



HAL
open science

Making Sense of the Biodiversity and Virulence of *Listeria monocytogenes*

Olivier Disson, Alexandra Moura, Marc Lecuit

► **To cite this version:**

Olivier Disson, Alexandra Moura, Marc Lecuit. Making Sense of the Biodiversity and Virulence of *Listeria monocytogenes*. Trends in Microbiology, 2021, 10.1016/j.tim.2021.01.008 . pasteur-03271346

HAL Id: pasteur-03271346

<https://pasteur.hal.science/pasteur-03271346>

Submitted on 25 Jun 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution - NonCommercial 4.0 International License

1 **Making sense of the biodiversity and virulence of *Listeria monocytogenes***

2

3 Olivier Disson^{1,2}, Alexandra Moura^{1,2,3} and Marc Lecuit^{1,2,3,4}

4

5

6 ¹ Institut Pasteur, Biology of Infection Unit, Paris-France

7 ² Inserm U1117, Paris-France

8 ³ Institut Pasteur, French National Reference Center and WHO Collaborating Center for *Listeria*,

9 Paris-France

10 ⁴ Université de Paris, Service des Maladies Infectieuses et Tropicales, Hôpital Universitaire Necker-

11 Enfants Malades, Institut Imagine, APHP, Paris-France

12

13 **Keywords:** *Listeria monocytogenes*, listeriosis, biodiversity, virulence, adaptation

14

15 **Correspondence to:**

16 Olivier Disson, olivier.disson@pasteur.fr & Marc Lecuit, marc.lecuit@pasteur.fr

17 Biology of Infection Unit, Institut Pasteur, Inserm U1117, 28 rue du Dr Roux, 75015 Paris,

18 France

19 Phone: +33 1 40 61 37 87

20 Fax: +33 1 40 61 34 21

21

22 **Abstract**

23 *Listeria monocytogenes* is a foodborne pathogen responsible for listeriosis, an infection which
24 can manifest in humans as bacteremia, meningoen­cephalitis in immunocompromised patients
25 and the elderly, and fetal-placental infection in pregnant women. Reference strains from this
26 facultative intracellular bacterium have been instrumental to investigate basic mechanisms in
27 microbiology, immunology and cell biology. The integration of bacterial population genomics
28 with environmental, epidemiological and clinical data allowed the uncovering of new factors
29 involved in the virulence of *L. monocytogenes* and its adaptation to different environments.
30 This review illustrates how these investigations have led to a better understanding of the
31 bacterium's virulence and the driving forces that shaped it.

32

33 **Highlights**

- 34 • *Listeria monocytogenes* is a foodborne opportunistic bacterial pathogen
- 35 • Classical genetic approaches in reference strains allowed identification of core
36 virulence genes
- 37 • Investigation of the species diversity of *L. monocytogenes* allowed uncovering hypo-
38 and hypervirulent strains and new virulence genes
- 39 • Hypervirulent strains are host-associated, whereas hypovirulent strains are associated
40 with food-processing environments
- 41 • Accumulation of loss-of-function mutations in virulence pathways leads to niche
42 restriction and obligate saprophytism

43

44 **Introduction**

45 *Listeria monocytogenes* is the causal agent of listeriosis, a foodborne infection associated with
46 high morbidity and mortality. The virulence of *L. monocytogenes* depends on its ability to
47 colonize the gut and disseminate in the host. This Gram-positive bacterium was first isolated
48 by G. Hülphers in Sweden, from rabbit liver necrosis [1] and from a human meningitis case in
49 1921 by J. Dumont [2]. It was then isolated and characterized by E.G.D. Murray in rabbits
50 and guinea pigs from a breeding facility in the UK in 1926, and J.H. Pirie in wild gerbils in
51 South Africa in 1927 [3, 4]. Human listeriosis is characterized by septicemia and central
52 nervous system (CNS) infections, which mostly occur in immunocompromised and/or elderly
53 individuals, and maternal-fetal infections, as well as rare anatomically localized infections [5-
54 12]. *L. monocytogenes* can also induce gastroenteritis in otherwise healthy individuals when
55 the ingested inoculum is large [13]. The foodborne origin of human listeriosis was formally
56 proven in the 1980s [14]. Although listeriosis is relatively rare compared to other foodborne
57 infections [15, 16], its high fatality rate (approximately 20% to 30%, despite effective
58 antibiotic treatment [5]) makes it a significant public health concern [17]. Recently, the
59 largest recorded outbreak occurred in South Africa, where 937 individuals were affected, with
60 a 27% mortality rate [18]. *L. monocytogenes* has also been detected in the feces of healthy
61 people, but the prevalence of *L. monocytogenes* fecal carriage in the general population is
62 unknown [19, 20]. *L. monocytogenes* is therefore mostly considered as an opportunistic
63 pathogen.

64 *L. monocytogenes* is not only an economical and public health issue, but is also a powerful
65 model to study basic microbiology, host biology including innate and adaptive immunity and
66 cell biology [21-24]. The identification of *L. monocytogenes* virulence factors in reference
67 strains in cell culture models allowed deciphering the underlying mechanisms of its
68 interactions with the host, and also addressing basic cell biology questions [22]. Further
69 studies combining epidemiologic data with comparative genomics analyses of food- and
70 clinically-associated strains allowed the uncovering of new genes involved in *L.*
71 *monocytogenes* virulence [25]. Current investigations indicate that virulence factors are key
72 for the adaptation of *L. monocytogenes* to its host and its optimal spread in the environment,
73 and may modify our understanding of how this bacterium has acquired and selected these
74 genes.

75

76 **Uncovering virulence genes of *L. monocytogenes* by mutagenesis of reference strains**

77 It was shown early on, using *in vitro* and *in vivo* approaches, that *L. monocytogenes* can enter
78 macrophages, survive and be transferred to adjacent cells [26]. *In vivo* experiments in guinea
79 pigs indicated that *L. monocytogenes* can be released in the cytoplasm of intestinal epithelial
80 cells, surrounded by electron dense material [27], which would later come to be identified as
81 polymerized actin. Entry of *L. monocytogenes* in non-phagocytic epithelial cells was reported
82 *in vitro* in the human enterocyte-like cell line Caco-2 [28]. Most of the factors involved in key
83 phenotypes of the cellular microbiology of *L. monocytogenes*, namely entry into host cells,
84 escape from the internalization vacuole and spread from cell to cell, were characterized in the
85 1980s and 1990s and shown to be critical for virulence *in vivo*. They were mostly discovered
86 by a combination of molecular and cell biology approaches, using transposon bacterial
87 mutagenesis, gene knock-out and complementation studies coupled with infection of cultured
88 macrophages and epithelial cell lines. These factors were characterized in the laboratory
89 reference strains EGD, 10403S or EGDe, which all belong to the serovar 1/2a but to two
90 distinct lineage II clonal complexes (CCs; see below). EGD (CC7) derives from a strain
91 isolated in guinea pig by E. G. D. Murray, hence its name [3]. 10403S (CC7) is a
92 streptomycin-resistant strain derived from 10403, a strain isolated from a human listeriosis
93 case [29]. The origin of EGDe (CC9) is more difficult to trace. It was supposed to be an EGD
94 isolate, but genome analyses have revealed that both strains are actually unrelated [30, 31].

95

96 LIPI-1 genes

97 A phenotypic characteristic of *L. monocytogenes* is its ability to completely lyse red blood
98 cells (beta hemolysis), due to the expression of a hemolysin encoded by *hly* and named
99 listeriolysin O (LLO) [32, 33]. Screening and characterization of nonhemolytic mutants
100 obtained by transposon mutagenesis indicated that hemolysin expression is critically needed
101 for *L. monocytogenes* virulence [34]. Further characterization of Δhly mutant in intestinal
102 Caco-2 cells showed that LLO's pore-forming toxin is involved in escape of *L.*
103 *monocytogenes* of its internalization vacuole [28]. Later analysis involved the characterization
104 of two phospholipases, PlcA and PlcB (activated by the metalloprotease Mpl), which are also
105 involved in vacuolar escape [35, 36]. Once in the cytosol, *L. monocytogenes* can polymerize
106 actin and form so-called "actin comet tails" via its surface protein ActA, which propel *L.*
107 *monocytogenes* in the cytoplasm, and allow its spread from cell to cell [37, 38]. The study of
108 *actA* and *plcA* transposon and deletion mutants indicated that they are both critical for *L.*
109 *monocytogenes* virulence [39, 40]. Analysis of an avirulent non-hemolytic variant of the
110 NCTC 7973 guinea pig strain (isolated by E.G.D. Murray) allowed the characterization of a

111 transcription factor involved in *hly* expression which was named PrfA for "Positive regulatory
112 factor of listeriolysin (*lisA*, the other designation for *hly*)" [41, 42]. PrfA was further shown to
113 be the major regulator of *L. monocytogenes* virulence genes *in vivo*, including *prfA*, *hly*, *actA*,
114 *plcA*, *mpl*, *plcB* and the nucleomoduline *orfX* [43], which are all encoded by the Listeria
115 pathogenicity island 1 (LIPI-1) [44] (Figure 1A). PrfA is also transcriptionally regulated by
116 σ^B , that controls the general stress response in *L. monocytogenes*, is regulated post-
117 transcriptionally and requires a glutathione co-factor for full activation [45, 46].

118

119 *inlAB*

120 Transposon mutagenesis was also used to screen for mutants that are unable to enter Caco-2
121 cells. This approach allowed for the identification and characterization of internalin (InlA), a
122 leucine-rich repeats (LRR)-containing protein anchored to the bacterial peptidoglycan by a
123 Leu-Pro-X-Thr-Gly (LPXTG) motif [47]. The receptor of InlA, namely E-cadherin (Ecad),
124 which is expressed on epithelial cells and forms *adherens* junctions, was identified by affinity
125 chromatography [48]. Further analyses indicated that the InlA-Ecad interaction is species-
126 specific, as mouse and rat Ecad are not a receptor for InlA, in contrast to human Ecad. Indeed,
127 InlA interaction with Ecad requires a proline in the 16th amino acid position of Ecad, which is
128 present in permissive species (human, rabbit, guinea pig, gerbil), and is replaced by a
129 glutamic acid in non-permissive murine species (mouse, rat), preventing InlA interaction with
130 Ecad in the latter [49]. In guinea pigs and transgenic mice expressing human Ecad, InlA-Ecad
131 interaction is critical for *L. monocytogenes* to cross the intestinal barrier, at the villus level, by
132 transcytosis across goblet cells [50, 51]. Live imaging in intestinal organoids has now shown
133 that *L. monocytogenes* hijacks E-cadherin recycling for its: (i) entry into; (ii) transepithelial
134 transfer across; and (iii) exocytosis from goblet cells, in a dynamin-, microtubule- and Rab11-
135 dependent manner, respectively [52]. *inlA* is located in an operon comprising a downstream
136 gene called *inlB* [47]. InlB is also an LRR-containing protein; however, it is not covalently
137 linked to the bacterial surface by an LPXTG motif, but loosely attached to it by its Gly-Trp
138 (GW) motif. InlB mediates the entry of *L. monocytogenes* in cultured cells by interacting with
139 c-Met/HGFR, which is ubiquitously expressed [53, 54]. InlB-c-Met interaction is also
140 species-specific: it occurs in humans, mice and gerbils but not in rabbits nor guinea pigs [55].
141 InlB is not involved in *L. monocytogenes* crossing of the intestinal barrier [55]. In order to
142 study the role of the InlA-Ecad and InlB-c-Met interactions in organs other than the intestine,
143 knock-in mice KIE16P, in which the glutamic acid in 16th position of the Ecad is replaced by
144 a proline, were generated [56]. In contrast to the intestinal barrier where only InlA is

145 involved, InlA and InlB are both required and act in a conjugated manner in *L.*
146 *monocytogenes* crossing of the placental barrier in gerbils and in KIE16P mice [56, 57].

147

148 **Uncovering virulence genes of *L. monocytogenes* by comparative *Listeria* genomics**

149 While classical genetics has identified major virulence factors in *L. monocytogenes*, whole
150 genome sequencing of pathogenic bacterial species, which was initiated with *Mycoplasma*
151 *genitalium* [58], opened a new avenue for identifying genes involved in virulence, by
152 comparison of the genomic content of pathogenic and phylogenetically closely related non-
153 pathogenic bacterial species. Indeed, comparison of the genome sequences of *L.*
154 *monocytogenes* EGDe and the nonpathogenic *Listeria innocua* CLIP 11262 reference strains
155 allowed identification of putative virulence factors present in *L. monocytogenes* but absent in
156 *L. innocua* [59] (Figure 1B).

157 Sequencing of EGDe revealed the presence of 25 genes that code for proteins of the internalin
158 family, characterized by the presence of an LRR domain. Seventeen of them had not been
159 described before, including InlJ, an LPXTG anchored protein involved in virulence, whose
160 exact function remains to be determined [60, 61] and InlP (*Imo2470*), a secreted protein
161 involved in the infection of the placenta, among other organs [62, 63].

162 Comparative genomics also revealed the presence of GW motif-containing proteins other than
163 InlB in EGDe (and which were absent in *L. innocua* CLIP 11262) that are involved in
164 virulence. As an example, the GW-containing protein Auto is an autolysin involved in cell
165 entry and virulence [64].

166 Another gene that is absent in *L. innocua*, named *bsh*, has been identified by this strategy.
167 This PrfA-regulated virulence gene encodes for a bile salt hydrolase, which increases
168 resistance of *L. monocytogenes* to bile, and enhances liver infection and bacterial faecal
169 carriage [65], an essential property for a foodborne pathogen.

170

171 **Intraspecies genomic studies and identification of novel virulence genes**

172 *The biodiversity of L. monocytogenes*

173 The biodiversity of *L. monocytogenes* was first investigated via serotyping, based on *Listeria*
174 somatic and flagellar antigens, which allowed identifying the four main serotypes (1/2a, 1/2b,
175 1/2c and 4b) associated to listeriosis [66]. However, this method does not allow establishing
176 reliable phylogenetic links between strains. Multilocus enzyme electrophoresis differentiated
177 two main lineages, I and II [67], while gene variation studies and multi-locus genotyping
178 assays have led to the delineation of two minor lineages III and IV [68, 69]. In order to

179 determine the population structure of *L. monocytogenes*, multi-locus sequence typing
180 (MLST), allowing to translate sequence diversity into numerical profiles, called sequence
181 types (STs), was set up with seven housekeeping genes [70]. Using this technique, isolates
182 having at least 6 alleles in common out of these seven genes are grouped into the same clonal
183 complex (CC). With advances in whole genome sequencing, this method has been extended
184 to larger sets of loci (typically thousands) to allow strain differentiation at the level of the
185 whole- or core genome (wgMLST or cgMLST, respectively) [71-75]. A standard
186 nomenclature system has been proposed based on the cgMLST scheme of 1,748 core loci
187 [75]. Isolates sharing less than 150 and 7 allelic mismatches are grouped into sublineages
188 (SL) and cgMLST types (CT), respectively, which allows for efficient inter-laboratory
189 communication, in particular for emerging strain types [76] (Figure 2A). In addition, the
190 compilation of genotypic data in dedicated databases, such as BIGSdb-*Lm*
191 (<http://bigsdb.pasteur.fr/listeria>) or GenomeTrakr (hosted at
192 <https://www.ncbi.nlm.nih.gov/pathogens>) allow for in-depth studies of the global spread and
193 biodiversity of *L. monocytogenes*.

194

195 *Hypervirulent strains belong to distinct CCs compared to hypovirulent strains*

196 *L. monocytogenes* can be divided into four evolutionary lineages [77], 13 serotypes [78]
197 grouped into 6 genosergroups [79] and more than 170 clonal complexes and 300 sublineages
198 (<http://bigsdb.pasteur.fr/listeria>). Of these, lineage I (serotypes 4b and 1/2b) and II (serotypes
199 1/2a and 1/2c) isolates represent the majority of clinical isolates. The most prevalent CCs and
200 SLs are geographically and temporally widespread [70, 80, 81]. However, an uneven
201 prevalence is observed among food vs. clinical cases [25, 75, 80], as previously noted at the
202 lineage and serotype levels [77]. While lineage II, in particularly sublineages SL9 (CC9) and
203 SL121 (CC121) are mostly associated with food, lineage I isolates, and in particular
204 sublineages SL1 (CC1), SL2 (CC2), SL4 (CC4) and SL6 (CC6) are strongly associated with
205 clinical cases in the Western world [25, 75, 82-87] (Figure 2B). In Asia, specifically in China
206 and Taiwan, the most prevalent sublineage in both food and clinical cases is SL87 (CC87,
207 lineage I, serotype 1/2b) [84, 88, 89], suggesting that food habits and/or local specificities are
208 involved in geographic diversity of *L. monocytogenes*.

209 A large study of 6,663 *L. monocytogenes* isolates of clinical and food origins prospectively
210 collected in France between 2005 and 2013 reported a higher prevalence of CC9 and CC121
211 (both of lineage II) isolates in food samples, while CC1, CC6, CC2 and CC4 (all of lineage I)
212 were the most prevalent in clinical cases [25]. In addition, CC1 and CC4 were found to be

213 statistically associated with more invasive forms of listeriosis (maternal-neonatal and CNS
214 infections). Similar results have been observed within farm environments, with CC1 being
215 also associated with CNS infection in ruminants in Switzerland [90]. This could reflect a
216 difference in virulence and/or in exposure of at-risk patients to these strains. Association of
217 CCs to clinical cases is inversely correlated to patients' immunosuppressive comorbidities,
218 suggesting that these are more virulent than CCs most associated with food [25]. *In vivo*
219 studies, which compared CC1, CC4, CC6, CC9 and CC121 isolates and the laboratory
220 reference strains EGDe and 10403S in a mouse model permissive for oral *L. monocytogenes*
221 infection, unambiguously showed that strains from CCs most associated with clinical cases
222 (CC1, CC4 and CC6) are hypervirulent when compared to laboratory reference strains EGDe
223 and 10403S, as they induced a more pronounced body weight decrease and are more
224 neuroinvasive [25]. In contrast, CC9 and CC121 are hypovirulent when compared to these
225 laboratory reference strains. Of note, CC9 and CC121 are not fully avirulent, since they can
226 still infect laboratory animals, although to a lower degree, as well as immunodeficient patients
227 [25]. This heterogeneity in virulence among lineages and CCs led to an investigation of the
228 putative molecular mechanisms involved in both hypo- and hyper-virulence.

229

230 Virulence factors found in the core genome

231 While *inlA* is present in all *L. monocytogenes* strains, some strains, including the reference
232 laboratory strain LO28 (CC9, lineage II), express a truncated and therefore nonfunctional
233 InlA [91]. Epidemiological analysis indicated that clinical strains, and most significantly
234 those from maternal-fetal infections, express full-length InlA more frequently than strains
235 from food origins [70, 92]. InlA is truncated in the majority of hypovirulent CC9 and CC121
236 isolates as well as in other lineage II clonal complexes (e. g. CC331, CC199 and CC321[75]).
237 Hypovirulence can therefore, at least in part, be accounted for by InlA truncation, leading to
238 decreased translocation across the intestinal barrier and placental infection [50, 56, 93].

239 Similarly to *inlAB*, LIPI-1 genes are present in all *L. monocytogenes* SLs and are therefore
240 core virulence genes (Figure 1A). As indicated above, non-pathogenic *L. innocua* lacks LIPI-
241 1 and *inlAB* genes, however rare *L. innocua* strains contain both LIPI-1 and *inlA* that are
242 functional *in vitro* and *in vivo* [94-99]. This indicates that *L. monocytogenes* and *L. innocua*
243 species likely evolved independently from a common pathogenic ancestor harboring both the
244 LIPI-1 and *inlAB* virulence loci. Speciation likely occurred upon differential environmental
245 selective pressures, with *L. innocua* evolving in an environment where virulence genes are not
246 needed [100]. In contrast, 0.1% of *L. monocytogenes* strains from both lineage I and II are

247 non-hemolytic. 93.3% of these non-hemolytic strains harbor at least one loss-of-function
248 mutation in *prfA*, and the remaining 6.7% harbor *hly* mutations leading to LLO inactivation.
249 In both of these cases, this results in a strong decrease in virulence, suggesting that niche
250 restriction leading to sub-speciation may still occur within *L. monocytogenes* species [101].

251

252 Virulence factors encoded by the accessory genome

253 LIPI-3

254 The LIPI-3 virulence cluster, which is absent in *L. monocytogenes* reference strains but
255 present in lineage I F2365 (SL1/CC1) and H7858 (SL6/CC6) outbreak strains, has also been
256 characterized (Figure 1A). It consists of 8 genes, including a hemolytic and cytotoxic factor,
257 listeriolysin S (LLS), involved in virulence [102]. LLS, encoded by *llsA*, acts as a bacteriocin
258 and alters microbiota composition, resulting in an increase in the luminal amount of *L.*
259 *monocytogenes*, while *llsB*, another gene of LIPI-3, plays a role in the systemic phase of the
260 infection [103, 104]. Functional LIPI-3 or its remnants can also be present in some *L. innocua*
261 strains [98, 105], where its role is not defined, reinforcing the hypothesis that the two species
262 share a common pathogenic ancestor. The presence of LIPI-3 in *Lm* sublineages SL1 (CC1),
263 SL4 (CC4) and SL6 (CC6), which are the most associated with clinical cases [75, 98], and its
264 absence in lineage II strains, suggests a possible role in their hypervirulence.

265

266 LIPI-4

267 Comparative genomic analyses derived from biodiversity studies allowed the characterization
268 of LIPI-4, a putative phosphotransferase system (PTS) present in SL4, a sublineage of the
269 lineage I, which strongly associated with a clinical origin (Figure 2B), particularly in CNS
270 and placental infections [25]. LIPI-4 is involved in placental and CNS infection in an *in vivo*
271 model of listeriosis, by a so far unknown mechanism (Figures 1A and 2A) [25]. Interestingly,
272 LIPI-4, is also present in the *L. monocytogenes* sublineage that is the most prevalent in Asia,
273 SL87 (CC87) [106], as well as in *L. innocua* [98]. Its actual role in such a non-virulent species
274 is currently unknown.

275

276 LIPI-2

277 LIPI-2 is a genomic region discovered in *Listeria ivanovii*, a *Listeria* species which is
278 pathogenic for mammals, mainly ovines and bovines, but also rarely humans [108, 109].
279 LIPI-2 was first described as an *L. ivanovii*-specific genomic region that encodes a
280 sphingomyelinase involved in phagosome disruption, as well as 10 genes of the internalin

281 family [110]. This region is involved in *L. ivanovii* virulence, together with the *L. ivanovii*
282 LIPI-1 orthologous region, both *in vitro* and in an *in vivo* sheep model of infection [110]. *L.*
283 *monocytogenes* isolates belonging to a new sublineage of the major lineage II (SL626/CC33,
284 serovar 4h) recovered from an ovine listeriosis outbreak in China have been characterized as
285 hypervirulent [107]. These isolates contain a partial LIPI-2 locus, comprising only the
286 sphingomyelinase *smcL* and the internalins *i-inlF* and *i-inlE* genes (Figure 1A),
287 likely by acquisition of exogenous DNA from *L. ivanovii*. The specific role of the partial
288 LIPI-2 in hypervirulence of this newly described *L. monocytogenes* sublineage remains to be
289 elucidated.

290

291 Heterogeneity in virulence as a reflection of differential host adaptation?

292 The distribution of *L. monocytogenes* is wide. It has been isolated in humans and animals in
293 all continents except Antarctica [80]. Linked to its ability to tolerate extreme temperatures,
294 acidity and osmolarity, *L. monocytogenes* can persist and grow in soil, plants, water, food and
295 silage [111-113]. However, the distribution of *L. monocytogenes* lineages and CCs is not
296 homogeneous in these ecosystems, indicating that CCs emergence may be driven by their
297 selective adaptation to the environment in which they are the most abundant [25, 77].

298 When investigating food samples, hypervirulent CC1, CC4 and CC6 were found to be
299 statistically associated with dairy products, whereas hypovirulent *L. monocytogenes* clones,
300 CC9 and CC121 were found associated with meat products [114], suggesting a differential
301 adaptation of the CCs to their environment. Interestingly, CC9 and CC121 harbor genes
302 involved in tolerance to benzalkonium chloride, a surface disinfectant commonly used in food
303 production environments, and are more efficient than their hypervirulent counterparts to form
304 biofilms in the presence of this disinfectant [114, 115]. Other stress resistance genes such as
305 the SSI-1 and SSI-2 stress survival islets [116, 117] are also typically present in lineage II,
306 and contribute to *L. monocytogenes* survival and growth in the stress conditions encountered
307 in food and food processing environments (e.g. high salt concentrations, oxidative stress).
308 Lineage I isolates typically display higher tolerance to low pH [118], which can be
309 advantageous for the survival of *L. monocytogenes* in the host's gastric acidic environment.
310 Consistent with these observations, strains of hypervirulent CCs colonize the gut more
311 efficiently than those of hypovirulent CCs [114], which may account for their strong
312 association with ruminant feces [119] and dairy products, which may result from milk fecal
313 contamination [114] (Figure 3).

314

315 **Concluding remarks**

316 *L. monocytogenes* is a major model pathogen which has allowed scientific breakthroughs in
317 microbiology, immunology, cell biology and host-pathogen interactions. It has also emerged
318 as a powerful model to study how microbes evolve from their natural habitat to anthropic
319 environments. While advances have been made in the discovery of genetic factors involved in
320 virulence and adaptation to host vs. food processing environments, the mechanisms by which
321 *L. monocytogenes* adapts to various environments remains to be fully deciphered. Seminal
322 studies have indicated how the transcriptome in the EGDe reference strain responds to the
323 switch from saprophytism to virulence [120]. Future large-scale transcriptomics and
324 metabolomics studies with representative strains from different clonal complexes will enrich
325 our understanding of how *L. monocytogenes* can adapt to the environments in which it thrives.
326 Further studies will also be needed to decipher the forces driving hypervirulence and its
327 relationship with adaptation to the host, and how this relates to capacity of *L. monocytogenes*
328 to be shed back in the environment and transmitted to new hosts, as suggested by data from
329 ruminants and animal models [114, 119]. The continuous analysis of biodiversity of *L.*
330 *monocytogenes*, together with environmental, epidemiological, clinical and experimental data
331 obtained from genome wide association studies will shed light into the mechanisms and
332 selective pressures that drive *Listeria* adaptation to diverse environments.

333

334 **Figure Legends**

335 **Figure 1. Virulence genes of *L. monocytogenes***

336 A) Typical genetic organization of *Listeria* pathogenicity islands (LIPI; colored arrows).
337 Flanking genes (gray arrows) are labeled based on their orthologs in *L. monocytogenes* EGDe,
338 when existing. *Lm*, *L. monocytogenes*; *Lin*, *L. innocua*; *Liv*, *L. ivanovii*; *, LIPI-2 genes
339 present in *Liv* atypical *L. monocytogenes* CC33 isolates are shown in darker yellow.

340 B) Genome maps of *L. monocytogenes* EGDe and *L. innocua* CLIP11262 [59]. Inner circles
341 represent the GC skew $((G - C)/(G + C))$ and GC content $((G+C)/(G+C+A+T))$ of *L.*
342 *monocytogenes* EGDe. Gaps in *L. innocua* chromosome (blue) represent genomic regions
343 absent in *L. monocytogenes* genome (red). The genomic positions of known *L.*
344 *monocytogenes* virulence genes are shown and their common and/or locus names indicated.
345 Internalin genes are highlighted in red. Triangles denote the genomic positions of LIPI
346 regions in either EGDe (black triangle) or other *Listeria* species (open triangles), as detailed
347 in panel A.

348

349 **Figure 2. Population structure of *L. monocytogenes***

350 A) Population structure of *L. monocytogenes* based on cgMLST profiles (adapted from [75];
351 $N=1,696$ isolates collected from food and clinical samples, mostly in North America and
352 Europe). Branches are colored by phylogenetic lineage (I-IV, shown in red, blue, green and
353 gray, respectively) and collapsed by sublineage (SL). The vertical dashed bar indicates the
354 established cgMLST cut-off to define SLs (i.e. 150 allelic differences out of 1748 loci). Main
355 SLs are labeled on the branches and corresponding MLST clonal complexes (CCs) are shown
356 in the first column, PCR serogroups, source types and continents of isolation are colored
357 according the key panels (upper left). Dark blue boxes denote the presence of selected genetic
358 traits: pathogenicity islands (LIPI-1 to -4), genes involved in teichoic acid biosynthesis
359 (*gltAB*, *tagB* and *gtcA*), genes coding for internalins (*inlABCEFGHJK*), autolysins (*aut* and
360 *autIVb*) and bile resistance (*bsh*), genes involved in benzalkonium resistance (*bcrABC*
361 operon, *ermC*, *Tn6188::ermC* and *ermE*), stress resistance islets (SSI-1 and SSI-2) and
362 biofilm formation and virulence (*comK*).

363 B) Frequency of the most prevalent *L. monocytogenes* MLST clonal complexes in clinical (y -
364 axis) versus food (x -axis) isolates, based on a exhaustively collected isolates dataset obtained
365 in the scope of listeriosis surveillance ($N=6,633$ isolates, France 2005-2013). (reproduced
366 with permission from [25]). Each circle represents a clonal complex, and is colored according
367 to its lineage. The size of each circle is proportional to the number of isolates it represents.
368 The clonal complexes of common reference lab strains (CC7: strains EGD and 10403S; CC9:
369 EGDe and LO28), are also indicated.

370

371 **Figure 3. Trade-off between host-associated and environmental lifestyles of *L.***
372 ***monocytogenes***

373 Properties of *L. monocytogenes* strains adapted to a host environment (in red) versus of those
374 adapted to saprophytism (natural and food-processing environments; in blue). Listeriosis
375 results from the interplay between host and microbial factors: the less immunocompromised is
376 the host, the more virulent *L. monocytogenes* strains need to be to cause disease. Thus,
377 immunocompromised hosts tend to be susceptible to even saprophytic (hypovirulent) *L.*
378 *monocytogenes*, whereas healthy hosts are usually only susceptible to host-associated
379 (hypervirulent) strains, as shown in [25].

380

381 **Acknowledgments**

382 Research in the Biology of Infection Unit is supported financially by Institut Pasteur, Inserm,
383 ANR and ERC. Marc Lecuit is a senior member of Institut Universitaire de France.

384 **References**

- 385 1. Hülphers, G. (1911) Lefvernekros hos kanin orsakad af en ej fovut beskريفuen bakterie.
386 Soensk Veterinar Tidskrift 16, 265-273.
- 387 2. Dumont, J. and Cotoni, L. (1921) Bacille semblable au bacille du rouget du porc rencontré
388 dans le liquide céphalo-rachidien d'un méningitique. Ann. Inst. Pasteur (Paris) 35 (10), 625-
389 33.
- 390 3. Murray, E.G.D. et al. (1926) A disease of rabbits characterised by a large mononuclear
391 leucocytosis caused by hitherto undescribed bacillus *Bacterium monocytogenes* (n. sp.). J.
392 Pathol. Bacteriol. 29, 407-439.
- 393 4. Pirie, J.H.H. (1927) A new disease of veld rodents, "Tiger river disease.". Pub South Africa
394 Inst Med Res 62, 163-186.
- 395 5. Charlier, C. et al. (2017) Clinical features and prognostic factors of listeriosis: the
396 MONALISA national prospective cohort study. Lancet Infect Dis 17 (5), 510-519.
- 397 6. Danion, F. et al. (2017) *Listeria monocytogenes* isolation from urine: a series of 15 cases
398 and review. Clinical microbiology and infection : the official publication of the European
399 Society of Clinical Microbiology and Infectious Diseases 23 (8), 583-585.
- 400 7. Chersich, M.F. et al. (2018) Diagnosis and Treatment of *Listeria monocytogenes*
401 Endophthalmitis: A Systematic Review. Ocular immunology and inflammation 26 (4), 508-
402 517.
- 403 8. Charlier, C. et al. (2012) *Listeria monocytogenes*-associated joint and bone infections: a
404 study of 43 consecutive cases. Clinical infectious diseases : an official publication of the
405 Infectious Diseases Society of America 54 (2), 240-8.
- 406 9. Charlier, C. et al. (2014) *Listeria monocytogenes*-associated biliary tract infections: a study
407 of 12 consecutive cases and review. Medicine 93 (18), e105.
- 408 10. Shoai-Tehrani, M. et al. (2019) *Listeria monocytogenes*-associated endovascular
409 infections: A study of 71 consecutive cases. The Journal of infection 79 (4), 322-331.
- 410 11. Morgand, M. et al. (2018) *Listeria monocytogenes*-associated respiratory infections: a
411 study of 38 consecutive cases. Clinical microbiology and infection : the official publication of
412 the European Society of Clinical Microbiology and Infectious Diseases 24 (12), 1339 e1-1339
413 e5.
- 414 12. Pilmis, B. et al. (2020) Cutaneous listeriosis, a case series of 16 consecutive patients over
415 25 years. The Journal of infection 80 (2), 232-254.
- 416 13. Dalton, C.B. et al. (1997) An outbreak of gastroenteritis and fever due to *Listeria*
417 *monocytogenes* in milk. N Engl J Med 336 (2), 100-5.
- 418 14. Schlech, W.F., 3rd et al. (1983) Epidemic listeriosis--evidence for transmission by food. N
419 Engl J Med 308 (4), 203-6.
- 420 15. ECDC (2018) Annual epidemiological report for 2016 Listeriosis. European Centre for
421 Disease Prevention and Control.
- 422 16. Pohl, A.M. et al. (2019) Differences Among Incidence Rates of Invasive Listeriosis in the
423 U.S. FoodNet Population by Age, Sex, Race/Ethnicity, and Pregnancy Status, 2008-2016.
424 Foodborne pathogens and disease 16 (4), 290-297.
- 425 17. European Food Safety, A. et al. (2019) The European Union One Health 2018 Zoonoses
426 Report. EFSA J 17 (12), e05926.
- 427 18. Thomas, J. et al. (2020) Outbreak of Listeriosis in South Africa Associated with Processed
428 Meat. N Engl J Med 382 (7), 632-643.
- 429 19. Sauders, B.D. et al. (2005) Low prevalence of *Listeria monocytogenes* in human stool. J
430 Food Prot 68 (1), 178-81.
- 431 20. Grif, K. et al. (2003) Incidence of fecal carriage of *Listeria monocytogenes* in three
432 healthy volunteers: a one-year prospective stool survey. Eur J Clin Microbiol Infect Dis 22
433 (1), 16-20.

- 434 21. Radoshevich, L. and Cossart, P. (2018) *Listeria monocytogenes*: towards a complete
435 picture of its physiology and pathogenesis. Nature reviews. Microbiology 16 (1), 32-46.
- 436 22. Lecuit, M. (2020) *Listeria monocytogenes*, a model in infection biology. Cellular
437 microbiology 22 (4), e13186.
- 438 23. Pamer, E.G. (2004) Immune responses to *Listeria monocytogenes*. Nature reviews.
439 Immunology 4 (10), 812-23.
- 440 24. Maudet, C. et al. (2020) Innate immune responses to *Listeria* in vivo. Current opinion in
441 microbiology In press.
- 442 25. Maury, M.M. et al. (2016) Uncovering *Listeria monocytogenes* hypervirulence by
443 harnessing its biodiversity. Nat Genet 48 (3), 308-313.
- 444 26. Mackaness, G.B. (1962) Cellular resistance to infection. J Exp Med 116, 381-406.
- 445 27. Racz, P. et al. (1972) Experimental *Listeria* enteritis. I. An electron microscopic study of
446 the epithelial phase in experimental listeria infection. Lab Invest 26 (6), 694-700.
- 447 28. Gaillard, J.L. et al. (1987) *In vitro* model of penetration and intracellular growth of
448 *Listeria monocytogenes* in the human enterocyte-like cell line Caco-2. Infect Immun 55 (11),
449 2822-9.
- 450 29. Edman, D.C. et al. (1968) *Listeria monocytogenes* L forms. I. Induction maintenance, and
451 biological characteristics. J Bacteriol 96 (2), 352-7.
- 452 30. Cantinelli, T. et al. (2013) "Epidemic clones" of *Listeria monocytogenes* are widespread
453 and ancient clonal groups. J Clin Microbiol 51 (11), 3770-9.
- 454 31. Becavin, C. et al. (2014) Comparison of widely used *Listeria monocytogenes* strains EGD,
455 10403S, and EGD-e highlights genomic variations underlying differences in pathogenicity.
456 mBio 5 (2), e00969-14.
- 457 32. Geoffroy, C. et al. (1987) Purification, characterization, and toxicity of the sulfhydryl-
458 activated hemolysin listeriolysin O from *Listeria monocytogenes*. Infect Immun 55 (7), 1641-
459 6.
- 460 33. Harvey, P.C. and Faber, J.E. (1941) Studies on the *Listerella* Group: I. Biochemical and
461 Hemolytic Reactions. J Bacteriol 42 (5), 677-87.
- 462 34. Gaillard, J.L. et al. (1986) Transposon mutagenesis as a tool to study the role of
463 hemolysin in the virulence of *Listeria monocytogenes*. Infect Immun 52 (1), 50-5.
- 464 35. Camilli, A. et al. (1993) Dual roles of *plcA* in *Listeria monocytogenes* pathogenesis. Mol
465 Microbiol 8 (1), 143-57.
- 466 36. Domann, E. et al. (1991) Molecular cloning, sequencing, and identification of a
467 metalloprotease gene from *Listeria monocytogenes* that is species specific and physically
468 linked to the listeriolysin gene. Infect Immun 59 (1), 65-72.
- 469 37. Kocks, C. et al. (1992) *L. monocytogenes*-induced actin assembly requires the *actA* gene
470 product, a surface protein. Cell 68 (3), 521-31.
- 471 38. Tilney, L.G. and Portnoy, D.A. (1989) Actin filaments and the growth, movement, and
472 spread of the intracellular bacterial parasite, *Listeria monocytogenes*. J Cell Biol 109 (4 Pt 1),
473 1597-608.
- 474 39. Camilli, A. et al. (1991) *Listeria monocytogenes* mutants lacking phosphatidylinositol-
475 specific phospholipase C are avirulent. J Exp Med 173 (3), 751-4.
- 476 40. Portnoy, D.A. et al. (1988) Role of hemolysin for the intracellular growth of *Listeria*
477 *monocytogenes*. J Exp Med 167 (4), 1459-71.
- 478 41. Leimeister-Wachter, M. et al. (1990) Identification of a gene that positively regulates
479 expression of listeriolysin, the major virulence factor of *Listeria monocytogenes*. Proc Natl
480 Acad Sci U S A 87 (21), 8336-40.
- 481 42. Mengaud, J. et al. (1991) Pleiotropic control of *Listeria monocytogenes* virulence factors
482 by a gene that is autoregulated. Mol Microbiol 5 (9), 2273-83.

- 483 43. Prokop, A. et al. (2017) OrfX, a Nucleomodulin Required for *Listeria monocytogenes*
484 Virulence. *mBio* 8 (5).
- 485 44. de las Heras, A. et al. (2011) Regulation of *Listeria* virulence: PrfA master and
486 commander. *Current opinion in microbiology* 14 (2), 118-27.
- 487 45. Reniere, M.L. et al. (2015) Glutathione activates virulence gene expression of an
488 intracellular pathogen. *Nature* 517 (7533), 170-3.
- 489 46. Tiensuu, T. et al. (2019) Flick of a switch: regulatory mechanisms allowing *Listeria*
490 *monocytogenes* to transition from a saprophyte to a killer. *Microbiology (Reading)* 165 (8),
491 819-833.
- 492 47. Gaillard, J.L. et al. (1991) Entry of *L. monocytogenes* into cells is mediated by internalin,
493 a repeat protein reminiscent of surface antigens from gram-positive cocci. *Cell* 65 (7), 1127-
494 41.
- 495 48. Mengaud, J. et al. (1996) E-cadherin is the receptor for internalin, a surface protein
496 required for entry of *L. monocytogenes* into epithelial cells. *Cell* 84 (6), 923-32.
- 497 49. Lecuit, M. et al. (1999) A single amino acid in E-cadherin responsible for host specificity
498 towards the human pathogen *Listeria monocytogenes*. *EMBO J* 18 (14), 3956-63.
- 499 50. Lecuit, M. et al. (2001) A transgenic model for listeriosis: role of internalin in crossing the
500 intestinal barrier. *Science* 292 (5522), 1722-5.
- 501 51. Nikitas, G. et al. (2011) Transcytosis of *Listeria monocytogenes* across the intestinal
502 barrier upon specific targeting of goblet cell accessible E-cadherin. *J Exp Med* 208 (11),
503 2263-77.
- 504 52. Kim, M. et al. (2020) Live Imaging Reveals *Listeria* Hijacking of E-Cadherin Recycling
505 as It Crosses the Intestinal Barrier. *Curr Biol*.
- 506 53. Dramsi, S. et al. (1995) Entry of *Listeria monocytogenes* into hepatocytes requires
507 expression of InIB, a surface protein of the internalin multigene family. *Mol Microbiol* 16 (2),
508 251-61.
- 509 54. Shen, Y. et al. (2000) InIB-dependent internalization of *Listeria* is mediated by the Met
510 receptor tyrosine kinase. *Cell* 103 (3), 501-10.
- 511 55. Khelef, N. et al. (2006) Species specificity of the *Listeria monocytogenes* InIB protein.
512 *Cell Microbiol* 8 (3), 457-70.
- 513 56. Disson, O. et al. (2008) Conjugated action of two species-specific invasion proteins for
514 fetoplacental listeriosis. *Nature* 455 (7216), 1114-8.
- 515 57. Gessain, G. et al. (2015) PI3-kinase activation is critical for host barrier permissiveness to
516 *Listeria monocytogenes*. *J Exp Med* 212 (2), 165-83.
- 517 58. Fraser, C.M. et al. (1995) The minimal gene complement of *Mycoplasma genitalium*.
518 *Science* 270 (5235), 397-403.
- 519 59. Glaser, P. et al. (2001) Comparative genomics of *Listeria* species. *Science* 294 (5543),
520 849-52.
- 521 60. Sabet, C. et al. (2005) LPXTG protein InIJ, a newly identified internalin involved in
522 *Listeria monocytogenes* virulence. *Infect Immun* 73 (10), 6912-22.
- 523 61. Sabet, C. et al. (2008) The *Listeria monocytogenes* virulence factor InIJ is specifically
524 expressed in vivo and behaves as an adhesin. *Infect Immun* 76 (4), 1368-78.
- 525 62. Faralla, C. et al. (2018) *Listeria monocytogenes* InIP interacts with afadin and facilitates
526 basement membrane crossing. *PLoS Pathog* 14 (5), e1007094.
- 527 63. Faralla, C. et al. (2016) InIP, a New Virulence Factor with Strong Placental Tropism.
528 *Infect Immun* 84 (12), 3584-3596.
- 529 64. Cabanes, D. et al. (2004) Auto, a surface associated autolysin of *Listeria monocytogenes*
530 required for entry into eukaryotic cells and virulence. *Mol Microbiol* 51 (6), 1601-14.

- 531 65. Dussurget, O. et al. (2002) *Listeria monocytogenes* bile salt hydrolase is a PrfA-regulated
532 virulence factor involved in the intestinal and hepatic phases of listeriosis. *Mol Microbiol* 45
533 (4), 1095-106.
- 534 66. Seeliger, H.P.R. and Höhne, K. (1979) Serotyping of *Listeria monocytogenes* and related
535 species. *Methods in Microbiology* New York: Academic Press, 33-48.
- 536 67. Piffaretti, J.C. et al. (1989) Genetic characterization of clones of the bacterium *Listeria*
537 *monocytogenes* causing epidemic disease. *Proceedings of the National Academy of Sciences*
538 of the United States of America 86 (10), 3818-22.
- 539 68. Rasmussen, O.F. et al. (1995) *Listeria monocytogenes* exists in at least three evolutionary
540 lines: evidence from flagellin, invasive associated protein and listeriolysin O genes.
541 *Microbiology* 141 (Pt 9), 2053-61.
- 542 69. Ward, T.J. et al. (2008) Multilocus genotyping assays for single nucleotide
543 polymorphism-based subtyping of *Listeria monocytogenes* isolates. *Appl Environ Microbiol*
544 74 (24), 7629-42.
- 545 70. Ragon, M. et al. (2008) A new perspective on *Listeria monocytogenes* evolution. *PLoS*
546 *Pathog* 4 (9), e1000146.
- 547 71. Jensen, A.K. et al. (2016) Molecular Typing and Epidemiology of Human Listeriosis
548 Cases, Denmark, 2002-2012. *Emerging infectious diseases* 22 (4), 625-33.
- 549 72. Chen, Y. et al. (2016) Core Genome Multilocus Sequence Typing for Identification of
550 Globally Distributed Clonal Groups and Differentiation of Outbreak Strains of *Listeria*
551 *monocytogenes*. *Applied and environmental microbiology* 82 (20), 6258-6272.
- 552 73. Ruppitsch, W. et al. (2015) Defining and Evaluating a Core Genome Multilocus Sequence
553 Typing Scheme for Whole-Genome Sequence-Based Typing of *Listeria monocytogenes*.
554 *Journal of clinical microbiology* 53 (9), 2869-76.
- 555 74. Pightling, A.W. et al. (2015) The *Listeria monocytogenes* Core-Genome Sequence Typer
556 (LmCGST): a bioinformatic pipeline for molecular characterization with next-generation
557 sequence data. *BMC microbiology* 15, 224.
- 558 75. Moura, A. et al. (2016) Whole genome-based population biology and epidemiological
559 surveillance of *Listeria monocytogenes*. *Nat Microbiol* 2, 16185.
- 560 76. Moura, A. et al. (2017) Real-Time Whole-Genome Sequencing for Surveillance of
561 *Listeria monocytogenes*, France. *Emerg Infect Dis* 23 (9), 1462-1470.
- 562 77. Orsi, R.H. et al. (2011) *Listeria monocytogenes* lineages: Genomics, evolution, ecology,
563 and phenotypic characteristics. *Int J Med Microbiol* 301 (2), 79-96.
- 564 78. Seeliger, H.P. and Langer, B. (1989) Serological analysis of the genus *Listeria*. Its values
565 and limitations. *Int J Food Microbiol* 8 (3), 245-8.
- 566 79. Doumith, M. et al. (2004) Differentiation of the major *Listeria monocytogenes* serovars by
567 multiplex PCR. *J Clin Microbiol* 42 (8), 3819-22.
- 568 80. Chenal-Francois, V. et al. (2011) Worldwide distribution of major clones of *Listeria*
569 *monocytogenes*. *Emerg Infect Dis* 17 (6), 1110-2.
- 570 81. Haase, J.K. et al. (2014) The ubiquitous nature of *Listeria monocytogenes* clones: a large-
571 scale Multilocus Sequence Typing study. *Environ Microbiol* 16 (2), 405-16.
- 572 82. Kwong, J.C. et al. (2016) Prospective Whole-Genome Sequencing Enhances National
573 Surveillance of *Listeria monocytogenes*. *J Clin Microbiol* 54 (2), 333-42.
- 574 83. Lee, S. et al. (2018) *Listeria monocytogenes* Source Distribution Analysis Indicates
575 Regional Heterogeneity and Ecological Niche Preference among Serotype 4b Clones. *mBio* 9
576 (2).
- 577 84. Wang, Y. et al. (2012) Genetic diversity and molecular typing of *Listeria monocytogenes*
578 in China. *BMC Microbiol* 12, 119.
- 579 85. Bertrand, S. et al. (2016) Diversity of *Listeria monocytogenes* Strains of Clinical and Food
580 Chain Origins in Belgium between 1985 and 2014. *PLoS One* 11 (10), e0164283.

581 86. Kuch, A. et al. (2018) Molecular diversity and antimicrobial susceptibility of *Listeria*
582 *monocytogenes* isolates from invasive infections in Poland (1997-2013). *Sci Rep* 8 (1), 14562.
583 87. Painset, A. et al. (2019) LiSEQ - whole-genome sequencing of a cross-sectional survey of
584 *Listeria monocytogenes* in ready-to-eat foods and human clinical cases in Europe. *Microb*
585 *Genom* 5 (2).
586 88. Huang, Y.T. et al. (2015) Disease Burden of Invasive Listeriosis and Molecular
587 Characterization of Clinical Isolates in Taiwan, 2000-2013. *PloS one* 10 (11), e0141241.
588 89. Wang, Y. et al. (2015) Characterization of *Listeria monocytogenes* isolated from human
589 Listeriosis cases in China. *Emerg Microbes Infect* 4 (8), e50.
590 90. Dreyer, M. et al. (2016) *Listeria monocytogenes* sequence type 1 is predominant in
591 ruminant rhombencephalitis. *Scientific reports* 6, 36419.
592 91. Jonquieres, R. et al. (1998) The *inlA* gene of *Listeria monocytogenes* LO28 harbors a
593 nonsense mutation resulting in release of internalin. *Infect Immun* 66 (7), 3420-2.
594 92. Jacquet, C. et al. (2004) A molecular marker for evaluating the pathogenic potential of
595 foodborne *Listeria monocytogenes*. *J Infect Dis* 189 (11), 2094-100.
596 93. Lecuit, M. et al. (2004) Targeting and crossing of the human maternofetal barrier by
597 *Listeria monocytogenes*: role of internalin interaction with trophoblast E-cadherin. *Proc Natl*
598 *Acad Sci U S A* 101 (16), 6152-7.
599 94. den Bakker, H.C. et al. (2010) Comparative genomics of the bacterial genus *Listeria*:
600 Genome evolution is characterized by limited gene acquisition and limited gene loss. *BMC*
601 *Genomics* 11, 688.
602 95. Johnson, J. et al. (2004) Natural atypical *Listeria innocua* strains with *Listeria*
603 *monocytogenes* pathogenicity island 1 genes. *Appl Environ Microbiol* 70 (7), 4256-66.
604 96. Milillo, S.R. et al. (2012) *Listeria monocytogenes* and hemolytic *Listeria innocua* in
605 poultry. *Poult Sci* 91 (9), 2158-63.
606 97. Moreno, L.Z. et al. (2012) Characterization of atypical *Listeria innocua* isolated from
607 swine slaughterhouses and meat markets. *Res Microbiol* 163 (4), 268-71.
608 98. Moura, A. et al. (2019) Atypical Hemolytic *Listeria innocua* Isolates Are Virulent, albeit
609 Less than *Listeria monocytogenes*. *Infect Immun* 87 (4).
610 99. Volokhov, D.V. et al. (2007) The presence of the internalin gene in natural atypically
611 hemolytic *Listeria innocua* strains suggests descent from *L. monocytogenes*. *Appl Environ*
612 *Microbiol* 73 (6), 1928-39.
613 100. Schmid, M.W. et al. (2005) Evolutionary history of the genus *Listeria* and its virulence
614 genes. *Syst Appl Microbiol* 28 (1), 1-18.
615 101. Maury, M.M. et al. (2017) Spontaneous Loss of Virulence in Natural Populations of
616 *Listeria monocytogenes*. *Infect Immun* 85 (11).
617 102. Cotter, P.D. et al. (2008) Listeriolysin S, a novel peptide haemolysin associated with a
618 subset of lineage I *Listeria monocytogenes*. *PLoS Pathog* 4 (9), e1000144.
619 103. Quereda, J.J. et al. (2016) Bacteriocin from epidemic *Listeria* strains alters the host
620 intestinal microbiota to favor infection. *Proc Natl Acad Sci U S A* 113 (20), 5706-11.
621 104. Quereda, J.J. et al. (2017) Listeriolysin S: A bacteriocin from epidemic *Listeria*
622 *monocytogenes* strains that targets the gut microbiota. *Gut Microbes* 8 (4), 384-391.
623 105. Clayton, E.M. et al. (2014) Atypical *Listeria innocua* strains possess an intact LIPI-3.
624 *BMC Microbiol* 14, 58.
625 106. Wang, Y. et al. (2019) Genomic dissection of the most prevalent *Listeria monocytogenes*
626 clone, sequence type ST87, in China. *BMC Genomics* 20 (1), 1014.
627 107. Yin, Y. et al. (2019) A hybrid sub-lineage of *Listeria monocytogenes* comprising
628 hypervirulent isolates. *Nat Commun* 10 (1), 4283.
629 108. Guillet, C. et al. (2010) Human listeriosis caused by *Listeria ivanovii*. *Emerg Infect Dis*
630 16 (1), 136-8.

631 109. Sergeant, E.S. et al. (1991) Abortions in sheep due to *Listeria ivanovii*. Australian
632 veterinary journal 68 (1), 39.

633 110. Dominguez-Bernal, G. et al. (2006) A spontaneous genomic deletion in *Listeria ivanovii*
634 identifies LIPI-2, a species-specific pathogenicity island encoding sphingomyelinase and
635 numerous internalins. Mol Microbiol 59 (2), 415-32.

636 111. Fenlon, D.R. (1985) Wild birds and silage as reservoirs of *Listeria* in the agricultural
637 environment. J Appl Bacteriol 59 (6), 537-43.

638 112. Nightingale, K.K. et al. (2004) Ecology and transmission of *Listeria monocytogenes*
639 infecting ruminants and in the farm environment. Appl Environ Microbiol 70 (8), 4458-67.

640 113. Weis, J. and Seeliger, H.P. (1975) Incidence of *Listeria monocytogenes* in nature. Appl
641 Microbiol 30 (1), 29-32.

642 114. Maury, M.M. et al. (2019) Hypervirulent *Listeria monocytogenes* clones' adaption to
643 mammalian gut accounts for their association with dairy products. Nat Commun 10 (1), 2488.

644 115. Ortiz, S. et al. (2016) The Connection between Persistent, Disinfectant-Resistant *Listeria*
645 *monocytogenes* Strains from Two Geographically Separate Iberian Pork Processing Plants:
646 Evidence from Comparative Genome Analysis. Appl Environ Microbiol 82 (1), 308-17.

647 116. Harter, E. et al. (2017) Stress Survival Islet 2, Predominantly Present in *Listeria*
648 *monocytogenes* Strains of Sequence Type 121, Is Involved in the Alkaline and Oxidative
649 Stress Responses. Appl Environ Microbiol 83 (16).

650 117. Ryan, S. et al. (2010) A five-gene stress survival islet (SSI-1) that contributes to the
651 growth of *Listeria monocytogenes* in suboptimal conditions. J Appl Microbiol 109 (3), 984-
652 95.

653 118. Hingston, P. et al. (2017) Genotypes Associated with *Listeria monocytogenes* Isolates
654 Displaying Impaired or Enhanced Tolerances to Cold, Salt, Acid, or Desiccation Stress. Front
655 Microbiol 8, 369.

656 119. Esteban, J.I. et al. (2009) Faecal shedding and strain diversity of *Listeria monocytogenes*
657 in healthy ruminants and swine in Northern Spain. BMC Vet Res 5, 2.

658 120. Toledo-Arana, A. et al. (2009) The *Listeria* transcriptional landscape from saprophytism
659 to virulence. Nature 459 (7249), 950-6.

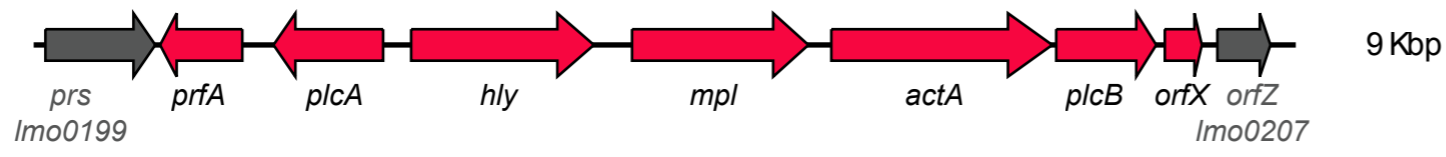
660

Outstanding questions

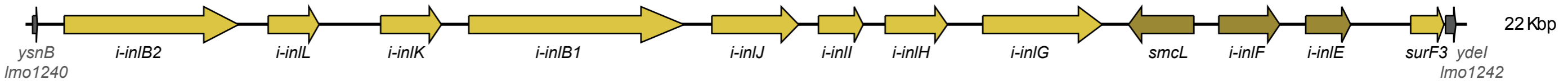
- What are the natural environments that have led to the current biodiversity of *L. monocytogenes*?
- What is the primary function of LIPI-1 genes, and other so-called “virulence genes”?
- What are the driving forces that have selected the virulence genes of *L. monocytogenes*?
- How does *L. monocytogenes* maintain its virulence since disease is not implicated in inter-host transmission?

(A)

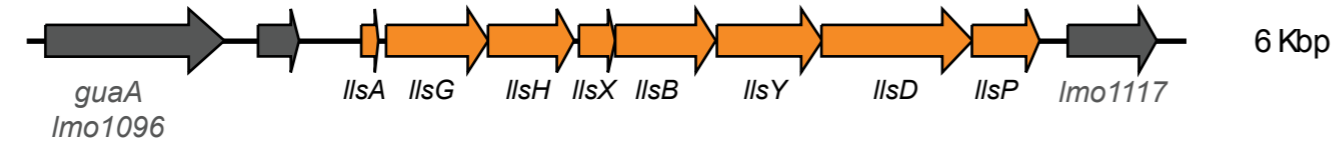
LIPI-1

Lm, *Liv* and atypical *Lin*

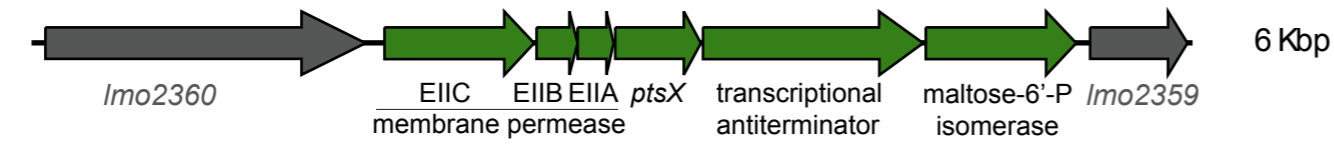
LIPI-2

Liv and atypical *Lm* CC33*

LIPI-3

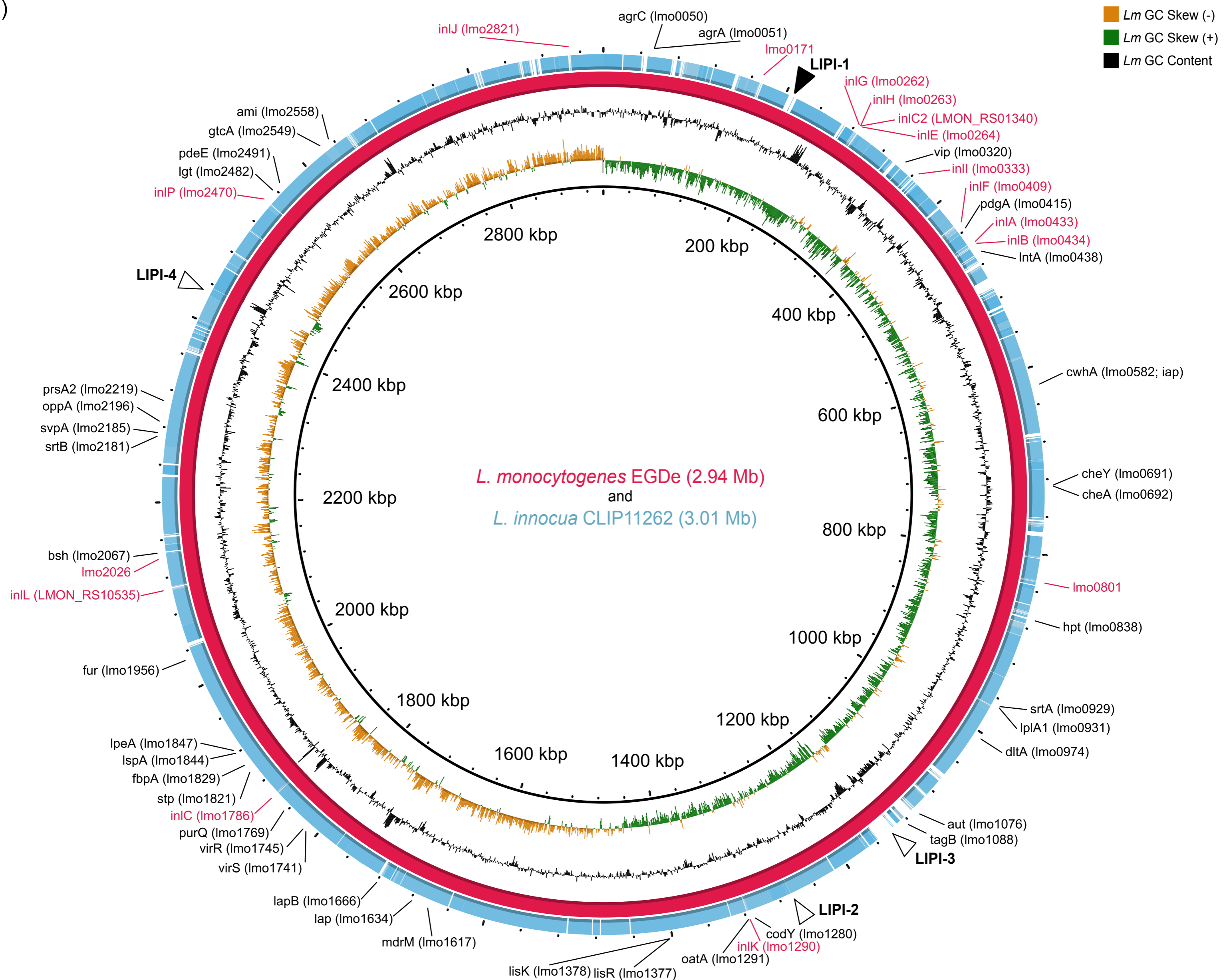
mainly *Lm* CC1, CC4, CC6 and some *Lin*

LIPI-4

mainly *Lm* CC4 and *Lin*

Scale: 1 Kbp

(B)



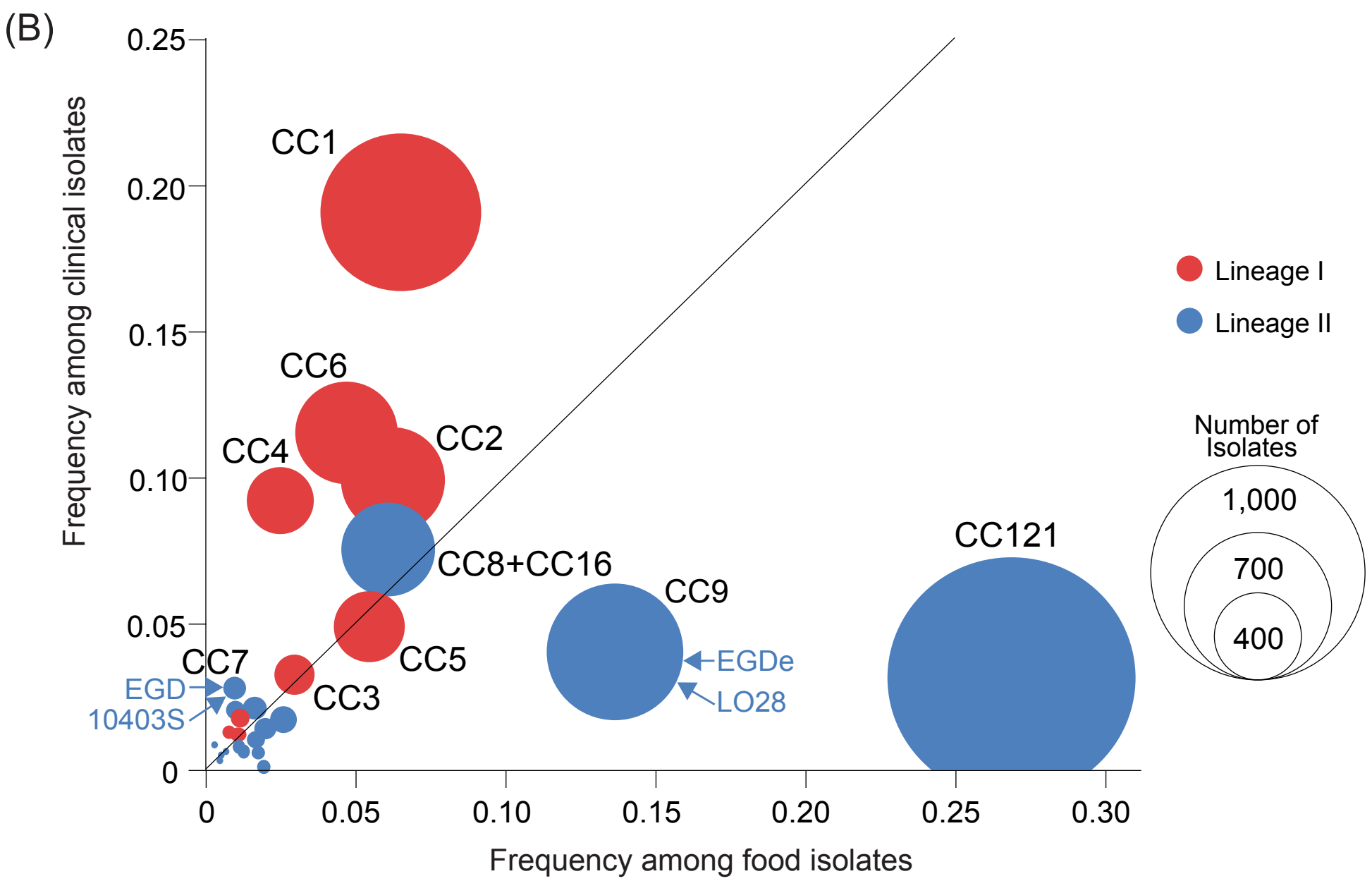
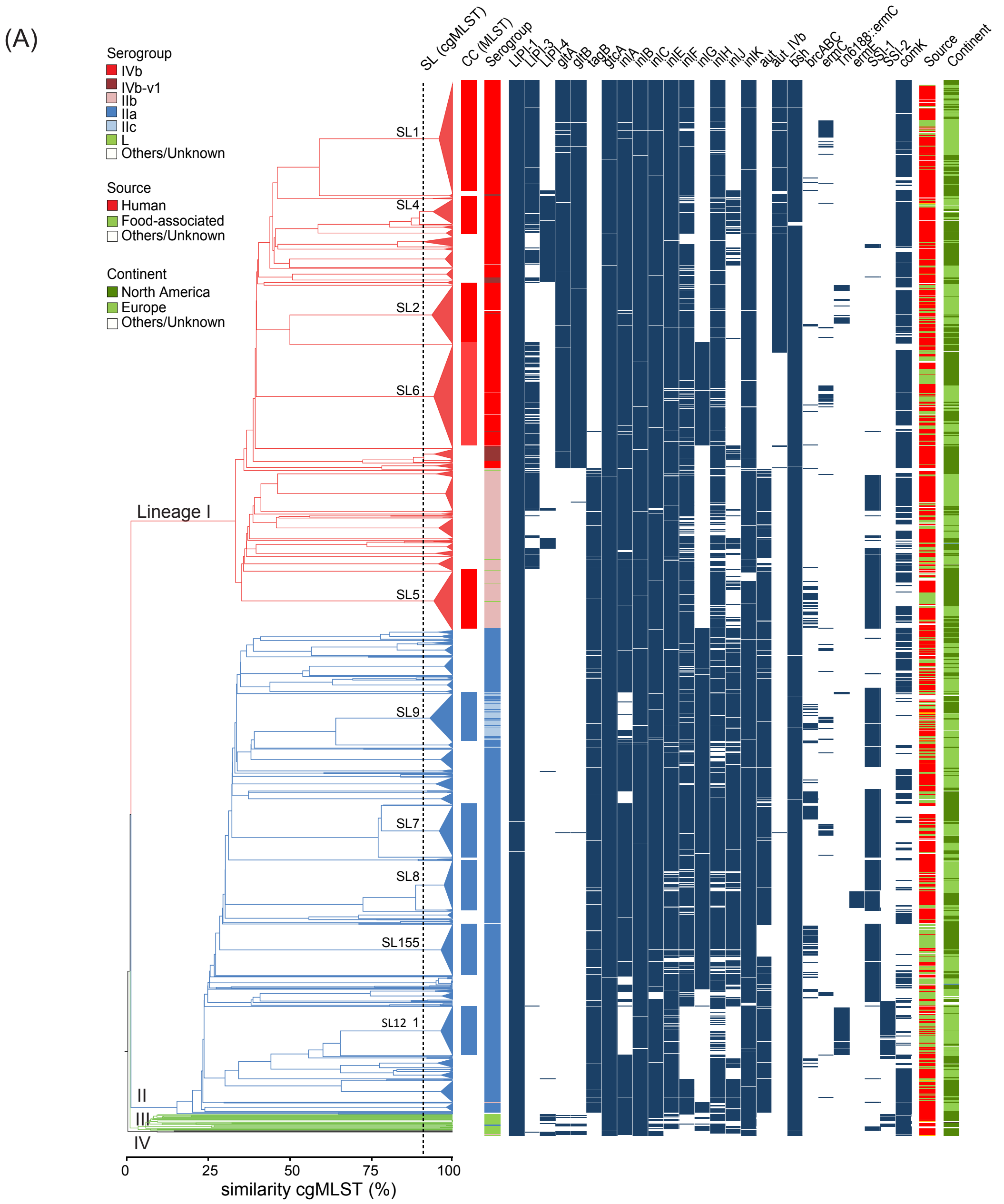


Figure 3



Host-association

Virulence
Active PrfA
Active LIPIs
Functional InIA
Fecal shedding
Transmission

Virulence

Immunodeficiency



Listeriosis depends on
microbial and host factors



Saprophytism

Loss of virulence
Inactive PrfA
Loss of LIPI-1 regulation
Truncation of InIA
Biofilm formation
Resistance to antiseptics