An outstanding bacterial membranome requires CRISPR-Cas systems to avoid the intervention of phages

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Abstract

Antimicrobial resistance is widely recognized as a serious global public health problem. To combat this threat, a thorough understanding of bacterial genomes is necessary. The current wide availability of bacterial genomes provides us with an in-depth understanding of the great variability of dispensable genes and their relationship with antimicrobials. Some of these accessory genes are those involved in CRISPR-Cas systems, which are acquired immunity systems that are present in part of bacterial genomes. They prevent viral infections through small DNA fragments called spacers. But the vast majority of these spacers have not yet been associated with the virus they recognize, and this has been named CRISPR dark matter (spacers without associated recognized sequence). By analyzing the spacers of tens of thousands of genomes from six bacterial species highly resistant to antibiotics, we have been able to reduce the CRISPR dark matter from 80-90% to 15% in some cases. In addition, we have observed that, when a genome presents CRISPR-Cas systems, this is accompanied by particular collections of membrane proteins. Our results suggest that when a bacterium presents membrane proteins that make it compete better in its environment, and these proteins are in turn receptors for specific phages, it would be forced to acquire CRISPR-Cas immunity systems to avoid infection by these phages.

Keywords

CRISPR-Cas, membrane, phage, pangenome, ESKAPE, machine learning

Introduction

The fight against antimicrobial resistant bacteria is one of the major challenges facing mankind in the near future (1). The World Health Organization prioritized a list of multidrug-resistant bacteria in 2017 to support research and development of effective drugs (2). The most relevant species in this list form the so-called ESKAPE group, whose acronym refers to two Gram-positive bacteria (*Enterococcus faecium* and *Staphylococcus aureus*) and four Gram-negative bacteria (*Klebsiella pneumoniae*, *Acinetobacter baumannii*, *Pseudomonas aeruginosa*, and *Enterobacter cloacae*). All of them cause infections, some of which are acquired in the hospital environment (3, 4). A major difference between Gram-positive and Gram-negative bacteria, which conditions the way to fight against them, is that Gram-negative bacteria have a second lipid membrane outside the cell wall, which gives them a richer variability of membrane proteins, such as the outer membrane proteins found in this second layer (5).

The CRISPR-Cas are adaptive immunity systems found in most archaea and in less than half of the bacteria sequenced (6). They provide acquired immune resistance against phages and other foreign

nucleic acid molecules such as plasmids, thus restricting gene transfer (7). There are different types of CRISPR-Cas systems based on genes that are part of the different steps of this immune system (adaptation or spacer integration, expression, and interference) and are generically called *cas* (CRISPR-associated genes). ESKAPE bacteria have CRISPR-Cas systems of the most common types, from I to IV, but only in a minimal number of strains, with frequencies ranging from less than 1% to 60% of genomes, depending on the species (8).

The acquired immunity of CRISPR-Cas systems is based on short nucleotide fragments, called spacers, cropped from the foreign nucleic acid sequence of a previous entry into the bacterial cell, called protospacers, which are inserted into the bacterial DNA next to the *cas* genes. These spacers are mostly similar to phage sequences and to a lesser extent to other extrachromosomal nucleic acid molecules. However, a large proportion of them have no known protospacer (over 80-90%), and these spacers of unknown origin are believed to come from phages not yet sequenced (9).

Bacterial genomes with CRISPR-Cas systems have *cas* genes along with the spacers, which are separated by repeats (short identical or nearly identical sequences). However, other genes have been linked with these systems because specific functionalities have been found in strains that have CRISPR-Cas systems. For example, a relationship has been demonstrated with the formation of multicellular structures called biofilms in *P. aeruginosa*, *Streptococcus mutant* and *Yersinia pestis* (10–12), connections with the regulation of outer membrane proteins have also been described in *Salmonella typhi* (13), and a specific relationship with virulence have also been shown in multiple bacterial species (14–16).

In earlier work with *A. baumannii*, we found that the group of strains that usually have CRISPR-Cas systems had genes involved in biofilm at a high frequency, and we also found genes encoding proteins with signal peptides and membrane lipoproteins (17). The analysis was based on a pangenome constructed from this species (all of the different genes found in the genomes of the species). The establishment of these pangenomes is now easier due to the large number of genomes available in public databases, and they allow to analyze the accessory genome, which is the set of genes that is not present in all the strains of a species (18) and are usually obtained by horizontal gene transfer (19). If strains with CRISPR-Cas systems have special functions not found in strains without these systems, accessory genes involved in these functions should appear almost exclusively in the former.

We have created a large pangenome for each ESKAPE species and compared strains with or without CRISPR-Cas systems to discover genes present with a significant frequency in the former. Then, these genes have been functionally analyzed and we found that they are enriched in genes encoding membrane proteins, which reveals a possible relationship with phages that infect bacteria and could take advantage of these membrane proteins as entry receptors.

Materials and Methods

Genome collection and annotation

The assembled sequences of ESKAPE species available in the National Center for Biotechnology Information (NCBI) Genome database on June 14, 2021, including complete and draft genomes, were collected (20). Genomes and metadata were downloaded with the tools datasets 12.1.0 and dataformat 12.4.0 (a total of 68,352 genomes). Genomes with a low number of total genes or a low average number of shared genes (>5 times the interquartile range) were removed on suspicion that they did not correspond to the species studied.

The protein-coding genes were predicted using Prokka version 1.14.5 (21), and the pangenome was created by Roary version 3.12.0 with an identity threshold of 90% and the -s parameter for not separating paralogs at this identity threshold (22). Protein sequences were functionally annotated using Sma3s v2 and the UniProt bacterial taxonomic division bacteria 2019_01 as the reference database (23). Gene names provided by Sma3s were preferentially assigned to each protein. When a gene name was repeated, a sequential number separated by two underscores was added. In cases where Sma3s did not assign a gene name, the one proposed by Prokka was taken, if available, preceded by an underscore. A gene was classified as a core gene if it appeared in ≥99% of the genomes of the species.

CRISPR-Cas systems and their specific types were assigned using CRISPRCasTyper 1.4.1 (24). Types I-Fa and I-Fb of *A. baumannii* were distinguished by looking for their different integration site. To discover the spacers of CRISPR-Cas systems, CRISPRCasFinder 4.2.20 was used with default parameters (25). Only CRISPR arrays with an evidence level equal to 4 were considered. Identical spacers were collapsed together, taking into account both chains. The number of sequences of each type for each species are available in Suppl. Table S1.

Search for specific gene groups in the pangenomes

Antibiotic resistance genes were found by AMRFinderPlus 3.10.1 using the databases of the 6 bacterial species analyzed (26). Virulence genes were found by performing a similarity search with BLASTP 2.9.0+ (27) against the VFDB database version December 2020 (Virulence Factor Database), requiring at least 90% sequence identity and 90% database sequence coverage (28).

To find genes encoding membrane proteins, we searched for genes that had the term 'membrane' in their functional annotation (within the Gene Ontology, UniProt keyword or Description annotation fields). Next, outer membrane proteins were added using a BLASTP similarity search against the OMPdb database release 2021, requiring at least 90% sequence identity and 90% database sequence coverage (29).

Genes from plasmids were searched using the annotation 'Plasmid' in the UniProt keyword field. Then, genes with ≥90% sequence identity and ≥90% query coverage with a sequence of the PLSDB database v2020_06_23_v2 were added (30). Viral genes were searched following the same protocol but using the IMG/VR database v3 (IMG_VR_2020-10-12_5.1) (31), and any of the following terms in the functional annotation: bacteriophage, prophage, phage, virus, virion, viral, but not "response to virus" or antiviral. Genes that were between 2 genes annotated as viral were also added. The Phaster web server was used to search for complete phages (32).

Search for protospacers

Protospacers, putative genes recognized by spacers, were searched by performing a similarity search with BLASTN and the blastn-short option turned on, using a threshold of ≥95% sequence identity and 100% spacer coverage. Those protospacers that were also found following the same strategy but using sequences from CRISPR repeats instead of the spacers were discarded to avoid misannotated sequences (33).

Searching for genes associated to CRISPR-Cas types with inference on random forests

We used inference on multiple iterations of random forests to search for genes associated with specific CRISPR-Cas types. For each species, we compiled one dataset containing all strains that have no CRISPR-Cas systems, and other datasets with the strains containing each CRISPR-Cas type present in the data, respectively. From 20 iterations with different random seeds, the most important

features were selected and counted. In this context the features are binary indicators of the presence of the genes for each strain. After all iterations, the count of a gene can indicate how often the random forest deemed it important for the difference between the respective CRISPR-containing and CRISPR-deficient genomes. *Cas* genes were removed from consideration, to be able to focus on non-directly related genes. Genes identified as important features in multiple iterations were considered to be associated with CRISPR-Cas systems, when they were more abundant in CRISPR-Cas containing genomes than in non-CRISPR-Cas containing genomes.

The random forest implementation was done in Python with the scikit-learn package 1.0.2 (34). With each iteration, random parts of the datasets were divided into train and test sets with the ratio of 0.8 to 0.2. For all six species and their CRISPR-Cas systems, the random forests achieved average accuracies higher than 0.93 over all iterations. The trained random forest object has the resulting feature importance by the mean decrease in impurity available as a parameter. The permutation feature importance may be more informative for high cardinality features, but since we only have two values for each feature, the mean decrease in impurity feature importance measurement is sufficient.

Functional enrichment analysis

To discover the functional enrichment of genes associated with specific CRISPR-Cas types, we used the R package TopGO version 2.40.0 (35), which uses GO terms from a specific ontology. The GO terms used were those annotated by Sma3s. Figures were created using the R ggplot2 library in a custom script.

MLST assignment, gene profiles and molecular phylogenies

MLST numbers were assigned to each genome by compiling the genes used in the species-specific schemes in PubMLST 23 Nov 2021 (36) and searching them in the genome sequences using the mlst program (https://github.com/tseemann/mlst). The MLST number assigned to each genome, along with the CRISPR-Cas systems it has is available in Suppl. Table S3.

MLST phylogenetic trees were constructed using the MLST sequences. Nucleotide sequences were aligned with mafft v7.271 using the G-INS-I option (37). The phylogeny was constructed with RAxML v8.2.9 with the GTRCAT model and bootstrap of 1000 (38). The model was selected with ModelFinder implemented in IQ-TREE (39). The phylogeny in Suppl. Fig. S6 was constructed with the same protocol but using all genomes, the PROTGAMMAWAG model and 500 core proteins.

The gene profiles for each species were constructed using a binary representation of the bacterial genome, where a gene is either absent or present in a strain, without accounting for the number of paralogs. This data is condensed to MLST level, by assigning 0 or 1 to the gene in the MLST group by majority vote of all strains in the group. The MLST groups are then subjected to a pairwise Jaccard distance measurement, resulting in a N x N matrix of Jaccard distances between MLST groups, with N equal to the number of MLST groups for the respective species dataset. The pairwise Jaccard distances were computed with scikit-learn.

These pairwise distances were used to construct a profile of genetic distances between the MLST groups for each species. We used ward-linkage and descending distance sort for the hierarchal clustering and the dendrogram. Dendrograms were produced using SciPy 1.6.2 (40), correlation plots were plotted with seaborn 0.11.2 and matplotlib 3.5.0 (41, 42).

Results

A large proportion of spacers in CRISPR-Cas systems can be annotated by pangenome analysis

The genomes of the different ESKAPE species were initially obtained and they were both structurally and functionally annotated with a special emphasis on the protein-coding genes and the elements that are part of the CRISPR-Cas systems. According to the number of genes, the smallest pangenome was found for *S. aureus* despite having started from a larger number of genomes (Fig. 1ab). The species with the largest number of genomes with CRISPR-Cas systems were *K. pneumoniae* and *P. aeruginosa*, which also had the largest pangenomes (along with the other Gram-negative species) and more than 3500 core genes, i.e., genes common to all genomes of the species. The difference between Gram-negative and Gram-positive bacteria is also evident when comparing the number of genes per genome, with *P. aeruginosa* showing both the largest number of genes and average number of shared genes (Fig. 1c). Finally, the major difference between genes per genome and average number of shared genes is found in *E. cloacae*, which could be explained by the low number of genomes used for this species (n=317).

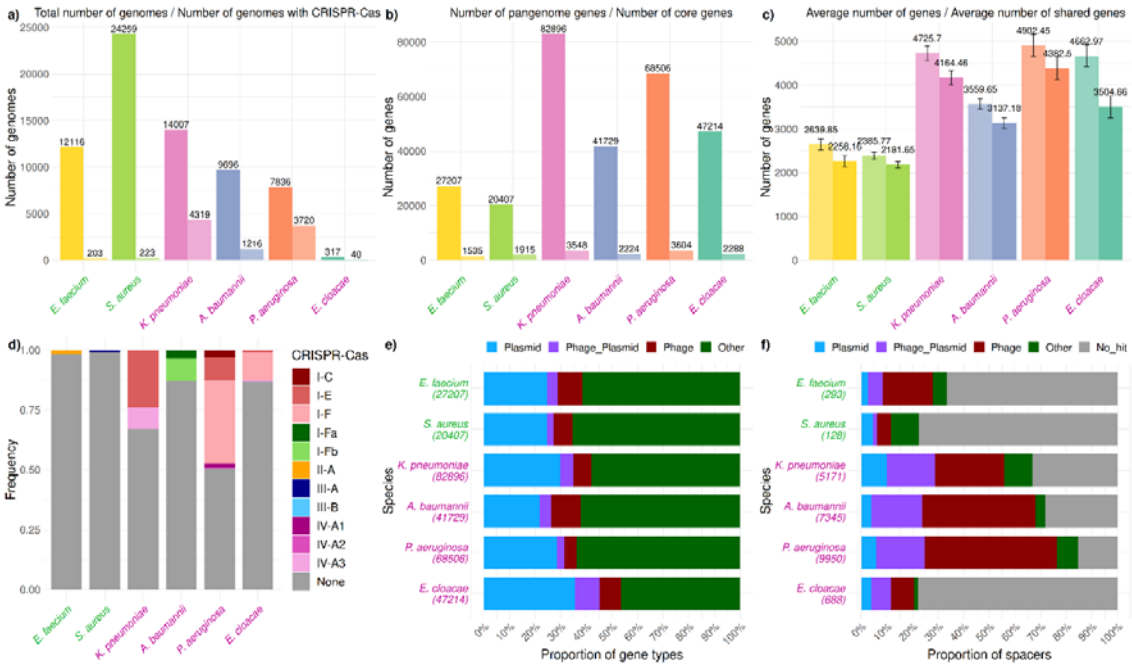

Fig. 1. Summary of analyzed genomes. a) Total number of genomes analyzed and number of genomes having CRISPR-Cas systems. b) Number of genes in the pangenome and number of core genes. c) Average number of genes per genome and average number of shared genes (average of the number of genes shared for each genome with the remaining genomes). d) Distribution of CRISPR-Cas systems among genomes. e) Proportion of gene types in the pangenome. For each species: the number of total genes is shown (below the species name), and the proportion of them that were annotated as belonging to plasmid sequences, phages, both plasmids and phages, and other genes not included in plasmids or phages. f) Spacers and assigned protospacer types by species. For each species: the number of total spacers is shown (below the species name), and the proportion of them that match with plasmid sequences, phages, sequences annotated as both plasmid and phage, other genes not included in plasmids or phages, and the spacers that do not match any gene (No_hit). The names of Gram-positive bacteria appear in green, and those of Gram-negative bacteria in purple.

The overall proportion of genomes exhibiting CRISPR-Cas systems is low, with Gram-positive bacteria having them in only 1% of their genomes, both *A. baumannii* and *E. cloacae* in 12%, and *K. pneumoniae* and *P. aeruginosa* in 30% and 47%, respectively (Fig. 1d). Since the spacers of CRISPR-Cas systems usually recognize mobile genetic elements, we first separated pangenome genes that could come from plasmids (an average of 28% of the genes) and bacteriophages (an average of 8% of the genes) (Fig. 1e). Interestingly, an average of 5% of the genes were annotated as originating from both genetic elements, which could come from what is known as phage-plasmids (43).

Next, the spacers were obtained and their cognate protospacers were searched for within the pangenome genes. The major number of different spacers was found in the three species with CRISPR-Cas IV and/or I-F types (Fig. 1f). Protospacers belonged in a much larger proportion to phage genes and, remarkably, it was possible to annotate more than 65% of them in the same three species mentioned, with a maximum of 85% in *P. aeruginosa*. The species with a lower number of annotated spacers (about 25%) were *E. cloacae* and *S. aureus*. It is noteworthy that about 18% of the spacers in CRISPR-Cas type I species appear to recognize genes annotated as phage-plasmids.

Since the presence of CRISPR-Cas systems prevents the entry of foreign DNA (including resistance and virulence plasmids) into the bacterium, the number of genes involved in these functions was compared between genomes with and without CRISPR-Cas systems. On average, genomes with CRISPR-Cas systems presented a lower number of resistance and virulence genes (Suppl. Fig. S1). The most significant difference was found with CRISPR-Cas type II and III, except for resistance genes in *S. aureus*. On the other hand, *P. aeruginosa* presented the highest number of virulence genes overall, although the genomes without CRISPR-Cas systems presented a lower number than those with CRISPR-Cas systems, a result that was repeated with the resistance genes. So, we cannot conclude that all genomes with CRISPR-Cas systems tend to carry a lower number of resistance and virulence genes.

CRISPR-Cas systems appear and disappear throughout the entire phylogeny

At this point we wanted to know if the CRISPR-Cas systems were linked to a branch of the phylogeny of the species studied. To define phylogenetic relationships between genomes with and without CRISPR-Cas systems, the multi-locus sequence typing was used (MLST). This is based on several house-keeping genes of each species (36), and two adjacent groups reflect genomes arising from a recent common ancestor. Except for *E. faecium* type II and some groups of *A. baumannii* type I genomes, CRISPR-Cas systems appear to be spread throughout the phylogenetic tree (Fig. 2, top), suggesting a possible gain by horizontal gene transfer in genomes for which it could provide an evolutionary advantage, and a possible subsequent loss when that advantage no longer exists.

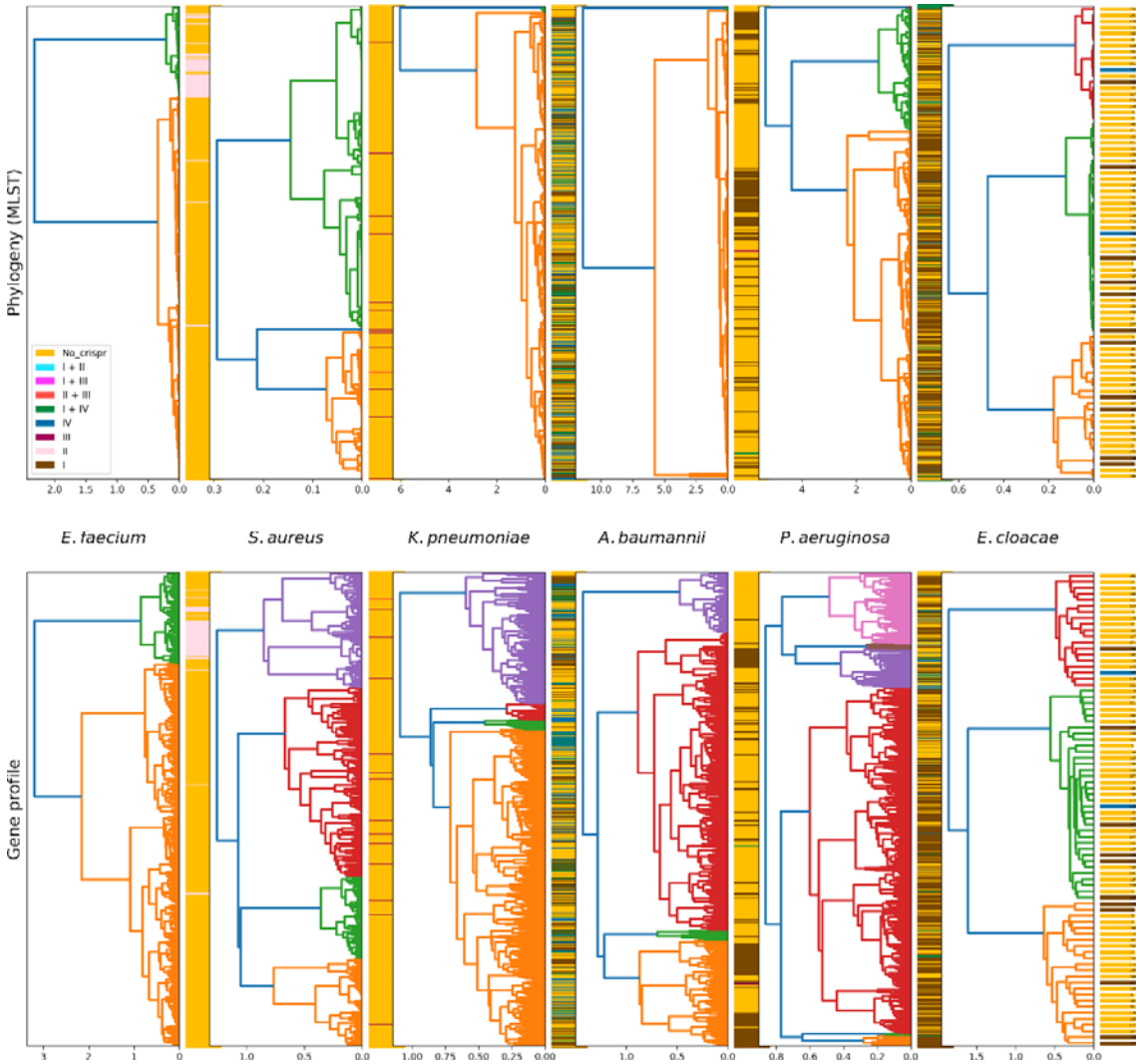

Fig. 2. Molecular phylogeny for each species based on MLST (top) and gene profile (bottom). The legend shows the different types of CRISPR-Cas (without specifying the subtype) with orange indicating the MLST group lacking any CRISPR-Cas system. The colors of the branches highlight different clades distanced from the rest.

If CRISPR-Cas systems are associated with other bacterial physiological functions, one possible hypothesis is that genomes with CRISPR-Cas systems always have a similar collection of accessory genes, regardless of *cas* genes. To test this idea, distance trees were constructed between the same MLST groups in the phylogeny, in this case based on the gene profile of the genomes (gene presence/absence matrix). These gene profiles showed a similar dispersion to that found with molecular phylogeny, except, again, for the case of *E. faecium* and the groups of *A. baumannii* genomes, suggesting that CRISPR-Cas systems do not appear in genomes with a fixed collection of accessory genes (Fig. 2, bottom). However, when the two types of relationships are compared, a high correlation is observed between distances of the molecular phylogeny and the gene profile in species with a smaller number of genomes with CRISPR-Cas systems (Suppl. Fig. S2). Nevertheless, this correlation decreases in those with a higher number of genomes with CRISPR-Cas systems. Taken together, this reinforces the idea that CRISPR-Cas systems do not appear to be linked to strains phylogenetically related or to specific accessory genomes. But this raised the question of

whether genomes with CRISPR-Cas systems presented particular genes at higher frequencies than genomes without these systems.

Genes associated with CRISPR-Cas systems mainly encode membrane proteins

We have already seen that CRISPR-Cas systems are associated with different accessory genomes and appear and disappear at any evolutionary time. However, assuming that CRISPR-Cas systems are associated with other physiological functions of bacteria, we should find some accessory genes more frequently in genomes having these systems. With this in mind, we searched for genes significantly associated with genomes showing CRISPR-Cas systems, excluding the *cas* genes themselves, and a mean of 147±62 genes per genome was found (Suppl. Table S2). To test whether these associated genes were significantly involved in any biological process or function, enrichment analysis was performed, and it was found that genes encoding membrane proteins were highly prominent (Fig. 3). These include the pili proteins of *P. aeruginosa* and *K. pneumoniae*, as well as outer membrane proteins of *A. baumannii* and *K. pneumoniae*, and other *A. baumannii* and *P. aeruginosa* proteins involved in type II secretion systems. This relationship with membrane proteins was especially relevant in CRISPR-Cas type I systems, while other genes involved in catabolic processes or DNA metabolism were found more notably in types II, III, and IV (Suppl. Fig. S3). In addition, types IV are also notable for the annotation "extrachromosomal DNA", reflecting the origin of these systems from mobile genetic elements (44).

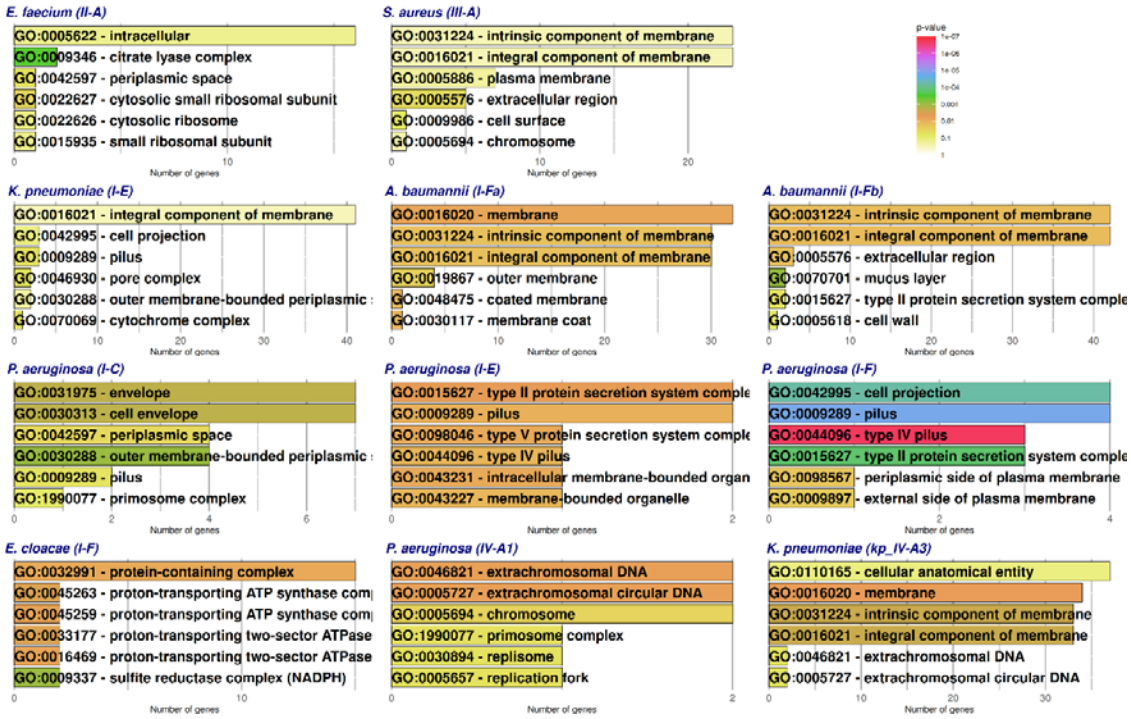

Fig. 3. Functional enrichment of genes associated with different CRISPR-Cas systems in the different species. Gene Ontology cellular component was used in these enrichments, using functional annotations of the pangenome obtained by Sma3s. Enrichment with Gene Ontology biological process and molecular function can be found in Suppl. Fig. S3.

Genomes with specific types of CRISPR-Cas systems have different sets of membrane proteins

Since the genomes bearing CRISPR-Cas systems presented specific types of membrane proteins, we wanted to know if all genomes with a specific CRISPR-Cas type had the same set of membrane

proteins. To assess this, genes encoding membrane proteins previously associated with CRISPR-Cas were searched for in genomes having these systems. In general, there were at least two clusters of genomes, especially when type I was analyzed, each one presenting a different collection of genes encoding membrane proteins (Fig. 4), except for type I-F in *E. cloacae*, as the number of genomes in this species is low. However, the separation between these two clusters of genomes were not as clear as with the other types (Suppl. Fig. S4). As an example, about half of the *A. baumannii* genomes with the CRISPR-Cas type I-Fb system carry the *opuD* and *betP* genes, involved in choline and glycine betaine transport, which could protect the bacterium from osmotic stress (45). And the other half of the genomes have porins such as *benP*, or a specific variant of the TonB-dependent siderophore receptor *bauA* (46).

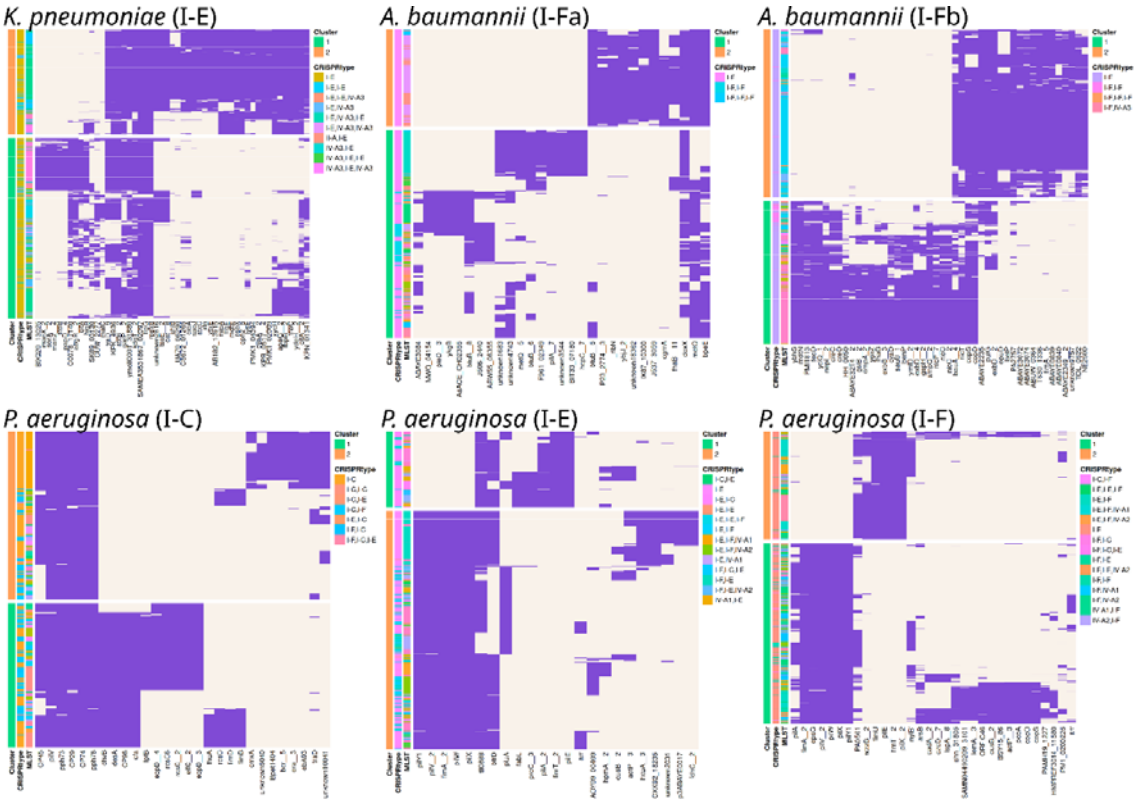

Fig. 4. Clusters of genomes with CRISPR-Cas type I systems according to the genes encoding membrane proteins that they have. The purple color indicates presence of the gene (X-axis) in the corresponding genome (Y-axis). On the left side of each plot, the cluster number is shown, along with the CRISPR-Cas type or combination of them and the MLST group.

Genomes with both CRISPR-Cas type I systems and specific membrane proteins show exclusive spacers and phage genes

Genomes with CRISPR-Cas type I systems show specific collections of membrane proteins. These proteins could provide the bacterium with important characteristics such as specific stress responses, or the detoxification that certain outer membrane proteins can offer. Since the majority of CRISPR-Cas systems studied here seem to recognize phage sequences (Fig. 1f), we hypothesized that these membrane proteins could be acting as receptors of specific phages, and genomes with them are forced to acquire CRISPR-Cas systems to defend against infection by these viruses, while maintaining the beneficial functions for the bacterium. Indeed, the *A. baumannii* cluster 1 for type I-

Fb includes the *ompA* gene, an outer membrane protein, the *P. aeruginosa* cluster 1 for type I-C shows the *fhuA* gene and the two *A. baumannii* clusters for I-Fa show different variants of the *btuB* gene, both encoding TonB-dependent proteins. These three genes have long been known to act as phage receptors in *Escherichia coli* and *Salmonella* (47).

To test the above hypothesis, we looked for spacers that recognized phage genes present in each of the two membrane protein-based clusters of genomes that did not appear (neither the spacer nor the phage gene) in the other cluster. We then searched for genomes in each cluster that might carry the phage genes recognized by these spacers. Thus, within a given cluster we could find genomes containing a cluster-specific spacer, or a phage gene recognized by one of these spacers. But it could also be the case that both elements appear in the same genome, and this would imply having two new alternatives: either a cluster-specific spacer and the viral gene recognized together, or a spacer and a viral gene recognized by one of the other cluster-specific spacers. When evaluated in CRISPR-Cas type I systems, the number of genomes with unique spacers for each cluster was high, with a predominance of genomes with unique spacers in *K. pneumoniae* I-E and *P. aeruginosa* I-C, with spacers and phage genes in *A. baumannii* I-F types, and with phage genes in *P. aeruginosa* I-E and I-F (Fig. 5).

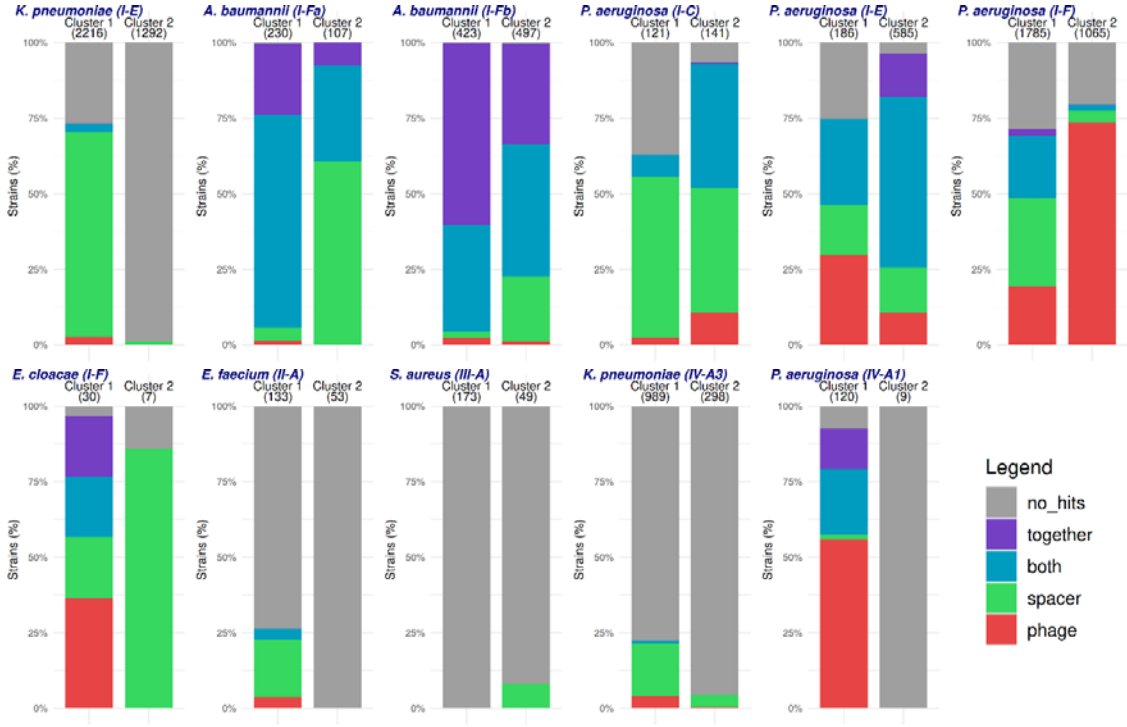

Fig. 5. Distribution of genomes in each membrane protein-based cluster with unique spacers and/or protospacers (phage genes). The chart shows the proportion of genomes from each cluster which contain: a phage gene recognized by a cluster-specific spacer (red), a cluster-specific spacer (green), both a spacer and a phage gene, but the latter is recognized by another cluster-specific spacer (blue), both a spacer and the recognized phage gene (purple), or neither the cluster-specific spacer nor the phage gene (grey). The types of CRISPR-Cas systems that defined the best clusters are shown in the row above (in the same order as in Fig. 4), and the others are shown in the row below (in the same order as in Suppl. Fig. S4). The number of genomes in each cluster is shown in parentheses.

In *A. baumannii*, cluster 2 of CRISPR-Cas I-Fb genomes (see Fig. 5) has specific spacers against a phage gene (*unknow5433*) that appears in 466 distinct genomes, with the phage gene itself also appearing in 68 of them (Fig. 6a). The complete phage is integrated near a tRNA-Arg gene and has a total of 59 proteins (Suppl. Fig. S5). In addition, the phage gene also appears in 283 genomes lacking CRISPR-Cas systems. When the specific membrane proteins of cluster 2 are also searched for in genomes lacking the CRISPR-Cas system but having the phage gene, the best match occurs with the *T630_1336* membrane protein (which we will refer to as *cam1* for CRISPR-associated membranome gene 1), while the rest of the membrane proteins appear more frequently in genomes lacking the phage gene (Fig. 6b). Specifically, 278 of the genomes that have *cam1* also have the phage gene (63%). In fact, of the 283 genomes without CRISPR-Cas systems that have the phage gene, only in 5 of them the *cam1* gene could not be found.

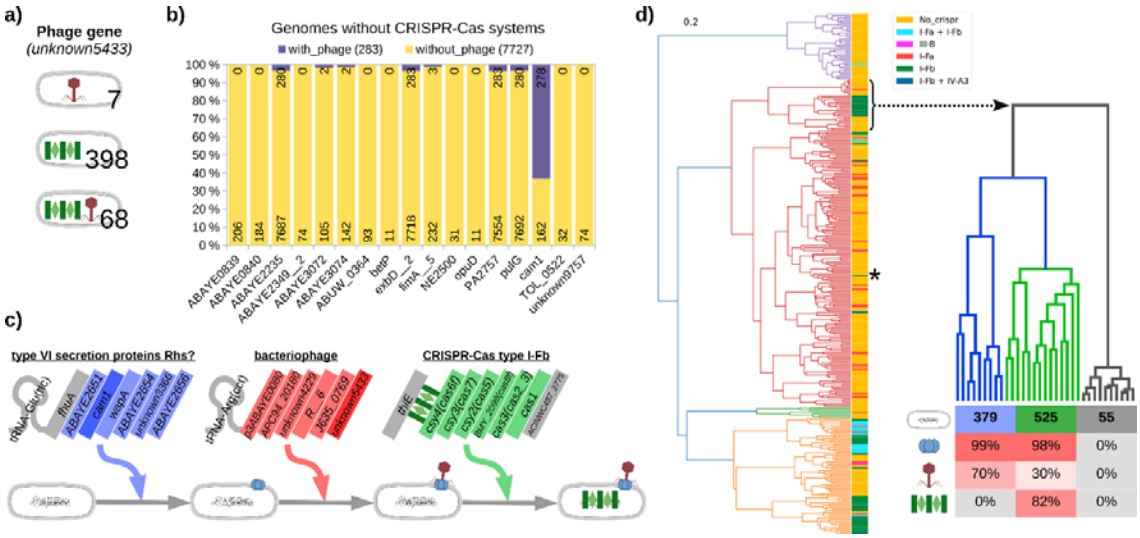

Fig. 6. Phage gene specific to cluster 2 of CRISPR-Cas I-Fb genomes and its co-occurrence with membrane genes. a) The phage gene *unknown5433* appears alone in only 7 genomes of cluster 2, but there are spacers against it in 398 genomes, and the phage gene appears together with these spacers in 68 other genomes. b) Frequency of cluster 2-specific membrane proteins in genomes lacking CRISPR-Cas systems, separated between those with and without the phage gene. The number of genomes with and without the phage gene is shown in parentheses. Note that the phage gene appears in most genomes having the membrane protein Cam1. c) Hypothesis of gain of gene groups from a genome lacking the membrane protein and the CRISPR-Cas system: first the region of 6 genes involved in a T6SS entered, which would allow virus entry through the Cam1 protein, and finally the CRISPR-Cas system would be obtained for protection against the phage. d) Hypothesis testing by the gene profile of *A. baumannii*. The grey clade (55 genomes) lacks CRISPR-Cas system, phage gene and membrane protein, the blue clade (379 genomes) has the membrane protein and the phage gene appears in 70% of genomes, and the green clade (525 genomes) shows CRISPR-Cas systems in the 82% of genomes and present the phage gene in a lower number of genomes (30%). The tree is consistent with the steps proposed in the hypothesis in c). *MLST734 which belongs to cluster 2 and appears separate from the rest of the genomes in the cluster, is composed of 6 genomes with the CRISPR-Cas system and the membrane protein, and the gene phage in only one of them. The rest of the MLSTs can be consulted in Suppl. Table S3.

Cam1 is a 349 amino acid protein that shows a transmembrane region followed by a Rhs repeat-associated core domain at its N-terminal half (InterPro:IPR022385). This domain appears in bacterial toxins involved in type VI secretion systems (T6SS), but when present in proteins of less than 400

amino acids, found in bacteria such as *Pseudomonas putida*, it has an unknown function (48). The gene encoding this membrane protein is part of a cluster of 6 genes that is integrated next to a tRNA-Glu and the gene *fhuA*, and close to this region there is another T6SS spike gene (*vgrG2__8*). Since the presence of this protein seems to be a *sine qua non* condition for the appearance of the phage, we could expect that the gain of the T6SS by the bacterium would imply that it would be exposed to infection by the phage, something that could be counteracted by the bacterium with the acquisition of a CRISPR-Cas system (Fig. 6c). This appears to be supported when analyzing the group of genomes containing this membrane protein, together with genomes that have a similar gene profile. In this group there is a set of 55 genomes without the membrane protein, phage, and CRISPR-Cas systems (Fig. 6d). Another set of 379 genomes with a similar gene profile carries the membrane protein and 70% of these genomes show the phage. Finally, there is a last set of 525 genomes carrying the membrane protein but showing a lower frequency of occurrence of the gene phage probably due to the CRISPR-Cas they present. All this is supported by a phylogeny in which 500 core genes from all the genomes of this cluster were used (Suppl. Fig. S6).

Discussion

A limited number of bacterial genomes have CRISPR-Cas systems to prevent the entry of foreign DNA. We have analyzed tens of thousands of genomes of bacterial species of the ESKAPE group and found that type I is the most frequent CRISPR-Cas system in the Gram-negative species of this group. Our results are consistent with previous reports that found, for example, a low proportion of genomes with CRISPR-Cas systems in *E. faecium* (49), and about fifty percent of those in *P. aeruginosa* (7). When a bacterial genome has a CRISPR-Cas system, it is expected to have fewer genes. Some of the missing genes may be those involved in antibiotic resistance and originating from plasmids or integrative and conjugative elements, or those involved in virulence and originating from phages. This was mostly found in our study with ESKAPE pangenomes, and coincides with previous studies with the same species, except for the fact that we show a more complete collection of type IV systems, and we have used twice as many genomes (8, 50).

Spacers of CRISPR-Cas systems usually recognize sequences originating from phages (9) and to a lesser extent from plasmids, such as we previously showed in *A. baumannii* (17). However, most spacers have unknown origin, as the corresponding protospacer cannot be found, and have been lumped into what is known as CRISPR dark matter (9, 51). This dark matter accounts for 80-90% of the spacers and is expected to recognize phage sequences that are still unknown or have diverged from known phage sequences. And it is believed that this percentage may decrease with the future increase of sequences in the databases (52). We show that by analyzing complete pangenomes, which can include all the variation of phages infecting the species, dark matter can be greatly minimized (Fig. 1f). Thus, we were able to annotate 85% of the 9950 *P. aeruginosa* spacers and 72% of the 7345 *A. baumannii* spacers, with approximately 70% of these corresponding to phages or phage-plasmids. These phage-plasmids include phages that can remain as extrachromosomal elements in the bacterium (43) and against which we have found more associated spacers than against plasmid genes.

We have also found that genomes with CRISPR-Cas systems do not appear phylogenetically restricted, nor do they have a unique accessory genome (Fig. 2). This suggests that bacteria acquire these systems when they provide an important evolutionary advantage. In a study carried out to measure the impact of CRISPR-Cas systems on horizontal gene transfer in bacteria, it was concluded that these systems would play an important role at the population level but not at the evolutionary scale (53). In fact, these systems are often recruited by mobile genetic elements on independent

phylogenetic times (54). All of this would support our results, positioning CRISPR-Cas systems as functional modules that are acquired and discarded under certain circumstances.

The association found between membrane proteins and CRISPR-Cas systems suggests that one such circumstance could be the defense against phages that use these proteins as receptors. A previous report had already suggested that misfolded membrane proteins may trigger an envelope stress response that activates a CRISPR-Cas system (55), and another relevant report found 79 genes with probable association to the CRISPR-Cas systems and some of them encoded integral membrane proteins (56). Many of these genes were related to type III systems, which is the type in *S. aureus,* where we found more than 20 genes annotated as "integral component of membrane" associated with its CRISPR-Cas system. However, we mainly found this association with membrane-related accessory genes among the different classes of type I CRISPR-Cas and propose that this may be related to the acquisition of beneficial functions for the bacterium that conversely make it more vulnerable to certain phages. These membrane proteins may help form biofilms or allow for certain virulence-related advantages (10–16), but at the same time, this membrane proteins can be receptors for specific phages.

It has been shown that phages can increase the virulence of the bacterium they infect when integrated into the bacterial chromosome, as they can carry toxins, resistance genes, or adhesion factors (57). CRISPR-Cas systems would prevent these phages from proliferating. However, we have found that in many cases spacers coexist with the cognate phage gene, especially in *A. baumannii*. These cases reflect that the phage would be integrated into the bacterial genome, suggesting that the immune system has not been fully efficient. Indeed, in *P. aeruginosa*, and partly in *K. pneumoniae*, we have seen that the number of virulence genes in strains carrying CRISPR-Cas systems may be higher than in those without (suppl. Fig. S1). The coexistence of the protospacer with the cognate spacer has been proposed as representing autoimmunity processes with a negative effect on the bacterium (58). However, the high number of cases found here would suggest some unknown function. This is supported by studies showing that CRISPR-Cas systems can prevent the lytic cycle of phages but tolerate the virus integration as a prophage, allowing the bacteria to co-opt the phage genes for possible use as virulence factors (59).

We observed that genomes with specific types of CRISPR-Cas systems can be separated into clusters based on the membrane proteins they have, and that the spacers of these CRISPR loci would recognize different non-overlapping phage genes (Fig. 4-5). By analyzing these relationships between membrane proteins, spacers and phage genes, we are able to propose a gene encoding a member of a putative T6SS as a putative receptor for a phage found in genomes with and without CRISPR-Cas systems. Phylogenetic data suggest that the gain of this immune system would protect the bacterium against this phage while allowing it to maintain the secretion system, which could be useful for intra- or interspecific competition (60). Interestingly, the gene cluster to which this membrane gene belongs is integrated next to the *fhuA* gene, and it is speculated that a class of specific receptors (TonB) could be critical for phage genome injection through their interaction with FhuA (61). Thus, this protein could help both the entry of the phage and the proper functioning of the T6SS system. Other reports have shown that secretion systems and CRISPR-Cas systems can depend on quorum sensing (62, 63), which enables the coordination of bacterial population growth and is therefore also related to biofilm formation. Thus, it could be hypothesized that the bacterium would express both systems at the same time, to avoid being exposed to phages that could take advantage of the activation of the T6SS system when the bacterial population and cell-to-cell contact is increased.

Conclusions

Most bacterial genomes do not have CRISPR-Cas systems, suggesting that they could be acquired by horizontal gene transfer under certain circumstances. Here, we describe a "Membrane protein-Phage-CRISPR" triad, in which the CRISPR-Cas systems might be especially necessary when the bacterium expresses accessory genes that encode for membrane proteins. This would give it a special advantage in that situation, but it would also represent a gateway for phages that use those proteins as receptors.

Data availability

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

The code used to analyze the data and the pangenomes as well as the phylogenetic distances can be found in our GitHub repository: https://github.com/UPOBioinfo/crispromeskape.

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Conflict of Interest

The authors declare that they have no conflict of interests.

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Supplementary data

Suppl. Fig. S1. Distribution of the number of genes involved in antibiotic resistance (top) and virulence (bottom) in groups of genomes with and without CRISPR-Cas systems. Asterisks highlight pairs of groups with significant differences in the median according to a Wilcoxon rank-sum test (from *≤0.05 to ****≤1e-03).

Suppl. Fig. S2. Correlation between molecular phylogeny and gene profile. Pearson correlation value is shown in parenthesis. Points show pairs of MLST groups without CRISPR-Cas systems (cyan), with one group having one of these systems (orange), and with the two groups having CRISPR-Cas systems (pink).

Suppl. Fig. S3. Functional enrichment of genes associated to CRISPR-Cas genomes (GO Biological Process and Molecular function).

Suppl. Fig. S4. Clusters of genomes with CRISPR-Cas types II, III and IV systems, together with the I-F of *E. cloacae* by the sets of membrane protein encoding genes that they present. The purple color indicates presence of the gene of the X-axis in the corresponding genomes of the Y-axis. At the left side of each heatmap the cluster number is shown, together with the CRISPR-Cas type or combination of them and the MLST group.

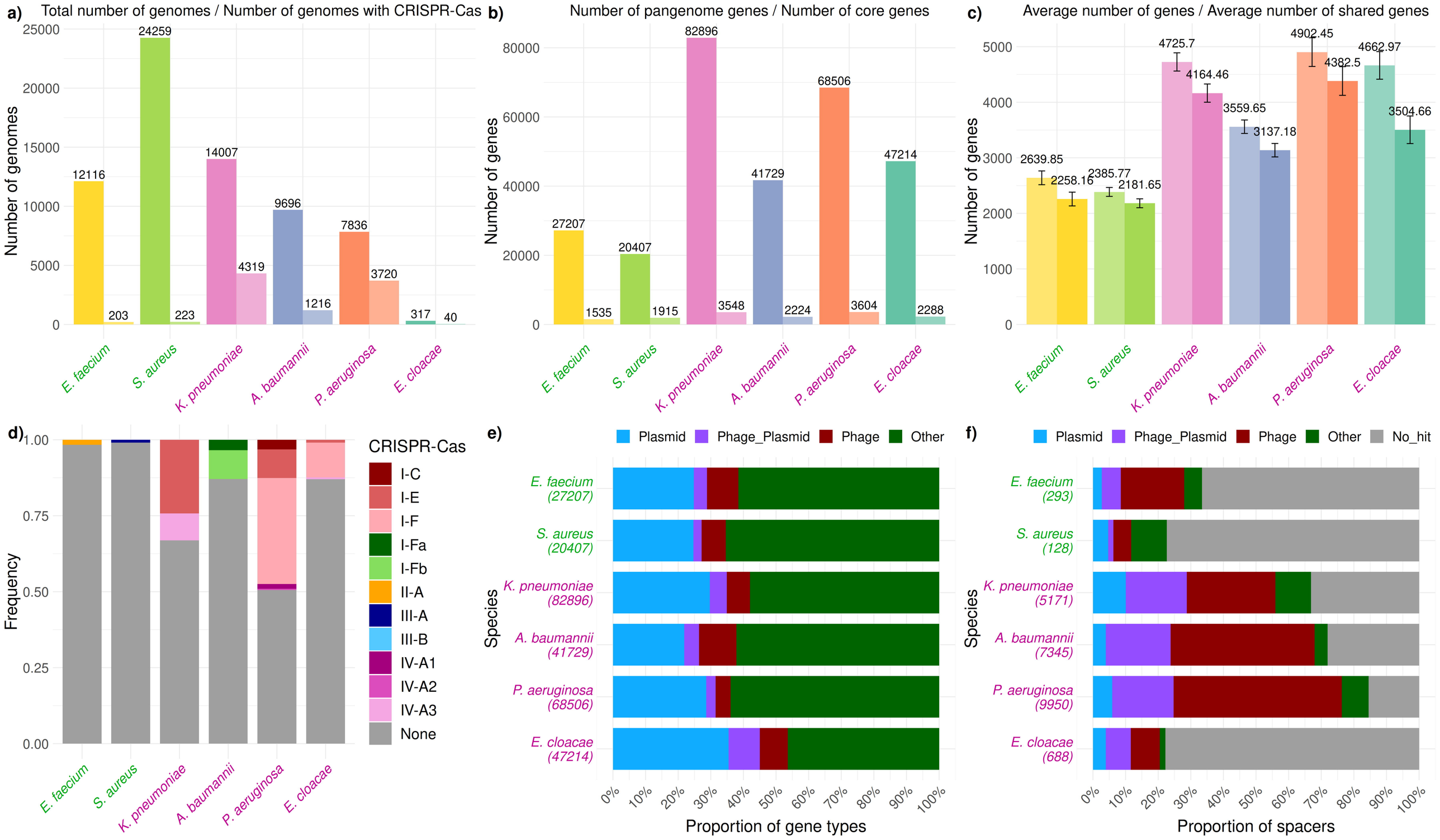
Suppl. Fig. S5. Phage with gene *unknown5433* integrated near the tRNA-Arg in *A. baumannii* (figure obtained with Phaster).

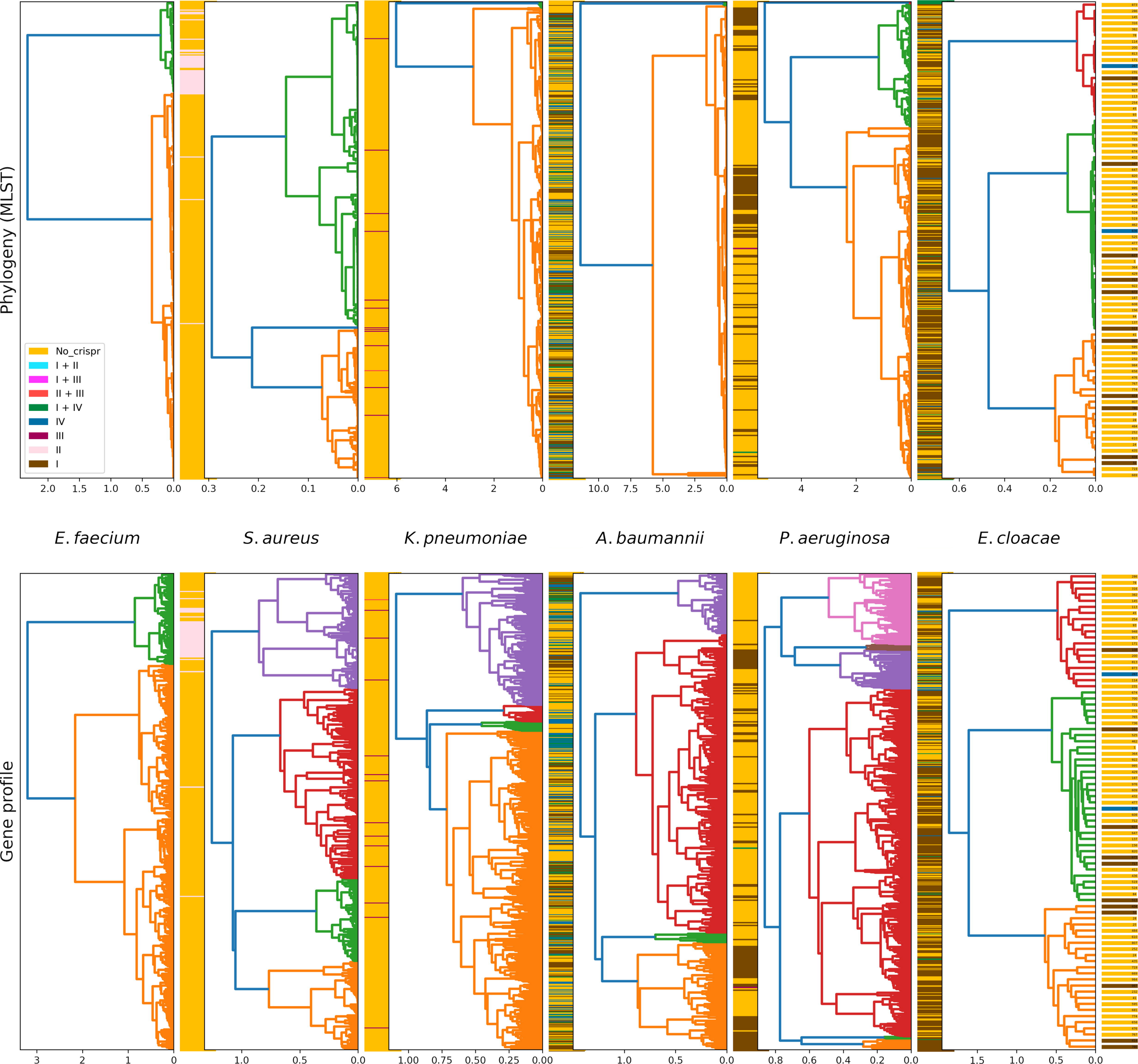
Suppl. Fig. S6. Molecular phylogeny of genomes in membrane protein-based cluster 2 of CRISPR-Cas type I-Fb. We used 500 genes that appeared in all genomes. A bootstrap of 1000 repeats was used, and nodes with a bootstrap of less than 70 % are marked with a red circle. Legend colors represent genomes with: the CRISPR-Cas type I-Fb (C), the membrane gene *cam1* (M), the phage gene *unknown5433* (P), or none of them (-).

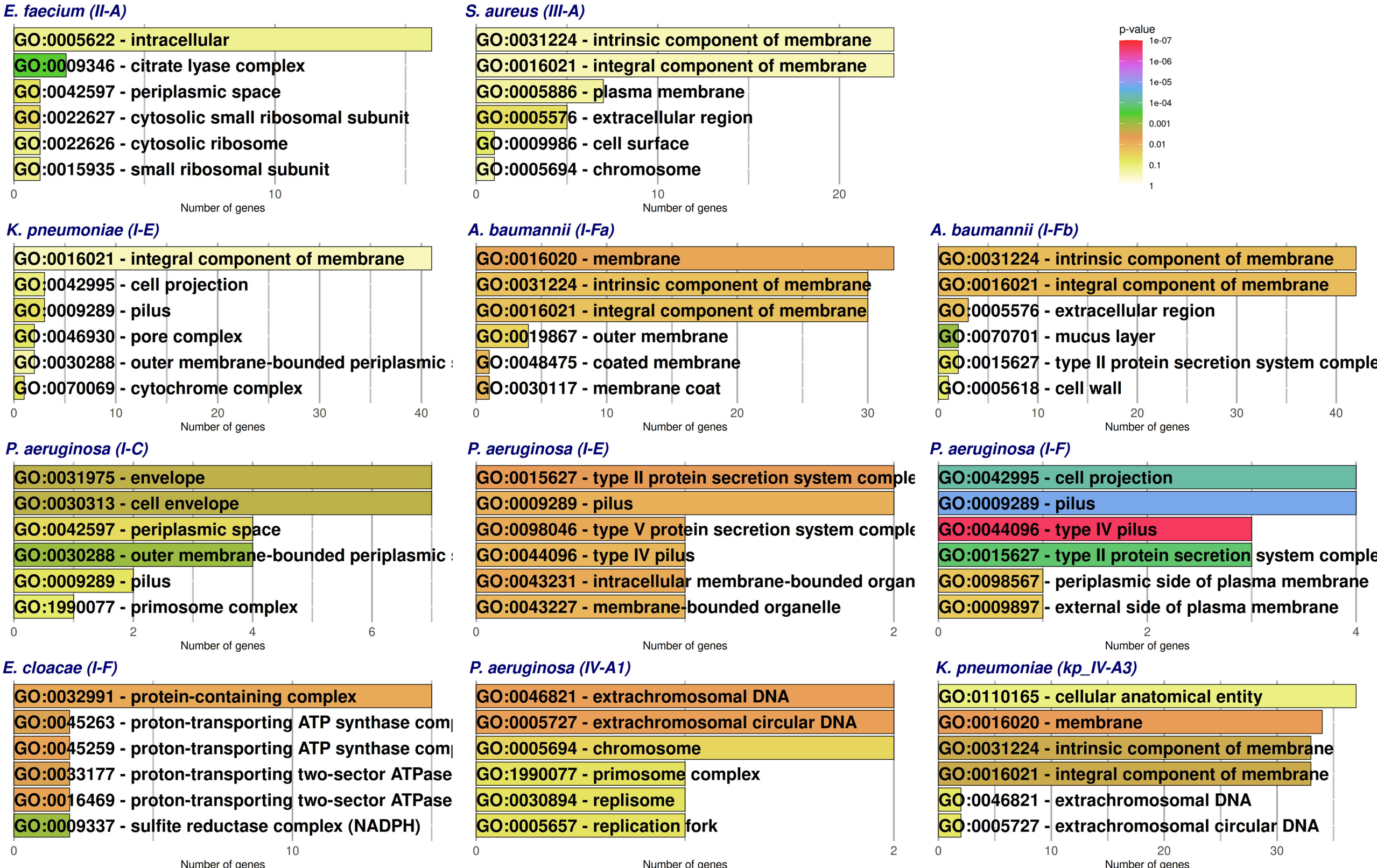
Suppl. Table S1. Distribution of CRISPR-Cas systems and spacers among all species.

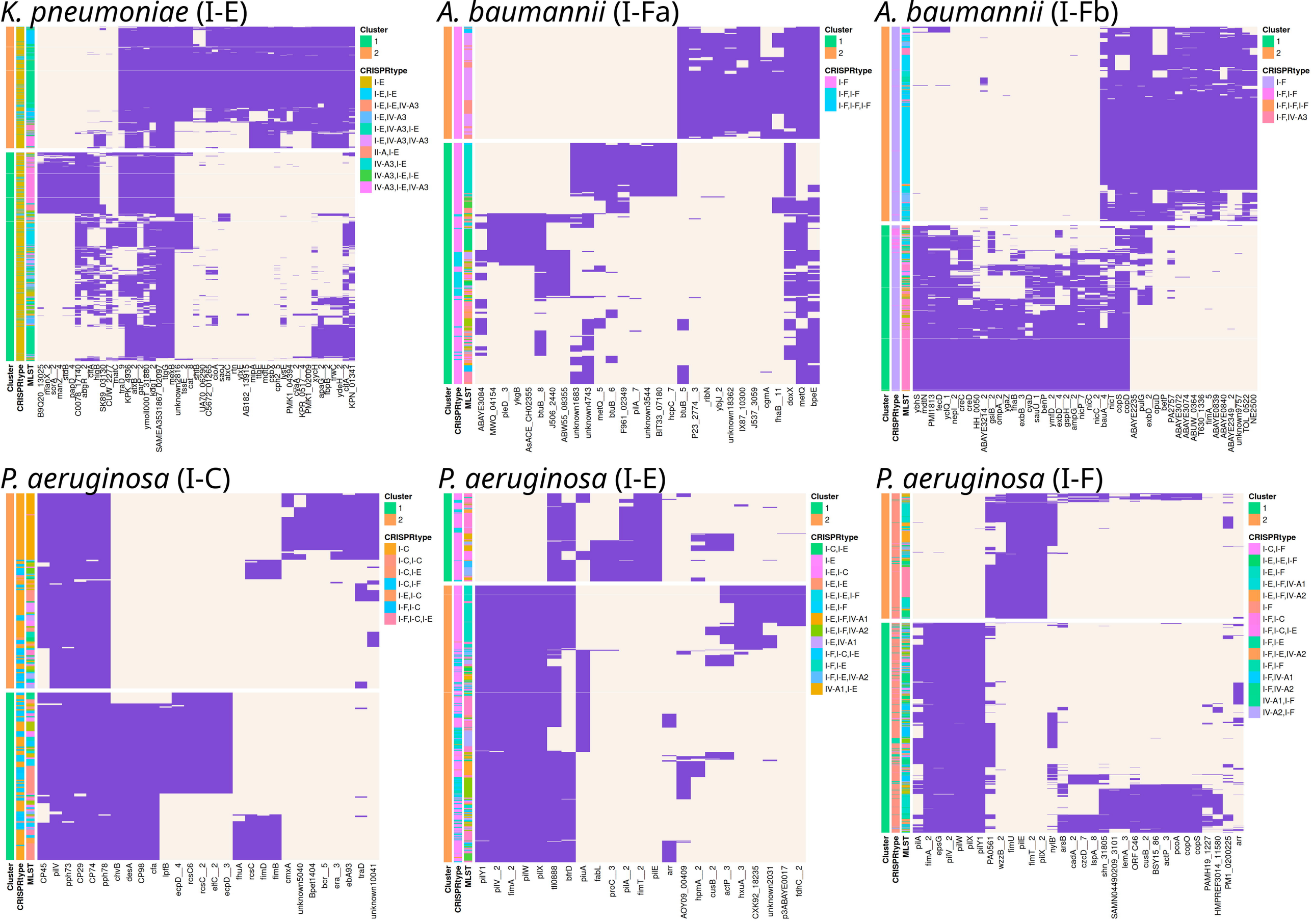
Suppl. Table S2. Genes associated to CRISPR-Cas genomes. Each worksheet represents the inference results of 20 iterations of a random forest for a species and the respective CRISPR-Cas systems. The first worksheet has a header explaining the columns as follows: genes (holds the encoded gene names); known_genes_retained (contains the count of how often the gene was in the top 30 most important features of the trained random forest); log2fc_gene and diff_gene, whic are the log2 fold change and the simple difference of means of the samples with and without a CRISPR-Cas system, respectively.

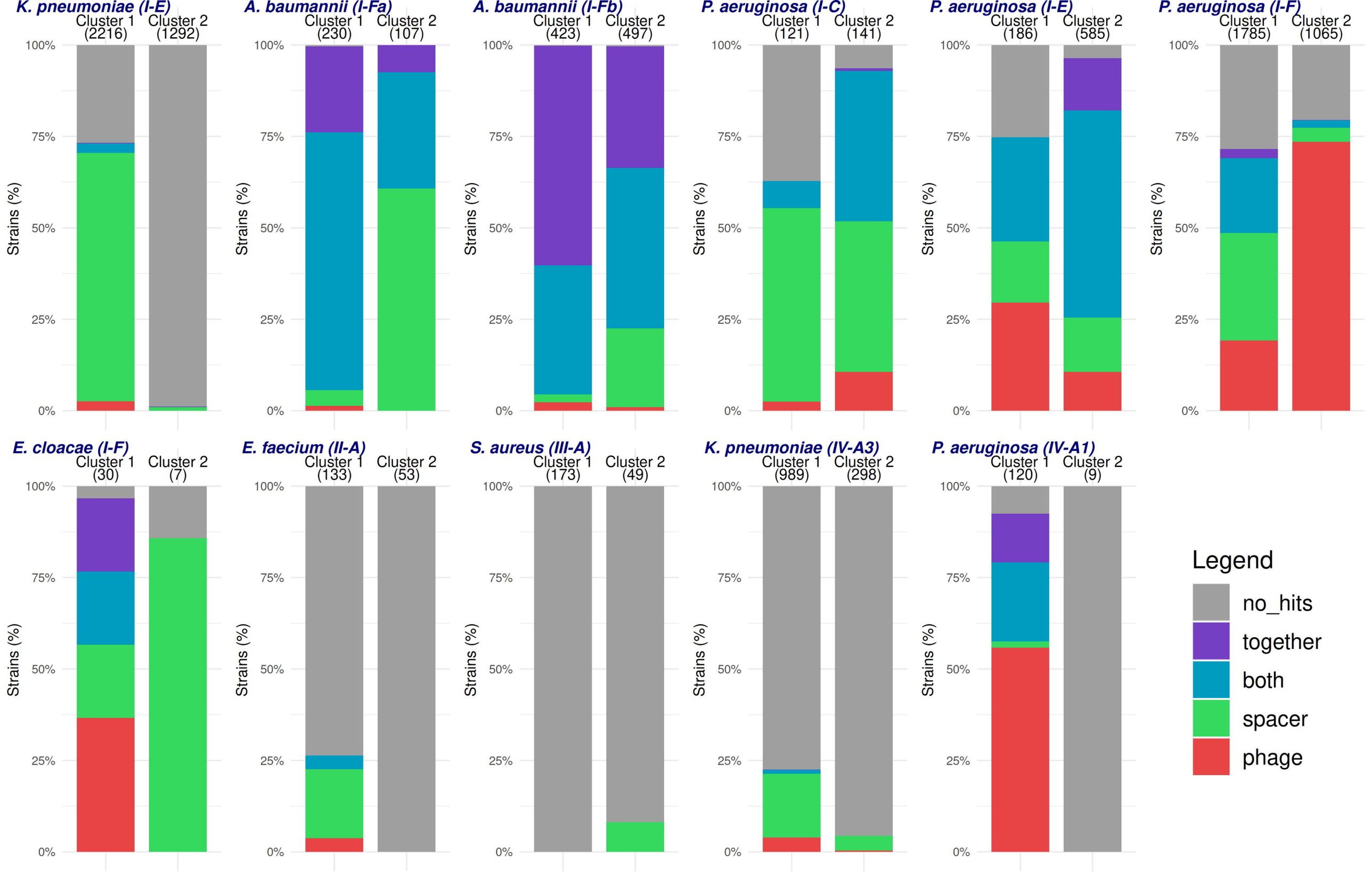
Suppl. Table S3. Genomes used for each species with the MLST to which they belong, and the CRISPR-Cas systems found in them. Incomplete CRISPR-Cas systems are labeled as "ambiguous". The *A. baumannii* I-Fb clusters have been added, along with the clade colors used in Fig. 6d.

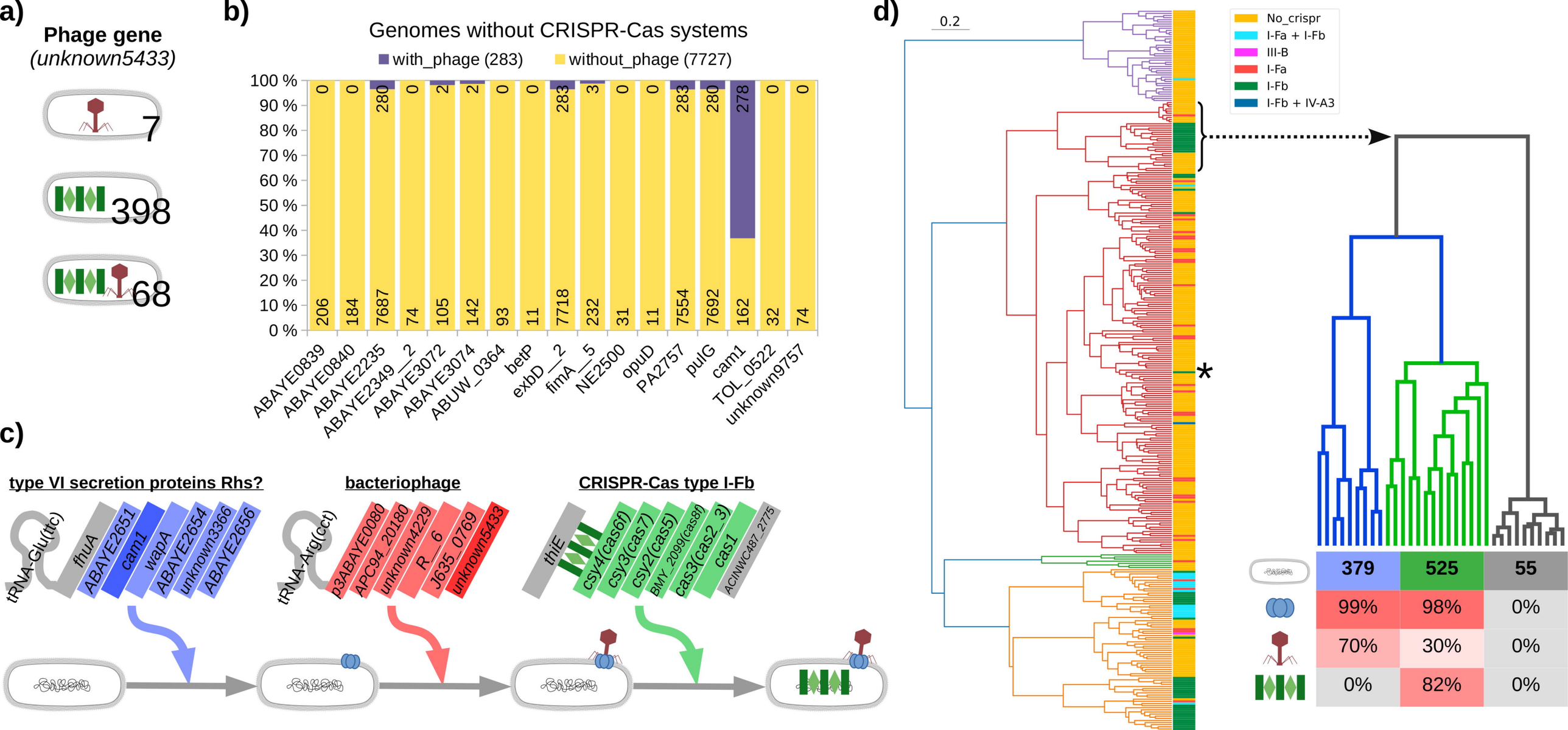

a)
Total number of genomes / Number of genomes with CRISPR-Cas
Number of genomes
12116
203
24259
223
14007
4319
9696
1216
7836
3720
317
40
E. faecium
S. aureus
K. pneumoniae
A. baumannii
P. aeruginosa
E. cloacae
b)
Number of pangenome genes / Number of core genes
Number of genes
27207
1535
20407
1915
82896
3548
41729
2224
68506
3604
47214
2288
c)
Average number of genes / Average number of shared genes
Number of genes
2639.85
2258.16
2385.77
2181.65
4725.7
4164.46
3559.65
3137.18
4902.45
4382.5
4662.97
3504.66
d)
Frequency
CRISPR-Cas
I-C
I-E
I-F
I-Fa
I-Fb
II-A
III-A
III-B
IV-A1
IV-A2
IV-A3
None
e)
Plasmid
Phage_Plasmid
Phage
Other
Species
E. faecium (27207)
S. aureus (20407)
K. pneumoniae (82896)
A. baumannii (41729)
P. aeruginosa (68506)
E. cloacae (47214)
Proportion of gene types
f)
Plasmid
Phage_Plasmid
Phage
Other
No_hit
Species
E. faecium (293)
S. aureus (128)
K. pneumoniae (5171)
A. baumannii (7345)
P. aeruginosa (9950)
E. cloacae (688)
Proportion of spacers


Phylogeny (MLST)
Gene profile
No_crispr
I + II
I + III
II + III
I + IV
IV
III
II
I
E. faecium
S. aureus
K. pneumoniae
A. baumannii
P. aeruginosa
E. cloacae


E. faecium (II-A)
GO:0005622 - intracellular
GO:0009346 - citrate lyase complex
GO:0042597 - periplasmic space
GO:0022627 - cytosolic small ribosomal subunit
GO:0022626 - cytosolic ribosome
GO:0015935 - small ribosomal subunit
0
10
Number of genes
S. aureus (III-A)
GO:0031224 - intrinsic component of membrane
GO:0016021 - integral component of membrane
GO:0005886 - plasma membrane
GO:0005576 - extracellular region
GO:0009986 - cell surface
GO:0005694 - chromosome
0
10
20
Number of genes
p-value
1e-07
1e-06
1e-05
1e-04
0.001
0.01
0.1
1
K. pneumoniae (I-E)
GO:0016021 - integral component of membrane
GO:0042995 - cell projection
GO:0009289 - pilus
GO:0046930 - pore complex
GO:0030288 - outer membrane-bounded periplasmic
GO:0070069 - cytochrome complex
0
10
20
30
40
Number of genes
A. baumannii (I-Fa)
GO:0016020 - membrane
GO:0031224 - intrinsic component of membrane
GO:0016021 - integral component of membrane
GO:0019867 - outer membrane
GO:0048475 - coated membrane
GO:0030117 - membrane coat
0
10
20
30
Number of genes
A. baumannii (I-Fb)
GO:0031224 - intrinsic component of membrane
GO:0016021 - integral component of membrane
GO:0005576 - extracellular region
GO:0070701 - mucus layer
GO:0015627 - type II protein secretion system comple
GO:0005618 - cell wall
0
10
20
30
40
Number of genes
P. aeruginosa (I-C)
GO:0031975 - envelope
GO:0030313 - cell envelope
GO:0042597 - periplasmic space
GO:0030288 - outer membrane-bounded periplasmic
GO:0009289 - pilus
GO:1990077 - primosome complex
0
2
4
6
Number of genes
P. aeruginosa (I-E)
GO:0015627 - type II protein secretion system comple
GO:0009289 - pilus
GO:0098046 - type V protein secretion system comple
GO:0044096 - type IV pilus
GO:0043231 - intracellular membrane-bounded organ
GO:0043227 - membrane-bounded organelle
0
2
Number of genes
P. aeruginosa (I-F)
GO:0042995 - cell projection
GO:0009289 - pilus
GO:0044096 - type IV pilus
GO:0015627 - type II protein secretion system comple
GO:0098567 - periplasmic side of plasma membrane
GO:0009897 - external side of plasma membrane
0
2
4
Number of genes
E. cloacae (I-F)
GO:0032991 - protein-containing complex
GO:0045263 - proton-transporting ATP synthase com
GO:0045259 - proton-transporting ATP synthase com
GO:0033177 - proton-transporting two-sector ATPase
GO:0016469 - proton-transporting two-sector ATPase
GO:0009337 - sulfite reductase complex (NADPH)
0
10
Number of genes
P. aeruginosa (IV-A1)
GO:0046821 - extrachromosomal DNA
GO:0005727 - extrachromosomal circular DNA
GO:0005694 - chromosome
GO:1990077 - primosome complex
GO:0030894 - replisome
GO:0005657 - replication fork
0
2
Number of genes
K. pneumoniae (kp_IV-A3)
GO:0110165 - cellular anatomical entity
GO:0016020 - membrane
GO:0031224 - intrinsic component of membrane
GO:0016021 - integral component of membrane
GO:0046821 - extrachromosomal DNA
GO:0005727 - extrachromosomal circular DNA
0
10
20
30
Number of genes


K. pneumoniae (I-E)
Cluster
1
2
CRISPRtype
I-E
I-E,I-E
I-E,I-E,IV-A3
I-E,IV-A3
I-E,IV-A3,I-E
I-E,IV-A3,IV-A3
II-A,I-E
IV-A3,I-E
IV-A3,I-E,I-E
IV-A3,I-E,IV-A3
Cluster
CRISPRtype
MLST
B9Q20_13025
manX__2
sorA__2
manZ__4
stdB
papD__4
C0078_17140
abgR__2
citA
higB
SK89_03130
CUW_2277
matC
traD__9
KPK_4936
atxB__2
garP__2
ymoll0001_31880
kdgT__2
SAMEA3531867_02097
ttgG
mexB
unknown2816
tssE__2
cat__8
shlB
UA70_06290
C5672_01265
cioA
saoJ
atxC
rtn
vtxE
AB182_13915
mlpA
ttgE
mdtE
cpsB
cpsE
lysE
PMK1_04394
cvaA__2
KPR_0517__4
PMK1_02009
xycH
apaG__2
fppB__2
frwC
ydeH__2
citA__2
KPN_01341
A. baumannii (I-Fa)
Cluster
1
2
CRISPRtype
I-F
I-F,I-F
I-F,I-F,I-F
Cluster
CRISPRtype
MLST
ABAYE3084
MWQ_04154
pleD__3
ykgB
AsACE_CH02355
btuB__8
J506_2440
ABW55_08355
unknown1683
unknown4743
metQ__5
btuB__6
F961_02349
pilA__7
unknown3544
BIT33_07180
hcpC__7
btuB__5
P23_2774__3
_ribN
ybjJ_2
unknown18362
IX87_10300
J537_3059
cgmA
fhaB__11
doxX
metQ
bpeE
A. baumannii (I-Fb)
Cluster
1
2
CRISPRtype
I-F
I-F,I-F
I-F,I-F,I-F
I-F,IV-A3
Cluster
CRISPRtype
MLST
ybhS
mdtN
PMI1813
fecD
yclQ__1
nepI__2
creC
creD
HH_0050
ABAYE3214__2
gsiB__2
ompA__2
ygaZ
fhaB
exbB__3
cyaD
sauU__1
benP
ymfD__2
exbD__4
gspH__2
ampG__2
nicP__7
nicC
nicC__2
bauA__4
nicT
copS
copD
ABAYE2235
pulG
exbD__2
opuD
betP
PA2757
ABAYE3072
ABAYE3074
ABUW_0364
T630_1336
fimA__5
ABAYE0839
ABAYE0840
ABAYE2349__2
unknown9757
TOL_0522
NE2500
P. aeruginosa (I-C)
Cluster
1
2
CRISPRtype
I-C
I-C,I-C
I-C,I-E
I-C,I-F
I-E,I-C
I-F,I-C
I-F,I-C,I-E
Cluster
CRISPRtype
MLST
CP45
pilV
pph73
CP29
CP74
pph78
chvB
desA
CP98
cfa
lpfB
ecpD__4
rcsC6
rcsC__2
elfC__2
ecpD__3
fhuA
rcsC
fimD
fimB
cmxA
unknown5040
Bpet1404
bcr__5
era__3
ebA93
traD
unknown10041
P. aeruginosa (I-E)
Cluster
1
2
CRISPRtype
I-C,I-E
I-E
I-E,I-C
I-E,I-E
I-E,I-E,I-F
I-E,I-F
I-E,I-F,IV-A1
I-E,I-F,IV-A2
I-E,IV-A1
I-F,I-C,I-E
I-F,I-E
I-F,I-E,IV-A2
IV-A1,I-E
Cluster
CRISPRtype
MLST
pilY1
pilV__2
fimA__2
pilW
pilX
tll0888
bfrD
piuA
fabL
proC__3
pilA__2
fimT__2
pilE
arr
AOY09_00409
hpmA__2
cusB__2
actP__3
hxuA__3
CXK92_18235
unknown2031
p3ABAYE0017
fdhC__2
P. aeruginosa (I-F)
Cluster
1
2
CRISPRtype
I-C,I-F
I-E,I-E,I-F
I-E,I-F
I-E,I-F,IV-A1
I-E,I-F,IV-A2
I-F
I-F,I-C
I-F,I-C,I-E
I-F,I-E
I-F,I-E,IV-A2
I-F,I-F
I-F,IV-A1
I-F,IV-A2
IV-A1,I-F
IV-A2,I-F
Cluster
CRISPRtype
MLST
pilA
fimA__2
epsG
pilV__2
pilW
pilX
pilY1
PA0561
wzzB__2
fimU
pilE
fimT__2
pilX__2
nylB'
arsB
cadA__2
czcD__7
lspA__8
shn_31805
SAMN04490209_3101
lemA__3
ORF C46
cusB__2
BSY15_86
actP__3
pcoA
copO
copS
PAMH19_1227
HMPREF3014_11580
PM1_0200225
arr


K. pneumoniae (I-E)
Cluster 1 (2216)
Cluster 2 (1292)
A. baumannii (I-Fa)
Cluster 1 (230)
Cluster 2 (107)
A. baumannii (I-Fb)
Cluster 1 (423)
Cluster 2 (497)
P. aeruginosa (I-C)
Cluster 1 (121)
Cluster 2 (141)
P. aeruginosa (I-E)
Cluster 1 (186)
Cluster 2 (585)
P. aeruginosa (I-F)
Cluster 1 (1785)
Cluster 2 (1065)
E. cloacae (I-F)
Cluster 1 (30)
Cluster 2 (7)
E. faecium (II-A)
Cluster 1 (133)
Cluster 2 (53)
S. aureus (III-A)
Cluster 1 (173)
Cluster 2 (49)
K. pneumoniae (IV-A3)
Cluster 1 (989)
Cluster 2 (298)
P. aeruginosa (IV-A1)
Cluster 1 (120)
Cluster 2 (9)
Strains (%)
100%
75%
50%
25%
0%
Legend
no_hits
together
both
spacer
phage


a)
Phage gene
(unknown5433)
7
398
68
b)
Genomes without CRISPR-Cas systems
with_phage (283)
without_phage (7727)
100 %
90 %
80 %
70 %
60 %
50 %
40 %
30 %
20 %
10 %
0 %
ABAYE0839 206 0
ABAYE0840 184 0
ABAYE2235 7687 280
ABAYE2349_2 74 0
ABAYE3072 105 2
ABAYE3074 142 2
ABUW_0364 93 0
betP 11 0
exbD_2 7718 283
fimA_5 232 3
NE2500 31 0
opuD 11 0
PA2757 7554 283
pulG 7692 280
cam1 162 278
TOL_0522 32 0
unknown9757 74 0
c)
type VI secretion proteins Rhs?
tRNA-Glu(ttc)
fhuA
ABAYE2651
cam1
wapA
ABAYE2654
unknown3366
ABAYE2656
bacteriophage
tRNA-Arg(cct)
p3ABAYE0080
APC94_20180
unknown4229
R_6
J635_0769
unknown5433
CRISPR-Cas type I-Fb
thiE
csy4(cas6f)
csy3(cas7)
csy2(cas5)
BMY_2099(cas8f)
cas3(cas2_3)
cas1
ACINWCA87_2775
d)
0.2
No_crispr
I-Fa + I-Fb
III-B
I-Fa
I-Fb
I-Fb + IV-A3
*
379 525 55
99% 98% 0%
70% 30% 0%
0% 82% 0%