Antibiotic resistance plasmids spread among natural isolates of *Escherichia coli* in spite of CRISPR elements

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Clustered, regularly interspaced, short palindromic repeats (CRISPRs) are implicated in defence against foreign DNA in various archaeal and bacterial species. They have also been associated with a slower spread of antibiotic resistance. However, experimental and evolutionary studies raise doubts about the role of CRISPRs as a sort of immune system in *Escherichia coli*. We studied a collection of 263 natural *E. coli* isolates from human and animal hosts, representative of the phylogenetic and lifestyle diversity of the species and exhibiting various levels of plasmid-encoded antibiotic resistance. We characterized the strains in terms of CRISPRs, performed replicon typing of the plasmids and tested for class 1 integrons to explore the possible association between CRISPRs and the absence of plasmids and mobile antibiotic resistance determinants. We found no meaningful association between the presence/absence of the *cas* genes, reflecting the activity of the CRISPRs, and the presence of plasmids, integrons or antibiotic resistance. No CRISPR in the collection contained a spacer that matched an antibiotic resistance gene or element involved in antibiotic resistance gene mobilization, and 79.8 % (210/263) of the strains lacked spacers matching sequences in the 2282 plasmid genomes available. Hence, *E. coli* CRISPRs do not seem to be efficient barriers to the spread of plasmids and antibiotic resistance, consistent with what has been reported for phages, and contrary to reports concerning other species.

INTRODUCTION

Clustered, regularly interspaced, short palindromic repeats (CRISPRs) and their CRISPR-associated (*cas*) genes have been identified in most archaea and many bacteria. CRISPRs typically consist of short (23–47 bp) and highly conserved direct repeats interspersed with non-repetitive sequences called spacers (Sorek *et al.*, 2008). The CRISPRs form peculiar genetic loci, which may provide acquired immunity against viruses and plasmids by targeting nucleic acids in a sequence-specific manner (Horvath & Barrangou, 2010; Wiedenheft *et al.*, 2012). CRISPR-encoded immunity is provided by transcription of the repeat-spacer array, followed by transcript processing into small CRISPR RNAs, which are then used in...
combination with Cas proteins to interfere with invasive DNA or RNA (Sapranuska et al., 2011). However, the number, sequence and length of CRISPR arrays and the number and sequence of the Cas proteins are highly variable between bacterial species (Makarova et al., 2011). In Escherichia coli, CRISPRs map in two largely independent pairs of loci, CRISPR1 and 2 located at 62 min on the chromosome, and CRISPR3 and 4 at 20 min, each including a different type of cas gene (subtype I-E and subtype I-F, respectively) (Diez-Villaseñor et al., 2010; Touchon & Rocha, 2010). The functional roles of the CRISPRs in E. coli remain unclear. Effective CRISPR-based protection against phages and/or plasmids cannot be observed in wild-type strains, and its demonstration requires complex genetic manipulation such as H-NS inactivation (Edgar & Qimron, 2010; Pougach et al., 2010; Pul et al., 2010), LuoO overexpression (Wexta et al., 2010), expression of a plasmid-encoded protein targeted for export in cells that lack DnaK (Perez-Rodriguez et al., 2011), or high-temperature protein G expression (Yosef et al., 2011). It is doubtful that such conditions are relevant in the natural environment. Molecular phylogeny analysis of several hundred natural E. coli isolates has shown a strong phylogenetic inertia of CRISPR loci, with CRISPR being almost absent from strains of the major B2 phylogenetic group, and a very slow spacer composition evolution; these observations are at odds with expectations for an active immune system (Delannoy et al., 2012; Diez-Villaseñor et al., 2010; Touchon & Rocha, 2010; Touchon et al., 2011). Lastly, cas genes evolve under purifying selection, without any evidence of positive selection, indicating that there is no evidence that they are involved in an arms race between bacterial and archaeal hosts and invading virus or plasmid DNA (Takeuchi et al., 2012). These various observations led to the suggestion that in E. coli, because CRISPRs are maintained in most lineages, the CRISPR/Cas system may have functions in addition to defence against foreign DNA (Babu et al., 2011).

The aim of this work was to gain further insights into the role of CRISPRs in E. coli by analysing a collection of 263 natural E. coli isolates from human and animal hosts, representative of the phylogenetic and lifestyle diversity of the species and exhibiting various levels of plasmid-encoded antibiotic resistance. In this collection, we characterized the CRISPR content and the presence of mobile elements, i.e. plasmids and integrons, and explored the putative link between the presence/absence of the CRISPRs/Cas systems, the presence of mobile elements and antibiotic resistance. We also searched for CRISPR spacers that target nucleic acid sequences in relation to antibiotic resistance and the acquisition or dissemination of plasmids and integrons.

METHODS

Bacterial strains and antibiotic susceptibility. The collection of 263 natural E. coli isolates used for this study has been described in Touchon et al. (2011). Briefly, they were isolated from various hosts [humans (n=217), domestic animals (n=22), wild animals (n=24)] over a 25-year period (1980–2005) (Branger et al., 2005; Lefort et al., 2011; Picard et al., 1999; Skurnik et al., 2006). The strains were either commensal (119 strains) or associated with extraintestinal infections (144 strains). The CRISPR sequences and the phylogroups (Moissetet et al., 2010; Tenailleon et al., 2010) as determined by specific PCR (Clermont et al., 2000; Clermont et al., 2011a) and multilocus sequence typing (Clermont et al., 2011b; Jauregy et al., 2008) were available for all the strains (Touchon et al., 2011). The proportions of the phylogroups in the collection are: A (23.5%), B1 (17.1%), B2 (33.8%), C (5.5%), D (12.5%), F (4.9%), Escherichia clades (0.7%) and ungrouped (2.6%). Extended-spectrum β-lactamases (ESBLs) were produced by 125 of the strains, including 53, 22 and 50 strains producing a TEM-, SHV- and CTX-M-type ESBL, respectively, all encoded by plasmid genes (Branger et al., 2005; Marcade et al., 2009).

For this study, resistance (mainly plasmid-encoded) to seven antibiotics (ampicillin, cefotaxime, streptomycin, kanamycin, tetracycline, chloramphenicol and sulphonamide) was determined for all the strains. The strains were also tested for penicillinase by PCR or isoelectric focusing (Branger et al., 1997, 2005): 32 non-ESBL- and 59 ESBL-producing strains were penicillinase-positive.

The main characteristics of the strains are given in Table S1 available with the online version of this paper.

Plasmid replicon typing and integron detection. Plasmid replicons were determined using the PCR-based replicon typing scheme, described by Carattoli et al. (2005), which detects 18 plasmid replicons frequently found in Enterobacteriaceae. This technique involves the detection of the coding sequence of repA, the cis-repeats of the origin of replication or the countertranscript RNAI (Carattoli et al., 2005), all elements defining the basic replicons of plasmids.

The strains were screened for the presence of class 1 integrons, the most prevalent integrons in E. coli (Lefort et al., 2011; Skurnik et al., 2005), by PCR amplification of an internal fragment of the integrase gene with a previously described primer pair (intI1L and intI1R) (Ploy et al., 2000).

CRISPR spacer analysis. To identify the spacer sequences matching sequences from plasmids (proto-spacers), the spacers were subjected to the standard BLASTN search (e-value threshold, 1.10⁻5) against 2282 plasmid genomes available in GenBank. Proto-spacers were defined by identifying homologous plasmid sequences with an e-value <1.10⁻5 and less than 10% difference in sequence length.

Statistical analysis. The Pearson’s χ² or the Fisher’s exact test was used, when appropriate, for comparisons. The Student’s t test was used for comparing means. A P value <0.05 was considered significant for all the tests. For correlation analyses, we used the Spearman rho correlation coefficient. Factorial analysis of correspondence (FAC) was used to describe associations between variables. FAC uses a covariance matrix based on χ² distances. The computation determines a plane defined by two principal axes of the analysis; the first axis, F1, accounts for most of the variance, and the second axis, F2, which is orthogonal to F1, accounts for the largest part of the variance not accounted for by F1 (Greenacre, 1992). SPAD.N software (Cisia) was used for FAC with a two-way table. The table had 263 rows, corresponding to the 263 E. coli strains studied, and 30 columns, corresponding to the 30 variables: the seven phylogenetic groups (A, B1, B2, D, C, F and the Escherichia clades), the ungrouped strains, the presence of ESBL and of penicillinase, the three resistance scores (absence of resistance, resistance to one to three antibiotics, resistance to four to seven antibiotics), the presence of class 1 integrons, the presence of five types of replicons [IncF (including IncFI, IncFIA, IncFIB and IncFIC), IncB/O, IncI1, broad-host-range (BHR) replicons including IncL/M, IncA/C, IncN and IncP, and other
replicons (IncK, IncH1, IncH2 and IncY), the absence of described replicons, the presence of two types of cas gene (subtype I-E and subtype I-F), the absence of cas genes, the presence of the four CRISPR loci (CRISPR1–4), and the total number of repeats classified into three categories (2–9, 10–24 and 25–54). For each strain, each variable was coded as a binary code: present=1, absent=0.

RESULTS AND DISCUSSION

Characterization of the plasmid and integron content of the collection

We used a PCR-based replicon typing scheme to analyse the plasmid content of the strains (Carattoli et al., 2005). Although some plasmids are not typable with this scheme (Carattoli, 2009; Marcadé et al., 2009), the results of such PCR typing correlate well with those of S1 nuclease plasmid analysis in a collection of natural E. coli and can therefore be considered to be a good approximation of plasmid content (Bengtsson et al., 2012).

Replicon typing detected 506 replicons in 221 strains (84%) (mean 1.92, range 0–6): 55 strains had one replicon, 78 strains had two, 63 strains had three, 21 strains had four, two strains had five and two strains had six. Fourteen different incompatibility groups were detected: IncF (n=186) [IncFII (n=155), IncFIB (n=142), IncFIA (n=48) and IncFIC (n=10)], IncI (n=37), IncB/O (n=14), BHR replicons (n=70) [IncA/C (n=46), IncN (n=15), IncP (n=9) and IncL/M (n=4)], other replicons (n=25) [IncY (n=9), IncK (n=8), IncH1 (n=5) and IncH2 (n=4)].

As expected, replicons were more prevalent among resistant strains than susceptible strains (89.2% versus 72.7%, respectively; P=0.002). Among strains with plasmids, resistant strains had more different replicons than susceptible strains (mean 2.4 versus 1.92, respectively; P=0.003); similarly, more ESBL-producing strains than non-ESBL-producing strains carried replicons ([92.8% versus 76.81%, respectively; P=0.0003] (mean number of replicons 2.44 versus 2.12, respectively; P=0.02)). There were no such differences between non-ESBL-producing strains producing and not producing penicillinase.

BHR replicons were detected only in resistant strains (70/186 strains) (P<0.0001), which were mostly ESBL-producing strains (n=61) (P<0.0001); most of the non-ESBL strains produced penicillinase (n=7) (P<0.0001). In the ESBL-producing strains, most of these replicons are on the same plasmid as the ESBL genes (Marcadé et al., 2009). IncI1 replicons were found more frequently among ESBL-producing strains than among all other strains (19.2% versus 10.1%; P=0.036). However, combining all resistance markers into one group, there was no difference in the frequency of IncI1 replicons between resistant and susceptible strains. The carriage of the IncF or IncB/O replicons was not associated with the antibiotic resistance of the strains. The remaining replicons were generally more frequent in the ESBL-producing than other strains (P=0.029).

We screened the collection for class 1 integrons. Integrons are mobile elements associated with antibiotic resistance, and class 1 integrons are the most prevalent (Mazel, 2006). Integrons were found in 135 strains, all of which were resistant to antibiotics and most of which produced an ESBL (120 strains) (96% of the ESBL-producing strains versus 10.8% of the non-ESBL producing strains, P<0.0001); seven (21.8%) of the non-ESBL-producing strains produced penicillinase and eight (7.5%) did not (P=0.02).

Comparison of CRISPRs/cas genes according to the presence of replicons and integrons in E. coli strains

Cas proteins are necessary for the functioning of the CRISPR/Cas immune system and are indicators of the system’s activity (Horvath & Barrangou, 2010). To test for an association between the presence and the activity of CRISPRs and the presence of replicons and integrons, we compared the presence of cas genes (both subtype I-E and I-F) between E. coli strains with and without replicons or integrons. There were no significant differences in the prevalence of cas genes between the replicon- or integron-positive and -negative E. coli strains (Table 1).

Plasmids of the IncF, IncB/O and IncI1 groups have narrow host ranges and are commonly found in Enterobacteriaceae (Johnson et al., 2007); IncF plasmids are particularly well adapted to E. coli (Boyd et al., 1996). Other groups of plasmids (IncL/M, IncA/C, IncN and IncP) are considered to have BHRs (Toukdarian, 2004). We analysed the data according to the type of replicon, because CRISPRs may be particularly involved in the defence against BHR replicons. We found no significant differences in the prevalence of the cas system except for strains containing IncF replicons, which had fewer cas genes (28.4% versus 41.5%, P=0.039) (Table 2). These data generally indicate that there is no significant link between CRISPR activity and the presence of foreign DNA (plasmids or integrons).

CRISPRs and acquired antibiotic resistance

Recently, an inverse correlation between the presence of a CRISPR/Cas locus in enterococci and that of acquired antibiotic resistance genes has been reported (Palmer & Gilmore, 2010). Thus, we compared CRISPRs between susceptible and resistant strains, assuming that the observed resistances were largely due to plasmid-borne determinants. We also looked for differences in the production of plasmid-encoded enzymes conferring resistance to β-lactams, i.e. penicillinase and ESBL.

The presence of subtype I-E cas genes did not differ between the susceptible and resistant strains, or between those producing and not producing β-lactamases. The subtype I-F cas system was significantly more prevalent among susceptible strains [n=11 (14.2%) versus n=5 (2.6%); P=0.0003] (Table 1). However, in our collection of strains this system was carried only by a small number of
| Table 1. CRISPR characteristics of the *E. coli* strains studied according to antibiotic resistance, *β*-lactamase production and presence of integrons and replicons |
|--------------------------------------------------|----------------|----------------|----------------|----------------|----------------|
|                              | Antibiotic resistance | ESBL production | Non-ESBL production | Integron | Replicon |
|                              | Presence  | Absence | P     | Presence  | Absence | P     | Presence  | Absence | P     | Presence  | Absence | P     |
| No. of strains               | 186      | 77     |       | 125      | 138     |       | 32       | 106     |       | 135      | 128     |       |
| No. (%) of strains with:     |           |         |       |           |         |       |           |         |       |           |         |       |
| Subtype I-E                  | 118 (63.4) | 45 (58.4) | 0.44 | 78 (62.4) | 85 (61.5) | 0.81 | 18 (56.2) | 67 (63.2) | 0.31 | 87 (64.4) | 76 (59.3) | 0.39 |
| Subtype I-F                  | 5 (2.6)   | 11 (14.2) | 0.0003 | 5 (4.0)   | 11 (7.9)   | 0.17 | 0 (0)     | 11 (10.3) | 0.057 | 5 (3.7)   | 11 (8.5)   | 0.09 |
| Absence of *cas* gene        | 64 (34.4) | 21 (27.2) | 0.26 | 43 (34.4) | 42 (30.4) | 0.49 | 14 (43.7) | 28 (26.4) | 0.061 | 44 (32.5) | 41 (32)   | 0.9  |

*Pase, Penicillinase.

| Table 2. CRISPR characteristics of the *E. coli* strains studied according to the five different types of replicons |
|--------------------------------------------------|----------------|----------------|----------------|----------------|----------------|
|                              | IncF replicon | IncI1 replicon | IncB/O replicon | BHR replicons | Other replicons |
|                              | Presence  | Absence | P     | Presence  | Absence | P     | Presence  | Absence | P     | Presence  | Absence | P     |
| No. of strains               | 186      | 77     |       | 37       | 226     |       | 14       | 249     |       | 70       | 193     |       |
| No. (%) of strains with:     |           |         |       |           |         |       |           |         |       |           |         |       |
| Subtype I-E                  | 122 (65.5) | 41 (53) | 0.06  | 27 (73)  | 136 (60.1) | 0.13 | 8 (57.1) | 155 (62.2) | 0.70 | 48 (68.5) | 115 (59.5) | 0.27 |
| Subtype I-F                  | 12 (6.4)   | 4 (5.1)   | 0.69  | 2 (5.4)   | 14 (6.2)   | 0.85 | 1 (7.1)  | 15 (6.0)  | 0.86 | 2 (2.8)   | 14 (7.2)  | 0.18 |
| Absence of *cas* gene        | 53 (28.4) | 32 (41.5) | 0.039 | 9 (24.3)  | 76 (33.6) | 0.26 | 5 (35.7) | 80 (32.1) | 0.78 | 20 (28.6) | 65 (33.6) | 0.43 |

*Pase, Penicillinase.*
strains of the B2 group (15/89) and a single B1 strain; most group B2 strains with subtype I-F systems were susceptible to antibiotics (11/15).

**Comparisons excluding the phylogenetic group B2 strains**

The major phylogenetic group B2 is almost devoid of CRISPR (Diez-Villasen˜or et al., 2010; Touchon et al., 2011). This may bias the analysis of our collection, so we reanalysed the data, excluding the 89 group B2 strains (19 replicon-negative and 70 replicon-positive strains): almost all (92.5 %) of the 174 non-B2 group strains possessed cas genes.

The distributions of replicons, integrons, antibiotic resistance and β-lactamases in the 174 non-B2 group strains were similar to those in the total collection: 152 (87.3 % versus 84.4 %, respectively) had at least one replicon, 92 had an integron (52.8 % versus 51.3 %, respectively), 48 (27.5 % versus 29.2 %, respectively) were antibiotic susceptible, 83 (47.7 % versus 47.5 %, respectively) produced an ESBL, and among the 91 non-ESBL-producing strains, 20 (21.9 % versus 23.1 %, respectively) produced a penicillinase.

The findings for the non-B2 group strains were similar to those for the entire collection including group B2 strains: there was no correlation between the presence of the subtype I-E cas system and any characteristics of the strains including the type of replicons. The subtype I-F cas system is not found in non-B2 strains, with one exception, as previously reported (Diez-Villasen˜or et al., 2010; Touchon & Rocha, 2010). Thus, exclusion of B2 strains did not substantially change the results.

**Numbers of repeats according to the presence of replicons, integrons or antibiotic resistance**

Another indicator of CRISPR activity could be the median number of repeats. Repeat-spacer units can be both added and deleted, such that a CRISPR can evolve while maintaining an identical number of repeats, although species with a large number of repeats nevertheless tend to have high CRISPR activity (Delaney et al., 2012; Kuno et al., 2012; Tyson & Banfield, 2008). Using this measure, we found no evidence of any negative association between the presence of replicons, integrons or antibiotic resistance and the total number of repeats (data not shown).

**Multidimensional analysis of the data**

We conducted an FAC to have an overview of the data. The first two axes (planes F1 and F2) accounted for 33.56 % of the total variance. The first factor separated the presence of CRISPR1 and 2, large numbers of CRISPR repeats (10–24 and 25–54 categories) and the presence of subtype I-E cas genes from small numbers of CRISPR repeats (two to nine category) and the absence of subtype I-E cas genes (Fig. 1). Thus, factor F1, which accounts for 19.57 % of the total variance, may be associated with the activity of the CRISPR system. The phylogenetic groups were projected along this axis, and group B2 appeared on the opposite side from the others, illustrating the association between the species phylogeny and CRISPRs (Diez-Villasen˜or et al., 2010; Touchon et al., 2011). The second factor, F2, accounted for 13.99 % of the total variance; it appeared to characterize the resistance and the plasmid content of the strains. Thus, a high resistance score (resistance to four to seven antibiotics) and the presence of a class 1 integron and an ESBL were projected at opposite extremes to the absence of resistance, a low level of resistance (resistance to one to three antibiotics) and the presence of a penicillinase. The presence and types of replicons were projected along this axis, with BHR, the other replicons, the IncI1 replicons and the absence of replicons being opposed to the presence of IncB/O replicons and the absence of replicons. The subtype I-F cas genes and the CRISPR4 were closely linked, as previously reported (Diez-Villasen˜or et al., 2010; Touchon et al., 2011), and distinguished by the negative values of the two axes.

This FAC indicates that the strain phylogeny and the CRISPR activity are correlated (axis F1) and that the resistance to antibiotics and the presence of replicons and integrons are also correlated (axis F2), but that strain phylogeny and CRISPRs are independent of antibiotic resistance and the presence of integrons and replicons.

**Sequence similarities between the spacers and available plasmid sequences**

We searched for proto-spacers in 2282 sequenced prokaryote plasmids to assess the role of CRISPRs in the acquisition of plasmids. Most (210/263) strains lacked spacers that matched plasmid sequences. In 53 strains, we identified only 29 different spacers with significant matches (19 spacers had 100 % and 10 had 90.6–96.8 % identity to plasmid sequences); of these, 17 spacers matched sequences in genes encoding functions such as methylase, partition proteins, antirestriction proteins, and replication and transfer systems (Table S2). We found no occurrence of spacers that matched antibiotic resistance genes or elements involved in antibiotic resistance gene mobilization (e.g. Tn3, IScep1, intI1). We observed no association between spacers that matched plasmid sequences and ESBL production or the type of ESBL or replicon. The number of spacers matching plasmid genes in the 53 strains was relatively small: 77.3 % of the strains had one, 13.2 % had two, 7.5 % had three or four and one single strain (IAI48) had 15 (Tables S2, S3 and S4). These spacers were specific for each group of CRISPR. In CRISPR1 and CRISPR2, their diversity was low: only seven different spacers were found in the CRISPR1 of 27 strains and three in the CRISPR2 of 13 strains. For the CRISPR3–4/subtype I-F cas system, the diversity of the spacers was greater: eight different spacers were found in the CRISPR3 of 14 strains and 11 in CRISPR4 of seven
strains. Strains with the same spacers matching plasmids belonged to the same CRISPR spacer relatedness group as defined by Touchon et al. (2011), reflecting phylogenetic inertia (Tables S3 and S4).

Concluding remarks

We were not able to find any solid or meaningful correlation between the presence of plasmids, integrons, acquired antibiotic resistance and CRISPRs. This is surprising given the suggested role of CRISPRs in moderating plasmid transfer (Horvath & Barrangou, 2010); it is also contrary to findings in enterococci, in which CRISPR are negatively associated with antibiotic resistance (Palmer & Gilmore, 2010). Indeed, the probability of acquiring plasmids and integrons and thereby becoming multi-resistant should be higher for strains with less active CRISPRs, if such activity inhibits transfer. Our findings are consistent with the observation that the sensitivity of an E. coli collection, representative of the species diversity, to a set of 59 coliphages (Kutter, 2009) does not correlate with CRISPR content (Diez-Villasenor et al., 2010). The patterns of distribution that we report suggest that the presence of CRISPRs has little effect on the epidemiology of plasmids in E. coli or on the spread of antibiotic resistance. The 20% of the strains that retain spacers with identity to plasmids could indicate ancient contacts with plasmids, the CRISPRs being no longer active.

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