Timescale and genetic linkage explain the variable impact of defense systems on horizontal gene transfer

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Running title: Defense systems and horizontal gene transfer

Keywords: defense system, horizontal gene transfer, mobile genetic element, CRISPR-Cas, phage, plasmid, comparative genomics.

Abstract

Prokaryotes have evolved a wide repertoire of defense systems to prevent invasion by mobile genetic elements (MGE). However, because MGE are vehicles for the exchange of beneficial accessory genes, defense systems could consequently impede rapid adaptation in microbial populations. Here, we study how defense systems impact horizontal gene transfer (HGT) in the short and long terms. By combining comparative genomics and phylogeny-aware statistical methods, we quantified the association between the presence of 7 widespread defense systems and the abundance of MGE in the genomes of 196 bacterial and 1 archaeal species. We also calculated the differences in the rates of gene gain and loss between lineages that possess and lack each defense system. Our results show that the impact of defense systems on HGT is highly taxon- and system-dependent. CRISPR-Cas stands out as the defense system that most often associates with a decrease in the number of MGE and reduced gene acquisition. Timescale analysis reveals that defense systems must persist in a lineage for a relatively long time to exert an appreciable negative impact on HGT. In contrast, at short evolutionary times, defense systems, MGE, and gene gain rates tend to be positively correlated. Based on these results and given the high turnover rates experienced by defense systems, we propose that the inhibitory effect of most defense systems on HGT is masked by recent co-transfer events involving MGE.

Introduction

Gene exchange plays a key role in the adaption of microbes to changing environments, facilitating the spread of antibiotic resistance, pathogenicity factors, metabolic genes, and other accessory functions (Arnold et al. 2022). Over the last decade, there has been an increasing interest in assessing the ecological and genetic factors that control horizontal gene transfer (HGT) and determine the outcome of newly acquired genes in microbial populations (Soucy et al. 2015; Hall et al. 2020; Lee et al. 2022). HGT is often mediated by mobile genetic elements (MGE), such as phages, integrative and

38 conjugative elements, and plasmids, against which bacteria have evolved an elaborate repertoire of
39 defense systems (Doron et al. 2018; Botelho et al. 2023; Georjon and Bernheim 2023; Mayo-Munoz
40 et al. 2023; Shaw et al. 2023). As a result, large-scale patterns of HGT are shaped by an interplay of
41 ecological and genetic variables that underlie cross-strain and cross-species differences in
42 susceptibility to MGE (Haudiquet et al. 2022).

43 Recent studies have highlighted the role of CRISPR-Cas as widespread adaptive immunity systems
44 that protect archaea and bacteria against viruses and other MGE (van Vliet et al. 2021; Watson et al.
45 2024). While in vitro experiments have provided supportive evidence for this function (Marraffini and
46 Sontheimer 2008; O'Hara et al. 2017; Watson et al. 2018), questions remain about the extent to
47 which CRISPR-Cas systems constrain gene exchange in nature (Gophna et al. 2015; O'Meara and
48 Nunney 2019; Shehreen et al. 2019; Westra and Levin 2020; Wheatley and MacLean 2021; Pursey et
49 al. 2022). More generally, there is a paucity of research on how other defense systems, such as
50 restriction-modification (RM), abortive infection (Abi), and an expansive repertoire of recently
51 identified gene systems including Gabija, CBASS (cyclic oligonucleotide-based antiphage signaling
52 system), DMS (DNA modification-based systems), and DRT (defense-associated reverse
53 transcriptases) affect HGT in prokaryotes (Tesson et al. 2022; Costa et al. 2024).

54 In contrast with the expectation that defense systems restrict HGT by interfering with the
55 propagation of MGE, several empirical and theoretical observations suggest that the relation
56 between defense systems and HGT might be more complex. First, the selection pressure to maintain
57 defense systems in a population (and consequently their prevalence) generally increases with the
58 exposure to MGE (Oliveira et al. 2016; Meaden et al. 2022). Second, defense systems are mobilized
59 by MGE, which could lead to a trivial positive association with HGT rates. In a less trivial manner,
60 whole defense systems or parts of them are often encoded by MGE (Makarova et al. 2011; Pinilla-
61 Redondo et al. 2022; Botelho 2023), promoting the retention of the latter for their beneficial side-
62 effects in cellular defense (Koonin et al. 2020; Rocha and Bikard 2022).

63 Here, we investigated the association between 7 widespread defense systems and HGT rates in 197
64 prokaryotic species. By combining high-quality genomic data, phylogenomic methods, and
65 phylogeny-aware statistical inference, our study aimed to shed light on the nuanced consequences of
66 the interplay between defense systems and MGE on bacterial evolution across time scales.

67

68 **Results**

69 Association between defense systems and MGE abundance is MGE- and taxon-dependent

70 We used species-wise phylogenetic generalized linear mixed models (PGLMM) to study the
71 association between the presence or absence of the 7 most prevalent defense systems in the dataset
72 (RM, DMS, Abi, CRISPR-Cas, Gabija, DRT, and CBASS), genome size (measured as the total number of
73 genes), and the number of MGE per genome (Fig. 1, Tables S1-S4). Notably, the sign and magnitude
74 of the associations are strongly taxon-, system-, and MGE-dependent. Out of 197 species included in
75 the analysis, around 20-30 (depending on the defense system) displayed a statistically significant ($p <$
76 0.05) positive association between the presence of the defense system and genome size. In contrast,
77 statistically significant negative associations were only observed in 5-15 species (Fig. 1a, top row).
78 Because differences in the number of genomes per species could bias comparisons based on p-
79 values, we also explored an alternative criterion based on effect sizes to determine the number of
80 positive and negative associations (see Methods). Regardless of the criterion, CRISPR-Cas stood out
81 as the defense system that most often displays negative associations with genome size. In contrast,
82 other defense systems positively correlate with larger genomes in most of the species. The analysis
83 also reveals differences regarding the association between defense systems and distinct types of
84 MGE (Fig. 1a, middle rows). In all defense systems, negative associations with prophages are more
85 frequent than negative associations with plasmids. Once again, the clearest trend corresponds to
86 CRISPR-Cas, whose presence often correlates with a reduction in the number of prophages, but with
87 higher numbers of transposable elements and plasmids. Significant associations, when detected,

88 affect a sizeable fraction of the accessory genome, with 20-40% differences in MGE content between
89 genomes that do and do not harbor the defense system (Fig. 1b).

90 A more detailed analysis at the level of functional categories reveals that the presence of defense
91 systems is most often associated with changes in the number of genes from COG categories X
92 (mobilome), L (replication, recombination and repair), U (intracellular trafficking and secretion), K
93 (transcription), D (cell cycle control, cell division and chromosome partitioning), and V (defense) (Fig.
94 1 and S1). Genes from these functional categories are typically present in MGE, suggesting that
95 correlations (both positive and negative) between defense systems and genome size are primarily
96 due to differences in the abundance of MGE. We confirmed that by masking genomic regions that
97 correspond to MGE and rerunning the statistical analysis. As expected, 82% of the significant
98 associations disappeared after masking MGE (Fig S2). Although a few significant associations
99 persisted, a closer inspection revealed that those were often related to genes from prophages and
100 other MGE (such as phage satellites) that had not been originally identified as such and remained
101 unmasked.

102 The sign of the association between defense systems and MGE does not follow a clear taxonomic
103 trend (Fig. 2, S3 and S4), with opposite signs sometimes found in closely related species (see, for
104 example, the differences for CRISPR-Cas in *Phocaeicola vulgatus* and *P. dorei*). Moreover, the same
105 species often display opposite trends for different defense systems. For example, in *Pseudomonas*
106 *aeruginosa*, genomes with CRISPR-Cas contain fewer MGE, whereas genomes with CBASS, Gabija,
107 and RM systems are significantly enriched in MGE. Negative associations between CRISPR-Cas and
108 MGE are more abundant in the genus *Acinetobacter* (5 out of 8 species), the phylum *Bacteroidota*
109 (negative association in 8 species, positive association in a single species) and the class *Clostridia*, the
110 latter especially affecting prophages (negative association in 11 species, positive association in 2
111 species; Fig. S4). The genus *Acinetobacter* is also enriched in negative associations involving Abi (5
112 species) and CBASS (4 species). Furthermore, three almost non-overlapping groups of streptococci

113 display negative associations for different defense systems. These encompass *S. pyogenes*, *S.*
114 *gordonii*, *S. anginosus*, *S. mutans*, and *S. salivarius* in the case of CRISPR-Cas; *S. anginosus*, *S. oralis*, *S.*
115 *intermedius*, and *S. suis* in the case of RM and DMS; and *S. pyogenes*, *S. dysgalactiae*, *S. equi*, and *S.*
116 *uberis* in the case of Gabija.

117 The only archaeal species in the study (*Methanococcus maripaludis*) did not show any remarkable
118 trend, other than a relatively strong (but not statistically significant) positive association between
119 prophages and DMS/RM, and between transposons and Gabija, and a moderate negative association
120 between prophages and Gabija.

121 Associations between defense systems and MGE arise from differences in the rate of gene
122 acquisition

123 To investigate if correlations between defense systems and MGE involve cross-strain differences in
124 genome plasticity, we built high-resolution strain trees and identified clades that contain defense
125 systems (DEF^+). Then, we inferred the rates of gene gain and loss associated with different classes of
126 MGE in DEF^+ clades and in their respective sister groups lacking the defense system (DEF^-) using the
127 phylogenomic reconstruction tool GLOOME. We quantified the relative differences in gene gain and
128 loss by dividing the rates observed in DEF^+ clades by those from sister DEF^- clades. Finally, we
129 compared the resulting DEF^+/DEF^- gain and loss ratios between species in which MGE abundances
130 and defense systems are positively and negatively correlated. We found that DEF^+ and DEF^- clades
131 often differ in their gain rates but not in their loss rates (Fig. 3). Specifically, in species in which
132 defense systems are associated with increased MGE abundance, gene gain rates are typically higher
133 in clades that contain the defense system. The opposite (that is, a reduction of gene gain rates in
134 DEF^+ clades) is observed in species in which defense systems are associated with lower MGE
135 abundance. Among the latter, the biggest reductions in plasmid acquisition occur in association with
136 RM, DRT, DMS and Abi, whereas significant drops in prophage gain are observed for DRT, CBASS and
137 CRISPR-Cas. Taken together, these results confirm that correlations between defense systems and

138 MGE arise from cross-strain differences in gene gain rather than loss, as expected given the role of
139 MGE as facilitators of HGT.

140 Timescales and linkage determine the sign of associations between defense systems and HGT

141 Because prokaryotic defense systems are often located within or adjacent to MGE, we hypothesized
142 that positive associations between defense systems and MGE abundance could be explained, at least
143 in part, by recent co-transfer events (henceforth, we term this the "linkage hypothesis"). Uder this
144 hypothesis, positive associations would simply result from defense systems travelling together with
145 MGE.

146 To test the linkage hypothesis, we first verified that defense systems tend to be co-transferred with
147 MGE. We used ancestral reconstruction methods to identify the branches in which defense systems
148 and MGE were gained and lost along strain trees (Table S5). We found that >95% of defense
149 acquisition events occurred in branches in which MGE were also gained, and >90% of defense losses
150 occurred in branches in which MGE were also lost (Fig. 4a). The random expectation given the rates
151 of MGE gain and loss would be 71% and 63%, respectively (deviations with respect to these
152 expectations are statistically significant with $p < 10^{-8}$, binomial test). We also quantified the effect of
153 MGE gain and loss on the per-branch probability of gaining or losing defense systems. The probability
154 of acquiring a defense system is around 50-fold higher in branches in which MGE are gained than in
155 other branches (Fig. 4b-c; Fisher exact test $p < 10^{-8}$ for all defense systems). Similarly, the probability
156 of losing a defense system is 10- to 50-fold higher if MGE are also lost in the same branch (Fig. 4b-c;
157 Fisher exact test $p < 10^{-20}$ for all defense systems). These trends are observed even in very short
158 branches, spanning an evolutionary time of 10^{-6} substitutions per site in core genes (Fig. S5). To rule
159 out the possibility that these associations were due to the presence of incomplete genomes, we
160 separately considered terminal and internal tree branches. Because the dataset only includes
161 complete and nearly complete genomes, the absence of a small number of missing genes, if relevant,
162 would only affect the inference of gene gain and loss in terminal branches (those leading from the

163 immediate ancestor to each incomplete genome). Despite some quantitative differences, strong co-
164 acquisition (and co-loss) of MGE and defense occurs in both terminal and internal branches (Fig. S5),
165 confirming that these associations are genuine.

166 Although co-gain of MGE and defense systems along the same tree branch does not necessarily imply
167 a single event of joint gain, the strength of the association, the extremely short timespans, and the
168 fact that similar trends are observed for gene losses strongly suggest that concurrent gain and loss of
169 MGE and defense systems involve genetic linkage. (Alternative explanations in terms of strong
170 selective pressure to quickly acquire defense mechanisms upon exposure to MGE and lose them
171 once the MGE disappear might produce similar trends at the population level but not at the single-
172 genome level, whereas episodic increases in the overall rate of HGT leading to separate but
173 correlated acquisition of MGE and defense systems would not explain correlated losses.)

174 A major consequence of the co-transfer of MGE and defense systems is that the negative effects of
175 defense systems on HGT should be easier to detect at longer timescales or under evolutionary
176 conditions that weakened genetic linkage. To test that prediction, we studied how relative
177 differences in the rates of gene gain between DEF^+ and DEF^- clades depend on the depth of their last
178 common ancestor in the strain tree (note that the depth of the last common ancestor serves as an
179 upper limit for the time that the defense system has been retained in a lineage). As expected,
180 positive outliers (with much higher gene gain rate in the DEF^+ clade than in the DEF^- clade) almost
181 invariably correspond to very recent lineages (less than 10^{-5} substitutions per bp in nearly universal
182 core genes), indicating a very recent acquisition of the defense system (Fig. 5a and Table S6). In
183 contrast, negative outliers (with much lower gene gain rate in the DEF^+ clade than in the DEF^- clade)
184 generally correspond to deeper lineages (at least 0.001 substitutions per bp). We quantified these
185 trends by calculating the skewness of the distribution of DEF^+/DEF^- log-transformed gain ratios in
186 very recent and older lineages (Fig. 5b and Table S7). All the distributions show significant positive
187 skewness in recent lineages and significant negative skewness in older lineages (all $p < 0.001$ except

188 recent RM lineages, with $p = 0.023$; d'Agostino test for skewness). This quantitative analysis confirms
189 that differences in timescale affect all defense systems, with positive associations between defense
190 systems and gene gain being dominant in the short term and negative associations becoming more
191 frequent in the long term.

192 A second prediction of the linkage hypothesis is that the net effect of defense systems on HGT is not
193 only species specific, but also lineage specific. That is, defense systems may display positive, zero, or
194 negative association with HGT in different lineages depending on when the defense system was
195 acquired and how tight is the linkage to MGE. A more detailed study of gene gain rates in individual
196 species confirms that the effect of CRISPR-Cas is, indeed, lineage-specific, with the same species
197 encompassing recent CRISPR-Cas$^+$ lineages that display increased gene gain rates and older CRISPR-
198 Cas$^+$ lineages with reduced gene gain rates (Fig. 5c). This observation, combined with the very recent
199 acquisition of CRISPR-Cas in most lineages, could explain why many species show non-significant or
200 positive associations between CRISPR-Cas and MGE abundances. Indeed, of the six representative
201 species shown in Fig. 5c, only *Streptococcus anginosus* and *Acinetobacter radioresistens* show a net
202 negative association between CRISPR-Cas and MGE abundance in the PGLMM analysis.

203 The findings described so far indicate that, of the seven defense systems considered in this study,
204 CRISPR-Cas is the one that most often induces a net reduction in HGT rates. According to the linkage
205 hypothesis, this could be due to a comparatively weaker physical association between CRISPR-Cas
206 and MGE. To evaluate that possibility, we calculated the percentage of genes from CRISPR-Cas
207 located inside MGE and compared that to other defense systems (Fig. 5d). Consistent with the
208 linkage hypothesis, CRISPR-Cas is the defense system that is least often encoded by MGE (4.97% vs
209 8.12%, $p < 10^{-8}$, chi-squared test), followed by Gabija and DMS.

210 <u>Anti-CRISPR proteins modulate associations between CRISPR-Cas and HGT</u>

211 Anti-CRISPR proteins (Acr) have the potential to suppress the possible negative effect of CRISPR-Cas
212 on MGE-driven gene transfer (Mahendra et al. 2020). This, in turn, could contribute to explaining the

213 sign of the association between CRISPR-Cas and MGE in different lineages. To assess that possibility,
214 we searched all the genomes in the dataset for known Acr, finding 16,058 proteins. Then, we
215 compared the prevalence of Acr in species in which the presence of CRISPR-Cas is positively or
216 negatively correlated with the MGE content, separately considering genomes that do and do not
217 harbor CRISPR-Cas. Because Acr are generally encoded by MGE (Pinilla-Redondo et al. 2020) and the
218 prevalence of MGE systematically varies across groups acting as a confounding factor, we restricted
219 our comparisons to genomes that contain at least one prophage. Our results indicate that genomes
220 with and without CRISPR-Cas differ in the prevalence of Acr (Fig. S6). More importantly, these
221 differences have opposite directions in species in which CRISPR-Cas is positively and negatively
222 associated with MGE abundance. In the former, Acr are more prevalent in genomes that contain
223 CRISPR-Cas (p = 0.0012, chi-squared test). In the latter, Acr are less prevalent in genomes with
224 CRISPR-Cas ($p < 10^{-6}$, chi-squared test). In both groups, the prevalence of Acr in genomes without
225 CRISPR-Cas is similar. These results suggest that negative associations between CRISPR-Cas and HGT
226 could be dependent on (or at least facilitated by) low prevalence of Acr in the genome.

227

228 **Discussion**

229 Defense systems could have a significant impact on microbial evolution by effectively blocking the
230 transfer of MGE, reducing gene flow and limiting the spread of accessory genes. However, because
231 defense systems are often carried by MGE and these are the main vehicles of HGT, a net positive
232 association between defense systems and gene exchange cannot be ruled out *a priori*. We assessed
233 the relative weight of these two opposite scenarios by quantifying the association between defense
234 systems, MGE abundance, and gene acquisition rates in a phylogeny-aware comparative study of 197
235 prokaryotic species.

236 Our results shed light on previous, apparently contradictory findings concerning the effect of CRISPR-
237 Cas on genome evolution and diversification (Gophna et al. 2015; O'Meara and Nunney 2019;

238 Shehreen et al. 2019; Wheatley and MacLean 2021; Pursey et al. 2022). A pioneering study
239 conducted in 2015 found no evidence to support an overall association between CRISPR-Cas activity
240 and reduced gene acquisition via HGT at evolutionary time scales (Gophna et al. 2015). Such lack of
241 association was explained by several factors, including the high mobility of CRISPR-Cas systems, that
242 limits their long-term impact on host genomes, and the possibility that HGT is mediated by MGE that
243 escape (or are not targeted by) CRISPR-Cas immunity.

244 Our analyses support the general conclusion that CRISPR-Cas and other defense systems have little
245 overall impact on HGT in most bacterial species. Specifically, differences in the rates of gene
246 acquisition in lineages that do and do not harbor defense systems are centered around zero. That
247 said, we identified significant opposite trends at very short and intermediate evolutionary time
248 scales: positive associations between defense systems and HGT are more frequent at very short time
249 scales, whereas negative associations become dominant at longer time scales. These opposite trends
250 suggest that the actual effects of defense systems on gene exchange may be obscured by recent co-
251 transfer events involving MGE. As a result, the possible negative effects of defense systems on HGT
252 only become detectable if the defense system is maintained for long enough periods of time.

253 Besides this general picture, we identified some species in which the presence of defense systems
254 (especially CRISPR-Cas) significantly correlates with smaller genome sizes and MGE abundances.
255 Some of those associations had been previously described in *P. aeruginosa* and *Klebsiella*
256 *pneumoniae* (Wheatley and MacLean 2021; Botelho et al. 2023). And yet, these species represent
257 special cases rather than the rule, even in the context of host-associated bacteria. In fact, our results
258 underline that the association between defense systems and HGT is strongly system- and lineage-
259 dependent. This conclusion confirms and extends previous findings that showed that the impact of
260 CRISPR-Cas on the spread of antibiotic resistance is highly variable across species and its sign cannot
261 be easily explained by simple ecological, environmental, or genomic variables (Shehreen et al. 2019).

262 Among the 7 defense systems included in this study, CRISPR-Cas stands out for being the most
263 recurrently associated with reduced genome sizes and lower MGE (especially prophage) abundances.
264 In contrast, DMS, Abi, DRT, and CBASS are more often associated with higher numbers of MGE and
265 accessory genes. This finding is fully consistent with a recent study that compared 73 defense
266 systems in 12 bacterial species (Kogay et al. 2024). We propose that what makes CRISPR-Cas systems
267 different is their weaker (though still substantial) linkage with MGE. Compared to fully functional
268 CRISPR-Cas systems, other defense systems like Abi, Gabija, CBASS, DMS, and RM are more
269 frequently located within or next to MGE (Makarova et al. 2011; Benler et al. 2021; Rousset et al.
270 2022; Botelho 2023) and may have alternative functions related to MGE propagation. For example,
271 Abi systems have been identified in PICIs as accessory genes that facilitate their parasitic lifecycle
272 (Ibarra-Chavez et al. 2021). The narrow specificity of some defense systems may be another reason
273 why those systems do not significantly interfere with HGT. For instance, the GmrSD type IV RM
274 system selectively targets phages with glucosylated hydroxymethylcytosine (Bair and Black 2007) and
275 the Thoeris defense system only appears to be effective against myoviruses (Doron et al. 2018). This
276 caveat extends to any defense system based on epigenetic modifications, such as RM and DMS,
277 whose overall effect on HGT depend on the repertoire of epigenetic markers in the host and MGE
278 populations (Oliveira et al. 2016). Although selective targeting is often viewed as an outcome of
279 phage-host coevolution, it is tempting to speculate that it could have been evolutionarily favored by
280 the need to fight harmful genetic parasites while maintaining sufficiently high rates of HGT to
281 prevent population-level gene loss (Iranzo et al. 2016).

282 Defense systems sometimes exhibit synergistic interactions (Dupuis et al. 2013; Wu et al. 2024),
283 which could contribute to the heterogeneity of effects reported in this study. The vast number of
284 potential interactions and the limited availability of high-quality genomes made it unfeasible to
285 systematically account for the effect of interactions with the methods developed in this work.
286 Determining if and to what extent synergy among defense systems affects HGT remains a subject for
287 future investigation, possibly focused on a small set of experimentally validated interactions in highly

288 sequenced species. Another open question concerns which levels of detail, both taxonomic and
289 functional, best capture the effect of defense systems on HGT. From a taxonomic perspective,
290 working at or below the species level is a natural choice because species represent genetically
291 cohesive units (Bobay and Ochman 2017; Konstantinidis 2023; Conrad et al. 2024) and, as a result,
292 uncontrolled confounding factors are less likely to affect within-species than cross-species
293 comparisons. In contrast, more complex multi-level approaches would be required to detect trends
294 at higher taxonomic ranks. From a functional perspective, we grouped defense systems based on
295 their mechanism of action, under the assumption that functionally similar systems produce similar
296 effects on HGT. Though reasonable, this grouping criterion may not be optimal in systems in which
297 subtypes markedly differ in their eco-evolutionary dynamics and linkage with MGE. Moreover, fine-
298 grain dissection of highly abundant systems, such as RM and DMS, could help improve the sensitivity
299 of statistical tests by producing more balanced sets of strains with and without the subtypes of
300 interest.

301 All in all, we showed that some defense systems, especially CRISPR-Cas, can significantly reduce HGT,
302 although the effect is often masked by the fact that these systems travel together with MGE. Beyond
303 possible functional connections, the linkage between defense systems and MGE is an inevitable
304 consequence of the arms race between parasites and hosts. Because defense systems are costly and
305 their efficacy drops as parasites evolve, they are subject to rapid turnover and depend on HGT for
306 long-term persistence in microbial populations (van Houte et al. 2016; Iranzo et al. 2017; Koonin et
307 al. 2017; Puigbo et al. 2017). As a result, it is extremely challenging to disentangle the impact of
308 defense systems on gene flow from the causes that lead to their presence or absence, especially at
309 short evolutionary time scales. As more and more genomic data become available, we expect that
310 future research will overcome this challenge by quantifying the linkage between defense systems
311 and MGE, developing more realistic null models, and testing the role of defense systems on microbial
312 adaptation at different time scales.

313

314 **Methods**

315 Genome collection and identification of defense systems

316 We parsed the Genome Taxonomy Database (GTDB, https://gtdb.ecogenomic.org) release 202 (Parks
317 et al. 2020) to identify all high-quality genomes (according to the MIMAG criteria (Bowers et al.
318 2017)) with completeness >99%, contamination <1%, and contig count <500. The 82,595 genomes
319 that passed these filters were downloaded from the NCBI FTP site (https://ftp.ncbi.nlm.nih.gov).
320 CRISPR-Cas systems were identified with CRISPRCasTyper v1.2.4 (Russel et al. 2020) using default
321 parameters. We classified a genome as CRISPR-Cas$^+$ if it contains at least one high-confidence Cas
322 operon and CRISPR-Cas$^-$ otherwise. Only the species with >10 genomes and at least 5 CRISPR-Cas$^+$
323 genomes were further considered. To reduce the computational cost, we only considered a
324 maximum of 500 genomes per species. Species with >500 genomes were randomly subsampled to
325 keep at most 350 CRISPR-Cas$^+$ and 150 CRISPR-Cas$^-$ genomes. Other defense systems were identified
326 with Padloc v1.1.0 (db v1.4.0) (Payne et al. 2022). Our analysis focused on the most prevalent
327 defense systems: restriction-modification (RM), DMS, Abi, CRISPR-Cas, Gabija, DRT, and CBASS.

328 After applying these criteria, 19,323 genomes belonging to 196 bacterial and 1 archaeal species
329 (*sensu* GTDB) were included in the analysis (Tables S1 and S2). Of those, 2,964 correspond to
330 complete genomes and the rest to high-quality, nearly complete ones.

331 Gene prediction and annotation.

332 Open reading frames (ORF) were predicted with Prodigal v2.6.3, using codon table 11 (prokaryotic
333 genetic code) and "single" mode, as recommended for finished and draft quality genomes (Hyatt et
334 al. 2010). Orthologous ORF were then separately clustered for each species with Roary v3.13.0 (Page
335 et al. 2015) setting an 80% identity threshold for initial clustering followed by synteny-based
336 refinement (options '-t 11 -i 80'). The resulting gene clusters were functionally annotated by selecting

337 a representative sequence, arbitrarily chosen among those with length between 0.95 and 1.05 times
338 the average length of all sequences in the cluster. Representative sequences were functionally
339 annotated by mapping them to in-house profiles of the Clusters of Orthologous Genes (COG)
340 database (2020 release) (Galperin et al. 2021) with HMMER v3.1b2 (e-value < 0.001)
341 (http://hmmer.org). The 26 major prokaryotic functional categories defined in the COG database
342 were assigned to the annotated genes. Some functional categories (A, RNA processing and
343 modification; B, chromatin structure and dynamics; W, extracellular structures; T, signal
344 transduction; and Z, cytoskeleton) were excluded since they rarely or never occur in prokaryotic
345 genomes. The case-insensitive keywords "phage", "plasmid" and "transpos*" in the COG gene
346 annotations were used to identify genes associated with prophages, plasmids, and transposons,
347 respectively, and the resulting gene lists were manually curated to minimize false assignments. We
348 based our statistical analyses on marker gene counts rather than full MGE counts because the latter
349 are more susceptible to technical artifacts (e.g., different heuristics to deal with nested MGE will
350 affect the number of MGE, but not the number of MGE marker genes). Gene counts per genome and
351 functional category are listed in Table S8. Anti-defense proteins, including anti-CRISPR, were
352 identified by running HMMER v3.1b2 (e-value < 10^{-10}) against the dbAPIS database (Yan et al. 2024).

353 Identification of genomic regions containing mobile genetic elements

354 Prophages were detected with Phispy v4.2.21 using default options (Akhter et al. 2012). Short
355 transposons were identified based on the presence of isolated or paired genes annotated as
356 transposases. In the latter case, we allowed for up to one additional gene between two transposon-
357 related genes to account for the genetic architecture of some insertion sequences (Gomez et al.
358 2014). Due to the inherent difficulty to discriminate among plasmids, ICE, and IME (Botelho et al.
359 2023), we restricted the search for these elements to complete genomes. Plasmids were identified as
360 extra-chromosomal replicons that contain the "plasmid" label in their NCBI description lines. After
361 removing those replicons, ICEfinder (Liu et al. 2019) was used to identify ICE and IME. Finally, we ran

362 Phispy v4.2.21 with option '-phage_genes 0' to identify any other MGE, integrons, pathogenicity
363 islands, and fragments of MGE that could have been missed by the previous approaches. The MGE
364 identified through these approaches were masked from complete genomes to produce the gene
365 counts in Table S9 and the results shown in Fig S2.

366 Species trees

367 Phylogenetic trees were separately built for each species based on the set of 120 prokaryotic marker
368 genes (122 in the case of Archaea) proposed by the GTDB r202. For each species, only those marker
369 genes with prevalence >80% were used for phylogenetic reconstruction. We aligned the amino acid
370 sequences of each marker gene with mafft-linsi (L-INS-I algorithm, default options, MAFFT v7.475)
371 (Katoh and Standley 2013) and back-translated the amino acid alignments to nucleotide alignments
372 with pal2nal.pl v14 (Suyama et al. 2006) using codon table 11. After concatenating all nucleotide
373 alignments, we built preliminary trees with FastTree v2.1.10 (options '-gtr -nt -gamma -nosupport -
374 mlacc 2 -slownni') (Price et al. 2010). The tree topologies produced by FastTree were subsequently
375 provided to RaxML v8.2.12 (Stamatakis 2014) for branch length optimization (raxmlHPC with options
376 '-f e -m GTRGAMMA'). The final trees are included in Supplementary File S1.

377 To visualize trends across species (Fig. 2, S3, and S4), we used the online tool iTOL (Letunic and Bork
378 2021) and the multispecies tree from GTDB r202.

379 Phylogenetic generalized linear mixed models (PGLMM)

380 For each genome in the dataset, we collected the following response variables: the total number of
381 genes, the number of genes belonging to each functional category, and the number of genes
382 associated with prophages, plasmids, and transposons (Table S8). Genes that belong to the 7 defense
383 systems of interest were excluded when computing these values. Then, for each response variable,
384 we fitted a PGLMM with Poisson distribution and canonical link function, using the presence or
385 absence of each defense system as predictors and the species trees as guides to generate the
386 covariance matrix.

387 For each species, the PGLMM assumes that the response variable, Y_i, follows a Poisson distribution
388 with mean μ_i, that is, $Y_i \sim \mathrm{Poisson}(\mu_i)$. The expected gene abundance, μ_i, is modeled as
389 $\log \mu_i = \beta_0 + \sum \beta_j X_{ij} + \epsilon_i$, where β_0 is the intercept, β_j is the coefficient associated with the defense
390 system j, and $X_{ij} \in \{0,1\}$ denotes the absence or presence of defense system j in genome i. The
391 random effects ϵ_i follow a multivariate normal distribution, $\epsilon \sim \mathrm{Gaussian}(0, \sigma^2_{phy}\mathbf{C})$, where σ^2_{phy} is the
392 strength of the phylogenetic signal and $\mathbf{C}$ is a covariance matrix derived from the phylogenetic tree
393 under the assumption of Brownian motion evolution.

394 To extend the PGLMM to multiple species, we considered that the effect of defense systems on gene
395 content can be species-dependent (this was a reasonable assumption *a priori* and later confirmed by
396 the analysis). To account for that, the multi-species model must include an interaction term "defense
397 × species", whose coefficients are relevant per se, and, accordingly, modeled as fixed effects.
398 Moreover, because HGT rates are highly variable among species (Puigbò et al. 2014; Iranzo et al.
399 2019), it makes little sense to extend the phylogenetic correction beyond single species or assume
400 that the strength of the phylogenetic signal is the same for all species. These considerations are
401 captured by a phylogenetic covariance matrix with block-diagonal structure (one block per species),
402 and species-wise values of the phylogenetic coefficient (one for each block of the phylogenetic
403 covariance matrix). In practice, fitting a multi-species model with these specifications is formally
404 equivalent to fitting independent models for each species. The latter approach, that we adopted, has
405 the advantage of being more suitable for parallelization and requiring fewer computational
406 resources. Thus, for each species and response variable, we fitted a PGLMM with the function
407 pglmm_compare(response_variable ~ Abi + CBASS + CRISPR + DMS + DRT + Gabija + RM, family =
408 "poisson", data = SpData, phy = SpTree) from the R package phyr v1.1.0 (Li et al. 2020). In the
409 formula, Abi, CBASS, CRISPR, and so on, are binary variables representing the presence (1) or absence
410 (0) of each defense system. Table S4 presents the coefficients, p-values, and goodness of fit of the
411 model.

412 The model described above includes nine coefficients (intercept, seven defense systems, and the
413 phylogenetic signal), which could lead to overfitting in species in which the number of available
414 genomes is limited. As an alternative, we also fitted seven separate PGLMM, one for each defense
415 system, involving a single predictor and the phylogenetic random effect (Table S3). This approach
416 does not account for correlations among defense systems, but it has the advantage of not being
417 affected by overfitting. The figures in the manuscript are based on this second set of models,
418 although, in practice, both approaches produce very similar quantitative results.

419 For each class of PGLMM, we also fitted non-phylogenetic models in which the covariance matrix $\mathbf{C}$ of
420 the random effect was replaced by the identity matrix. The PGLMM were compared to their non-
421 phylogenetic counterparts using the conditional Akaike Information Criterion (cAIC) as previously
422 described (Greven and Kneib 2010; Säfken et al. 2021). Based on the cAIC, PGLMM performed better
423 than non-phylogenetic models in 65% of the species-response-defense triplets and were close to
424 non-phylogenetic models (ΔcAIC < 2) in another 25% of the triplets (in most of those cases, the
425 strength of the phylogenetic signal was close to zero, which made both phylogenetic and non-
426 phylogenetic models equivalent).

427 To account for the possibility that other (less abundant) defense systems could explain part of the
428 variability in the results, we explored a more complex set of models that included the total number
429 of other defense systems as an additional predictor. These models generally performed worse than
430 their simpler variants (ΔcAIC > 0 in 83% of the species) and were not considered for further analysis.

431 Large differences in sample size among species and defense systems translate into unequal precision
432 in the estimation of the model coefficients. To deal with that limitation, we used two different
433 criteria to identify species in which the presence of a defense system is positively or negatively
434 associated with gene numbers. For one option, we adopted a classical criterion of statistical
435 significance ($p < 0.05$) for the predictor variable in the PGLMM. For the other option, we applied an
436 alternative criterion based on effect size, aimed at comparing species with different sample sizes in

437 which p-values are not commensurable. Specifically, for each variable and defense system, we jointly
438 considered the PGLMM of all the species and determined the smallest effect size (in absolute value)
439 that reached statistical significance ($p < 0.05$) in any species. Then we used the smallest significant
440 effect size (SSES) as a threshold to classify associations as positive, negative, or null.

441 Inference of gene gain and loss

442 Gene gains and losses at each branch of each species tree were estimated with Gloome (Cohen and
443 Pupko 2010), using as inputs the gene presence/absence matrices previously generated by Roary and
444 the species trees. The parameter configuration file was set to optimize the likelihood of the observed
445 phyletic profiles under a genome evolution model with 4 categories of gamma-distributed gain and
446 loss rates and stationary frequencies at the root.

447 Comparison of gene gain and loss rates between DEF^+ and DEF^- clades

448 For each species tree, we defined DEF^+ clades as the narrowest possible clades such that at least 80%
449 of the leaves contain the defense system of interest. Candidate DEF^- clades were defined in an
450 analogous way, referring to leaves without the defense system. Next, we identified pairs of DEF^+ and
451 DEF^- clades that constitute sister groups. Sister $DEF^{+/-}$ pairs were excluded if both clades contained a
452 single genome. For each clade in a valid pair, we computed the overall gene gain and loss rates as the
453 expected number of gene gains (or losses) in that clade divided by the total branch length. Gain and
454 loss rates for different functional categories and MGE we calculated in an analogous way but
455 restricting the sum of gene gains and losses to the genes of interest. When calculating overall and
456 category-wise gain and loss rates, we did not take into account the contribution of species-wise
457 singletons (genes without homologs in other genomes of the same species), as they may represent
458 false gene predictions or genes that are replaced at unusually high rates (Wolf et al. 2016). To
459 account for the non-negative nature of gain and loss rates and their heavy-tailed distributions,
460 comparisons between DEF^+ and DEF^- sister branches were done based on log-transformed rate
461 estimates. To calculate the skewness of the distributions and their statistical significance, we use the

462 method proposed by D'Agostino (D'Agostino et al. 1990) as implemented by the
463 scipy.stats.skewtest() function in Python (Virtanen et al. 2020).

464 **Data access**

465 All the data generated or analyzed during this study are included in this published article and its
466 supplementary information files.

467 **Competing interests**

468 The authors declare no competing financial interests.

469 **Acknowledgements**

470 Y.L. is supported by China Scholarship Council (No.202008440425). J.B. is supported by the Maria
471 Zambrano grant of the Spanish Ministry of Universities (Grant No. UP2021-035), and the Severo
472 Ochoa Program for Centres of Excellence in R&D of the Agencia Estatal de Investigación of Spain
473 (Grant No. CEX2020-000999-S (2022–2025) to the CBGP). J.I is supported by the Ramón y Cajal
474 Programme of the Spanish Ministry of Science (Grant No. RYC-2017–22524); the Agencia Estatal de
475 Investigación of Spain (Grant Nos. PID2019-106618GA-I00 and CNS2023-145430), the Severo Ochoa
476 Programme for Centres of Excellence in R&D of the Agencia Estatal de Investigación of Spain (Grant
477 No. SEV-2016–0672 (2017–2021) to the CBGP); and the Comunidad de Madrid (through the call
478 Research Grants for Young Investigators from Universidad Politécnica de Madrid, Grant No.
479 M190020074JIIS).

480 We thank Jorge Calle-Espinosa for helpful discussions.

481

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652

653

654 **Figure legends**

655 **Figure 1: Association between 7 widespread defense systems, total number of genes, and MGE**
656 **abundance.** (a) Number of species displaying positive or negative associations in a phylogenetic
657 generalized linear mixed effects model (see Methods) according to two different criteria: $p < 0.05$
658 and absolute effect size greater than the smallest significant effect size (|ES|>SSES), separately
659 computed for each response variable and defense system. The top row (Total) indicates the
660 association with the total number of genes. The association with MGE was calculated based on
661 marker genes for prophages (Ph), plasmids (Pl), and transposons (Tr). Abbreviations of functional
662 categories: X (mobilome), L (replication, recombination and repair), U (intracellular trafficking and
663 secretion), K (transcription), D (cell cycle control, cell division and chromosome partitioning), and V
664 (defense). (b) Effect sizes, measured as relative differences in gene and MGE abundances. Each point
665 corresponds to one species. Species with values beyond the axis limits are collapsed in a single point
666 with size proportional to the number of species. Vertical lines indicate the median over all the
667 species that show positive or negative association according to the p-value and SSES criteria.

668 **Figure 2: Taxonomic distribution of species displaying positive and negative associations between**
669 **the presence of defense systems and the number of genes from the mobilome** (based on COG
670 annotations). Taxa discussed in the text are labeled: A, *Acinetobacter*; Pa, *Pseudomonas aeruginosa*;
671 Pv, *Phocaeicola vulgatus*; Pd, *Phocaeicola dorei*; S1, *Streptococcus pyogenes*, *S. dysgalactiae*, *S. equi*,
672 and *S. uberis*; S2, *S. gordonii*, *S. anginosus*, *S. mutans*, and *S. salivarius*; S3, *S. oralis*, *S. intermedius*,
673 and *S. suis*. The classes *Bacilli* and *Clostridia* are the major components of the Genome Taxonomy
674 Database phyla "Bacillota" and "Bacillota_A". See figure S3 for a larger version including all species
675 names.

676 **Figure 3: Relative differences in the rates of gene gain and loss between sister clades that do and**
677 **do not harbor defense systems** (DEF^+ and DEF^-, respectively). The boxplots represent the distribution
678 of the DEF^+/DEF^- ratio of gene gain (or loss) rates for each class of MGE, calculated for every pair of

679 sister clades. Values greater (or smaller) than 1 indicate increased (or reduced) gene flux in lineages
680 that contain the defense system. (a) Species that show a negative association between the defense
681 system and the number of marker genes for each class of MGE (PGLMM with smallest significant
682 effect size criterion). (b) Species that show a positive association between the defense system and
683 the number of marker genes for each class of MGE. In the boxplots, the central line indicates the
684 median, the box limits correspond to the 25 and 75 percentiles, and the whiskers extend to the
685 largest and smallest values not classified as outliers. P-values are based on Wilcoxon test with log-
686 ratio = 0 as null hypothesis.

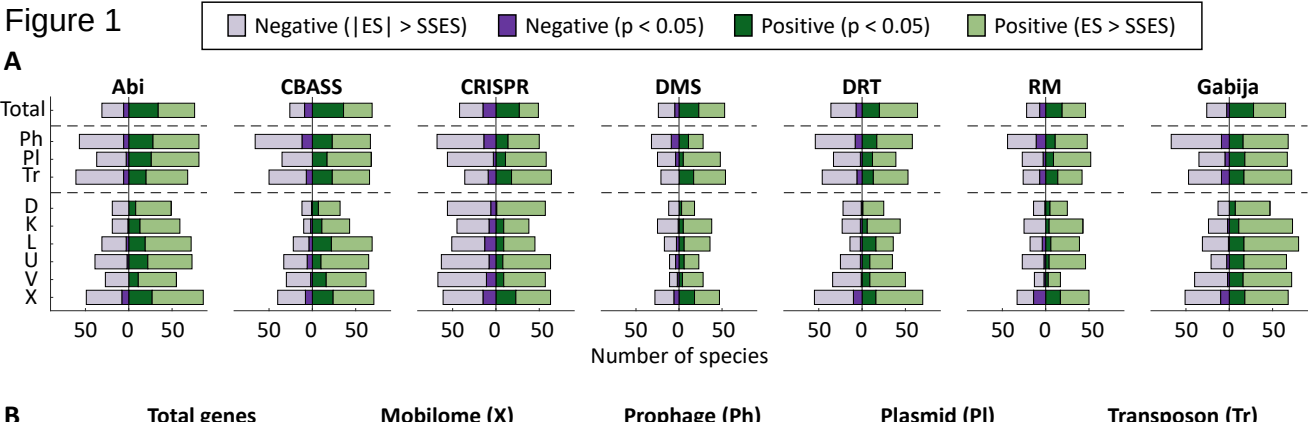
687 **Figure 4: Co-occurrence of defense system gains and losses and MGE gains and losses along the**
688 **phylogeny.** (a) Percentage of DEF gains (and losses) that occur in the same branch as an MGE gain (or
689 loss). (b) Conditional probability of gaining (or losing) a defense system provided that an MGE is also
690 gained (or lost) in the same branch, compared to the conditional probabilities when an MGE is not
691 acquired (or lost) in the same branch. (c) Effect of MGE gain (or loss) on the per branch probability to
692 acquire (or loss) a defense system, measured as a risk ratio. Error bars in (a) and (b) correspond to
693 95% confidence intervals based on the binomial distribution. Error bars in (c) indicate the 95%
694 confidence intervals for the relative risk (Morris and Gardner 1988).

695 **Figure 5: Association between defense systems and gene gain rates depends on the time scale.** (a)
696 Ratio of overall gene gain rates between sister clades that do and do not harbor defense systems
697 (DEF^+ and DEF^-, respectively). Values greater (or smaller) than 1 indicate increased (or reduced) gene
698 flux in lineages that contain the defense system. (b) Density distribution of the DEF^+/DEF^- gain ratios,
699 comparing clades that represent recent acquisitions of the defense system (depth < 10^{-5} substitutions
700 per site in core genes) and clades that have retained the defense system for longer times (depth > 10^{-}
701 ^{3} substitutions per site). The probability density function (y-axis) is represented in logarithmic scale to
702 facilitate the visualization the tails. The skewness of all distributions is statistically significant (Table
703 S7), with positive values for recent clades and negative values for deeper clades (c) Representative

704 examples of gene gain ratios in shallow and deeper sister groups from the same species. (d)
705 Percentage of genes from different defense systems located within known MGE. Whiskers represent
706 95% confidence intervals based on the binomial distribution.

707

Figure 1


Negative (|ES| > SSES)
Negative (p < 0.05)
Positive (p < 0.05)
Positive (ES > SSES)
A
Abi
CBASS
CRISPR
DMS
DRT
RM
Gabija
Total
Ph
Pl
Tr
D
K
L
U
V
X
50 0 50
Number of species

B
Total genes
Mobilome (X)
Prophage (Ph)
Plasmid (Pl)
Transposon (Tr)
Abi
CBASS
CRISPR
DMS
DRT
RM
Gabija
-10 -5 0 5 10
-40 -20 0 20 40
Relative change (%)

Figure 2

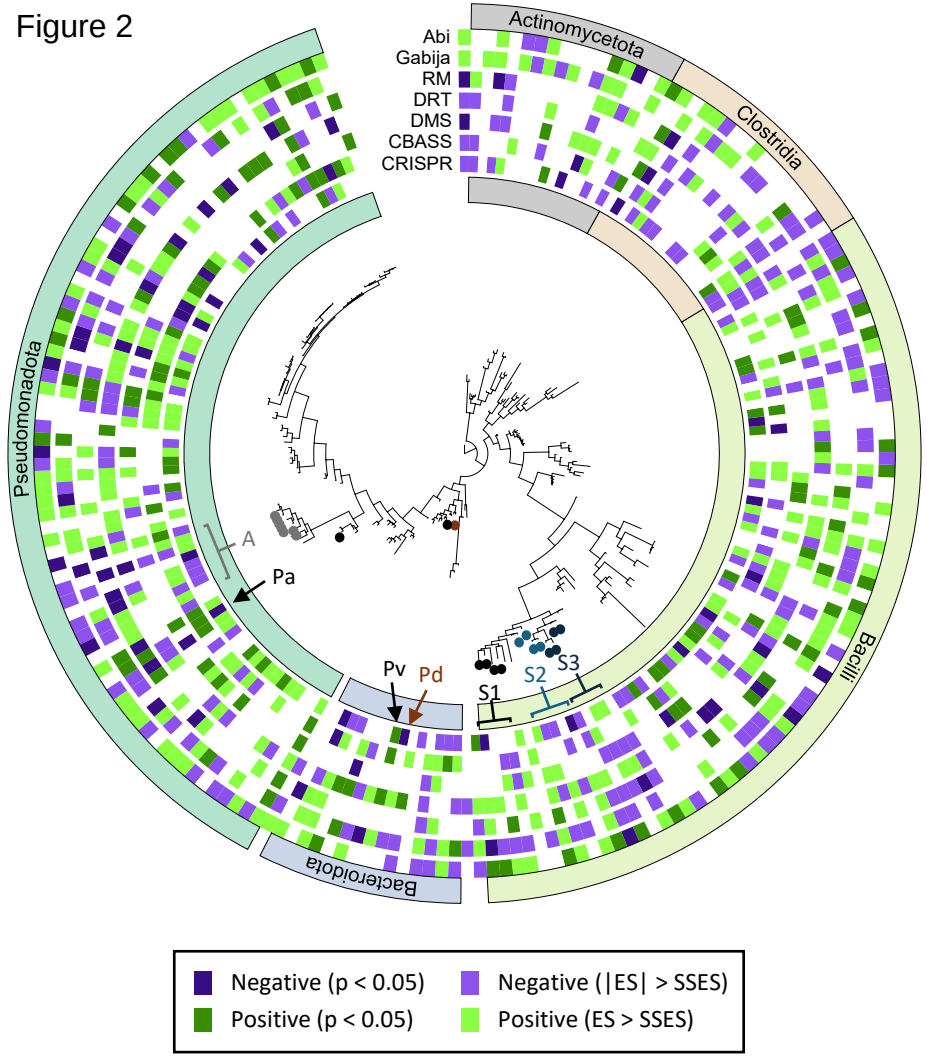

Abi
Gabija
RM
DRT
DMS
CBASS
CRISPR
Actinomycetota
Clostridia
Bacilli
Bacteroidota
Pseudomonadota
A
Pa
Pv
Pd
S1
S2
S3
Negative (p < 0.05)
Positive (p < 0.05)
Negative (|ES| > SSES)
Positive (ES > SSES)

Figure 3

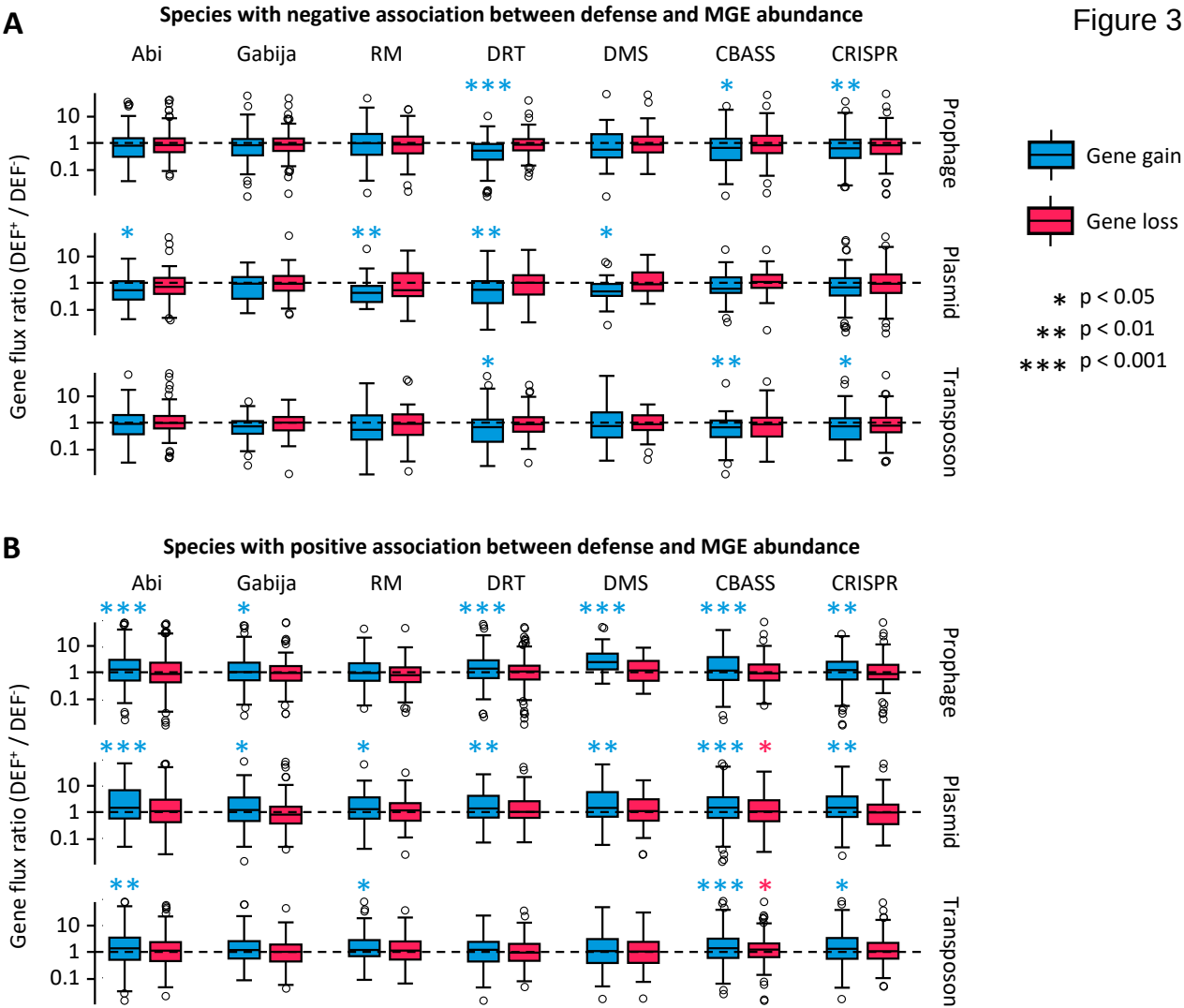

A
Species with negative association between defense and MGE abundance
Abi
Gabija
RM
DRT
DMS
CBASS
CRISPR
Prophage
Plasmid
Transposon
Gene flux ratio (DEF+ / DEF-)
10
1
0.1
Gene gain
Gene loss
* p < 0.05
** p < 0.01
*** p < 0.001
B
Species with positive association between defense and MGE abundance

Figure 4

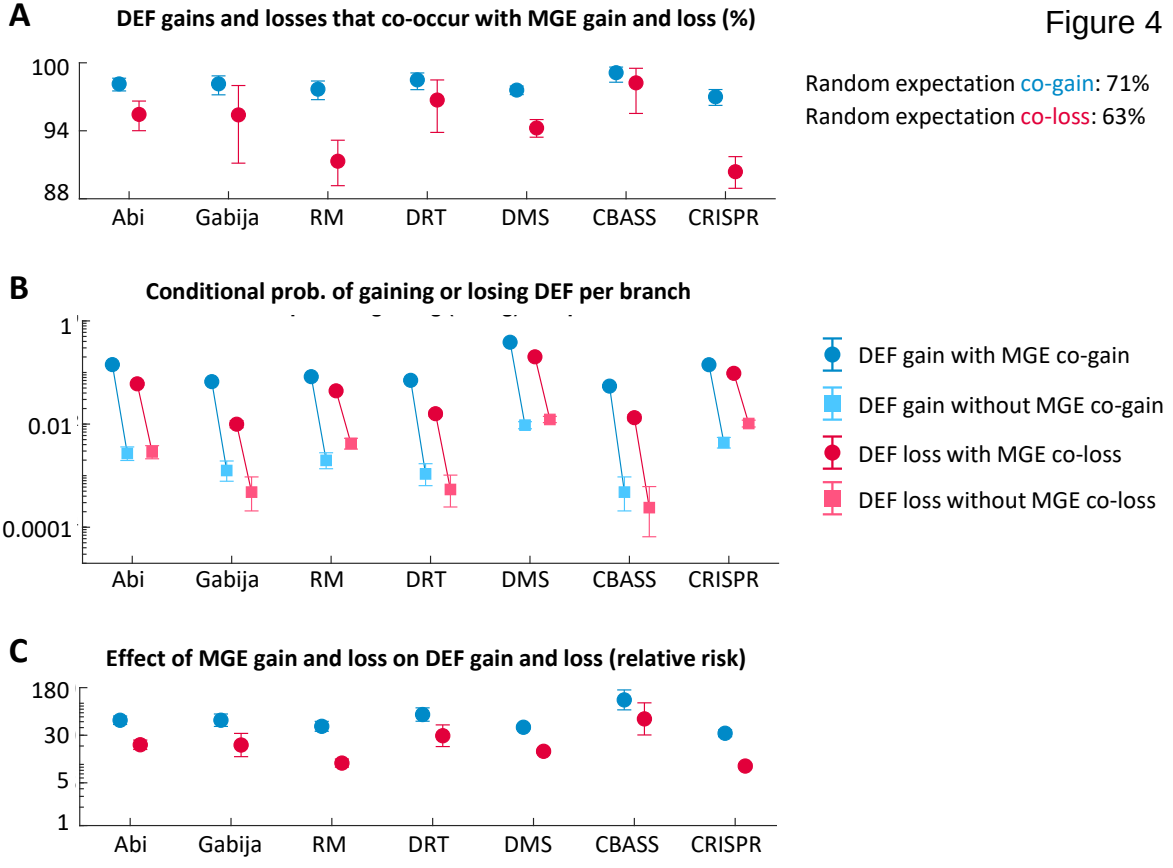
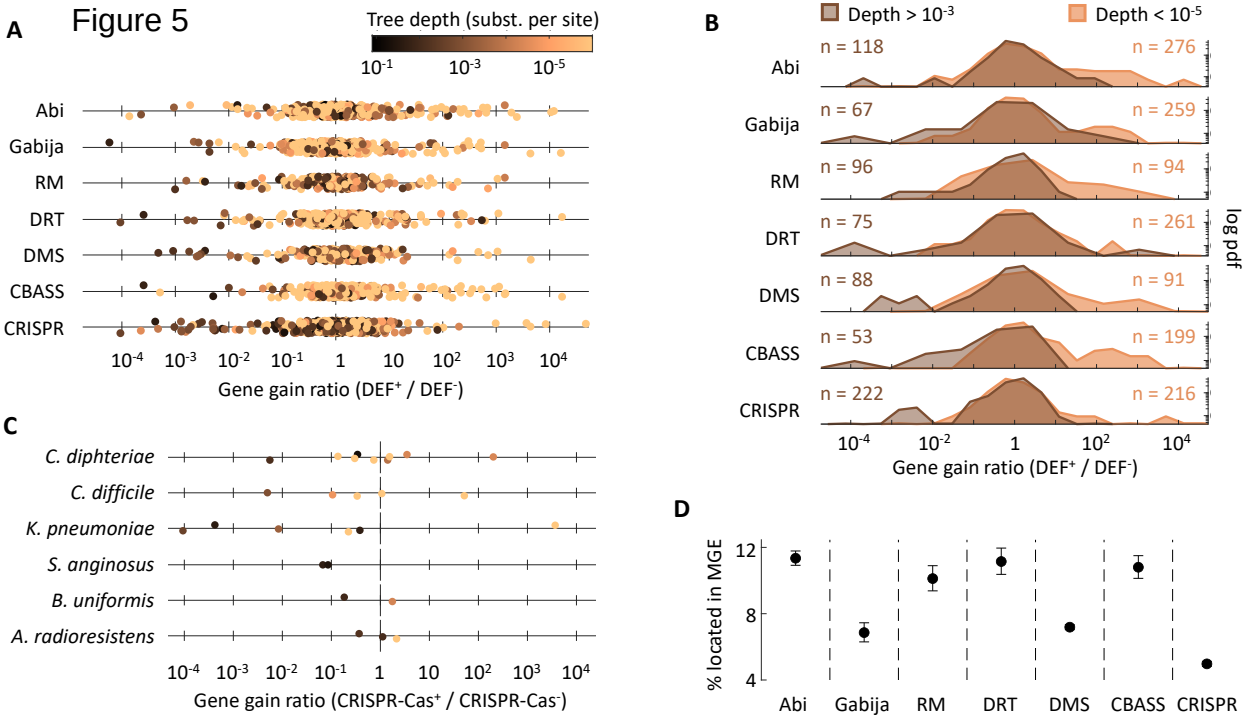

A
DEF gains and losses that co-occur with MGE gain and loss (%)
100
94
88
Abi
Gabija
RM
DRT
DMS
CBASS
CRISPR
Random expectation co-gain: 71%
Random expectation co-loss: 63%
B
Conditional prob. of gaining or losing DEF per branch
1
0.01
0.0001
Abi
Gabija
RM
DRT
DMS
CBASS
CRISPR
DEF gain with MGE co-gain
DEF gain without MGE co-gain
DEF loss with MGE co-loss
DEF loss without MGE co-loss
C
Effect of MGE gain and loss on DEF gain and loss (relative risk)
180
30
5
1
Abi
Gabija
RM
DRT
DMS
CBASS
CRISPR

Figure 5


A
Tree depth (subst. per site)
10-1
10-3
10-5
Abi
Gabija
RM
DRT
DMS
CBASS
CRISPR
10-4 10-3 10-2 10-1 1 10 102 103 104
Gene gain ratio (DEF+ / DEF-)
B
Depth > 10-3
Depth < 10-5
n = 118
n = 276
n = 67
n = 259
n = 96
n = 94
n = 75
n = 261
n = 88
n = 91
n = 53
n = 199
n = 222
n = 216
log pdf
10-4 10-2 1 102 104
Gene gain ratio (DEF+ / DEF-)
C
C. diphteriae
C. difficile
K. pneumoniae
S. anginosus
B. uniformis
A. radioresistens
10-4 10-3 10-2 10-1 1 10 102 103 104
Gene gain ratio (CRISPR-Cas+ / CRISPR-Cas-)
D
% located in MGE
12
8
4
Abi
Gabija
RM
DRT
DMS
CBASS
CRISPR