Azithromycin resistance levels and mechanisms in *Escherichia coli*

Cláudia Gomes¹, Lidia Ruiz-Roldán¹,⁴, Judit Mateu¹, Theresa J. Ochoa²,³ & Joaquim Ruiz¹

Despite azithromycin being used in some countries to treat infections caused by Gram-negative pathogens, no resistance breakpoint for *Escherichia coli* exists. The aim of this study was to analyse the levels and mechanisms of azithromycin resistance in *E. coli*. The presence of chromosomal (rplD, rplV and 23S rRNA) mutations, 10 macrolide resistance genes (MRGs) and efflux pump overexpression was determined in 343 *E. coli* isolates. Overall, 89 (25.9%) isolates had MICs ≥ 32 mg/L to azithromycin, decreasing to 42 (12.2%) when assayed in the presence of Phe-Arg-β-Napthylamide, with 35 of these 42 possessing at least one MRG. Efflux pumps played a role in azithromycin resistance affecting the Minimal Inhibitory Concentration (MIC) levels of 91.2% isolates whereas chromosomal alterations seem to have a minimal role. At least one MRG was found in 22.7% of the isolates with mph(A) being the most commonly found gene. The mph(A) gene plays the main role in the development of azithromycin resistance and 93% of the mph(A)-carrying isolates showed a MIC of 32 mg/L. In the absence of a specific resistance breakpoint our results suggest a MIC of 32 mg/L to be considered in order to detect isolates carrying mechanisms able to confer azithromycin resistance.

Infantile diarrhoea is a serious problem in developing countries and remains the second most common cause of death among children under five worldwide. In fact, it causes >800,000 deaths globally per year representing around 10–11% of the annual global child deaths¹⁻². *Escherichia coli* play a relevant role in the death of children by diarrhoea, being involved in more than 120,000 deaths annually of children under 5 years old³. The treatment approach to diarrhoea often does not require the use of antibacterial agents being frequently limited to the replacement of lost liquids and salts by means of Oral Rehydration Salts solutions in order to fight the dehydration risks³. However, according to the patient's nutritional status, the presence of comorbidities, the specific pathogen, illness severity and symptom duration, the use of antimicrobial agents may be required. Ampicillin and cotrimoxazole are the usual first line treatments in most low and middle-income countries⁴⁻⁵. Unfortunately, antimicrobial resistance has increased over time, and in different areas these antimicrobial agents are losing their usefulness as a treatment of diarrhoea⁴⁻⁷. Since antibiotic resistance is a severe health problem worldwide which can lead to inefficiency of antimicrobial agents and therapeutic failure⁸, surveillance of the development of antimicrobial resistance should be performed, establishing molecular mechanisms of resistance to thereby design alternative treatments.

Azithromycin and other macrolides have been largely used to treat Gram-positive infections and also possess good activity against different Gram-negative microorganisms, such as *Bartonella* spp., *Campylobacter* spp., *Haemophilus influenzae*, or *Neisseria gonorrhoeae*⁹⁻¹⁰. Classically, macrolides present low levels of activity against *Enterobacteriaceae* which have been related to the poor membrane penetration of these antimicrobial agents, preventing their use to treat *Enterobacteriaceae*⁹. Nonetheless, in comparison with other macrolides, azithromycin has a higher basic character⁹. Thus, while low permeability prevents the action of most of macrolide agents against *Enterobacteriaceae*⁹, this basic character confers to azithromycin a true role in the treatment of diarrhoeal infections related to different *Enterobacteriaceae*¹¹⁻¹². Thus, azithromycin is a promising alternative because of its excellent activity against most common diarrhoeagenic pathogens such as diarrhoeagenic *E. coli*, *Shigella* spp., *Salmonella* spp. or *Campylobacter* spp.¹³⁻¹⁶, and has been included in the considered armamentarium to fight against specific *Enterobacteriaceae*¹³⁻¹⁴.

Nonetheless, despite ranking amongst the most frequent etiological causes of diarrhoea¹⁵⁻¹⁶, and the association of some specific diarrhoeagenic pathotypes with high levels of children mortality¹⁸, at present no clinical breakpoint for resistance in *E. coli* has been established. However, a Minimal Inhibitory Concentration
(MIC) $\geq 32$ mg/L or a halo diameter $\leq 12$ mm have been proposed as the azithromycin resistance breakpoints in some Enterobacteriaceae\cite{17,18}. Furthermore, a series of questions on the use of azithromycin in the treatment of diarrhoeagenic Enterobacteriaceae remain to be fully answered. These include questions such as specific azithromycin resistance rates, azithromycin resistance mechanisms in circulation, as well as a more relevant question, such as the effect of different alterations on the final azithromycin MIC.

Chromosomal efflux pumps are bacterial systems involved in the extrusion of molecules from bacteria to the environment, including bacterial products such as siderophores as well as toxics and antibiotics\cite{19}. In this line chromosomal efflux pumps are involved in intrinsic and acquired azithromycin resistance\cite{9,20}. Additionally, target amino acid substitutions in the L4 (rplD) and L22 (rplV) ribosomal proteins and in 23S rRNA (rrlH) have also been involved in macrolide resistance\cite{5}.

Nonetheless, the most relevant mechanisms of azithromycin resistance in Enterobacteriaceae are those encoded in mobile elements\cite{8}. Different Macrolide Resistance Genes (MRGs) have been described, leading to resistance through different pathways such as target modifications produced by rRNA methylases encoded in ermA genes or macrolide-inactivation, mediated by esteras as those encoded by ere(A) or ere(B) genes or by phosphorylases such as those encoded in the mph(A) and mph(B) genes. Additionally, transferable genes such as msr(A), mef(A) or mef(B) have been reported to encode macrolide-efflux pumps\cite{5}.

This study aimed to evaluate the levels and the mechanisms of resistance to azithromycin in a collection of samples of E. coli from children with and without diarrhoea. In the absence of a specific azithromycin breakpoint for E. coli, we analyse the relationship between specific mechanisms of resistance and MIC levels.

### Results

#### Antibiotic susceptibility levels.

The MICs of azithromycin ranged between 0.06 mg/L and $>256$ mg/L, with a MICmin of 8 mg/L and MICmax of 128 mg/L (Table 1).

Overall, 140 (40.8%) and 89 isolates (25.9%) had a MIC $\geq 16$ and $\geq 32$ mg/L, respectively, while only 18.7% and 11.9% ($P<0.0001$ in both cases) remained with a MIC $>16$ and $>32$ mg/L respectively when Phe-Arg-$\beta$-Naphthylamide (PAβN) was added (Table 1, Figs 1 and 2). When the analysis was made comparing diarrhoeagenic and commensal E. coli no differences were observed. Nonetheless, when analysing the isolates by pathotypes the levels of resistance of enteroaggregative (EAEC) (48.7%) and diffuse-adhering (DAEC) (45%) were significantly higher than those of enterotoxigenic (ETEC) (17.1%) and enteropathogenic (EPEC) (10%). Moreover, the resistance levels of EAEC isolates were also significantly higher than those of commensal isolates ($P = 0.0007$) and ETEC isolates ($P = 0.0007$).

In the presence of PAβN all groups showed decreased levels of resistance, which was significant ($P = 0.0080$) amongst EAEC isolates (Table 1).

#### Effect of PAβN

In all cases the isolates were able to grow in the presence of PAβN. As mentioned above the addition of PAβN affected the azithromycin susceptibility levels (Tables 1 and 2, Figs 1 and 3). Overall, when the MIC was established in the presence of PAβN (MICPAβN) the effect of PAβN on the MIC levels was observed in 91.2% of the isolates, independently of the initial MIC (MICi) of azithromycin, with 256 being the maximum MIC/MICPAβN quotient (from MIC; of 64 mg/L to MICPAβN of 0.25 mg/L) (Table 2). In 47 out of 89 (52.8%) azithromycin-resistant isolates, the addition of PAβN resulted in a MIC within the range of susceptibility (Table 1, Fig. 1). On the other hand, 35 out of these 47 isolates (74.5%) possessed at least 1 MRG (unidentified in one case - see conjugation results below).

Two commensal and 4 diarrhoeagenic isolates presented a MICi $>256$ mg/L and a MICPAβN $\geq 256$ mg/L, thereby not allowing the effect of PAβN to be accurately established.

### Table 1. Analysis of azithromycin resistance by E. coli categories. PAβN: Phe-Arg-$\beta$-Naphthylamide; Com: Commensal, EPEC: Enteropathogenic; ETEC: Enterotoxigenic; EAEC: Enterocytogenic; DAEC: Diffuse-adhering; DEC: Diarrhoeagenic; MIC: Minimal Inhibitory Concentration (expressed in mg/L); R: Resistance (considering MIC $\geq 32$ mg/L); N: Without PAβN; Y: With PAβN; P: Differences between resistance levels in the absence and presence of PAβN (highlighted in bold the significant differences found). \(4^*$EAEC isolates were significantly more resistant than commensal ($P = 0.006$), EPEC ($P < 0.0001$) and ETEC isolates ($P = 0.0007$). \(5^*$DAEC isolates were significantly more resistant than EPEC ($P = 0.0004$) and ETEC ($P = 0.0302$).

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|---|---|---|---|---|---|---|---|
| MIC Range | PAβN | Com. (84) | EPEC (120) | ETEC (41) | EAEC (78) | DAEC (20) | Total DEC | Overall (343) |
| MIC<sub>i</sub> | N | 2 $\geq$ 256 | 0.06 $\geq$ 256 | 2 $\geq$ 256 | 2 $\geq$ 256 | 1 $\geq$ 256 | 0.06 $\geq$ 256 | 0.06 $\geq$ 256 |
| Y | 0.06 $\geq$ 256 | 0.25 $\geq$ 64 | 0.25 $\geq$ 256 | 0.25 $\geq$ 128 | 0.06 $\geq$ 256 | 0.06 $\geq$ 256 |
| MIC<sub>p</sub> | N | 16 | 6 | 4 | 16 | 16 | 8 | 8 |
| Y | 2 | 1 | 1 | 2 | 4 | 1 | 1 |
| MIC<sub>y</sub> | N | 128 | 16 | 64 | $>256$ | 128 | 128 | 128 |
| Y | 32 | 2 | 4 | 64 | 32 | 32 | 32 |
| R (Nos/%) | N | 23 (27.4) | 12 (10.0) | 7 (17.1) | 38 (48.7) | 9 (45.0) | 66 (25.5) | 89 (36.6) |
| Y | 13 (15.5) | 4 (3.3) | 1 (2.4) | 21 (26.9) | 3 (15.0) | 29 (10.8) | 42 (12.2) |
| P | 0.0897 | 0.0671 | 0.0571 | 0.0080 | 0.0824 | $<0.0001$ | $<0.0001$ |

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As a general rule the MIC$_{I}$/MIC$_{P\beta N}$ quotient ranged from 4 to 16 (267 isolates, 77.8% of total isolates). The MIC$_{I}$/MIC$_{P\beta N}$ mode was 8 (overall, and among commensal and diarrhoeagenic groups), while the mean effect was 12 (Table 2). When the diarrhoeagenic group was subdivided into pathotypes, only DAEC and EAEC showed slight differences (Tables 1 and 2).

![Figure 1](image1.png)

**Figure 1.** Analysis of Minimal Inhibitory Concentration (MIC) of 16 and 32 mg/L to detect the presence of specific macrolide-resistance mechanisms. R: Resistance; R$_{P\beta N}$: resistance in presence of PA$_{P\beta N}$. (a) Overall. (b) Isolates in which no sought mechanism of resistance was found. (c) Isolates carrying the mph(A) gene alone or with a target mutation. (d) Isolates carrying the mph(A) gene together other MRG. (e) Isolates carrying a MRG different that mph(A). (f) Isolates carrying only L4 and/or L22 amino acid changes. The single isolate (isolate 3491) which remains resistant after PA$_{P\beta N}$ addition possesses an unidentified MRG.

![Figure 2](image2.png)

**Figure 2.** Minimal Inhibitory Concentration (MIC) distribution. MRG: Macrolide resistance gene (other than mph(A)); wt: wild type. Any MIC category with ≥5% of the isolates is highlighted in dark grey. If a strain had a L4 and/or L22 mutation(s) and a MRG, then the isolates are included in either the mph(A) or MRG category.

|   | MIC$_{I}$/MIC$_{P\beta N}$ |   |
|---|--------------------------|---|
|   | 40.0% - 55 isolates | 8.7% - 12 isolates |
|   | 28.5% - 72 isolates | 7.5% - 10 isolates |
|   | 11.1% - 20 isolates | 2.8% - 3 isolates |
|   | 0.4% - 1 isolate | 0.0% - 0 isolates |

As a general rule the MIC$_{I}$/MIC$_{P\beta N}$ quotient ranged from 4 to 16 (267 isolates, 77.8% of total isolates). The MIC$_{I}$/MIC$_{P\beta N}$ mode was 8 (overall, and among commensal and diarrhoeagenic groups), while the mean effect was 12 (Table 2). When the diarrhoeagenic group was subdivided into pathotypes, only DAEC and EAEC showed slight differences (Tables 1 and 2).

Analysing the effect of PA$_{P\beta N}$ in 255 diarrhoeagenic and 82 commensal isolates, a non-significant trend of a higher number of affected commensal isolates was observed ($P = 0.0810$). Thus, the effect of PA$_{P\beta N}$ was not observed in 8.6% and 2.4% diarrhoeagenic and commensal isolates respectively. Despite the significant effect of PA$_{P\beta N}$ on the MIC of EAEC isolates, 11 (14.7%) were not affected by PA$_{P\beta N}$. Interestingly, 10 out of these 11 isolates presented MIC$_{I}$ of 64–32 mg/L and MIC$_{P\beta N}$ of 32–16 mg/L, with MRGs being detected in only 2 cases. In
addition, 3 DAEC isolates (15%) were also not affected by PAβN presenting borderline significant differences with commensal isolates.

Only 17 out of 263 isolates analysed (6.5%) presented mutations in the rplD or rplV genes. Thus, 6 isolates had mutations in the rplD gene and 7 in the rplV gene, while 4 isolates presented amino...
Acid codon alterations concomitantly in both genes. Thirteen of these had a MIC≥32 mg/L (including 3 presenting mutations in both of the targets analysed), but only one (isolate 3491), in which an unidentified MRG was detected by conjugation (see below), remained resistant when the MICPAβN was established. In 4 cases were detected concomitant MRGs (Table 4). None of the isolates analysed had mutations in the 23S rRNA gene.

Macrolide resistance genes. Seventy-eight isolates (22.7%) possessed at least one MRG (Table 5). The MRG most frequently found was mph(A), which was present in 53 isolates (67.9% of isolates possessing MRG) belonging to all the groups analysed. In 43 cases no other MRG was detected, while in the remaining 10 cases mph(A) was detected together with the erm(A) gene in 4 cases, the erm(B) gene in 3 cases and the mef(A) and ere(A) gene in 2 and 1 cases, respectively. When more than one MRG was identified within the same isolate the mph(A) gene was always present.

Macrolide resistant genes (MRGs) absence (265) presence 1 MRG (68) 2 MRGs

| MIC Range | PAβN | Absence (265) | Presence mph(A) (43) | Other (25) | Overall (10)* |
|-----------|------|--------------|------------------|------------|--------------|
| Y         | N    | 0.06–>256    | 8–>256           | 4–64       | 64–>256 |
|           | Y    | 0.06–128     | 0.25–>256        | 0.25–32    | 16–>256 |
| MIC<sub>cp</sub> | N | 8 | 8 | 256 |
|           | Y | 1 | 32 | 1 | 32 |
| MIC<sub>cp</sub> | N | 32 | >256 | 64 | >256 |
|           | Y | 4 | 256 | 16 | 128 |
| R (N/%)   | N | 81/30.6% | 41/95.4%** | 8/32% | 10/100% |
|           | Y | 21/7.9% | 29/67.4% | 4/16% | 10/100% |
| P         | <0.0001 | 0.003 | 0.3209 | 1.000 |

Table 3. Analysis of azithromycin resistance in the presence and absence of macrolide resistance genes. PAβN: Phe-Arg-β-Naphtylamde; MRG: Macrolide resistance gene; N: Absence of PAβN; Y: Presence of PAβN. MIC: Minimal Inhibitory Concentration (expressed in mg/L). P: Azithromycin resistance (considering MIC≥32 mg/L). A non-identified conjugative mechanism of resistance was detected (isolate 3491).

| E. coli | L4 | L22 | MRG | MIC ± PAβN |
|---------|----|-----|-----|------------|
|         |    |     |     |            |
| Commensal | V52I | I4L + L6Q + T72A | — | 16 0.125 |
| Commensal | A37S + V52L | wt | — | 16 1 |
| Commensal | wt | I4L + K6Q + T72A | — | 16 1 |
| Commensal | V52I + D91E + T173N | wt | — | 16 2 |
| Commensal | wt | S101T + I103L | — | 64 2 |
| DAEC | wt | K83N + D94H + K98N | mph(A) | 64 4 |
| EAEC | wt | V17I | — | 2 1 |
| EAEC | A37T + K74T | wt | — | 4 1 |
| EAEC | V120I | wt | — | 4 1 |
| EAEC | wt | L46Q | mph(A) | 64 16 |
| EAEC | K123S | I4L + K6Q | — | >256 128 |
| EPEC | A190V | wt | mcr(A) | 8 0.5 |
| EPEC | D154E | wt | — | 8 1 |
| EPEC | V52I + T173N | I4L + K6Q + T72A | mph(B) | 16 4 |
| EPEC | K123S | I4L + K6Q + T72A | — | 32 1 |
| ETEC | wt | L46Q | — | 64 16 |
| ETEC | wt | L46Q | — | 128 2 |

Table 4. L4 (rpI/D) and L22 (rpI/V) amino acid substitutions. PAβN: Phe-Arg-β-Naphtylamde; MRG: Macrolide resistance gene; wt: wild type. N: Absence of PAβN; Y: Presence of PAβN. *A non-identified conjugative mechanism of resistance was detected (isolate 3491).
| E. coli | N | Phosphotransferases | Methylesses | Esterases | Efflux Pumps | Overall |
|--------|----|---------------------|------------|-----------|--------------|---------|
|        |     | mph(A) | mph(B) | ermA | ermB | ermC | ere(A) | mef(A) | mef(B) | mcr(A) | mcr(D) | N | % | Genes |
| EAEC   | 78  | 21     | 0      | 1      | 3a    | 3b   | 1     | 3a     | 3b     | 0     | 0    | 1  | 29b | 39.8 | 37 |
| EPEC   | 120 | 6      | 1      | 0      | 2     | 0    | 0     | 0      | 1      | 1     | 2    | 13 | 10.8 | 13 |
| ETEC   | 41  | 2      | 0      | 0      | 1     | 2    | 2     | 0      | 0      | 1     | 0    | 6  | 14.6 | 6  |
| DAEC   | 20  | 10     | 0      | 0      | 1     | 0    | 2     | 0      | 0      | 1     | 0    | 6  | 14.6 | 6  |
| DEC    | 259 | 39     | 1      | 5      | 4     | 4    | 5     | 3      | 1      | 2     | 3   | 58 | 23.2 | 67 |
| Comm.  | 84  | 14     | 0      | 1      | 2a    | 0    | 2     | 0      | 1      | 1     | 0    | 20 | 23.8 | 21 |
| Overall| 343 | 53     | 1      | 6      | 6     | 4    | 7     | 3      | 2      | 3     | 3   | 78 | 23.3 | 89 |

Table 5. Macrolide resistance genes. EAEC: Enteroaggregative; EPEC: Enteropathogenic; ETEC: Enterotoxigenic; DAEC: Diffusely Adherent; DEC: Diarrhoeagenic; Comm.: Commensal. 1 of them concomitantly with mph(A); 2 of them concomitantly with mph(A); 3 of them concomitantly with mph(A); 4 of them concomitantly with mph(A). Overall the EAEC isolates possess more MRGs than EPEC (P < 0.0001) and ETEC (P = 0.0113). Overall the DAEC isolates possess more MRGs than EPEC (P < 0.0001), ETEC (P = 0.0053) and commensal (P = 0.0283).

The presence of the mph(A) gene was correlated with higher MIC levels (Table 3, Figs 1, 2 and 3), while the presence of other MRGs alone seemed to have a lesser effect. In fact, 40 out of 43 isolates presenting the mph(A) gene as a single MRG had MICs ≥ 32 mg/L. Interestingly, those isolates presenting the mph(A) gene together with another MRG exhibited slightly higher MIC values than those possessing only the mph(A) gene (Fig. 1). The effect of PAβN on the 25 isolates carrying any other MRG was significantly higher (P < 0.0001) than in those isolates with the mph(A) gene. Thus, only 2 ermA, 1 ere(A) and 1 ermC carrying isolates were classified as non-wt when PAβN was added.

**Conjugation assay.** Transconjugants with MICs ≥ 32 mg/L were observed in 16 (24.2%) out of 66 isolates analysed. The mph(A) gene was transferred in 14 cases and the ermA gene in 3 cases (2 together with mph(A)). Finally, 1 transconjugant was obtained from a parental isolate (strain 125: MIC < 256 mg/L; MICPAβN = 128 mg/L, carrying amino acid changes in L4 [K123S] and L22 [I4L, K6Q]) in which no MRG was previously detected.

**Wt/non-wt phenotypes and MIC levels.** Overall, 22 out of 78 (28.2%) isolates carrying at least one MRG presented MIC levels < 32 mg/L. Of these, 3 isolates harbouring the mph(A) gene alone (7% of isolates carrying the mph(A) gene alone; 3.8% of isolates carrying MRG) and 19 carrying other MRGs other than mph(A) alone (76% of isolates carrying other MRGs; 86.4% of isolates with MIC < 32 carrying any MRG) having a MIC < 32 mg/L. No isolates possessing more than one MRG presented a MIC < 32 mg/L (Figs 1 and 2). The cumulative MIC curves of wt isolates and those presenting a MRG other than mph(A) were similar. The cumulative MIC curves of the isolates possessing target mutations, mph(A) and mph(A) plus other MRG were sequentially displaced towards higher MIC levels. When the cumulative MICs were established in presence of PAβN the results showed that those belonging to wt isolates, and those presenting MRG or L4/L22 amino acid substitutions were close similar, while only a spurious displacement towards high MIC levels of those non-wt, while isolates possessing mph(A) and mph(A) plus other MRG were sequentially displaced towards higher MIC levels in a clear manner (Fig. 3).

**Discussion**

Diarrhoea-related deaths in children remain among the most relevant health challenges worldwide, being of special concern in low- and middle-income countries. In these countries, antibiotic therapy when needed may be crucial to achieve a successful outcome. However, antibiotic resistance to commonly used antibacterial agents is dramatically increasing requiring new alternatives.

Regarding the feasibility to consider azithromycin as an alternative to treat diarrhoeagenic E. coli in the studied areas, the present study showed moderate azithromycin resistance levels highlighting some concerns about its usefulness as treatment in the absence of antibiotic susceptibility data, especially when EAEC or DAEC isolates are present.

In accordance with what has been previously described, the relevant role of PAβN-inhibitable efflux pumps in azithromycin resistance has been demonstrated once more. However, differences related to the specific bacteria groups were observed. The presence of a series of EAEC isolates in which no PAβN-effect was observed opens the door to different options, including the presence of alterations in the outer membrane composition which results in a possible azithromycin impaired permeability leading to an increase in the basal azithromycin resistance levels, combined with lesser efflux pump activity, at least in regard to PAβN-inhibitable efflux pumps. Another possibility is the presence of different patterns of overexpressed efflux pumps. In this line, selecting azithromycin resistant mutants in the presence of PAβN a similar scenario was observed (MIC of 32–16 mg/L with no further PAβN effect). In all these mutants the presence of an overexpressed OmpW was observed. In fact, OmpW has been associated with EmrE, an efflux pump belonging to the small multidrug resistance (SMR) family. Furthermore, the overexpression of EmrE has been related to E. coli grown in the presence of erythromycin.

In agreement with the presence of up to 7 gene copies and the subsequent need for multiple mutated alleles to visualize an effect on macrolide resistance, in the present study no mutations in the 23S rRNA gene were...
observed in the 66 isolates analysed. Regarding L4 and L22, the alterations detected seem to have a minor role in the development of azithromycin resistance, and most might be gene polymorphisms without antibiotic resistance relevance. Regarding the alterations at L4 and L22 observed, to our knowledge only the alterations at amino acid codon K82, K98 and K98 of L22 have previously been described in vitro obtained E. coli macrolide-resistant mutants but always concomitantly with other L22 amino acid alterations. The L22 alteration L46Q was present in 3 cases, all having a MIC ≥ 32 mg/L. Although in one case the addition of PA3N resulted in a MIC of 2 mg/L, and another was concomitantly present with the mph(A) gene, a possible slight effect of this alteration on macrolide susceptibility cannot be ruled out.

Regarding MRGs, in our series the relevant role of Mph(A) is undoubtable. This finding is in accordance with what has been previously described in E. coli and other Enterobacteriaceae. Those isolates with the mph(A) gene presented the highest percentages of azithromycin resistance both in the presence and the absence of PA3N. Nonetheless, relevant differences were observed in the MIC levels among isolates carrying the mph(A) gene. Thus, while 2 mph(A)-carrying isolates had a MIC of 8 mg/L which decreased to MICPA of 0.25 and 1 mg/L, another 11 isolates in which no other MRG was detected had a MIC of 256 mg/L which in no case decreased below the breakpoint considered in the presence of PA3N. This heterogeneity may be observed on analysing together different studies performed either in E. coli or other closely related Enterobacteriaceae. Different explanations may be proposed, including differences related to expression levels which may be due to the number of copies of the gene related to its genetic environment (e.g.: plasmids with different sizes and copy numbers), with alterations at the promotor sequence or with the presence of other undetected MRGs.

The remaining MRGs, seemed to have a marginal role in azithromycin resistance. In fact, the cumulative MIC curve of these isolates was close to that of wt microorganisms. Nonetheless, those isolates presenting the mph(A) gene together with another MRG ranked among those most resistant and less affected by the addition of PA3N, suggesting a slight contribution of other MRGs to final MIC levels when mph(A) gene is present. This finding was also showed when cumulative MICs were established.

Of these MRGs, among Enterobacteriaceae, the Msr(A) has only been described in E. coli and Enterobacter spp. In the present study, the mrr(A) gene was detected in isolates having MIC of 8 mg/L, supporting the loss of activity of this gene when cloned in E. coli. The other ATP binding transporter studied, Msr(D), was detected independently of the presence of Mef(A). Moreover, in no case the mef(A) gene and the mrr(D) genes were detected together. To our knowledge this is the first description of the mrr(D) gene alone, since it has always been described concomitantly with mef(A). Nevertheless, the presence of polymorphisms in the mef(A) primers annealing region cannot be ruled out. While the effect of Msr(D) on the final MIC levels was within the range of those previously described, this dissociation might result in impaired Mef(A). Contrary to what was observed in the present study, Mef(A) has been described to be frequent in Enterobacteriaceae. This difference may be related to the geographical origin of the samples.

This is the first description of Erm(A) in Enterobacteriaceae. While no data on erm(A) functionality in Enterobacteriaceae has been found, previous studies have described an impairment in the expression levels of erm(C), which, if combined with a limited gene copy number, might result in a marginal influence on azithromycin MIC levels such as detected in present study. Regarding Erm(B), the concomitant presence with mph(A) detected here in 3 isolates, has also been previously described. Also Ere(A) had a minimal role in the resistance to azithromycin in the present isolates. This finding is in accordance with the proposed lack of activity of Ere(A) in azithromycin.

There is controversy about the ability of Mph(B) to hydrolyse azithromycin. Thus, while Chesneau and colleagues have described its inability to confer azithromycin resistance, other authors have established a similar activity on hydrolysing erythromycin and azithromycin. The only isolate of our study that possessed the mph(B) gene exhibited an azithromycin MIC of 16 mg/L in the absence of PA3N. Despite this marginal role of most MRGs in the final azithromycin MIC, the detection of 6 out of 10 MRGs among commensal E. coli is noteworthy because of their role as a gene-reservoir. Conjugation studies showed that only the mph(A) or erm(B) genes were transferred alone or together. Additionally, in one case in which no MRG was previously detected, transconjugants were obtained showing the presence of an undetermined MRG. In fact other MRGs have been described in E. coli. Nonetheless, the presence of sporadic E. coli isolates possessing Mph(A) with MIC values of 8–16 mg/L was also showed. Therefore studies are needed to determine the possible need for more conservative breakpoint.

In summary, the present data demonstrate the presence of azithromycin resistance among intestinal, either pathogenic or not, E. coli from the area of Lima, highlighting the need for susceptibility data to adequately use this antimicrobial agent. Moreover, the relevant and hidden role of efflux pumps in the intrinsic levels of azithromycin
in the study) were used to validate the results. Additionally random selected positive PCRs were sequenced.

isolates by PCR (Table 6). In all cases negative and positive controls (microorganisms carrying the MRGs included colleagues as a quality control.

Madison, Wi) following the manufacturer's instructions and thereafter sequenced (Macrogen, Seoul, Korea).

obtained from parents and/or children legal guardians and all experiments were performed in accordance with relevant guidelines and regulations.

To study. The 

isolates from faeces samples collected in previous studies from children under 5 years of age in periurban areas of Lima (Peru) were recovered from frozen stocks 

E. coli 

Bacterial strains. Three hundred forty-three diarrhoeagenic (259 isolates, including 78 EAEC, 41 ETEC, 20 DAEC and 66 EPEC) or commensal (84 isolates) 

E. coli isolates from faeces samples collected in previous studies from children under 5 years of age in periurban areas of Lima (Peru) were recovered from frozen stocks to be included in the study. The uidA gene of all grown isolates was amplified as previously described by Walk and colleagues as a quality control.

In all cases the previous studies in which were collected the E. coli isolates were approved by the Ethical Committee of the Universidad Peruana Cayetano Heredia, faeces were sampled after informed consent was obtained from parents and/or children legal guardians and all experiments were performed in accordance with relevant guidelines and regulations.

Antimicrobial susceptibility testing. The MIC of azithromycin was determined by the agar dilution method in accordance with the CLSI guidelines in the absence (MIC<sub>IC<sub>50</sub></sub>) and presence (MIC<sub>IC<sub>90</sub></sub>) of 20 mg/L of PA<sub>N</sub> in the statistical analysis.

The effect of 20 mg/L of PA<sub>N</sub> on the viability of microorganisms was also assessed. The PA<sub>N</sub> effect on the MIC levels was considered when MIC<sub>IC<sub>90</sub></sub>/MIC<sub>IC<sub>50</sub></sub> > 2. The isolates with a MIC > 256 mg/L that remained unaltered or decreased to 256 mg/L when PA<sub>N</sub> was added were not considered in the statistical analysis.

Ribosomal target gene amplification and DNA sequencing. In a random selected subset of 263 (rplD and rplV genes) and 66 samples (23S rRNA) the presence of point mutations was established by PCR (Table 6), as previously described. The amplified products were recovered with Wizard SV Gel and the PCR Clean Up System (Promega, Madison, WI) following the manufacturer's instructions and thereafter sequenced (Macrogen, Seoul, Korea).

Transferable azithromycin resistance mechanism detection. The presence of 10 established MRGs (erm(A), erm(B), erm(C), ere(A), mph(A), mph(B), msr(A), msr(D), mef(A) and mef(B) genes) was sought in all isolates by PCR (Table 6). In all cases negative and positive controls (microorganisms carrying the MRGs included in the study) were used to validate the results. Additionally random selected positive PCRs were sequenced.

Conjugation assays. A total of 66 isolates with a MIC ≥ 32 mg/L were selected to determine the transferability of the MRGs. The conjugation was carried out in Luria-Bertani broth (Conda, Madrid, Spain) with azide-resistant E. coli J53 as a recipient strain. Transconjugants were selected in plates containing 150 mg/L of sodium azide and 32 mg/L of azithromycin. In order to avoid considering possible contaminations the relationship of transconjugants and the respective recipient strain was established by REP-PCR. The amplification of the MRGs present in the donor and derived transconjugant strains was performed by PCR as mentioned previously.

| Target | Gene | Prot | Primers | Size (bp) | Ann. (°C) | Ref. |
|--------|------|------|---------|----------|----------|------|
| Macrolide Resistance Genes | erm(A) | EreA | GCGGGTGCTCATGAACCTTGGAG | 420 | 60 | 29 |
| | erm(B) | ErmA | TCTAAAAGCAGTTAAAAGAAA | 533 | 52 | 29 |
| | erm(C) | ErmC | CGATATCTTTTTGTACCTTTC | 639 | 45 | 29 |
| | mef(A) | MeA | AAAAAAGTACTACAACCAATA | 345 | 54 | 29 |
| | mef(B) | MeB | TCTCTCTGCTATCATAAGTGG | 430 | 60 | 29 |
| | mph(A) | MphA | GCTGAGGAGACCTGGCGG | 403 | 60 | 29 |
| | mph(B) | MphB | ATTAAACAGTATCAGGATAGC | 868 | 50 | 29 |
| | msr(A) | MsrA | GCCATTATGTGGAGTAATG | 384 | 58 | 29 |
| | msr(D) | MsrD | CCCCAGTTGAGCAGGAAAT | 781 | 50 | 29 |
| Macrolide Chromosomal Targets | rplD | L4 | GCCGAAGAAATGCAGGATGCG | 845 | 56 | 25 |
| | rplV | L22 | CCGTGGAAGACCCGAGAACAGAAGAACCC | 925 | 56 | 25 |
| | rpiF | — | TAAGGTAGCGAAATTCCTTGTCG | 756 | 61 | 25 |
| | rep<sup>a</sup> | — | GCCGGCCGACATGGCGCATT | MB | 40 | 23 |
| | uidA | — | CATTACGGCAAATGTGCTAATC | 658 | 55 | 22 |

Table 6. Oligonucleotides used in the study. DNA: Amplified gene or DNA fragment; Prot: Encoded protein; Size: Amplified product size; Ann: Annealing temperature; MB: Multiband (having different and no related sizes). <sup>a</sup>Encode the 23S rRNA; <sup>b</sup>Primer designed to amplify the space between Repetitive Extragenic Palindromic (REP) sequences.
Statistical analysis. The Fisher exact test was used for statistical analysis. P values ≤ 0.05 were considered significant. A microorganism was considered “wt” when no sought mechanism of resistance other than PA/IN inhibitory efflux pumps was identified.

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Author Contributions
C.G., T.J.O. and J.R. designed the experiments; C.G., L.R.-R. and J.M. developed the laboratory studies; C.G. and J.R. analysed the data; C.G. and J.R. wrote the manuscript draft. All authors read the manuscript critically, provide suggestions and approved the final version.

Additional Information
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