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Molecular screening of pathogenic *Escherichia coli* strains isolated from dairy neonatal calves in Cordoba province, Argentina


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**KEYWORDS**
*Escherichia coli*; Calves; Diarrhea; PCR; Virulence genes; Argentina

**Abstract** The aim of this study was to perform a current molecular characterization of bovine pathogenic *Escherichia coli* strains isolated from random samplings in Argentinean dairy farms. Rectal swabs were obtained from 395 (63.7%) healthy and 225 (36.3%) diarrheic calves, belonging to 45 dairy farms in Cordoba Province, Argentina. *E. coli* isolates were examined for virulence genes (f5, f41, f17, sta, stb, lt, eae, vt) using PCR and the prevalence of *E. coli* virulence profiles was spatially described in terms of spatial distribution. A total of 30.1% isolates were found to be positive for at least one of the virulence genes. Depending on the different gene combinations present, 11 virulence profiles were found. Most of the isolates analyzed had a single gene, and no combination of fimbrial and enterotoxin gene was predominant. There was no association between the frequency and distribution of *E. coli* virulence genes and calf health status. Most of the virulence profiles were compatible with ETEC strains and showed a homogeneous distribution over the sampled area. A clustering pattern for *E. coli* virulence profiles could not be recognized. This work provides updated information on the molecular characterization of pathogenic *E. coli* strains from dairy herds in Cordoba, Argentina. These findings would be important to formulate prevention programs and effective therapies for diarrhea in calves caused by *E. coli*.

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**Introduction**

Neonatal calf diarrhea is an important cause of morbidity and mortality worldwide in newborn calves. This multifactorial disease involves the calf immune status, environmental factors and farm management practices (housing, feeding and hygienic conditions) as well as the interaction of different pathogens such as bacteria, viruses and protozoa.

*Escherichia coli* is the predominant aerobic organism in the normal intestinal microbiota of mammals; here it plays an important role in host metabolism, immunity and nutrition. However, a reduced number of highly adapted pathogenic strains are capable of causing intestinal or extraintestinal diseases and great economic losses.

Pathogenic *E. coli* strains have different virulence factors that allow them to colonize the host’s small intestine, avoiding the immune response and stimulating the deleterious inflammatory response to produce diarrhea. These virulence factors include the antigens of colonization or adhesion (F2–F6, F17, F18, F41 fimbriae and intimin) and exotoxins (heat-labile enterotoxin [LT], heat-stable enterotoxins [STa and STb] and verotoxins [VT]) only the most successful combinations of virulence factors have persisted to become specific *E. coli* pathotypes that are capable of causing disease in healthy individuals.

Farm animal and human diarrhea are frequently due to infection by one or several of *E. coli* pathotypes: enterotoxigenic *E. coli* (ETEC), enteropathogenic *E. coli* (EPEC), enteroinvasive *E. coli* (EIEC). Vero toxin-producing/Shiga toxin-producing *E. coli* (VT/STEC) which include its well-known subgroup enterohemorrhagic *E. coli* (EHEC), enteraggregative *E. coli* (EAggEC) and enteroadherent *E. coli* (EAdEC). ETEC infection is the most common type of colibacillosis in young animals especially in calves and piglets.

In Argentina, neonatal calf diarrhea is a severe and common disease, affecting both beef and dairy herds with a morbidity above 60%. Previous reports have shown the critical role of pathogenic *E. coli* in calf diarrhea. However, these studies are not recent and have generally been restricted to few animals or herds.

The aim of this study was to perform a current molecular characterization of pathogenic *E. coli* strains isolated from a random sampling of neonatal calves.

**Materials and methods**

**Calves and data collection**

Forty-five farms from Villa María (Córdoba) dairy area, with a herd size of 100–250 cows were randomly selected from a roster provided by the producers’ association. Such herd size strata represent 80% of the dairy operations in Villa Maria and 10% of Argentina’s dairies. A cross-sectional study involving Holstein calves less than 10 days of age, both healthy and diarrheic, was carried out between February and October 2008. The identification, age and gender of each animal were recorded. A clinical examination of each calf was performed, and the clinical parameters related to diarrhea were registered.

**Bacterial isolates**

Individual rectal swabs (*n* = 620) were collected and transported in Stuart’s transport medium (Britania, Argentina) to the Bacteriology laboratory. The swabs were plated
in MacConkey agar plates (Oxoid, UK) and Gram staining was performed on isolated lactose-positive colonies. Gram-negative bacteria were biochemically characterized as belonging to the Enterobacteriaceae family, Escherichia genus, according to the Bergey’s Manual\textsuperscript{19}. Colonies compatible with E. coli were transferred to tubes containing minimal agar medium\textsuperscript{24}.

Molecular typing by PCR

E. coli-positive strains were randomly selected and cultured on MacConkey agar plates.

Following overnight incubation, a loop was taken from the bacterial confluent growth zone, suspended in 300 μl of distilled water, boiled for 5 min, and centrifuged at 11,000 rpm for 2 min. The supernatant was used for PCR. From each PCR-positive sample, 10 colonies were streaked on MacConkey agar plates, pooled and processed by PCR. If a positive result was obtained, individual colonies were tested by PCR to identify positive isolates.

PCR technique was performed as described by Siqueira et al.\textsuperscript{39}, with minor modifications. Briefly, the amplification reaction was carried out with 7 μl of template, 6 μl of reaction buffer (5× Green Go Taq\textsuperscript{®}, Promega) containing MgCl\textsubscript{2} (1.5 mM), 0.6 μl of dNTP (10 mM, Promega), 0.5 μl of primer (300 ng/μl, Ruralex Fago) and 0.2 μl of Taq DNA polymerase (5 U/μl, Go Taq\textsuperscript{®} DNA Polimerasa, Promega). Volumes were adjusted to 30 μl with sterile distilled water. Templates were screened for the presence of E. coli virulence factors coding gene by conventional PCR, using the primers described in Table 1. PCR cycling conditions consisted of initial denaturation at 94 °C for 2 min, followed by 30 cycles of amplification (denaturation at 93 °C for 1 min, annealing for 1 min, extension at 72 °C for 1 min) and final elongation at 72 °C for 7 min. All PCR reactions were performed in an MJ Research Thermocycler Bio-Rad model PTC-220. The amplified PCR products were separated by electrophoresis on a 2% agarose gel with Tris–acetate–EDTA (TAE) buffer (Sigma–Aldrich\textregistered). A 100-bp DNA ladder (CienMarker, Biodynamics) was used as molecular weight marker. Control E. coli strains for each gene (Table 2) and distilled water were used as positive and negative controls respectively. The gel was stained with ethidium bromide (Sigma–Aldrich\textregistered) and visualized under a UV-transilluminator at 302 nm.

The E. coli strains were classified into different pathotypes, based on their specific virulence genes\textsuperscript{26}.

### Data analysis and mapping

All data analyses were performed using the commercially available statistical software, SPSS for Windows (Version 10.0, Chicago, IL, USA). Differences were considered significant at the level of p < 0.05. Herd geographic coordinates were recorded using a Global Positioning System (GPS). The prevalence of E. coli pathotypes was spatially described using a dot map displayed in Arcview 3.2 (ESRI, Redlands, CA, USA).

#### Table 1: Characteristics of the oligonucleotide primers used for the PCR reaction.

<table>
<thead>
<tr>
<th>Target gene</th>
<th>Primer</th>
<th>Primer sequence (5′–3′)</th>
<th>Amplicon size (bp)</th>
<th>Annealing temperature</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>vt</td>
<td>VT VT-F</td>
<td>GAGCGAAATATTATATATAT</td>
<td>322</td>
<td>48 °C</td>
<td>36</td>
</tr>
<tr>
<td></td>
<td>VT VT-R</td>
<td>CCGAATCCTAATATATATAT</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>LT LT-B</td>
<td>CCGAATCTGTTATATATATGTC</td>
<td>696</td>
<td>52 °C</td>
<td>6</td>
</tr>
<tr>
<td></td>
<td>LT LT-FN</td>
<td>GGGCAGCATATAACGTGC</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>stb</td>
<td>Stb Stb-F</td>
<td>ATCCGATTTCCTTGATCAT</td>
<td>175</td>
<td>55 °C</td>
<td>6</td>
</tr>
<tr>
<td></td>
<td>Stb Stb-R</td>
<td>GGGCCCAAGACTGTCCTC</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>sta</td>
<td>Sta Sta-A</td>
<td>ATTTTTATTCTGTATGTTCTTT</td>
<td>176</td>
<td>48 °C</td>
<td>14</td>
</tr>
<tr>
<td></td>
<td>Sta Sta-B</td>
<td>GATTTACACAGCATACACAGCTG</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>eae</td>
<td>EaeA-F</td>
<td>AGGCCTTGCCACTAGTTG</td>
<td>570</td>
<td>55 °C</td>
<td>8</td>
</tr>
<tr>
<td></td>
<td>EaeA-R</td>
<td>CCATCTCACCAGAGGA</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>f17</td>
<td>F17 F17-F</td>
<td>GGGGTGACAGAGGAGGTGGGGG</td>
<td>411</td>
<td>60 °C</td>
<td>29</td>
</tr>
<tr>
<td></td>
<td>F17 F17-R</td>
<td>CCGGGCGAACATTCATCACGG</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>f5</td>
<td>K99 K99-A</td>
<td>CACGCGCGCGCGAGTAATGACTG</td>
<td>278</td>
<td>60 °C</td>
<td>14</td>
</tr>
<tr>
<td></td>
<td>K99 K99-B</td>
<td>CCACTATTGAGGGAGCCGCGGG</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>f41</td>
<td>F41 F41-A</td>
<td>GGCATGAAAGACGAGAGGAGG</td>
<td>551</td>
<td>55 °C</td>
<td>14</td>
</tr>
<tr>
<td></td>
<td>F41 F41-R</td>
<td>GGGGTGACTGAGGTCTACCC</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

#### Table 2: Characteristics of E. coli strains used as positive control for virulence genes in PCR reaction.

<table>
<thead>
<tr>
<th>Identification</th>
<th>Origin</th>
<th>Virulence profiles</th>
</tr>
</thead>
<tbody>
<tr>
<td>FV-10185</td>
<td>OVINE</td>
<td>vt1, vt2</td>
</tr>
<tr>
<td>FV-10186</td>
<td>PORCINE</td>
<td>eae</td>
</tr>
<tr>
<td>FV-10187</td>
<td>PORCINE</td>
<td>vt2, sta, stb, f18</td>
</tr>
<tr>
<td>FV-10188</td>
<td>PORCINE</td>
<td>lt, stb, f4</td>
</tr>
<tr>
<td>FV-10189</td>
<td>PORCINE</td>
<td>lt, stb, f18</td>
</tr>
<tr>
<td>FV-10190</td>
<td>PORCINE</td>
<td>sta, f6</td>
</tr>
<tr>
<td>FV-10191</td>
<td>BOVINE</td>
<td>sta, f41, f5</td>
</tr>
<tr>
<td>FV-10192</td>
<td>OVINE</td>
<td>f17</td>
</tr>
</tbody>
</table>

*Note: These control E. coli strains were kindly provided by the Animal Health Group (EEA INTA Balcarce) and come from the strain collection of Dr. Jorge Blanco (E. coli Reference Laboratory, University of Santiago de Compostela, Lugo, Spain).*
A spatial scan statistic was used to identify and locate significant spatial clusters of *E. coli* virulence profiles within the study area. The null hypothesis to test was that herds within a particular window have the same prevalence level as herds outside the window. All analyses were performed by using SaTScan® software (Version 9.0, http://www.satscan.org).

**Results**

**Diarrhea prevalence**

Overall, 620 calves from 45 dairy farms were sampled. The average number of calves sampled per herd was 14 ± 7.95 (media ± standard deviation). At the time of sampling, 36.3% of calves (n = 225) presented diarrhea. Diarrhea prevalence found in dairy farms markedly varied from 0% to 71.4% with a median of 35.7% (percentile 25: 20.0% and percentile 75: 50.0%). There were no cases of diarrhea in 2 farms, while prevalence of diarrhea was higher than 50% in 8 farms (Fig. 1).

**Molecular characterization of *E. coli* isolates**

A total of 156 samples from 363 *E. coli*-positive isolates were analyzed by PCR to detect 8 virulence genes. A total of 109 samples (69.9%) were negative for all genes investigated (probably non-pathogenic strains) and 47 (30.1%) were found to be positive for at least one of the virulence genes. A total of 39 isolates were obtained from the PCR-positive samples. Depending on the different gene combinations present, 11 different virulence profiles were found (Table 3). Most of the isolates analyzed had a single gene. The prevalent virulence gene was *f17*, while the only enterotoxin gene observed was *sta*; *lt* and *stb* genes were not detected, while *vt* and *f41* genes were only found in combination with other genes. No combination for both fimbriae and toxins was prevalent.

Regarding the frequency and distribution of *E. coli* virulence genes, no significant differences (p = 0.3008, Chi Square) were observed between healthy and diarrheic calves (Table 3).

**Identification and spatial distribution of *E. coli* genetic profiles**

The map with the locations of all the farms visited and the spatial distribution of *E. coli* virulence profiles is farm is shown in Fig. 2. Most of the virulence profiles were compatible with ETEC strains and showed a homogeneous distribution over the sampled area. A clustering pattern for *E. coli* virulence profiles could not be recognized.

**Discussion**

Diarrhea is a frequent and growing concern in young calves, especially in the first week of age. Regarding the main enteropathogens causing diarrhea, ETEC, Rotavirus, Cryptosporidium parvum have been reported in 75–95% of cases of intestinal infections in young calves. In general, pathogenic *E. coli* appears to be less important compared with other diarrheagenic pathogens, since it is shed during a short period of time, resulting in a low prevalence. However, this bacterium plays an important role in the occurrence of enterotoxigenic and septicemic colibacillosis, and its incidence and impact on morbidity/mortality of newborn calves might be relatively high.

In Argentina, the role of pathogenic *E. coli* isolated from healthy and diarrheic calves has been previously investigated. However, these investigations have been limited to the study of few animals or herds and were conducted over thirty years ago. Odeón and Barrandeguy *et al.*, demonstrated that ETEC strains were commonly found in beef and dairy herds. Conversely, Bellinzoni *et al.*, reported that pathogenic *E. coli* was not detected in any calf. In this study, pathogenic *E. coli* prevalence was 30.1%. The differences in these findings could be due to regional variations, management and hygienic conditions, the age of the animals, or to the fact that the analysis of F5 fimbriae in the previous reports was determined only by serology,
Pathogenic *E. coli* in neonatal calves from Argentina

Table 3  Virulence profiles detected in the *E. coli* isolates (n = 39).

<table>
<thead>
<tr>
<th>Number of genes</th>
<th>Virulence profiles</th>
<th>Number of isolates</th>
<th>Frequencies (%)</th>
<th>Dairy farms (Calf health status)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>f17</td>
<td>16</td>
<td>41</td>
<td>11(H), 12(D), 18(D), 29(H), 32(H), 41(H), 42(D), 43(H), 75(D), 72(H), 76(H), 80(D), 85(H), 86(D), 93(H), 632(H)</td>
</tr>
<tr>
<td>1</td>
<td>eae</td>
<td>5</td>
<td>12.7</td>
<td>2(D), 25(D), 57(D), 63(H), 632(H)</td>
</tr>
<tr>
<td>1</td>
<td>sta</td>
<td>3</td>
<td>7.7</td>
<td>10(D), 90(H), 632(D)</td>
</tr>
<tr>
<td>1</td>
<td>f5</td>
<td>1</td>
<td>2.6</td>
<td>11(D)</td>
</tr>
<tr>
<td>2</td>
<td>f17 vt</td>
<td>3</td>
<td>7.7</td>
<td>11(H), 93(H), 43(H)</td>
</tr>
<tr>
<td>2</td>
<td>eae vt</td>
<td>3</td>
<td>7.7</td>
<td>2(D), 57(D), 86(D)</td>
</tr>
<tr>
<td>2</td>
<td>f17 sta</td>
<td>3</td>
<td>7.7</td>
<td>25(D), 90(H), 92(H)</td>
</tr>
<tr>
<td>2</td>
<td>f17 f5</td>
<td>2</td>
<td>5.1</td>
<td>1(H), 15(H)</td>
</tr>
<tr>
<td>2</td>
<td>sta eae</td>
<td>1</td>
<td>2.6</td>
<td>632(D)</td>
</tr>
<tr>
<td>3</td>
<td>sta f5 f41</td>
<td>1</td>
<td>2.6</td>
<td>42(D)</td>
</tr>
<tr>
<td>3</td>
<td>sta f5 f17</td>
<td>1</td>
<td>2.6</td>
<td>25(D)</td>
</tr>
</tbody>
</table>

* D, diarrheic calf; H, healthy calf.

without searching for virulence genes with molecular methods. The present work involved a great number of dairy farms and calves, and was focused on the molecular characterization of pathogenic *E. coli* strains among dairy farms from Cordoba province, Argentina. Previous studies in the same dairy farms of Cordoba province investigated the role of *Cryptosporidium* spp. and *Giardia* spp. producing calf diarrhea and the participation of *E. coli* causing subclinical mastitis. However, investigations about the molecular characterization of pathogenic *E. coli* causing diarrhea in neonatal calves have not been conducted. The results of the present study provide new and complementary information about one of the pathogens causing diarrhea in neonatal calves in this dairy area.

In order to categorize *E. coli* pathotypes, the presence of virulence characteristics needs to be identified. The use of PCR-based technology to identify virulence genes has become widely adopted to distinguish pathogenic *E. coli* strains from normal gut flora. In this study, ETEC-compatible strains were the most prevalent strains determined by the PCR technique. Nagy and Fekete reported that diarrhea caused by ETEC is considered to be the major infectious disease of newborn calves during the pre-weaning and weaning periods. Infected animals are important reservoirs of ETEC and their feces are the major source of environmental contamination with the bacteria, which are acquired by newborn calves soon after birth. Although bovine ETEC strains do not represent a real hazard to man and cannot be regarded as zoonotic agents, the high circulation of such strains affects the productive capacity and efficiency of calves in artificial breeding.

ETEC bacteria are known to adhere to the small intestinal epithelium without inducing significant morphological changes and to secrete enterotoxins that alter the functions of enterocytes by increasing secretion and reducing absorption. The main colonization factors or fimbriae described in bovine pathogenic *E. coli* are F5 (K99), F17 and F41. F5 fimbriae has been associated with most ETEC strains and is responsible for most cases of ETEC infection in newborn calves. In this study, the prevalence of ETEC F5+ was 12.9%. Similar results were obtained by Younis et al., in Egypt (10.4%), Ok et al., in Turkey (13.4%) and Pourtaghi et al., in Iran (14.1%). A higher prevalence of ETEC F5+ strains was reported in Egypt and Israel (23%), Mozambique (25%) and Brazil (35%). On the contrary, a lower prevalence of *E. coli* F5+ was found in the United States, the Netherlands, Brazil and Turkey, with rates of 1.8%, 2.6%, 5.8% and 9.4% respectively.

F17 fimbriae have been found on some bovine ETEC strains and mediate attachment of these bacteria to the intestinal epithelium. We found that the most prevalent virulence factor in both diarrheic and healthy calves was individual F17 fimbria (41%). The observation that a high number of *E. coli* strains carried genes for fimbriae but no toxin genes has been reported in others studies. Güler et al., found similar results (44%) for F17 fimbriae in *E. coli* isolates from calves. Nguyen et al., showed a low prevalence (16%) of F17 fimbriae among *E. coli* strains isolated from diarrheic calves. It is important to take into account that *E. coli* F17+ strains have also been associated with extraintestinal disease and usually show other virulence factors associated with the ability to cause septicemia. In Iran, Ghanbarpour and Oswald found strains of *E. coli* F17+ isolated from septicemic calves (29%) and from bovine mastitis (20.4%). Thus, in our study, an additional PCR screening of extraintestinal virulence factors would be necessary to confirm the likely presence of non-enterotoxigenic F17+ strains.

After adherence to the intestinal mucosa, ETEC strains produce one or both enterotoxins, LT and ST, which are responsible for hypersecretion of electrolytes and water into the small intestine. In this study, we found that sta was the only enterotoxin gene detected, while *lt* and *stb* genes were not detected in any of the examined...
strains. These results agree with Nagy and Fekete, who reported that the STa enterotoxin is produced in porcine and bovine ETEC strains, while the LT and STb enterotoxins are predominantly produced by human and porcine ETEC strains. Conversely, there are some reports describing the isolation of LT and STb enterotoxins from cows and calves. In Brazil, Salvadori et al., recorded 3.9% ETEC possessing st and lt genes from diarrheic calves and Rigobelo et al., reported a prevalence rate of E. coli isolated from diarrheic calves carrying st (25.4%) and lt (13.2%) genes. In China, Huasai et al., reported that 15.5% of ETEC strains isolates from healthy cows were positive for lt and st genes. In Italy, it has been found that all the ETEC strains isolated from diarrheic water buffalo calves had the lt gene but lacked st gene. Finally, Acha et al., reported that the sta gene was not detected in any of the E. coli F5 strains isolated from diarrheic calves in Mozambique. Since ETEC enterotoxins are plasmid-encoded secreted proteins, the variation in the prevalence of these toxins could be explained by the existence of conjugative plasmids widely distributed among bovine ETEC strains.

Regarding the distribution of E. coli virulence profiles, most of them were compatible with ETEC strains and were widely distributed throughout the examined geographical area. This pattern of spatial distribution was similar to the one reported for Giardia spp. in the same dairy farms, but disagrees with the primary cluster shown by Cryptosporidium spp. The ETEC virulence gene distribution on the farms showed different combinations but no profile was prevalent. There was no association between the virulence profiles and dairy farm location.

In conclusion, this study showed a high frequency of pathogenic E. coli strains that were widely distributed either among diarrheic or healthy calves in the examined farms. Mainly, F17 and F5 fimbriae were found to be the most common virulence factors of E. coli strains isolated from dairy calves. This work provides updated information on the molecular characterization of pathogenic E. coli strains obtained from a random sampling of dairy herds in Cordoba, Argentina. These findings would be important to formulate prevention programs and effective therapies for calf diarrhea caused by pathogenic E. coli.

**Ethical disclosures**

**Protection of human and animal subjects.** The authors declare that no experiments were performed on humans or animals for this study.
Confidentiality of data. The authors declare that no patient data appear in this article.

Right to privacy and informed consent. The authors declare that no patient data appear in this article.

Conflict of interest

The authors declare that they have no conflicts of interest.

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