and cause nosocomial/opportunistic infections.

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183   **Figure 1.** The diagram depicts the occurrence top terms (i.e., the number of species in which it

184   occurred). This occurrence ranged from 2 to 10. Here, the most common terms are summarized,

185   whose occurrence is not less than 4. The "human-related" and "vertebrate-related" terms are

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186   depicted in **red**, the "plant-related" terms are depicted in **green**, "public health threat" group of
187   terms is depicted in **yellow**, and the "marine-related" terms are depicted in **blue**. *Created with MS*
188   *Excel 2019*

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190   The network analysis results (**Figure 2 a, b**) show that the entire network can be divided
191   in some clusters of terms:
192   1) a "plant-related" cluster comprising "plant-related" terms;
193   2) a "human and public health threat" cluster comprising both "human related" and
194   "public health threat" term groups;
195   3) a "marine-related" cluster comprising "marine-related" terms;
196   The only "insect-related" term was tightly clustered with "human-related" terms suggesting
197   the common mechanisms of affecting insects and humans.

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199 **Figure 2a.** The full network of all terms in our dataset.

200 The nodes depict terms, the edges depict their co-occurrence, the edge widths depict their weights.

201 The "human-related" and "vertebrate-related" terms are depicted in **red**, the "plant-related" terms

202 are depicted in **green**, and the "public health threat" group of terms is depicted in **yellow**, the

203 "insect-related" terms are **magenta**, and the "marine-related" terms are **blue**. *Created with Gephi*

204 *0.9*

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208 **Figure 2b.** The "backbone" of our term network depicting only edges with the weight not less

209 than 2 and the corresponding nodes only (i.e., only the terms occurring in the same LOX-carrying

210 species not less than two times). This graph better depicts the network-forming connections and

211 hubs of the network. Notably, the yellow nodes (the "public health threat" group terms) are the

212 network-forming hubs providing connectivity to the entire networks, as well as the "human-

213 related" group terms".

The nodes depict terms, the edges depict their occurrence in the same bacterial species. The "human-related" and "vertebrate-related" terms are depicted in **red**, the "plant-related" terms are depicted in **green**, the "public health threat" term group is depicted in **yellow**, the "insect-related" terms are **magenta**, and the "marine-related" terms are **blue**. *Created with Gephi 0.9*

The "plant-related" cluster had many connections to the "human and public health threat" cluster presumably *via* the network-forming hubs. These hubs represented the same terms from the "human-related" and the "public health threat" groups that were included in **Figure 1**. Thus, the "AMR" term had direct connections to 82% of all nodes including more than 50% of "plant-related" terms. The "nosocomial" term had direct connections to 68% of all nodes including the same proportion of "plant-related" terms and is tightly clustered with the "emerging" term. This fact corresponds to the most common term prevalence and confirms the fact that the human infections caused by LOX-carrying bacteria have predominantly an "emerging" status. Moreover, the term "emerging" is a hub connected to 73% of all nodes including almost all "human-related" and "plant-related" clusters (but almost not connected to the "marine-related" cluster) and reflecting the fact that pathogenic LOX carriers are predominantly characterized in the literature as emerging pathogens.

When analyzing the more specific terms, it appeared that the "cystic fibrosis" term had connections to 55% of all nodes, including 42% of "plant-related" terms such as "endophytic" and "leaves". It is a very interesting association: the endophytic dwelling capacity in plants is associated with affecting cystic fibrosis patients in LOX-carrying bacteria. The organ-denominating term "lung" was also a hub having connections to 70% of all plant organ terms such as "leaves" and "root". So, the same bacterial LOXs are probably associated with affecting lungs in humans and affecting roots in plants.

The results of this analysis also show that the plant pathogenesis terms (such as "rot") and plant symbiosis terms (such as "growth-promoting") lie within the same "plant-related" cluster. This shows that bacterial LOXs are associated both with plant pathogenicity and plant symbionts. In contrast, we do not see such an association in the "human" cluster where all terms are pathogenesis-related.

We may also conclude that this network analysis strongly supports our previous hypothesis that cross-kingdom host jump ability is a common trait of the LOX-carrying bacteria. They are mainly represented by plant-human host jumps. Conversely, "marine-related" cluster occupied the peripheral position and had a significantly lower number of connections to other clusters indicating that plant-marine or human marine host jumps are less common. It is particularly evident when analyzing only the graph "backbone" consisting of edges with weight

249   more than or equal to 2 (**Fig. 2b**). The only "backbone" hub connecting the "marine-related"

250   cluster with all other clusters is "gut". This fact represents that both in marine animals and in

251   terrestrial animals, LOX-carrying bacteria tend to dwell in gut (intestines). It is a common trait of

252   many bacteria, and its relevance to lipoxygenases is dubious. So, "marine-related" ecological

253   functions of LOX-carrying bacteria seem to be largely isolated from other ecological functions —

254   such as colonizing plants, insects, and vertebrates.

255   The full list of term co-occurrences is provided in **Table S3**.

256   *Phylogenetic analysis results correspond with the collective ecological profile of LOX-carrying*

257   *bacteria*

258   The most important finding was that the results of bacterial LOX phylogenetic analysis

259   corresponded not with the phylogeny of the bacterial species themselves rather with their

260   ecological profile described above.

261   All LOXs analyzed in the current study can be divided into 4 clusters. Two clusters,

262   "Cluster 1" and "Cluster 2", (**Figure 3**, **Figure S1**) comprised the LOXs from phylogenetically

263   distinct species that are: (1) human/vertebrate pathogens; (2) plant pathogens; (3) both — are able

264   to affect both plants and humans. They reflected two independent series of the ancient horizontal

265   transfers of the LOX genes between plant symbionts, versatile (plant/human) pathogens (including

266   *Pseudomonas aeruginosa*), and human/vertebrate pathogens. Thus, these clusters share the

267   common ecological function pattern with the networks in the previous section where the terms for

268   plant pathogenicity, plant symbiosis, and vertebrate pathogenicity are tightly connected.
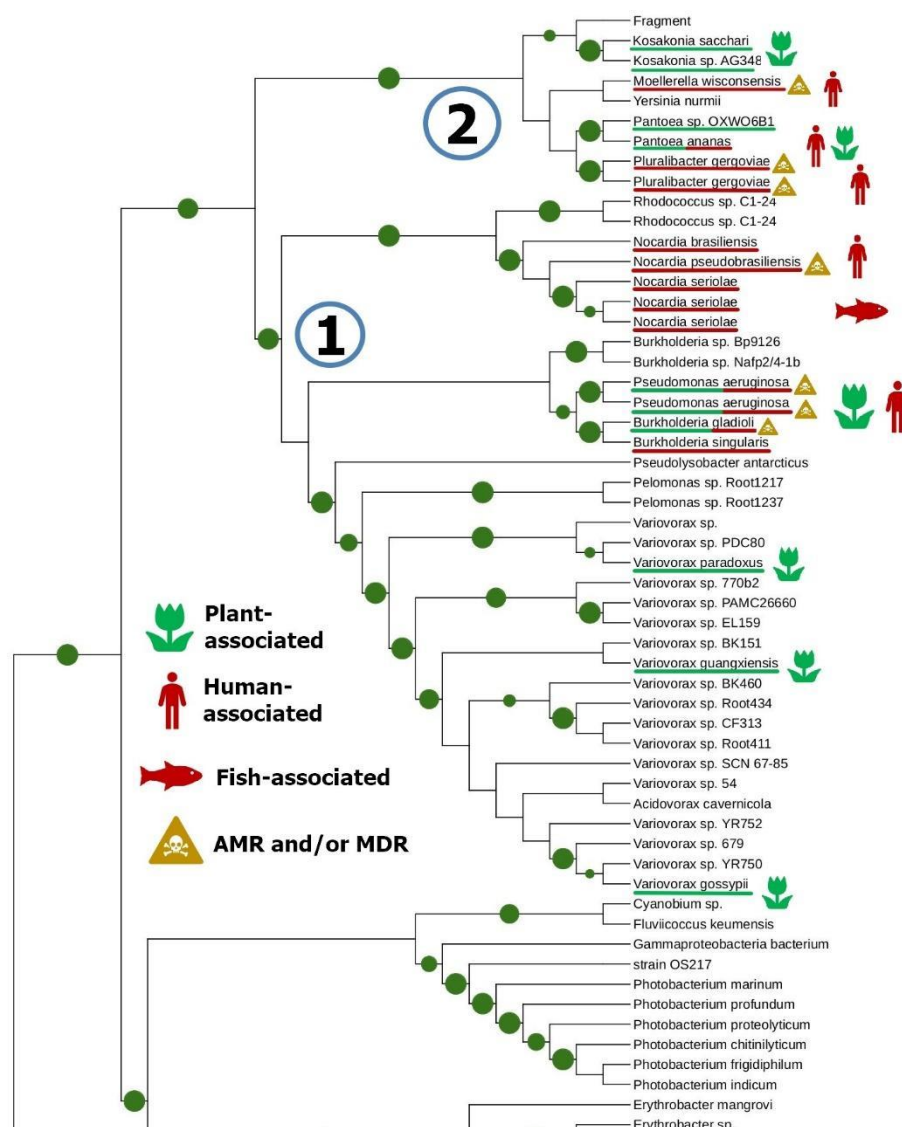
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Figure 3. Clusters 1 and 2 of bacterial LOXs reflect two independent series of horizontal gene transfers between plant symbionts, versatile pathogens, and vertebrate/human pathogens. Here, only a fragment of the phylogenetic tree is represented (the full tree is depicted in the **Figure S1**). The circle diameters depict the bootstrap values (display range of 0.5 to 1.0). *Created with iTOL*

In contrast, the LOXs of bacteria associated with marine invertebrates were predominantly separated to the distinct cluster (we call it "Cluster 4", **Figure 4**, **Figure S2**) and a distinct subcluster within the "Cluster 3" along with plant and human/vertebrate pathogens. Notably, no versatile (cross-kingdom) pathogens were included in the Cluster 3 (**Figure 5**, **Figure S3**), and it is the only cluster where the possible "terrestrial-marine" LOX gene transfer was observed. So, this cluster is significantly dissimilar from the Clusters 1 and 2, although this phylogenetic picture still corresponds to the collective ecological profile described above in terms

283   of low connectivity between "marine-related" terms and "plant-related"/"human-related" term
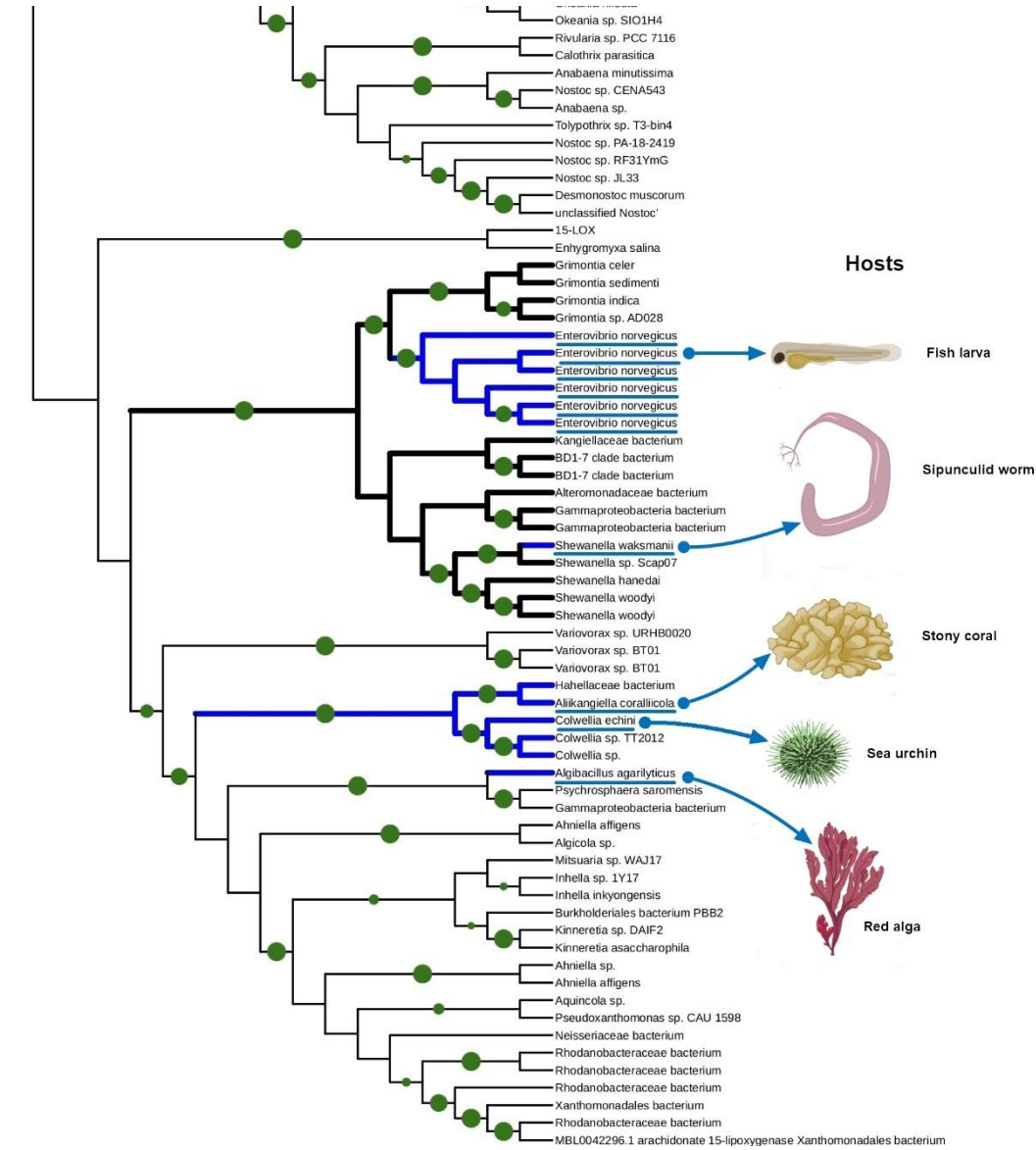
284   groups.

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287   **Figure 4.** The fragment of phylogenetic tree depicting the "Cluster 4" of bacterial LOXs. This

288   cluster encompasses bacteria isolated from different aquatic organisms: a red alga, a sea urchin, a

289   stony coral, a sipunculid worm, and a fish larva. The circle diameters depict the bootstrap values

290   (display range of 0.7 to 1.0). *Created with iTOL and BioRender.com*

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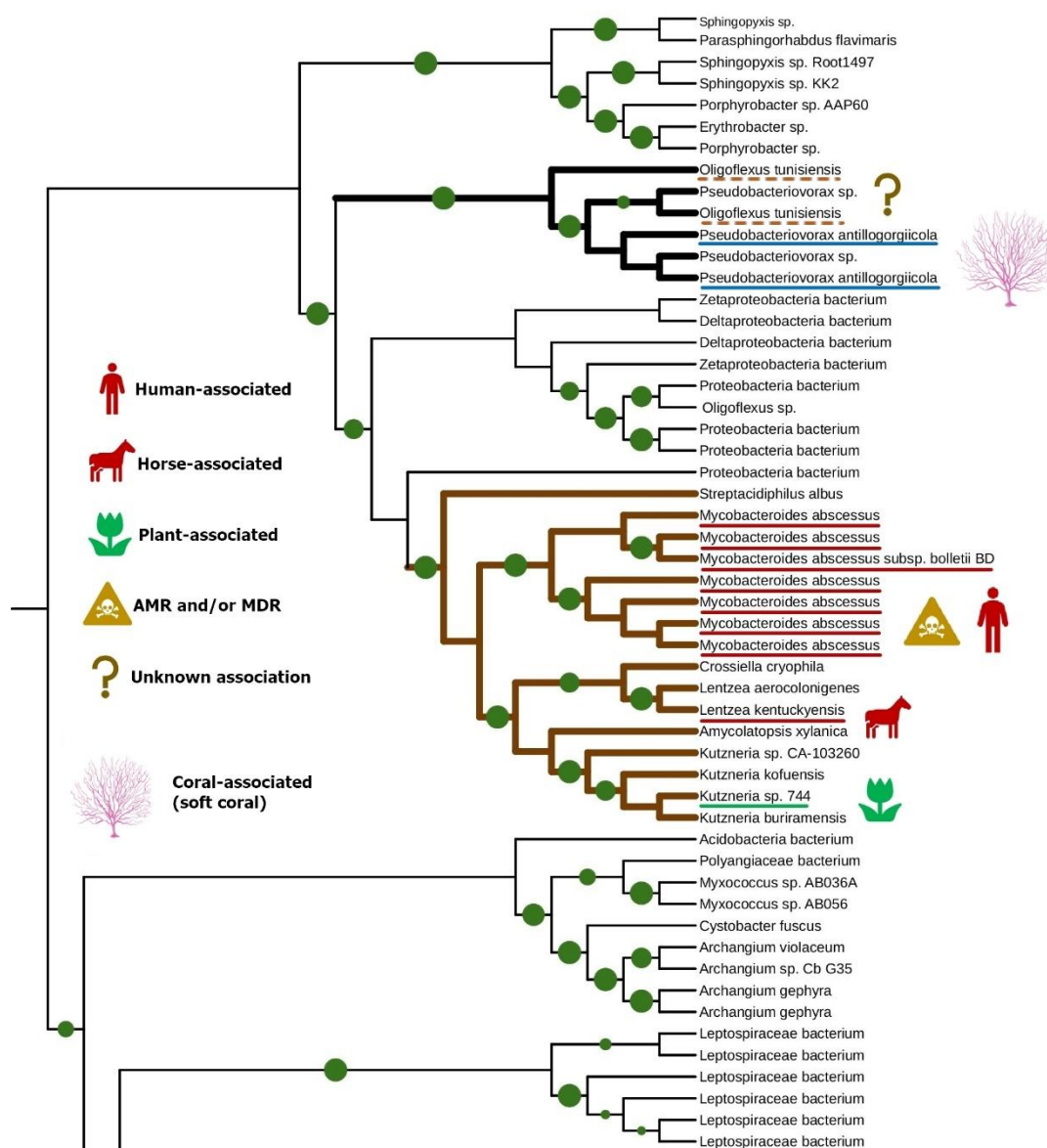
Figure 5. This fragment of a phylogenetic tree depicts the "Cluster 3" of bacteria LOXs — the only cluster, where the transfer between plant- and vertebrate associated bacteria and a coral-associated bacteria was recorded. The circle diameters depict the bootstrap values (display range of 0.7 to 1.0). *Created with iTOL and BioRender.com*

Regarding the phylogenetic relationship of these four clusters, our data are only partially conclusive due to the insufficient number of gap-free sites while analyzing the whole dataset. However, we constructed the phylogenetic network including subsets of Clusters 1, 2, and 3, which remained after applying MaxAlign procedure to make the number of gap-free-sites sufficient. This network also included outgroups, such a cyanobacteria and myxobacteria. To construct this phylogenetic network, we used the same sequence subset as for the phylogenetic tree depicting Cluster 1 and 2. It did not include Cluster 4 due to significant distance with it — it was included in another subset. However, even at the network construction stage, we had indirect evidence that

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306   the phylogenetic distance between Clusters 1, 2, and 3 is significantly less than the phylogenetic

307   distance between each of these clusters and Cluster 4.

308   The resulting network showed that Clusters 1, 2, and 3 were located closely to each other

309   and formed a kind of "supercluster" (**Fig. 6**). Within this study, this was the first evidence that

310   lipoxygenases of Cluster 3 are closely related to lipoxygenases of Clusters 1 and 2.
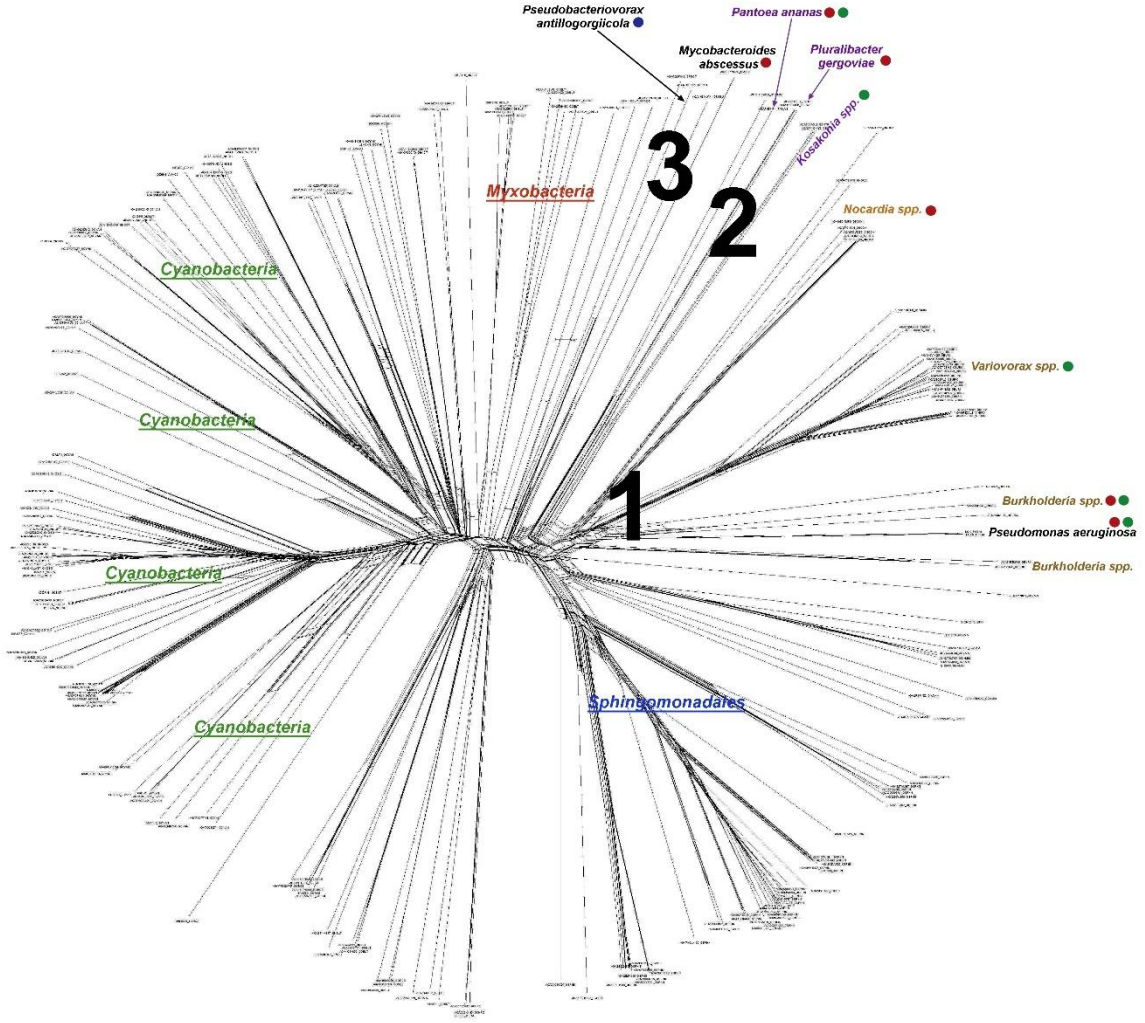

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Figure 6. SplitsTree phylogenetic network for lipoxygenases of Clusters 1, 2, and 3 shows their close location. Cyanobacteria, Myxobacteria, and Sphingomonadales were added as large outgroups. The cluster numbering and the color coding of ecological functions correspond to those in those in the Figures 3 and 5. *Created with SplitsTree.*

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317   The "supercluster" (comprising Clusters 1, 2, 3) and Cluster 4 seem to be located

318   distantly, and this result corroborates our earlier results and results of other researchers. In our

319   previous paper, this "supercluster" is also evident in the phylogenetic network of the larger dataset

320   of bacterial and eukaryotic LOXs. However, due to larger dataset and more diverse sequences,

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321 MaxAlign procedure excluded more sequences, including all Cluster 2. But Cluster 1 and Cluster

322 3 are still located closely together in the phylogenetic network in our previous paper (**Figure 7**).
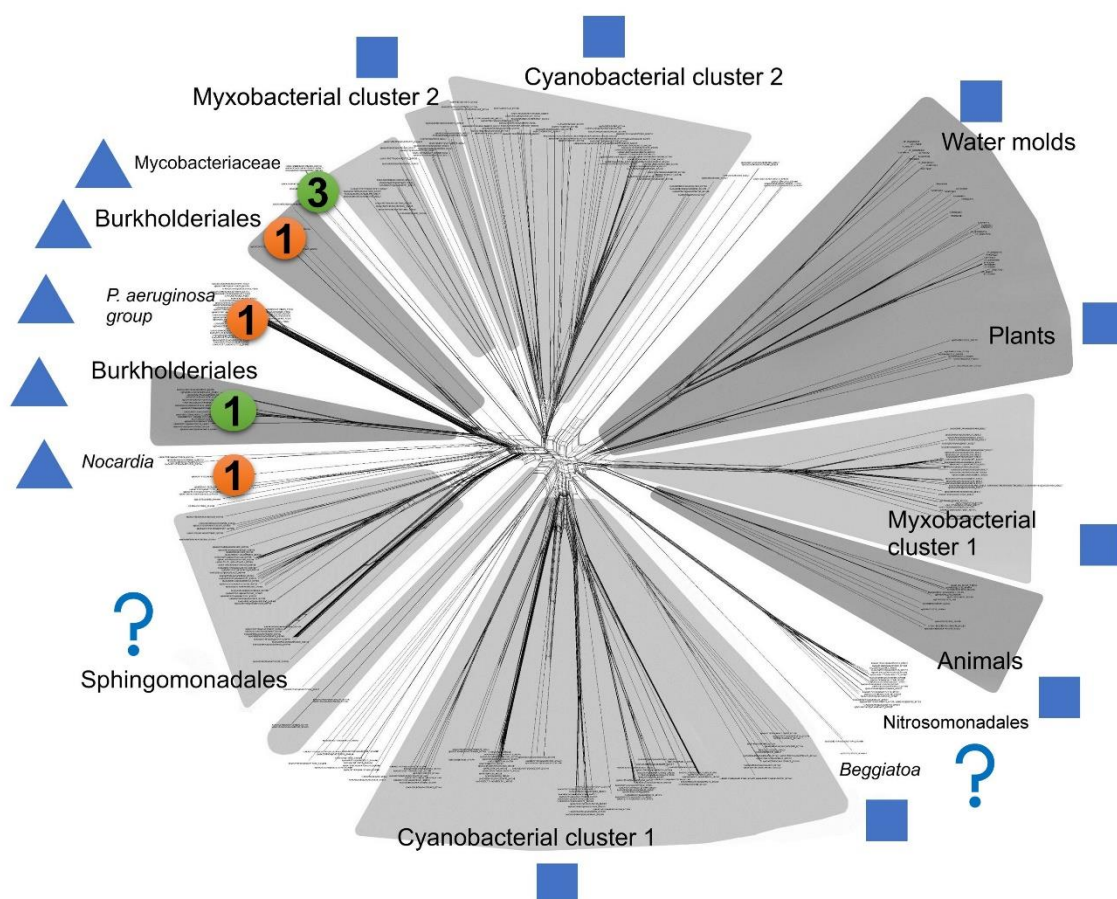
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324

325 **Figure 7.** *SplitsTree* phylogenetic network from our previous paper [8] with markers added.

326 Squares depict multicellular taxa, and triangles depict taxa involved in any kind of host-microbe

327 interaction. It is evident that the diagram is divided by these two ecological functions. However,

328 there are taxa for which multicellularity or host-microbe interactions have not been described; they

329 are marked with question mark.

330 All taxa marked with triangle cluster closely together despite they include the representatives of

331 "Cluster 1" (the number "1" in a round) and "Cluster 3" (the number "3" in a round). This

332 relationship is similar to the one presented in the Fig. 6. This suggests that the Clusters 1, 2, and 3

333 are closely related. *Image credit: Kurakin G. et al. (2020) [8], amended.*

334

335 These results corroborate the aforementioned fact that lipoxygenases of "marine-related"

336 bacteria are phylogenetically distant from their counterparts in "plant-related" and "human-

337 related" bacteria, and these later groups, in turn, are phylogenetically close to each other. Taken

338 together, these data indicate that the bacterial LOX phylogeny is associated with host-microbe

339 interactions in several independent clusters. This means that LOXs probably faced host-associated

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340   selection pressure. In turn, this means that bacterial LOXs are biochemically involved in host-

341   microbe interaction in different host types: plants, vertebrates, and invertebrates.

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Binding site analysis reveals that (ω-6)S-specificity of bacterial LOXs may contribute to plant-human host jumps

345   Our analysis also suggests that in the course of evolution, bacterial LOXs might swiftly

346   switch their function from plant-microbe interaction to human-microbe interaction and *vice versa*.

347   Moreover, such switches can ever occur in the same species without significant evolution of the

348   LOX sequence — it is the case of versatile pathogens capable of cross-kingdom host jumps.

349   Conversely, function switches between plant- or human-microbe interaction and interaction with

350   marine hosts are significantly less common. It could be explained by the environmental difference:

351   both plants and humans live in the terrestrial environment and have much more chances to contact

352   each other than to contact any organism from the aquatic environment. However, we decided to

353   check if any biochemical properties of these LOXs determine and explain their ecology.

354   We used ConSurf and additional statistical analysis to find any differences in key

355   residues determining the substrate insertion direction, stereospecificity, and regiospecificity.

356   These residues had been already characterized in papers [148, 149]. We have not found any

357   significant differences between studied bacterial lipoxygenases in the terms of insertion-

358   determining residues (probably the "tail first" insertion in all lipoxygenases) and Coffa residue

359   (**Ala** in all our LOXs, conservation score = 9, which means that all LOXs in our dataset have S-

360   stereospecificity).

361   LOX regiospecificity is determined by the bottom triad of the substrate-binding site —

362   3 amino acid residues forming the binding site bottom. We calculated the total volume of these

363   residues (as the sum of residue volumes) and performed Mann-Whitney test for three groups of

364   LOXs:

365   1) LOXs of human- and vertebrate-associated bacteria (n=12);

366   2) LOXs of plant-associated bacteria (n=10);

367   3) LOXs of marine-associated bacteria (n=6)

368   (the LOXs of cross-kingdom pathogens were included both in the "plant" and the "human-

369   vertebrate" samples).

370   There was no statistically significant difference between the "human/vertebrate group"

371   and the "plant group" (U=31). But there were statistically significant differences between each of

372   the above groups and the "marine group" (U=6 and U=2, respectively) even when checking the

373   null hypothesis against p=0.01 (it approximately corresponds to p=0.05 with Bonferroni

374   correction).

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375   Indeed, the LOXs of human-associated bacteria and the LOXs of plant-associated

376 bacteria have on average almost the same total volumes of the binding site bottom triad

377 (mean=475.24×10^{-3} nm^3, σ=30.45×10^{-3} nm^3 and mean=497.27×10^{-3} nm^3, σ=30.82×10^{-3} nm^3,

378 respectively). The bottom volumes of the LOXs of marine-associated bacteria were much lower

379 (mean=411.87×10^{-3} nm^3, σ=21.72×10^{-3} nm^3). This is graphically shown in the box plot (**Figure**

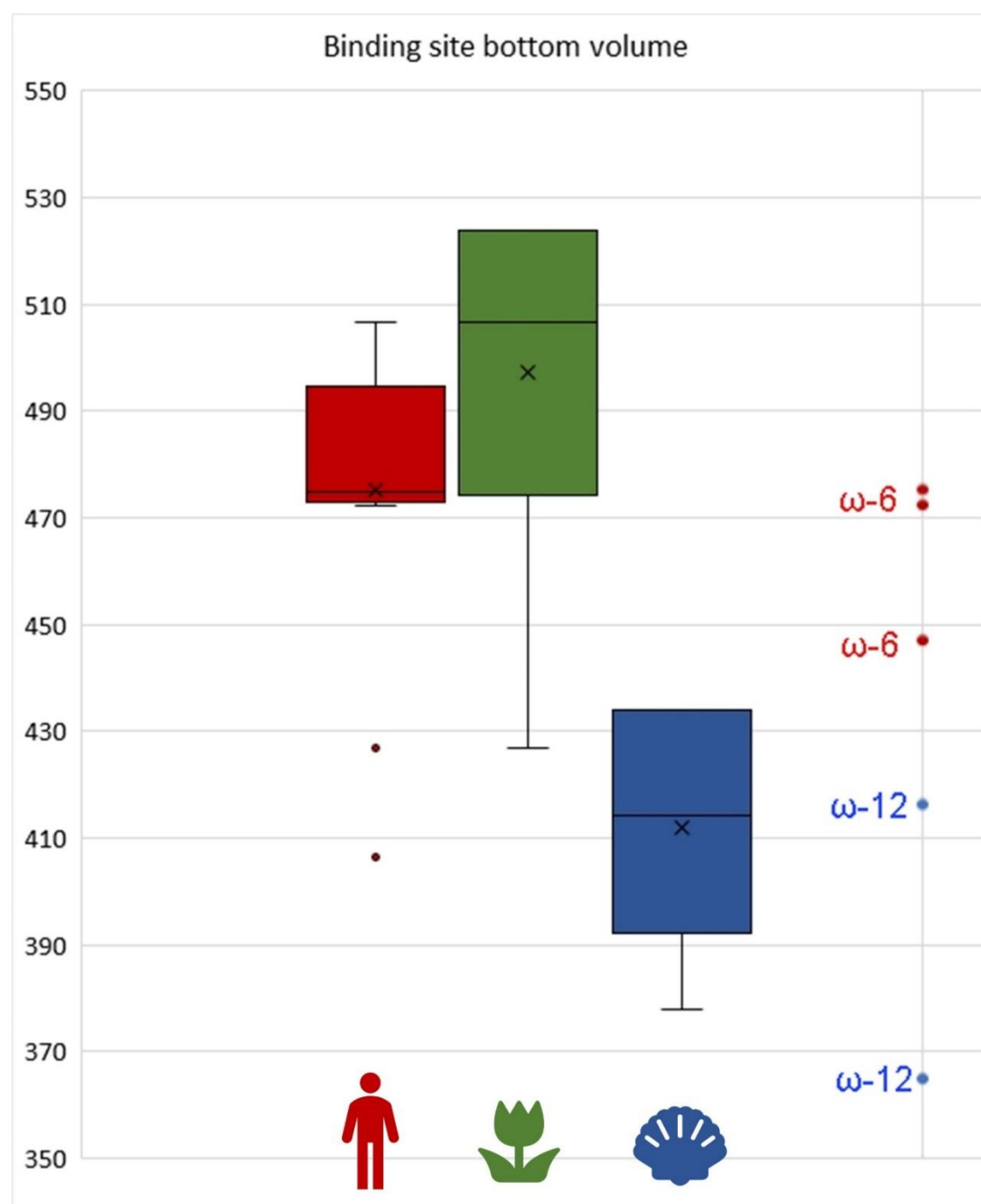380 **8**). The data for all bacteria included in this statistical analysis are provided in **Table S4**.

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383 **Figure 8.** The box plot of total LOX ligand binding site bottom volumes for "human/vertebrate-

384 associated" (**red**), "plant-associated" (**green**), and "marine-associated" (**blue**) groups. All volumes

385 are given in 10^{-3} nm^3 according to Perkins (1986) [146]. LOXs of "human/vertebrate-associated"

386 and "plant-associated" groups have total bottom volumes at almost the same level; in contrast,
387 LOXs of the "marine-associated" group have significantly lower bottom volumes ($p < 0.05$ with
388 Bonferroni correction).

389 The bar at the right shows total binding site bottom volumes of experimentally characterized LOXs
390 [149, 151-156] along with their activities. "Human-related" and "plant-related" groups correspond
391 (ω-6)-LOX activity while "marine-related" group is shifted towards (ω-12)-LOX activity.

393 The box plot (**Figure 8**) also graphically shows the presence of two outliers in the
394 "human-vertebrate" group with extremely low total bottom volume. But they fully correspond to
395 the average values for the "marine" group and lie within 1σ from its mean. These LOXs
396 phylogenetically belong to the Cluster 3 — the only cluster where the ancient terrestrial-marine
397 LOX gene transfer was recorded.

398 Compared with the binding site bottom volumes of experimentally characterized LOXs
399 [148, 150-155] (calculated by the same method), the "human-related" and "plant-related" LOX
400 groups appear to correspond to (ω-6)-LOXs. The *Pseudomonas aeruginosa* and *Burkholderia*
401 *thailandensis* LOXs (both belonging to the "human/vertebrate" group in our statistics) have total
402 binding site bottom volumes (472.2×10^{-3} nm^3 and 475×10^{-3} nm^3, respectively) near to the mean of
403 the "human-related" group. The mean of the "plant-related" group is even more. It must also
404 correspond to the (ω-6)S-LOX activity because it requires the minimally profound substrate
405 penetration into the active site (and, thus, the maximal volume of the site bottom). So, we may
406 conclude that (ω-6)S-LOX activity is required for plant-vertebrate host jumps. Conversely, the
407 "marine-related" group is significantly shifted towards (ω-12)-LOXs. It means that different
408 ecological groups of bacterial LOXs statistically correspond to different biochemical activities.

Discussion

Possible mechanism of the LOX-mediated plant-human host jumps

411 We have for the first time confirmed that pathogen and symbiont LOXs are associated
412 with a broad host range and cross-kingdom host jump ability. The correspondence between the
413 LOX phylogeny and the host-microbe interactions also confirms that these enzymes are involved
414 in pathogenesis and symbiosis and probably interact with a host.

415 One of the pathogen LOXs in our dataset – *Pseudomonas aeruginosa* LoxA — was
416 earlier experimentally characterized providing the pathophysiological insights into this interaction
417 [150, 156, 157]. This LOX is represented in our dataset, our statistical analysis sample and in the
418 Cluster 1 of the phylogenetic tree and attributed to be a LOX of a cross-kingdom pathogen because

419   of a broad host range of *Pseudomonas aeruginosa*. However, the available experimental data
420   regard only the role of *Pseudomonas aeruginosa* LOX in interactions with a human organism.

421   In addition to its direct destructive action on invaded tissues [156, 157], this LOX is
422   involved in the complicated signalling crosstalk with human leukocytes. When invading human
423   tissues, this pathogen secretes a 15S-LOX which converts the host's arachidonic acid to 15S-
424   HETE which is further metabolized by the human leukocytes to anti-inflammatory mediators such
425   as lipoxin A_4 (**Figure 9**). Their action, in turn, leads to decrease of immune cell recruiting and
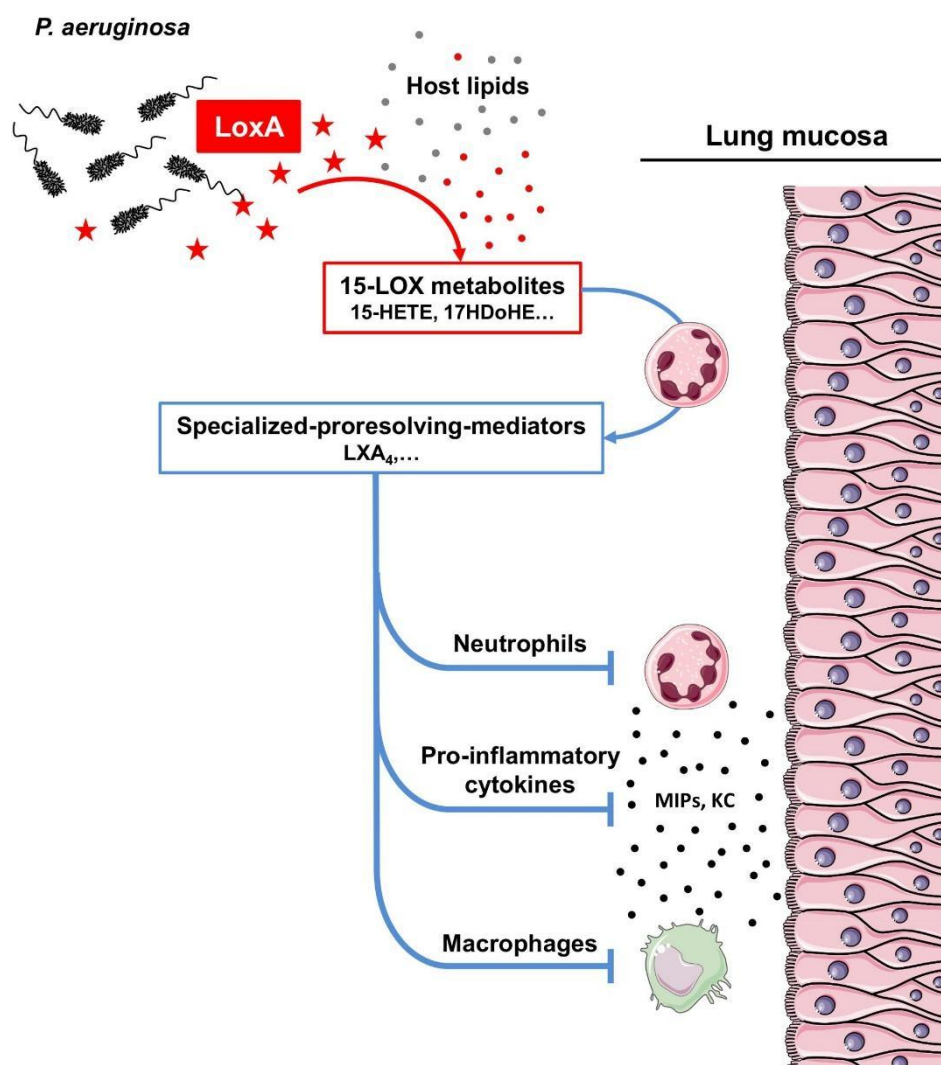426   immune response itself, and, finally, to the invasion facilitation [150].

427

428   **Figure 9.** The graphical outline of *Pseudomonas aeruginosa* LOX action to facilitate a host's
429   tissue invasion. ***Image credit:*** Morello, E. et al. (2019) *Frontiers in Microbiology* [150], CC BY
430   4.0

431

432   Host-pathogen oxylipin crosstalk is not restricted to humans and is widely exploited by
433   bacterial and fungal plant pathogens, but the exact biochemical mechanisms differ case by case

434 [158]. Our results suggest that some bacteria share similar LOXs and biochemical pathways for

435 interacting with different hosts — from plants to corals.

436 Moreover, we could now assume that both plant pathogenesis and plant symbiosis may

437 be driven by similar bacterial LOXs. This is the finding of particular interest because plant-

438 symbiont oxylipin signalling remains poorly elucidated: one of the most comprehensive reviews

439 on this topic by Beccacioli et al. [158] cites only one example of such interaction — the paper on

440 beneficial interaction between the fungus *Trichoderma virens* and maize (*Zea mays*) [159]. No

441 instances of oxylipin-mediated crosstalk between plants and their bacterial symbionts were cited.

442 Our results highlight the possible way of such oxylipin signalling.

443 Our data also show that LOX genes can make horizontal transfers thus finding new

444 ecological niche for themselves (like Richard Dawkins's "selfish genes" [160]) and enable their

445 carriers to make cross-kingdom host jumps. Such species are discussed above as versatile

446 pathogens.

447 According to our data, versatile LOX-carrying bacteria affect plants, vertebrates,

448 sometimes insects, but never affect marine invertebrates. Furthermore, LOX transfers between

449 plant-associated and vertebrate-associated pathogens are the most frequent events according to our

450 phylogeny (at least 3 independent series of plant-human transfers), while LOX transfers between

451 plant- or human-associated and marine-associated pathogens are rarer (only one even according to

452 our phylogeny). In our ecological networks, a small "marine-related" cluster is located apart from

453 the "plant-related" and "human-related" clusters. It leads to an assumption that bacterial

454 lipoxygenases can easily switch from a plant host to a vertebrate host (and *vice versa*), but a switch

455 to a marine host is very uncommon. The simplest explanation could be the ecological isolation:

456 both plants and humans live in the terrestrial environment and have much more tight contact

457 between each other than with any aquatic organism. According to this hypothesis, if humans were

458 aquatic mammals (like dolphins) and lived in a coral reef ecosystem, we could observe multiple

459 "coral-human" host jumps. This hypothesis assumes full "compatibility" of pathogen/symbiont

460 LOXs with any organism: vertebrate, insect, plant, or marine invertebrate.

461 However, we have found that LOXs themselves have structural differences in the ligand-

462 binding site which are linked to the host type. The "plant-compatible" and the "vertebrate-

463 compatible" LOXs appeared to have no statistically significant differences in the binding site

464 structure in contrast to the "marine-compatible" LOXs. It leads to an assumption that bacterial

465 LOXs face some biochemical "compatibility requirements" for successful host-microbe

466 interactions. According to this hypothesis, bacterial LOXs are capable of cross-kingdom switches

467 because plant and animal hosts require the same regio- and stereospecificity.

468   Matching statistical data on LOX binding site bottom volumes to experimental data for

469   biochemically characterized bacterial LOXs enabled us to infer the LOX specificity needed both

470   for colonizing plants and humans. It must correspond to the (ω-6)S-LOX activity.

471   In the case of human (and other vertebrates), any (ω-6)S-LOX of a pathogen is expected

472   to behave like *Pseudomonas aeruginosa* LOX: for arachidonic acid prevalent in the vertebrate

473   pool of polyunsaturated fatty acids (PUFAs), (ω-6)S-LOX activity means 15S-LOX activity,

474   which contributes to the lipoxin biosynthetic pathway and suppresses the host's immune response

475   (which is fulfilled in the case of *Pseudomonas aeruginosa*).

476   But what about plants? In contrast to humans, we have no direct experimental data on

477   the interaction of any bacterial LOX with them. However, we could biochemically infer this

478   interaction. In the case of linolenic acid (abundant in plant cells), any (ω-6)S-LOX could be

479   predicted to produce 13S-HpOT, which is a normal precursor of jasmonates. So, if (ω-6)S-LOX

480   enhances lipoxin production in the human tissues, they should enhance jasmonate production in

481   the plant body.

482   Here, the "explanative gap" is fully closed because jasmonate signalling hijacking is a

483   well-known trick of some plant pathogens. They increase a plant's susceptibility by inappropriate

484   activation of jasmonate signalling pathway. However, almost all instances of such pathogenesis

485   known up do date involve the use of toxin mimicking the natural jasmonate or acting as

486   prohormone. The best-known example is *Pseudomonas syringae* that uses coronatine activating

487   the jasmonate receptor JAZ-COI1 [161] to facilitate the invasion. A grapevine pathogen

488   *Lasiodiplodia mediterranea* facilitates its invasion by the prohormone toxin lasiojasmonate A

489   [162]. The current study provides computational evidence that bacteria might use lipoxygenase to

490   "spoof" the plant immunity with natural jasmonates rather than jasmonate-mimicking toxin. By

491   accident or by the convergent evolution, the same LOX activity appeared to be "compatible" with

492   a human lipoxin pathway which enabled LOX-carrying bacteria to be versatile pathogens (**Figure**
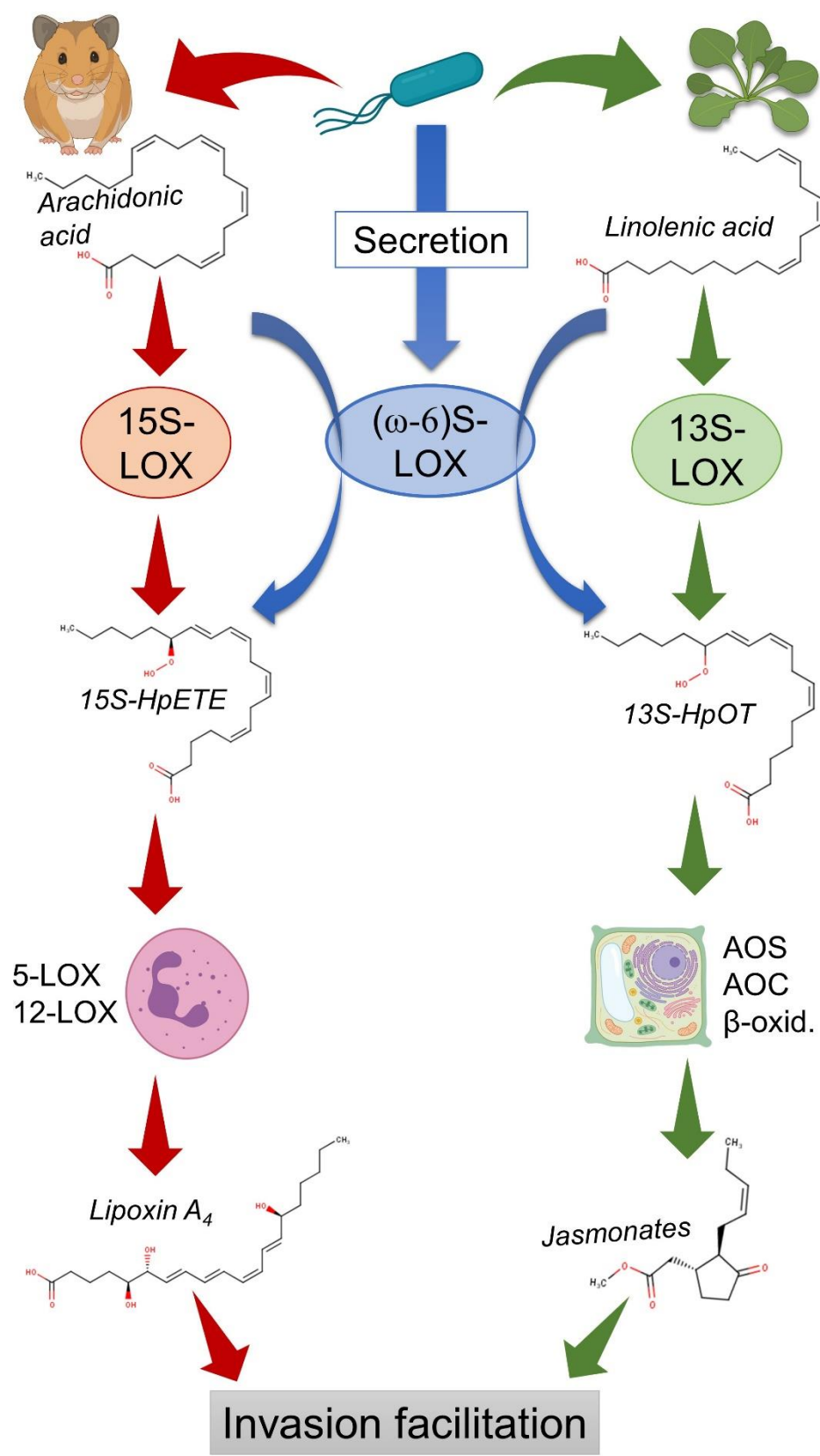
493   **10**).

494

Figure 10. Hypothetical mechanism of plant-animal host jumps provided by bacteria LOXs. In humans and vertebrates, a secreted bacterial (ω-6)S-LOX oxidizes arachidonic acid to form 15S-HpETE, which is further converted into lipoxins in leukocytes by other LOX isozymes. Lipoxins

499   suppress inflammation and immune response thus facilitating the invasion. *Created with*

500   *BioRender.com*

501

502   The immune response suppression hypothesis partially explains why this feature is

503   shared by plant pathogens and plant symbionts (as well as insect pathogens and symbionts): the

504   both groups need to evade the host's immune response to survive in its body. However, this raises

505   an additional question: why do human symbionts not use this strategy?

506   It is not the only specific thing of interaction between LOX-carrying bacteria and a

507   human (or a vertebrate) organism. As discussed above, the ability of such pathogens to affect

508   humans is extremely strongly associated with nosocomial or opportunistic traits. It means that

509   these bacteria are prone to affect immunocompromised people (the terms "compromised" and

510   "immunocompromised" are also present in the "human-related" cluster, but have relatively low

511   connectivity compared to hubs described above). Conversely, in the case of plants and insects, we

512   did not observe such associations: LOX carriers are capable of full-fledged pathogenesis and

513   symbiosis.

514   These differences are tightly connected: if a pathogen oxylipin "spoofing" is insufficient

515   to suppress the healthy human's immune response, there is no reason to expect that it will be

516   sufficient for human symbionts to evade the immune response and survive. Evidently, there is

517   something in the human (or, more broadly, vertebrate) immune system that makes it more resilient

518   to oxylipin signalling hijacks. Here, we come on shaky grounds: we have insufficient data to

519   explain what it could be. But it is reasonable to suggest that it is the presence of adaptive immunity.

520   Indeed, one of the mechanisms used by human microbiota to evade immune response is immune

521   tolerance — the acquired silencing of a host's adaptive immunity to the bacterium's antigens [163].

522   *Public health risks and molecular epidemiology surveillance*

523   First of all, we should downgrade the alert regarding possible "superspreading" of

524   lipoxygenases between nosocomial bacteria which we postulated in our previous work [8]. The

525   "*Pseudomonas aeruginosa* group" which comprised *Klebsiella pneumoniae*, *Enterobacter*

526   *cloacae*, *Acinetobacter baumannii,* and other bacteria putatively linked by fast horizontal transfers

527   of LOX genes proved to be an artifact. All LOX sequences of this group excepting *Pseudomonas*

528   *aeruginosa* sequences were subsequently removed from the databases due to probable

529   contamination. Here, we have found new evidence for the role of horizontal LOX gene transfers

530   in pathogenicity evolution, but we suggest that these processes occur on longer timescales.

531   The fact that the terms "emerging", "AMR", and "MDR" are prevalent in the collective

532   ecological profile of LOX carriers and serve as hubs in the term network representing this profile

533   should raise concerns. Moreover, antimicrobial-resistant and multi-drug resistant bacteria are

24

534   represented in each of 3 phylogenetic clusters of the pathogen and symbiont LOXs. It means that

535   bacterial LOXs are associated with emerging status, antimicrobial resistance and some other public

536   health threats. Furthermore, multiple connections between these traits and plant or human

537   pathogenicity entitle us to say that emerging pathogens use oxylipin signalling for plant-human

538   (or human-plant) host jumps.

539   But this conclusion should be considered carefully. Firstly, we have mentioned above

540   that the traits "AMR" and "MDR" could have overestimated weights due to normalization bias.

541   We are sure that this possible bias has limited effect because the used term "AMR" corresponds

542   to the real trait of studied bacteria. Moreover, the term "emerging" has also a very big weight, but

543   cannot be subjected to normalization bias. But on the visual level, it could produce a formidable

544   picture and lead to the emergence of news about "new deadly pathogens" or a "new global

545   epidemic". We caution all science journalists reading this article against doomsaying.

546   Secondly, our data provide strong evidence that LOX-carrying pathogens have low

547   epidemic and pandemic potential. As we discussed above, they are prone to affect

548   immunocompromised people or people with specific diseases (e.g., cystic fibrosis), and cannot

549   fully overcome the immune barrier of a healthy human despite multiple recorded host jump events.

550   So, we find no reason to expect a full-fledged epidemic or pandemic caused by LOX-carrying

551   pathogens.

552   However, all statistical signs of "public health threat" are connected with the extreme

553   danger they pose to specific populations — immunocompromised people or people with

554   comorbidities, especially with cystic fibrosis. Regarding these populations, we cannot rule out the

555   possibility of limited outbreaks. Sporadic cases are also dangerous for these people and could lead

556   to higher mortality and decrease in the quality of life. Thus, bacterial lipoxygenases could be a

557   useful molecular epidemiology marker for oncology, hematology, transplantology, and cystic

558   fibrosis medicine.

559   We continue updating our "pathogen blacklist" — the list of LOX-carriyng pathogens

560   that can pose danger to humans — on the grounds of publicly available data. The last update before

561   this article was published in English in the *Nature Portfolio Microbiology Community* blog [164]

562   and in Russian in *Priroda* [9]. In this work, this list is incorporated into Table **1**. But the full-

563   fledged molecular epidemiology surveillance requires much more strain-dependent data. We are

564   open for further collaboration for this purpose.

565   *Possible implications for marine biology*

566   In contrast to plants and humans, our knowledge about LOX-carrying bacteria in marine

567   organisms is limited. In most cases, we suggested the association with marine organisms only on

568   the basis of isolation of respective bacteria from a particular organism. So, the ecological functions

569   of LOX-carrying bacteria (pathogen/symbiont) are unknown for all "marine-associated" bacteria

570   in our dataset. But we can suggest that in the underwater world, more intriguing host jumps of

571   LOX-carrying bacteria occur. They involve corals, sea urchins, algae, sipunculids, and fishes. We

572   have found that these host jumps require other LOX regiospecificity, than for plants and humans.

573   We cannot infer any mechanism of their actions because of scarce data on oxylipin signalling in

574   marine invertebrates, but can suggest that bacterial LOXs could be a useful tool to study it.

575

Conclusions

577   In this article, we have proposed a mechanism of cross-kingdom host jumps by

578   interacting with evolutionary distant immune systems. The analysis of available data shows that

579   some bacteria may exploit the hosts' oxylipin signalling systems which are widespread across

580   different kingdoms. The most prevalent are plant-human host jumps because of a biochemical

581   "intercompatibility" of plant and human LOX activities needed for immune response suppression.

582   This finding could be interesting for basic scientists in a wide range of areas — from

583   immunology to marine biology. But it could also be useful for medical microbiology and public

584   health. Bacterial LOXs in a nosocomial environment could indicate an emerging pathogen which

585   could be dangerous for some vulnerable groups of patients. Bacterial LOX carriers isolated from

586   patients may require further consideration in terms of antimicrobial therapy and the particular role

587   of this bacteria in the patient's condition pathogenesis (as is the case of cystic fibrosis). It is evident

588   that pathogen and symbiont LOXs require further investigation in both basic and applied life

589   sciences.

590

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605

Ethics statement

607 This research did not involve any experimental animals or humans as a research object.

608

Conflict of interests

610 The author declares no conflict of interests. This research did not require any special funding.

611

Data availability

613 The unedited versions of phylogenetic trees and the full versions of tables are included in this
614 paper as **Supplementary Figures** and **Supplementary Tables**, respectively. Datasets,
615 spreadsheets, interactive data and any other data generated for this research are available by
616 request to the author's email.

617

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