Research Article

Pseudomonas aeruginosa Isolates from Spanish Children: Occurrence in Faecal Samples, Antimicrobial Resistance, Virulence, and Molecular Typing

Lidia Ruiz-Roldán,1,1 Alba Bellés,2 Jessica Bueno,2 José Manuel Azcona-Gutiérrez,3 Beatriz Rojo-Bezares,1 Carmen Torres,1,4 F. Javier Castillo,2,5 Yolanda Sáenz,1 and Cristina Seral2,5

1 Área de Microbiología Molecular, Centro de Investigación Biomédica de La Rioja (CIBIR), Logroño, Spain
2 Servicio de Microbiología, IIS Aragón, Hospital Clínico Universitario Lozano Blesa, Zaragoza, Spain
3 Departamento de Diagnóstico Biomédico, Laboratorio de Microbiología, Hospital San Pedro, Logroño, Spain
4 Área de Bioquímica y Biología Molecular, Universidad de La Rioja, Logroño, Spain
5 Departamento de Microbiología, Facultad de Medicina, Universidad de Zaragoza, Zaragoza, Spain

Correspondence should be addressed to Yolanda Saenz; ysaeenz@riojasalud.es

Received 23 January 2018; Accepted 2 May 2018; Published 11 June 2018

Academic Editor: Philippe Lanotte

Copyright © 2018 Lidia Ruiz-Roldán et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Pseudomonas aeruginosa is a major opportunistic human pathogen, responsible for nosocomial infections and infections in patients with impaired immune systems. Little data exist about the faecal colonisation by P. aeruginosa isolates in healthy humans. The occurrence, antimicrobial resistance phenotype, virulence genotype, and genetic lineages of P. aeruginosa from faecal samples of children from two different Spanish regions were characterised. Seventy-two P. aeruginosa were isolated from 1,443 faecal samples. Low antimicrobial resistance levels were detected: ceftazidime (8%), cefepime (7%), aztreonam (7%), gentamicin (3%), ciprofloxacin (1%), and imipenem (1%); susceptibility to meropenem, amikacin, tobramycin, levofloxacin, and colistin. Four multidrug-resistant strains were found. Important differences were detected between both geographical regions. Forty-one sequence types were detected among the 48 tested strains. Virulence and quorum sensing genes were analysed and 13 virulotypes were detected, being 26 exoU-positive strains. Alteration in protein OprD showed eight different patterns. The unique imipenem-resistant strain showed a premature stop codon in OprD. Intestinal colonisation by P. aeruginosa, mainly by international clones (as ST244, ST253, and ST274), is an important factor for the systemic infections development and the environmental dissemination. Periodic active surveillance is useful to identify these community human reservoirs and to control the evolution of antibiotic resistance and virulence activity.

1. Introduction

Pseudomonas aeruginosa is a ubiquitous Gram-negative bacteria present in many diverse environmental settings, including different living sources, such as animals and humans. The ability of P. aeruginosa to survive on minimal nutritional requirements and to tolerate a variety of physical conditions has allowed this organism to persist in both community and hospital settings [1]. This species is a major opportunistic human pathogen, responsible for worldwide nosocomial infections with increasing medical and veterinary importance [1] and causing infections mostly in patients with impaired immune systems [2]. P. aeruginosa shows high intrinsic resistance to antibiotics and has an extraordinary capacity to acquire new resistance mechanisms [3–5]. The increasing prevalence of multidrug-resistant (MDR) P. aeruginosa isolates is a global health problem, because of the limitation in clinical treatment options [5, 6].

This microorganism is seldom a member of the normal microbiota in humans. Colonisation rates in the range of
2.6 to 24% have been reported in human faecal samples [1, 3, 7]. Nevertheless, other studies show a high prevalence of *P. aeruginosa* in small intestinal and faecal samples from irritable bowel syndrome patients [7]. Relatively little data exist so far about the faecal colonisation by *P. aeruginosa* isolates in healthy humans, as well as about their antimicrobial resistance and virulence characteristics [1, 3, 7, 8]. Children tend to be exposed to more disease-causing bacteria through diary activities such as childcare and mouthing behaviours; they are more vulnerable to bacterial illness than adults, and they develop many nonfatal bacterial infections that require antimicrobial treatments, whereas no descriptions exist about the faecal carriage and characterisation of *P. aeruginosa* isolates from children. For that reason, the aim of this study was to determine the occurrence, the antimicrobial resistance phenotypes, the virulence genotypes, and the genetic lineages of *P. aeruginosa* in faecal samples of children from two Spanish regions (La Rioja and Aragón).

2. Material and Methods

2.1. Bacterial Strains. A total of 966 faecal samples of children were recovered in the Hospital Clínico Universitario Lozano Blesa (HLB) (Zaragoza, Aragón, Spain) from June to October 2013 and 477 samples in the Hospital San Pedro (HSP) (Logroño, La Rioja, Spain) from June to August 2014. La Rioja and Aragón are close regions but have different epidemiology settings. All children under 15 years old, who attended the hospital primary cares with gastroenteritis, food allergies, or disorders and with a microbiological study of their stools, were included in the study. The recruitment was the same in both hospitals. Samples were recovered from unique patients without repeated sampling from same patients. Samples were streaked onto cetrimide agar plates. After incubation at 37°C during 24h, one colony per sample with *P. aeruginosa* morphology was selected. Identification of the isolates was performed by classical biochemical methods and MicroScan® WalkAway (Siemens) microdilution system.

2.2. Antimicrobial Susceptibility Testing. In vitro antimicrobial susceptibility testing was performed by MicroScan WalkAway (Siemens) microdilution system following the Clinical and Laboratory Standards Institute (CLSI) guidelines [9]. Thirteen antimicrobial agents were tested, including (range of tested Minimal Inhibitory Concentration in μg/mL, MIC): ticarcillin (8–32), piperacillin-tazobactam (8/4–64/4), aztreonam (1–16), cefepime (1–16), ceftazidime (1–32), meropenem (1–8), imipenem (1–8), tobramycin (2–8), gentamicin (1–8), amikacin (8–32), levofloxacin (1–4), ciprofloxacin (0.5–2), and colistin (2–4). Extended-spectrum-beta-lactamase (ESBL), metallo-beta-lactamase (MBL), and class A carbapenemase phenotypes were determined by double-disc synergy tests [10–12].

2.3. Molecular Typing. The clonal relationship among the recovered isolates was determined by pulsed-field gel electrophoresis (PFGE) with *SpeI* restriction enzyme to digest genomic DNA [3].

Multilocus sequence typing (MLST) for *P. aeruginosa* was performed by PCR and sequencing of seven housekeeping genes (*acsA, aroE, guaA, mutL, nuoD, ppsA*, and *rpeE*) [13] (see Table S1 Supplementary Material). Allelic profiles and sequence types (STs) were compared with the PubMLST database (http://pubmlst.org/paeruginosa/). New STs were submitted to the PubMLST website.

2.4. Virulence Marker Genes. The following virulence and quorum sensing genes were analysed by PCR [14]: *exoU, exoS, exoy, exoT*, *lasA, lasB, exoA, aprA, rhlAB, rhl, rhlR, lasl*, and *lasR* (see Table S1 Supplementary Material).

2.5. Detection and Characterisation of Integrons. The presence of genes encoding type 1, 2, and 3 integrases, and 3’-conserved segment of class 1 integrons (*qacEΔ1+sul1*) was studied by PCR. The characterisation of class 1 integron variable regions was carried out by PCR mapping and sequencing [15] (see Table S1 Supplementary Material).

2.6. Characterisation of OprD Porin. The amino acid changes of the porin protein OprD were analysed by PCR and sequencing in all *P. aeruginosa* strains [16, 17] (see Table S1 Supplementary Material). The mutations were determined by comparison with the sequence of the control strain *P. aeruginosa* PAO1 (GenBank accession number AE004091).

The outer membrane proteins of selected strains were obtained and visualized by SDS-PAGE (Sodium Dodecyl Sulphate-PolyAcrylamide Gel Electrophoresis) in a Bio-Rad apparatus (Mini-Protean II) and were stained with Coomassie Brilliant Blue as previously described [18]. The *P. aeruginosa* PAO1 and PAO1 lacking-OprD were included as control strains.

3. Results

3.1. *P. aeruginosa* Isolates and Antimicrobial Susceptibility. A total of 72 *P. aeruginosa* were isolated from 1,443 faecal samples of children (5%): 42 isolates were obtained in HLB (4.3%) and the 30 remaining ones in HSP (6.3%). *Campylobacter, Salmonella, Giardia*, adenovirus, or rotavirus were the microorganisms involved in the diagnosis of 12 out of the 72 *P. aeruginosa*-positive children. Only 10 of the 72 children showed any underlying disease (number of children): atop dermatitis (2), weight delay (2), lactose intolerance (2), Williams syndrome (1), anorexia (1), cystic fibrosis (1), or psychomotor disturbance/delay (1). No relation was detected between the clinical background of the children and the *P. aeruginosa* isolates (see Table S2 Supplementary Material).

All isolates from HSP were susceptible to all antimicrobial agents tested, with the exception of one isolate that was cefazidime resistant. On the other hand, the 42 *P. aeruginosa* isolates from HLB showed the following resistance percentages to cefepime, 11.9%; piperacillin-tazobactam, 11.9%; ticarcillin, 11.9%; atazanavir, 11.9%; ceftazidime, 11.9%; gentamicin, 4.7%; imipenem, 2.4%; ciprofloxacin, 2.4% and were susceptible to meropenem, amikacin, tobramycin, levofloxacin, and colistin. Only 4 multidrug-resistant isolates were...
found (Pc31, Pc33, Pc40, and Pc43). None of the 72 isolates showed class A carbapenemase, MBL, or ESBL phenotypes.

Only two out of the 72 P. aeruginosa–harbouring children had received antibiotic treatment in the three months previous to sampling: i.e., one cystic fibrosis child (individual 44) who took amoxicillin-clavulanic acid due to a previous to sampling: i.e., one cystic fibrosis child (individual 44) who took amoxicillin-clavulanic acid due to previous infected infection and took meropenem, whereas the P. aeruginosa isolated from both patients were piperacillin-tazobactam-resistant and susceptible, respectively.

3.2. Clonal Relationship and Molecular Typing. Sixty-eight different PFGE patterns were detected among the 72 P. aeruginosa: 39 different PFGE patterns among the 42 isolates from HLB, and 29 among the 30 P. aeruginosa isolates from HSP. No relationship was found among strains from HLB and those from HSP.

The sequence types (ST) were determined among 48 P. aeruginosa strains using MLST method. Forty-one different sequence types were observed, five of them (ST2221, ST2222, ST2223, ST2241, and ST2242) being firstly described in this study and were annotated in the MLST database. Six ST were repeated more than once: ST27 (n=2 strains), ST253 (n=2), ST313 (n=2), ST390 (n=2), and ST667 (n=2) (see Table S2 Supplementary Material).

3.3. Detection of Virulence Markers. The distribution of thirteen genes involved virulence, type III secretion system (T3SS), and quorum sensing system was investigated among the 72 P. aeruginosa strains, and thirteen different profiles were obtained (Table 1).

The T3SS exoU gene was detected in 26 P. aeruginosa strains and the exoS in 47 strains. Both genes were amplified in two strains that were confirmed by three independent experiments and sequencing of the amplicons. On the other hand, neither exoU nor exoS genes were amplified in one strain that also lacked exoY, exoT, exoA, rhlI, and rhlR genes. At least one of the genes involved in quorum sensing (lasI, lasR, rhlR, or rhlR) was not amplified in six strains. However, lasA, lasB, aprA, and rhlAB genes were amplified in all strains (Table 1). The lasR amplicon sized higher than 2,000 bp in three strains (Pc5, Pc10, and Ps519). After sequencing, the insertion sequences IS246, IS246, and IS1394, respectively, were detected surrounding lasR gene. IS246 was detected upstream of lasR gene in Pc5 strain. IS246 was observed truncating lasR gene at position 660 in Pc10 strain, and IS1394 at position 639 in Ps519 strain; therefore those sequences were submitted in GenBank database with accession numbers MH050328 and MG815635, respectively.

3.4. Detection and Characterisation of Integrons. No class 1, 2, or 3 integrases were detected among the 72 P. aeruginosa strains.

3.5. Characterisation of Porin OprD. The oprD gene was amplified in all P. aeruginosa strains, and their OprD amino acid sequences (443 amino acids) were compared with the OprD sequence of reference P. aeruginosa PAO1. Table 2 shows the polymorphism detected in OprD sequences. Eight OprD profiles were distinguished among the 72 P. aeruginosa strains, all but one strain being imipenem-susceptible ones. Only four strains had the same pattern as P. aeruginosa PAO1, and the pattern B was the most frequently detected (31 strains). Among the amino acid changes detected in OprD loop 1 (44 to 61 amino acid (aa)), the most frequent ones were D43N, S57E, and S59R (31 strains, 43%). Alterations in loop 2 (93 to 127 aa) were observed in 51% of the P. aeruginosa strains, T103S and K115T being the most frequent in 21 strains and V127L in 16 strains. The E185Q, P186G, and V189T amino acid changes were the most common described in loop 3 (153 to 192 aa) followed by F170L. On the other hand, E230K
change in the loop 4 (221 to 233 aa) was detected in 47 of the strains. The N262T and A267S amino acid changes in loop L5 (260 to 274 aa) and R310E and A315G in loop 6 (304 to 317 aa) were identified. Thirty-one strains showed the G425A or Q424E substitutions in loop 8 (418 to 431 aa). In the region from 372 to 383 aa of the loop 7, the deletion of two amino acids (loop L7-short), which encoded a protein OprD of 441 amino acids, was identified in 46 strains (64%).

Only the imipenem-resistant strain (Pc1) showed a premature stop codon, due to a point mutation which modifies the expected OprD protein (pattern H in Table 2).

A total of 6 P. aeruginosa strains (Ps514, Ps518, Ps487, Ps499, Ps486, and Ps506) with different protein OprD profiles were selected to study their outer membrane proteins by SDS-PAGE. OprD band was detected in all tested strains.

<table>
<thead>
<tr>
<th>Strains (n* total)*</th>
<th>OprD Size</th>
<th>OprD Pattern</th>
<th>Amino acid Changes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ps514, Pc6, Pc10, Pc15 (4)</td>
<td>443</td>
<td>A</td>
<td>D43N; S57E; S59R; E202Q; I210A; E230K; S240T; N262T; A267S; A281G; K296Q; Q301E; R310G; V359L; Loop L7-short</td>
</tr>
<tr>
<td>Ps488, Ps492, Ps493, Ps494, Ps495, Ps496, Ps497, Ps501, Ps502, Ps507, Ps510, Ps515, Ps516, Ps518, Ps520, Ps521, Pc3, Pc8, Pc11, Pc12, Pc13, Pc17, Pc19, Pc22, Pc24, Pc26, Pc27, Pc28, Pc35, Pc39, Pc43 (31)</td>
<td>441</td>
<td>B</td>
<td>T103S; K115T; F170L; E185Q; P186G; V189T; R310E; A315G; G425A</td>
</tr>
<tr>
<td>Ps487, Ps498, Ps500, Ps517, Ps519, Ps522, Pc4, Pc5, Pc7, Pc14, Pc18, Pc20, Pc29, Pc30, Pc40, Pc42 (16)</td>
<td>443</td>
<td>C</td>
<td>S57E; S59R; V127I; E185Q; P186G; V189T; E202Q; I210A; E230K; S240T; N262T; T276A; A281G; K296Q; Q301E; R310E; A315G; L347M; Loop L7-short; S403A; Q424E</td>
</tr>
<tr>
<td>Ps489, Ps491, Ps499, Ps505, Ps512, Pc2, Pc23, Pc33, Pc37, Pc38, Pc41 (11)</td>
<td>441</td>
<td>D</td>
<td>V127I; E185Q; P186G; V189T; E202Q; I210A; E230K; S240T; N262T; T276A; A281G; K296Q; Q301E; R310E; A315G; L347M; Loop L7-short; S403A; Q424E</td>
</tr>
<tr>
<td>Pc16, Pc31, Pc32, Pc36 (4)</td>
<td>441</td>
<td>E</td>
<td>T103S; K115T; F170L; D43N; T103S; K115T; F170L</td>
</tr>
<tr>
<td>Ps486, Pc9, Pc21, Pc25 (4)</td>
<td>443</td>
<td>F</td>
<td>G60R; T105A; V127I; E185Q; P186G; V189T; E202Q; I210A; E230K; S240T; S248E; N262T; A267S; W277STOP</td>
</tr>
<tr>
<td>Ps506 (1)</td>
<td>443</td>
<td>G</td>
<td>I210A; E230K; S240T; S248E; N262T; A267S; W277STOP</td>
</tr>
<tr>
<td>Pc1 (1)</td>
<td>277</td>
<td>H</td>
<td>I210A; E230K; S240T; S248E; N262T; A267S; W277STOP</td>
</tr>
</tbody>
</table>

4. Discussion

This study intended to gain new insights into the P. aeruginosa in non-clinical environments, analysing the occurrence, and characteristics of P. aeruginosa from faecal samples of children. In fact, the occurrence of P. aeruginosa isolates in this study (5%) (HSP 6.3% and HLB 4.3%) was in the range detected in other studies (2.6-24%) [3, 7]. The presence of P. aeruginosa in stool constitutes a digestive reservoir of the bacteria that may be of importance in P. aeruginosa pathophysiology. In our case, there is not any association between diagnosis or clinical symptoms of the children and the P. aeruginosa detection.

Additionally, low antimicrobial resistance levels were detected for ceftazidime (8%), cefepime (7%), aztreonam (7%), gentamicin (3%), ciprofloxacin (1%), or imipenem (1%) among the isolates tested. Only four multidrug-resistant strains were found. High susceptibility rates were detected among the P. aeruginosa isolates from Logroño (La Rioja). In a previous work performed in the same region by our group, all eight P. aeruginosa isolated from healthy humans were susceptible to all tested antibiotics [3]. All these results contrast with the frequent multidrug-resistant P. aeruginosa detected in studies carried out with clinical strains [5, 6]. Additionally, in our study, only one imipenem-resistant strain was found and this resistance was associated with the loss of porin OprD by a premature stop codon (277 amino acids). Porin OprD serves as the imipenem entryway, and the presence of premature stop codons or certain insertions/deletions in the oprD sequence involves the OprD loss and confers carbapenem resistance in P. aeruginosa [1, 6, 16]. On the other hand, a wide variety of amino acid changes were detected in OprD of the remaining 71 P. aeruginosa strains of our study, even if they were susceptible to imipenem or meropenem. Thirty-one of our strains (43%) contained the same changes in protein OprD (D43N, S57E, S59R, E202Q, I210A, E230K, S240T, N262T, A267S, A281G, K296Q, Q301E, R310G, V359L, and loop L7-short). Some of these amino acid changes had been shown by other authors, in both carbapenem-susceptible and -resistant strains [19–22]. It is known that L2 and L3 loops are involved in binding to imipenem [23–25]. However, most of our strains showed substitutions in both loops and were carbapenem-susceptible, as other studies previously described [22]. The deletion of two amino acids, namely loop L7-short, identified in 48 of these strains has been previously described in both MBL-producing and non-MBL-producing isolates [6]. Summarising, there is a high polymorphism in the oprD gene of our P. aeruginosa strains, but neither the loop L7-short nor amino acid changes, with the exception of the premature stop codon, were associated with carbapenem resistance.
Regarding molecular typing, a wide variety of sequence types has been found in this study (41 different ST among the 48 *P. aeruginosa* strains tested), in contrast to other studies with clinical *P. aeruginosa* isolates, in which mainly appeared the “high-risk clones” ST11, ST175, and ST235. Some works showed that a high genetic diversity is associated with low rates of antimicrobial resistance, as our work has also demonstrated [8, 26]. So, the spread of these different STs among the community is independent from the antimicrobial resistance, but other features could be involved such as host adaptation (virulence, biofilm activity, ...). Among the diversities of STs found in this work, intercontinental clones were encountered: ST27, ST244, ST274, ST277, ST395, ST446, and ST560 [27]. ST244 that was detected in three strains from both regions is a global *P. aeruginosa* clone identified in several countries, among different origins (clinical, animal, and environmental), and associated with different antimicrobial resistance phenotypes (MDR or susceptible strains) and genotypes (carbapenemase-positive or negative) [27–30]. *P. aeruginosa* belonging to ST313 and ST274 have been already described as intestinal colonisers in healthy humans [8], although ST274 has been also previously described in carbapenem-susceptible *P. aeruginosa* isolates [3, 8], associated with cystic fibrosis patients, and is considered as an epidemic clone circulating in Spain [29, 31, 32]. *P. aeruginosa* strains ascended to ST27, ST252, ST253 (clonal complex PA14), ST395, and ST560 have been previously observed in clinical and environmental samples [14, 27, 31, 33, 34].

Considering *P. aeruginosa* pathogenicity, this species possesses at least two well-defined and interrelated quorum sensing (QS) systems, las and rhl. These signalling mechanisms are used to regulate gene expression through the production and secretion of autoinducers called N-acylhomoserine lactones, produced by lasI and rhlI genes, which activate LuxR family regulators, LasR and RhlR, respectively [35]. These systems control the production of different virulence factors, including elastases (LasB and LasA), alkaline protease (AprA), hydrogen cyanide, exotoxin A, pyocyanin, lectins, rhamnolipids (RhlA and RhlB), and superoxide dismutase [36–38]. In our study, the genes implicated in QS system, las and rhl, were amplified in most of the *P. aeruginosa* isolates. However, rhlI and rhlR genes were absent in one strain, rhlI gene were in other three strains, and lasI and lasR genes were not amplified in two more strains. Moreover, the insertion sequence ISI394 was found truncating lasR gene in one *P. aeruginosa* strain. Several reports have mentioned the frequency of lasR mutations among clinical and environmental isolates [39, 40] and have demonstrated that a lasR mutation does not lead to loss of virulence factors thanks to the regulation mediated by rhlR gene [41].

One important determinant of virulence is the T3SS, which is present in several Gram-negative bacilli, including *Salmonella*, *Shigella*, and *Yersinia* spp. In *P. aeruginosa*, the T3SS is a major virulence weapon that contributes to cytotoxicity and acute infections [42]. *P. aeruginosa* is able to produce and secrete virulence factors directly. This secretion system injects potent ExoS and ExoT (two ADP-ribosyltransferase enzymes), ExoU (acute cytolytic factor), and ExoY (an adenylate cyclase) exotoxins into the cytoplasm of the host cells by a cell contact-mediated mechanism [43–48]. ExoU is one of the most important virulence determinants of *P. aeruginosa* and is associated with an increased risk of early clinical mortality [49–51]. In our study, two strains showed the infrequent exoU/exoS-positive genotype, which has been rarely reported [52, 53]. Additionally, other 24 of our strains amplified the exoU gene (33% of total) that belonged to important clinical STs, such as ST253 or ST560, among others. This is a high percentage of exoU presence, considering that these strains were recovered from healthy children. Defined clonal lineages are expected to be linked to specific T3SS genotypes. Indeed, in a Spanish multicenter study, the vast majority of the extensively drug resistant (XDR) isolates belonged to the high-risk clones ST175 and ST111, which showed in all cases an exoU-negative/exoS-positive genotype, and to the ST235 clone that showed an exoU-positive/exoS-negative genotype [54].

5. Conclusion

This work assesses for the first time occurrence of *P. aeruginosa* in faecal samples of children from two different geographical Spanish regions, providing fundamental information about antimicrobial resistance, virulence, and the genetic lineages of the recovered isolates. These faecal *P. aeruginosa* isolates were characterised by their high clonal diversity, great variety of genetic lineages, low antimicrobial resistance percentages, and variability of virulence patterns. Intestinal colonisation by *P. aeruginosa*, mainly belonging to international clones (such as ST244, ST253, or ST274), is an important factor for the development of systemic infections and the environmental dissemination. Periodic active surveillance is useful to identify these community human reservoirs and control the evolution of antibiotic resistance carriage and virulence activity over time.

Data Availability

All data are included in the manuscript and Supplementary Material. The new sequence detected of ISI394 truncating lasR gene in Ps519 strain was submitted in GenBank database with accession number MG815635.

Conflicts of Interest

The authors declare that there are no conflicts of interest regarding the publication of this article.

Authors’ Contributions

Lidia Ruiz-Roldán and Alba Bellés contributed equally to this work.

Acknowledgments

This publication made use of the *Pseudomonas aeruginosa* MLST website (https://pubmlst.org/paeruginosa/) developed by Keith Jolley and sited at the University of Oxford (Jolley & Maiden 2010, BMC Bioinformatics, 11:595). The
development of this site has been funded by the Wellcome Trust. Part of this study was presented at the 26th European Congress of Clinical Microbiology and Infectious Diseases (ECCMID) (Abstract N° 5218, Amsterdam, Netherlands, 9–12th April 2016), XX Congreso Nacional Sociedad Española de Enfermedades Infecciosas y Microbiología Clínica (SEIMC) (Abstract N° 370, Barcelona, Spain, 26–28th May 2016), and the 16th International Congress on Pseudomonas (Abstract N° PI14, Liverpool, UK, 5–9th September 2017). Lidia Ruiz-Roldán has a predoctoral fellowship from the Consejería de Industria, Innovación y Empleo, Gobierno de La Rioja, Spain. This work was partially supported by the Instituto de Salud Carlos III of Spain (ISCIII) [Projects FIS PI12/01276 and PI16/01381] (cofunded by European Regional Development Fund (FEDER) “Away to make Europe”).

Supplementary Materials

Table S1: primers and annealing temperatures used in the PCR reactions of this study. Table S2: characteristics of P. aeruginosa isolates recovered from faecal samples of children. (Supplementary Materials)

References


[52] M. Maatallah, J. Cheriaux, A. Backhouf et al., “Population structure of *Pseudomonas aeruginosa* from five Mediterranean


Submit your manuscripts at www.hindawi.com