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Dissection of the Role of Pili and Type 2 and 3 Secretion Systems in Adherence and Biofilm Formation of an Atypical Enteropathogenic *Escherichia coli* Strain

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Atypical enteropathogenic *Escherichia coli* (aEPEC) strains are diarrheal pathogens that lack bundle-forming pilus production but possess the virulence-associated locus of enterocyte effacement. aEPEC strain 1551-2 produces localized adherence (LA) on HeLa cells; however, its isogenic intimin (*eae*) mutant produces a diffuse-adherence (DA) pattern. In this study, we aimed to identify the DA-associated adhesin of the 1551-2 *eae* mutant. Electron microscopy of 1551-2 identified rigid rod-like pili composed of an 18-kDa protein, which was identified as the major pilin subunit of type 1 pilus (T1P) by mass spectrometry analysis. Deletion of *fimA* in 1551-2 affected biofilm formation but had no effect on adherence properties. Analysis of secreted proteins in supernatants of this strain identified a 150-kDa protein corresponding to SslE, a type 2 secreted protein that was recently reported to be involved in biofilm formation of rabbit and human EPEC strains. However, neither adherence nor biofilm formation was affected in a 1551-2 *sslE* mutant. We then investigated the role of the EspA filament associated with the type 3 secretion system (T3SS) in DA by generating a double *eae espA* mutant. This strain was no longer adherent, strongly suggesting that the T3SS translocon is the DA adhesin. In agreement with these results, specific anti-EspA antibodies blocked adherence of the 1551-2 *eae* mutant. Our data support a role for intimin in LA, for the T3SS translocon in DA, and for T1P in biofilm formation, all of which may act in concert to facilitate host intestinal colonization by aEPEC strains.

Enteropathogenic *Escherichia coli* (EPEC) is a common cause of infant diarrhea in both developing and developed countries (1, 2). The hallmark of EPEC pathogenesis is the formation of attaching and effacing (AE) lesions characterized by intimate bacterial adherence, microvillus effacement, and accumulation of polymerized actin and many other cytoskeleton elements, resulting in pedestal-like structures underneath adherent bacteria. The virulence factors necessary for the establishment of the AE lesion are encoded in a chromosomal pathogenicity island called the locus of enterocyte effacement (LEE) (3).

The EPEC pathotype is divided into typical (tEPEC) and atypical EPEC (aEPEC). This classification is based on the occurrence of the virulence-associated EAF (EPEC adherence factor) plasmid (pEAF) in tEPEC and its absence in aEPEC (4). Although tEPEC strains were major causative agents of acute diarrhea in very young children in Brazil until the 1990s (5), there is currently a clear decrease in their frequency (6, 7). Retrospective studies indicate that aEPEC strains have existed as important agents of diarrhea in developed countries since the 1960s and are currently emerging pathogens that produce acute and persistent diarrhea affecting children and adults worldwide (2, 8, 9).

The first step in the establishment of diarrheal disease caused by the various pathotypes of diarrheagenic *E. coli* (DEC) is the colonization of the gastrointestinal tract, which is mediated by specific fimbrial and nonfimbrial adherence factors (10). A feature of tEPEC is the production of a characteristic pattern of adherence, termed localized adherence (LA), in which bacteria bind to localized areas of the cultured epithelial cell surface, forming compact microcolonies that are visualized after 3 h of infection (11). Generally, most aEPEC strains are weakly adherent and display a LA-like (LAL) phenotype manifested as loose cell-associated bac-

terial microcolonies; less common are aEPEC strains that display the typical LA pattern. These patterns can be observed only in adherence assays performed for extended periods of infection (e.g., 6 h). Some strains however, do exhibit diffuse (DA) or aggregative (AA) adherence (12–15).

Upon infection of host cells EPEC translocates a range of effector proteins to the eukaryotic cell by means of the type 3 secretion system (T3SS) (16, 17). The EspA protein forms a long filamentous extension of the T3SS needle complex, and EspB and EspD are coupled to the end of the EspA filament to generate a pore in the host cell membrane through which effectors are injected into the host cell cytosol (18, 19).

The LEE-encoded intimin adhesin is required for intimate adhesion to epithelial cells and cytoskeletal reorganization (20). The translocated intimin receptor (Tir) (21) is injected via T3SS and inserted in the plasma membrane of eukaryotic cells and is crucial for development of AE lesions. So far, intimin has been recognized as the main adhesin in aEPEC strains (22, 23).

Our previous studies on the adherence mechanisms of aEPEC strains isolated from clinical cases in Brazil identified a strain

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TABLE 1 *Escherichia coli* strains and plasmids used in this study

<i>E. coli</i> strain or plasmid	Relevant characteristics ^a	Reference or source
Strains		
1551-2	aEPEC (ONT:H ⁻) isolated from a child with diarrhea in Brazil (Nal ^r)	13
1551-2 <i>eae</i> ::Km	1551-2 harboring the suicide vector pJP5603 inserted into the <i>eae</i> gene (Nal ^r Km ^r)	24
1551-2 Δ <i>fimA</i>	<i>fimA</i> ::zeo (Nal ^r Zeo ^r)	This study
1551-2 <i>ssIE</i> ::Km	1551-2 harboring the suicide vector pJP5603 inserted into <i>ssIE</i> gene (Nal ^r Km ^r)	This study
1551-2 <i>escN</i> ::Km	1551-2 harboring the suicide vector pJP5603 inserted into <i>escN</i> gene (Nal ^r Km ^r)	This study
1551-2 <i>eae</i> ::Km Δ <i>tir</i>	Double intimin/Tir mutant in aEPEC 1551-2 (Nal ^r Km ^r Zeo ^r)	This study
1551-2 <i>eae</i> ::Km Δ <i>fimA</i>	Double intimin/T1P mutant in aEPEC 1551-2 (Nal ^r Km ^r Zeo ^r)	This study
1551-2 <i>eae</i> ::Km Δ <i>espA</i>	Double intimin/EspA mutant in aEPEC 1551-2 (Nal ^r Km ^r Zeo ^r)	This study
Plasmids		
pEscN	pACYC184 vector harboring the <i>escN</i> gene from EPEC prototype E2348/69	31
pJP5603	3.1-kb R6K-based suicide vector, Km ^r	29
pKOBEG-Apra	Red recombinase system plasmid, Apra ^r	44
1711-4 Δ <i>flhC</i>	Source of zeocin cassette	44
pGEM-T Easy	Cloning vector	Promega
pBAD/Myc-His A	Cloning vector	Invitrogen
pFimA	pBAD/Myc-His A vector harboring the <i>fimA</i> gene from aEPEC 1551-2	This study
pInt	pBAD/Myc-His A vector harboring the <i>eae</i> gene from aEPEC 1551-2	This study
pEspA	pBAD/Myc-His A vector harboring the <i>espA</i> gene from aEPEC 1551-2	This study

^a Nal^r, nalidixic acid resistant; Km^r, kanamycin resistant; Zeo^r, zeocin resistant; Apra^r, apramycin resistant.

(1551-2, ONT:H⁻) that forms LA on cultured epithelial cells. However, an isogenic intimin mutant strain adheres to HeLa cells exhibiting a diffuse-adherence (DA) pattern (24). Our main interest in the present study was to elucidate the nature of the aEPEC 1551-2 adhesin responsible for the DA phenotype.

MATERIALS AND METHODS

Bacterial strains, plasmids, and growth conditions. All strains and plasmids used in this study are listed in Table 1. aEPEC 1551-2 was isolated from a child with diarrhea in the absence of other recognized pathogens in an epidemiological study conducted in São Paulo, Brazil (13). Strains were grown aerobically in Luria-Bertani (LB) medium or Dulbecco's modified Eagle's medium (DMEM) (Invitrogen) at 37°C. Recombinant DNA and molecular biology techniques were performed as previously described (25).

Epithelial cell adherence. The assay described by Cravioto et al. (26) was used, with some adaptations, to determine the adherence pattern of the strains. HeLa cells at 70% confluence were cultivated in 24-well tissue culture plates containing glass coverslips and DMEM supplemented with 10% bovine fetal serum (Gibco Invitrogen, USA) and 1% antibiotics (Gibco Invitrogen, USA). After one wash with phosphate-buffered saline (PBS), pH 7.4, 1.0 ml of fresh medium (DMEM supplemented with 2% fetal bovine serum) was added to the cell monolayers. *E. coli* strains were grown overnight in LB broth without shaking and then diluted 1:50 in the medium contained in the microplates. After an incubation period of 3 h at 37°C, the monolayers were washed once with PBS and fresh medium was added to the wells, followed by an additional incubation period of 3 h for a total of 6 h. After six washes with PBS, the preparations were fixed with methanol, stained with May Grünwald-Giemsa stain, and examined by light microscopy.

For quantitative experiments, after the adherence assay (performed as described above) the washed monolayers were lysed in 1% Triton X-100 for 30 min at room temperature. Following cell lysis, adherent bacteria were quantified by plating serial dilutions onto MacConkey agar plates to obtain the total number of cell-associated bacteria. Assays were carried out in triplicate, and the results represent the means $\pm$ standard errors from at least three independent experiments.

Adherence inhibition experiments with EspA antibody. For adherence inhibition experiments, samples (approximately 10⁷ CFU) of an

overnight culture of the 1551-2 *eae*::Km mutant strain grown in LB broth were incubated with 1:100, 1:50, and 1:10 dilutions of anti-EspA antibody in DMEM at room temperature for 30 min and then added to the HeLa cell monolayers. After incubation for 6 h, the cells were lysed with PBS–0.1% Triton X-100, diluted, and plated onto MacConkey agar to estimate the number of bacteria adhering to the cell monolayers (27). All inhibition experiments were performed at least three times in triplicate on different days to ensure reproducibility.

Gene amplification. For PCR, the GoTaq Green master mix (Promega, Madison, WI) was used with 1 μ M each of the primers (Table 2).

Mutant construction and complementation. Derivative mutants were constructed by the one-step allelic exchange recombination method, using the thermosensitive plasmid pKOBEG-Apra, which carries lambda red recombination genes (28). Primers containing a 40-bp region homologous to the 5' and 3' extremities of the target genes (*fimA*, *tir* and *espA*) and specific sequence for the zeocin (Zeo) resistance-encoding gene were used to amplify the Zeo cassette. Amplicons were purified from the agarose gel and quantified (BioPhotometer; Eppendorf) before they were electroporated into competent aEPEC 1551-2 or 1551-2 *eae*::Km cells containing the pKOBEG-Apra plasmid. Recombinant bacteria were selected on Zeo-containing LB agar plates (60 μ g/ml). Primers employed for mutagenesis are described in Table 2. The loss of the target gene in the isogenic mutants was confirmed by PCR, and all mutants were checked for susceptibility to apramycin, indicative of the loss of pKOBEG-Apra.

For directed insertional mutagenesis, internal segments of the *escN* or *ssIE* gene were prepared by PCR and cloned into the pGEM-T Easy vector (Promega). This recombinant plasmid was digested with EcoRI, and the segment of interest was cloned into the corresponding sites in suicide vector pJP5603 (29). After transformation of recombinant plasmids into DH5 α (λ pir), transformants were selected on LB agar plates containing kanamycin (50 μ g/ml), X-Gal (5-bromo-4-chloro-3-indolyl- β -D-galactopyranoside; 80 μ g/ml), and IPTG (isopropyl- β -D-thiogalactopyranoside; 0.5 mM).

Plasmid DNA of transformants harboring the correct insert was transformed into strain S17-1(λ pir) for mobilization into a nalidixic acid-resistant derivative of strain 1551-2 via filter mating on cellulose nitrate membranes. Transconjugants were selected on L agar plates containing kanamycin and nalidixic acid as previously described (30). The integration of suicide vector pJP5603 in the *escN* gene was confirmed by PCR

TABLE 2 Primers used for PCR amplification

Purpose and designation	Primer sequence (5' → 3')
Allelic exchange	
tir-zeo(F)	ATGCCTATTGGTAATCTTGGTCATAATCCCAATGTGAGTGGTCATCGCTTGCATTAGAAAGG
tir-zeo(R)	TTAAACGAAACGATTGGATCCCGGCACTGGTGGGTATTTCGAATGATGCAGAGATGTAAG
fimA-zeo(F)	ATGAAAATTAAAACCTCTGGCAATTGTTGTTCTGTCGGCTCGTCATCGCTTGCATTAGAAAGG
fimA-zeo(R)	TTATTGATACTGAACCTTGAAGGTTGCATCCGCGTTAGCTGAATGATGCAGAGATGTAAG
espA-zeo(F)	ATGGATACACCAAATGCAACATCCGTTGCTAGTGCAGCGGTCATCGCTTGCATTAGAAAGG
espA-zeo(R)	TTATTTACCAAGGGATATTGCCGAATAGTCTGTATTGCGAATGATGCAGAGATGTAAG
Gene amplification by PCR	
espA-F	GGGGTACCTCGGTGTTTTTCAGGCTGC
espA-R	CGAGCTCATTTCAAGCTGGCTATTATT
fimA-F	GTCGGTTTTAACATTCAACTGAATG
fimA-R	AATAACGCGCCTGGAACGGAAT
fimA-NcoI-F	AGCAGCCCATGGAAATTTAACTCTGG
fimA-HindIII-R	GGTAGGAAGCTTTTATTGATACTGAACCTT
eae-NcoI-F	ATACCATGGTTACTCATGGTTGTTATACCCGG
eae-HindIII-R	ATAAAGCTTTCTCACACAGACTCGATAGGCATAATTACC
espA-NcoI-F	ATACCATGGATACACCAAATGCAACATCCGTTGC
espA-HindIII-R	ATAAAGCTTTTACCAAGGGATATTGCCGAAATAG
tir-F	CACCTATAAGGCAGTCTGTTGCTG
tir-R	GGTTTCAGTTGCACTTGCAGCAG
escN-F	CGACGACTATTGCAGAGT
escN-R	GCCTTATCTGCTTCAGGA
EscN-B-19-F	CGCGGATCCTCTAAGGGAATAATATCGAACTTAAAG
EscN-S-20-R	ACGCGTCGACTCAGGCAACCACTTTGAATAGGC
ssIE-F	GATTCTGATGGCGCGTAATGATAA
ssIE-R	CTGATACAGACGCTGAATGTTGG
ssIE-F2	AGGGCGTTATCAACAAGCTGTG
ssIE-R2	CAGCGGTTTTTCCATATAGTTGAG
PJP-F	CCCAGTCACGACGTTGTAAAACG
PJP-R	AGCGGATAACAATTCACACAGG

analysis using primers EscN-B-19-F and PJP-R to verify the right junction and primers EscN-S-20-R and PJP-F to verify the left junction (Table 2). The *escN* mutant was complemented with the pEscN plasmid carrying the *escN* gene from the prototype tEPEC E2648/69 in the pACYC184 vector (31).

The integration of suicide vector pJP5603 in the *ssIE* gene was also confirmed by PCR using primers ssIE-F2 and PJP-R to verify the right junction and primers ssIE-R2 and PJP-F to verify the left junction. For complementation, the *fimA* gene from aEPEC 1551-2 was amplified by PCR using primers fimA-NcoI-F and fimA-HindIII-R (Table 2). The PCR product was then digested with NcoI and HindIII and cloned in the respective site of the pBAD/Myc-His A vector, originating recombinant plasmid pFimA, and transformed into 1551-2 Δ fimA. T1P production in the complemented strain was induced with 0.01% L-arabinose. The *eae* and the *espA* genes from the 1551-2 strain were cloned in the pBAD/Myc-His A vector, in order to complement 1551-2 *eae::Km* and 1551-2 *eae::Km* Δ espA. Experiments with the complemented strains carrying the pInt (*eae* gene in pBAD/Myc-His A) or the pEspA (*espA* in pBAD/Myc-His A) plasmids were carried out in the presence of 0.01% L-arabinose.

All mutant derivatives obtained in this study were confirmed to grow at the same rate as the wild-type strain under the conditions tested in this work (data not shown).

Biofilm formation assay on abiotic surface. Adhesion to abiotic surface (polystyrene) was analyzed using 24-well plates as described before (32). Overnight cultures of bacteria grown in LB broth were adjusted to an optical density at 600 nm (OD_{600}) of 1.0 (corresponding to ca. 10^8 CFU/ml), and 40- μ l aliquots were added to the wells containing 1 ml of LB and incubated at 37°C for 48 h. Unbound bacteria were removed after washing the cultures three times with PBS, and biofilms were fixed for 20 min with

2% formalin, washed, and then stained with 1% crystal violet (CV) for 20 min. Wells were thoroughly rinsed three times with PBS, and the dye was solubilized in 1 ml of methanol. Finally, the amount of extracted crystal violet was determined by measuring the OD_{600} by using an enzyme-linked immunosorbent assay (ELISA) Multiskan plate reader.

Pellicle formation. Strains were grown for 48 h in LB at 37°C with shaking (225 rpm) in glass tubes. Following incubation, the pellicles formed at the air-liquid interface were washed three times with PBS and then stained with CV (1%) for 20 min. After staining, pellicles were washed thoroughly with distilled water and photographed (33).

Agglutination of yeast cells. Surface-expressed T1P on aEPEC strains was assayed by the ability to agglutinate mannan-rich yeast (*Saccharomyces cerevisiae*) cells on glass slides as previously described (34). Equal volumes of bacterial and yeast suspensions were mixed on a glass slide and rocked for 1 to 2 min until the aggregates were visualized by naked eye.

Pilus purification and characterization. To purify the pilus structures, aEPEC 1551-2 growing in 35 plates (150 by 15 mm) of MacConkey agar at 37°C was harvested, and the pili were purified by mechanical shearing and differential centrifugation in order to remove outer membranes and bacterial debris (35). Pilus purification was monitored by transmission electron microscopy (TEM) and SDS-PAGE, as previously described (36). The pilus-containing supernatant was denatured with HCl (37) and resolved by SDS-PAGE (38). An 18-kDa protein was excised from the gel and subjected to mass spectrometry analysis after trypsin digestion for protein identification.

Heat extraction of surface-associated proteins. Toward identifying a putative adhesin, outer membrane proteins from aEPEC 1551-2 were heat extracted as follows. Briefly, 10 ml of an overnight bacterial culture in LB

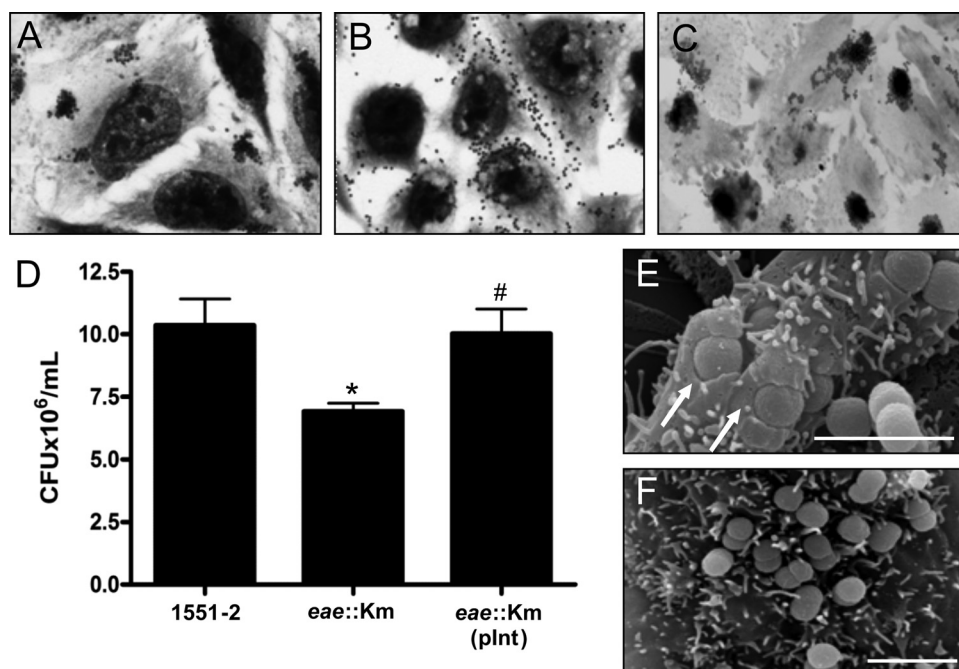


FIG 1 Atypical EPEC 1551-2 interactions with HeLa cells. (A, B, and C) Light microscopy images demonstrating the differences in the adherence patterns between 1551-2 (LA), the derivative *eae* mutant (DA), and the complemented strain (LA), respectively (original magnification, $\times 1,000$). (D) Quantitative adherence assay demonstrating a reduction of approximately 33.7% in adherence of 1551-2 *eae::Km* compared with the wild-type strain. *, $P < 0.05$; #, significantly more adherent than 1551-2 *eae::Km* ($P < 0.05$). (E and F) SEM images showing the presence or absence of AE lesions (arrows) after 6 h of contact by 1551-2 and 1551-2 *eae::Km*, respectively. Bar = 2 μ m.

or DMEM was harvested, resuspended in 100 μ l of PBS (pH 7.4), and heated at 65°C for 1 h (39). This treatment was followed by centrifugation at 10,000 $\times$ g at 4°C, and the supernatant was analyzed by SDS-PAGE in 12% polyacrylamide gels (38).

Analysis of aEPEC secreted proteins. Determination of aEPEC secreted proteins was done as previously described, with minor modifications (40). Briefly, strains were grown overnight in LB at 37°C in a shaker at 225 rpm and then subcultured 1:50 into 5 ml of DMEM. After 6 h of growth, 1.5 ml of each culture was harvested by centrifugation at 16,000 $\times$ g for 5 min, and the supernatant was filtered through a 0.45- μ m filter (Millipore). Trichloroacetic acid was added to a final concentration of 10% (vol/vol) to precipitate the secreted proteins. After overnight incubation at 4°C, the Eppendorf tubes were centrifuged for 20 min at 4°C, the supernatant was discarded, and the pellet was dissolved in 20 μ l of SDS-PAGE sample buffer. Saturated Tris was added to neutralize the residual trichloroacetic acid. The secreted proteins were used for SDS-PAGE (12%) analysis.

Ultrastructural studies. The presence of pili on 1551-2 was visualized by negative staining with 1% phosphotungstic acid and electron microscopy (EM) in a Philips CM12 electron microscope at 80 kV as previously described (27, 36). Glass coverslips containing fixed HeLa cells with adhering bacteria were prepared for scanning electron microscopy (SEM) as described previously (27) and visualized by using a Hitachi S-4500 scanning electron microscope (Hitachi, Tokyo, Japan).

Statistical analysis. All analyses were performed by using Student's *t* test. A *P* value of ≤ 0.05 by two-tailed analysis was taken to indicate statistical significance.

RESULTS

aEPEC 1551-2 continues to adhere to HeLa cells in the absence of intimin. We have previously demonstrated that deletion of the *eae* gene completely abolished the ability of 1551-2 to display the LA pattern in HeLa cells (24). Surprisingly, 1551-2 *eae::Km* re-

mained adherent to HeLa cells and no longer displayed LA but DA (Fig. 1A and B). Quantitatively, the *eae* mutant was approximately 33.7% less adherent than 1551-2 ($P < 0.05$) (Fig. 1D). Complementation of 1551-2 *eae::Km* in *trans* with *eae* in pInt restored the ability of this strain to produce the original LA pattern of 1551-2 (Fig. 1C) and at comparable adherence levels ($P > 0.05$) (Fig. 1D). The complemented strain was significantly more adherent than the *eae* mutant ($P < 0.05$). As expected, 1551-2 *eae::Km* lost the ability to produce the AE lesions observed in the wild-type strain (Fig. 1E and F).

Identification and role of T1P in cell adherence of aEPEC 1551-2. We set out to investigate the production of fimbrial adhesins in 1551-2 propagated at 37°C using different culture media (e.g., MacConkey, Luria-Bertani, Minca, 1% tryptone, colonization factor agar [CFA]) by electron microscopy analyses. Bacteria recovered from all the different media tested showed the presence of rigid rod-like pili (Fig. 2A), and MacConkey agar was selected for pilus purification (Fig. 2B). The purified pili dissociated into monomers of 18 kDa in 16% polyacrylamide gels under denaturing conditions with HCl treatment (Fig. 1C). Mass spectrometry analyses of the 18-kDa protein showed the presence of peptides whose amino acid sequence matched the sequence of the T1P major pilin subunit FimA (data not shown). In order to investigate the presence of functional T1P in 1551-2, the standard yeast agglutination assay was employed. 1551-2 recovered from the supernatant of infected HeLa cells after 6 h of incubation agglutinated yeast cells through recognition of mannosidic residues. This phenotype could be abolished in the presence of 3% D-mannose (data not shown).

The contribution of T1P in the adherence process of 1551-2

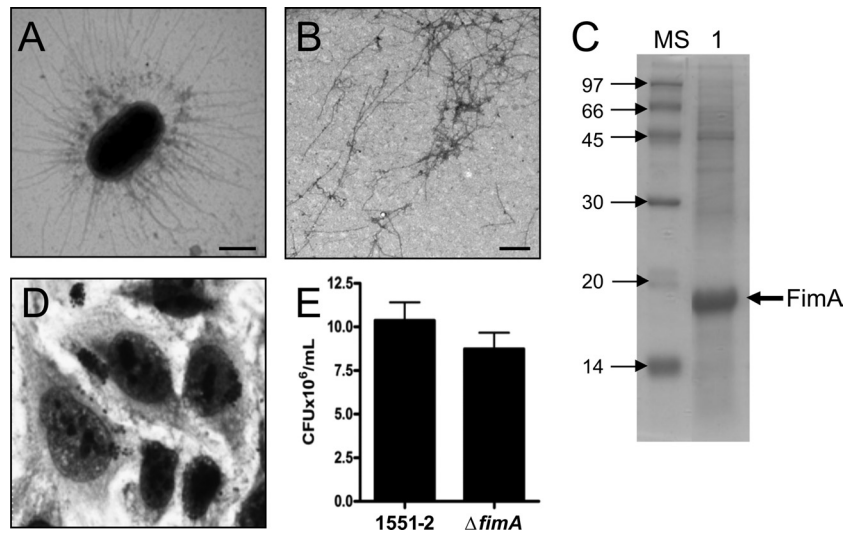


FIG 2 Identification and purification of T1P from 1551-2 and analysis of its role in HeLa cell adherence. (A) Micrograph of piliated 1551-2 obtained from the supernatant of infected cells after 6 h of incubation and visualized by TEM. (B) TEM micrograph showing pili purified from 1551-2. (C) Depolymerization of purified pili in a 16% SDS-PAGE gel showing an 18-kDa protein (lane 1) whose mass spectrometry analysis showed identity to the FimA pilin subunit protein. MS, mass standards. (D) Light microscopy micrograph showing the LA pattern exhibited by 1551-2 $\Delta fimA$ after 6 h of incubation with HeLa cells (original magnification, $\times 1,000$). (E) Quantification of adhering bacteria showing only minor differences in the level of adherence between the wild type and the T1P mutant ($P > 0.05$). Bar = 500 nm.

was investigated by mutating the *fimA* gene. The resulting *fimA* mutant lost its capacity to agglutinate yeasts, confirming the absence of T1P. However, the *fimA* mutant retained the ability to adhere to HeLa cells in an LA pattern (Fig. 2D). Although a reduction of approximately 15% could be observed between the *fimA* mutant and the wild-type strain (Fig. 2E), this difference could not be considered statistically significant ($P > 0.05$).

Identification and characterization of SsIE, a type 2 secretion substrate, in aEPEC 1551-2. A protein of approximately 150 kDa was identified in the heat extracts and secreted protein fractions analyzed by SDS-PAGE (Fig. 3). Mass spectrometry analyses of this protein showed the presence of peptides whose amino acid

sequence matched the sequence of a recently reported type 2 secreted protein termed SsIE (secreted and surface-associated lipoprotein from *E. coli*) (41). We observed that this protein is produced in both LB medium and DMEM, at 37 and 28°C, and in an atmosphere supplemented with 5% CO₂ (Fig. 3A). Given that SsIE was suggested to be involved in biofilm formation in the tEPEC E2348/69 and rabbit EPEC strains, we investigated the contribution of this protein in the adherence of 1551-2 by deleting the *ssIE* gene. Absence of the SsIE corresponding band in the mutant strain can be observed in Fig. 3B. Qualitative and quantitative adherence assays did not indicate any contribution of this protein to the ability of 1551-2 to adhere to HeLa cells or to form LA (Fig. 3C and D).

Contribution of the T3SS proteins to aEPEC 1551-2 adherence. In order to evaluate if the DA phenotype observed in the 1551-2 *eae::Km* strain is dependent on proteins secreted by the T3SS, we deleted the *escN* gene in the aEPEC 1551-2 strain. This gene encodes a cytoplasmic ATPase protein responsible for energizing the secretion machinery (31). The adherence of the 1551-2 *escN::Km* strain was drastically reduced (approximately 99.3%) compared with wild-type 1551-2, with only a few isolated bacteria adhering to HeLa cells ($P < 0.005$), strongly supporting the involvement of the T3SS translocon in the adherence of 1551-2 (Fig. 4A). Trans-complementation of the 1551-2 *escN::Km* strain with pEscN restored the LA pattern after 6 h of incubation (Fig. 4B and C).

The DA phenotype is dependent on the T3SS needle complex. Aiming to confirm the involvement of the T3SS needle complex in promoting DA, we constructed a double intimin/EspA mutant. In contrast to 1551-2 *eae::Km*, which displays the DA pattern in contact with HeLa cells, the intimin/EspA double mutant strain became nonadherent (Fig. 5A and B). Complementation of this mutant with the pEspA plasmid restored the ability of the mutant to adhere in a DA manner (Fig. 5C). Comparative quantitative as-

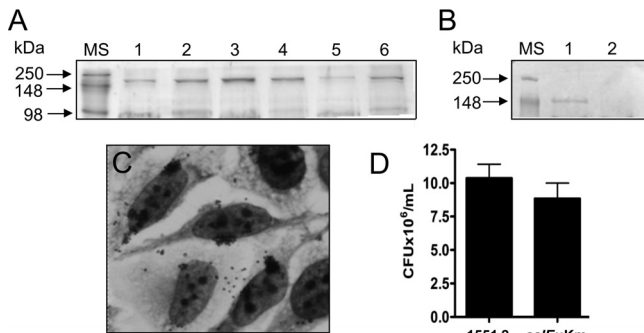


FIG 3 Identification and characterization of the T2SS substrate SsIE in supernatants of 1551-2. (A) Heat extract of the 1551-2 strain grown in DMEM (lanes 1, 3, and 5) and LB (lanes 2, 4, and 6) at 37°C with shaking (lanes 1 and 2), at 28°C with shaking (lanes 3 and 4), and at 37°C under a 5% CO₂ atmosphere without shaking (lanes 5 and 6). (B) Secreted protein extract showing the presence of the T2SS SsIE in the wild type (lane 1) and absence in the *ssIE* mutant (1551-2 *ssIE::Km*) (lane 2). (C) Light microscopy image showing LA by the *ssIE* mutant after 6 h of incubation with HeLa cells (original magnification, $\times 1,000$). (D) Quantitative comparison of adherence between the wild type and *ssIE* mutant showing no statistical difference ($P > 0.05$).

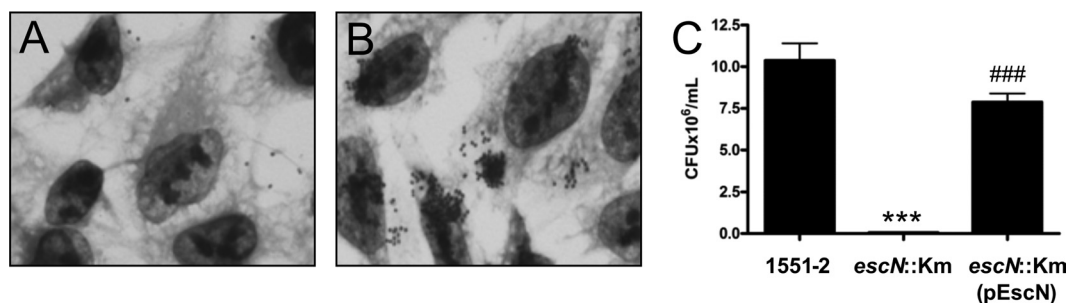


FIG 4 Contribution of T3SS to the adherence of aEPEC 1551-2 to HeLa cells. (A and B) Adherence of the T3SS-ATPase-deficient mutant (1551-2 *escN::Km*) and the complemented strain after 6-h assay (original magnification, $\times 1,000$). (C) Quantitative comparison of adherence between the indicated strains demonstrating that adherence of 1551-2 to HeLa is dependent on a functional T3SS. ***, significantly less adherent than 1551-2 ($P < 0.005$); ###, significantly more adherent than the 1551-2 *escN::Km* mutant ($P < 0.001$).

says demonstrated that adherence of the 1551-2 *eae::Km* Δ *espA* double mutant was reduced approximately by 99.4% with respect to the intimin mutant ($P < 0.0001$) (Fig. 5D). Similar results were observed in the adhesion inhibition assay with EspA antibodies (Fig. 5E), confirming the involvement of the T3SS translocon in the DA phenotype observed in the *eae* mutant strain. Complementation with pEspA plasmid quantitatively restored the adherence of 1551-2 *eae::Km* Δ *espA* compared with 1551-2 *eae::Km* ($P < 0.005$) (Fig. 5D). However, no statistical difference was observed between the 1551-2 *eae::Km* strain and the double mutant complemented strain ($P > 0.05$).

Additionally, two other double mutants were generated, intimin/Tir (1551-2 *eae::Km* Δ *tir*) and intimin/T1P (1551-2 *eae::Km* Δ *fimA*) mutants, that were approximately 16.3 and 9.1% less adherent, respectively, than the intimin mutant (Fig. 5D). These mutants were still able to display DA on HeLa cells, discarding the involvement of T1P and the intimin-Tir interaction in the DA phenotype (data not shown).

T1P mediated biofilm formation in aEPEC 1551-2. The ability of 1551-2 and its derivative mutants to develop biofilm on an abiotic surface (polystyrene) was investigated. In comparison with the wild-type strain, the mutation in the *fimA* gene drastically

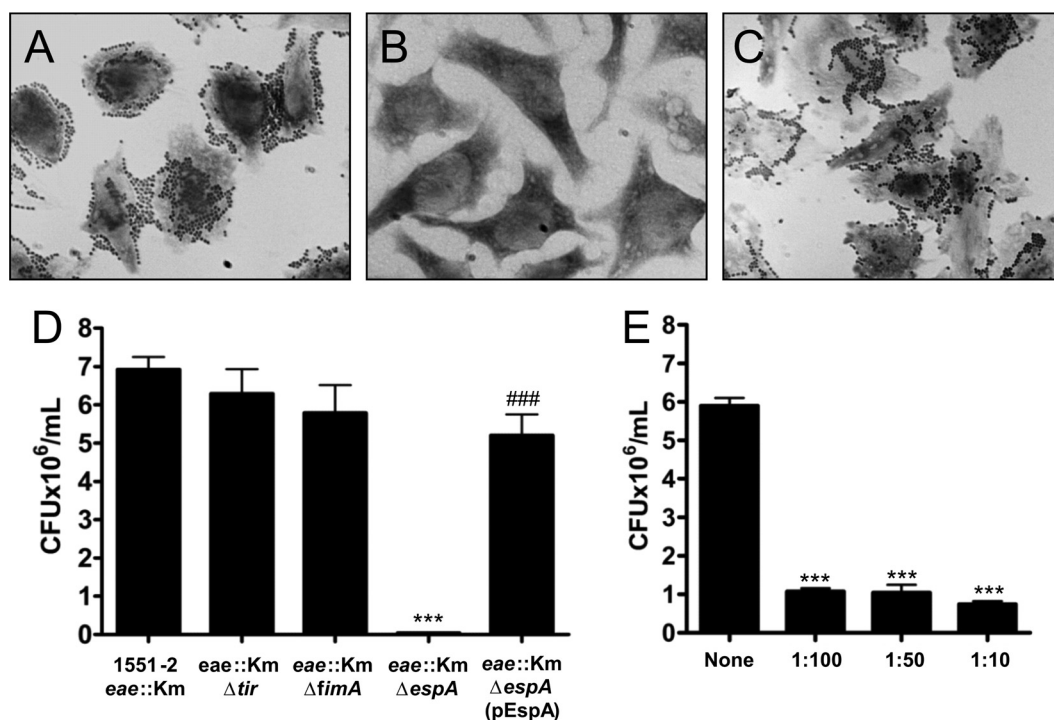


FIG 5 The DA phenotype is dependent on the T3SS needle complex. (A, B, and C) Light microscopy images showing the adherence patterns observed in 1551-2 *eae::Km*, the double *eae espA* mutant, and the complemented strain, respectively (original magnification, $\times 1,000$). (D) Effect of mutations in T3SS-related and *fimA* genes in the DA phenotype of 1551-2 *eae::Km* to HeLa cells. While mutations in *tir* and *fimA* genes did not affect the adherence ability of the *eae* mutant, deletion of *espA* significantly impaired its adherence. ***, significantly less adherent than 1551-2 *eae::Km* ($P < 0.0001$); ###, significantly more adherent than 1551-2 *eae::Km* Δ *espA* ($P < 0.005$). (E) Quantification of adherence of the 1551-2 *eae::Km* strain in the presence of three dilutions of the anti-EspA (1:100, 1:50, and 1:10). ***, statistically significant with respect to the values obtained in the absence (none) of the anti-EspA antibody ($P < 0.0001$).

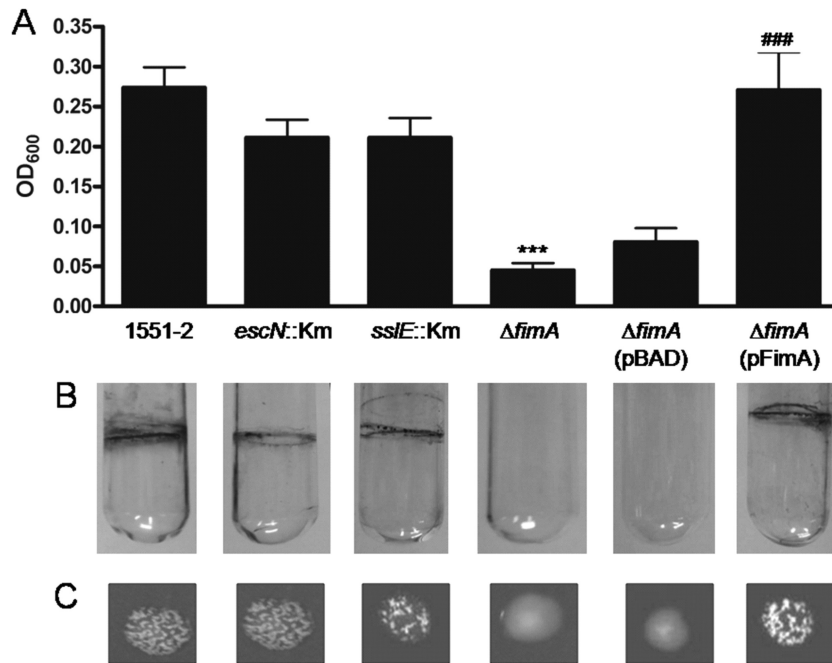


FIG 6 Biofilm formation by aEPEC 1551-2 and derived mutants. (A) Quantitative determination of crystal violet adsorption upon biofilm formation by the indicated strains. Biofilm formation by 1551-2 is dependent on T1P production but not on T3SS or the T2SS substrate. (B) Pellicle formation at the air-liquid interface upon bacterial growth in glass tubes. This phenotype was dependent on T1P production. (C) Images comparing yeast agglutination by the indicated strains. ***, significantly less adherent than 1551-2 ($P < 0.0001$); ###, significantly more adherent than 1551-2 $\Delta fimA$ ($P < 0.005$).

affected the ability of 1551-2 to form biofilm on plastic (Fig. 6). The quantitative biofilm assay confirmed the difference in adherence efficiency between the wild-type and the *fimA* mutant strains ($P < 0.0001$). Our results indicate that T3SS proteins as well as the T2SS substrate SslE did not interfere with the biofilm formation phenotype in strain 1551-2 (Fig. 6A).

aEPEC 1551-2 formed a pellicle at the air-liquid interface when grown with shaking in LB for 48 h at 37°C (Fig. 6B). Interestingly, the *fimA* mutant lost the ability to adhere to the glass tube wall in a ring-like format, while the *escN* and the *sslE* mutants preserved their biofilm-producing traits. The association of pellicle formation at the air-liquid interface with T1P production could be confirmed by demonstrating the close association between the presence of T1P, investigated by the ability to agglutinate yeast cells, and the ring-like formation in glass tube (Fig. 6C). Deletion of the *fimA* gene, confirmed by loss of the ability to agglutinate yeast, completely abolished the ability to form a pellicle at the air-liquid interface and to develop biofilm on a polystyrene surface.

The introduction of pFimA into the *fimA* mutant resulted in regaining of the capacity to adhere to abiotic surfaces (polystyrene and glass) and express T1P, as confirmed by the yeast agglutination assay (Fig. 6C). The pBADMyc-His A plasmid vector was employed as a negative control, confirming that the plasmid caused no alteration in the phenotype of the *fimA* mutated strain.

DISCUSSION

The importance of intimin as an adhesin has been extensively documented in tEPEC and aEPEC strains (20, 22, 23, 42). Although it is clear that intimin is an important adhesin of aEPEC, it is also evident that intimin is not the only adhesion factor present in this group of strains (24, 43, 44). We previously reported that

aEPEC strain 1551-2 forms LA 6 h postinfection and that an isogenic *eae* mutant shows DA (24). These data indicated that the switch from LA to DA pattern was intimin dependent but that the DA phenotype was intimin-Tir independent. This prompted us to search in 1551-2, using a PCR-based approach, for adhesin genes previously shown to be involved in DA of other DEC strain adhesin genes, namely, the genes for F1845, AIDA-I, and LDA (locus for diffuse adherence). We also surveyed in 1551-2 many other fimbrial genes present in other pathogenic *E. coli* strains, for example, the genes for AAF/I-II, LPF, HCP, curli, T1P, ECP, and P and S fimbriae (8, 13, 45, 46). Except for *fimE*, none of the genes encoding these adhesins were found in 1551-2 (46). Thus, in this study, in order to identify the elusive DA adhesin, we undertook a different approach: using TEM, we sought novel pili upon growth of the bacteria in different growth conditions, we analyzed supernatants for new secreted proteins, and we studied the role of the T3SS translocon.

Much to our surprise, the pili produced by and purified from the 1551-2 strain turned out to be T1P. Of note, bacteria harvested from the 6-h adherence assay strongly agglutinated yeast cells, demonstrating that T1P is produced in the presence of HeLa cells. T1P are ubiquitous among the *Enterobacteriaceae* and have been vastly studied and implicated in various virulence-associated events of pathogenic *E. coli* strains, e.g., invasion and persistence in host cells, biofilm formation, induction of proinflammatory molecules, and cell signaling (reviewed in reference 47). T1P enable uropathogenic *E. coli* (UPEC) strains to attach to mannose residues on the surface of urinary epithelium cells and play a significant role in the colonization and invasion of bladder epithelial cells as the first critical step in the establishment of cystitis and urinary tract infections (48, 49). T1P of prototypic tEPEC strain

E2348/69 were not associated with cell adherence (30) but were considered important for development of the AA pattern of enteroaggregative *E. coli* (EAEC) strain 042 (50). In the present study, 1551-2 Δ *fimA* showed the same level of adherence to HeLa cells as the wild-type strain, indicating that these pili were not the DA adhesins. Discrepancies in the role of T1P in cell adherence by different *E. coli* strains could be attributed to nucleotide variations in the tip adhesin *fimH* gene resulting in differences in their receptor-binding capabilities (51, 52). Studies showing variations in *fimH* and possible implications of these variations in aEPEC adherence are missing. Likewise, the fact that in our initial PCR 1551-2 was *fimA* negative could be explained by differences in the flanking regions of the gene that were not totally compatible with the sequence of the primers employed. Thus, it is possible that variants of *fimA* also exist among aEPEC.

The search for new potentially relevant surface-associated (heat extracts) or secreted proteins in supernatants of 1551-2 *eae::Km* cultures obtained in LB broth or DMEM led us to identify a large protein of approximately 150 kDa. Mass spectrometry analyses of this protein revealed that it was a product of the *yghJ* chromosomal gene, which encodes a predicted lipoprotein. YghJ production was demonstrated in bacteria grown at 28° and 37°C in an atmosphere supplemented or not with 5% CO₂. The fact that YghJ is produced under conditions similar to those that induce production of the BFP and T3S effectors in tEPEC (e.g., 37°C in DMEM) might indicate an important biological significance.

While this work was in progress, a study by Baldi et al. (41) became available demonstrating that tEPEC E2348/69 secretes an outer membrane lipoprotein via T2SS, which they called SslE (for secreted and surface-associated lipoprotein from *E. coli*) and found to be important for biofilm maturation but not for cell adherence (41). SslE is an ortholog protein of the accessory colonization factor D (AcfD) of *Vibrio cholerae*, which is required for efficient colonization of infant mouse intestine (53). It was apparent that the YghJ protein of aEPEC 1551-2 and the SslE of E2348/69 were the same protein.

In order to establish the function of this outer membrane lipoprotein in the adherence processes of strain 1551-2, we constructed an *sslE* mutant. The mutation in the *sslE* gene led to the absence of SslE in supernatants but did not affect the secretion of EspA, EspB, or EspD, which are dependent on the T3SS (data not shown). The 1551-2 *sslE::Km* mutant did not lose the ability to produce LA on HeLa cells, did not show a significant reduction in the number of bacteria associated with the epithelial cells ($P > 0.05$), and was not associated with biofilm formation. Thus, the role of SslE in the pathogenicity of 1551-2 or other aEPEC strains in general remains unclear.

After ruling out the involvement of T1P and SslE as DA adhesin, we investigated if the DA phenotype would be dependent on a functional T3SS. We created an insertional mutation in *escN*; this gene encodes the ATPase required for secretion of the T3SS assembly components (31). 1551-2 *escN::Km* was no longer adherent in HeLa cells, providing compelling evidence of the role of the T3SS in DA of 1551-2 *eae::Km*. We also tested the *escN* mutant in differentiated Caco-2 cells. As observed in HeLa cells, the ability of 1551-2 *escN::Km* to adhere to this enterocyte cell line model was significantly reduced (approximately 91.4%) in comparison with that of the wild-type 1551-2 strain ($P < 0.0001$) (data not shown). The role of the T3SS in the virulence properties of EPEC and EHEC has been well established (54). In particular, the EspA fila-

ment, or translocon, was shown to link the bacteria to the host cell membrane and to be responsible for injection of proteins into the cytosol. Previous studies carried out with tEPEC E2348/69 and an EHEC strain 413/89-1 (serotype O26:H⁻) pointed to a possible involvement of the T3SS filaments in adherence to epithelial cells (42, 55, 56). The T3SS apparatus was also demonstrated to be responsible for attachment to leaves in EHEC O157 and O26 strains (57, 58). To specifically define the role of the EspA translocon in DA, we then constructed a double intimin/EspA mutant. As predicted, this mutant totally lost its adherence capabilities. Thus, this event is intimin-Tir independent, as the single *eae* and the double *eae tir* mutants displayed DA. Further, the DA pattern was inhibited with anti-EspA antibodies. All together, these data strongly indicated that the T3SS translocon is responsible for the DA displayed by 1551-2 *eae::Km*. Nonetheless, future studies are needed to elucidate whether the EspA filament *per se* or the tip-associated pore-forming proteins, EspB and EspD, are responsible for mediating host cell adherence.

Biofilm formation is a property not fully explored among aEPEC isolates. The ability of an aEPEC strain of serotype O55:H7 to produce biofilm at low temperatures was associated with curli production (33). While curli biogenesis genes are commonly found among aEPEC isolates, production of curli is a rare event (46). The EspA filament of tEPEC E2348/69 was suggested to be responsible for biofilm development on plastic under static conditions (59). In this study, we showed that the T3SS translocon is not associated with biofilm in 1551-2, since the *escN* mutant strain is still able to produce the same level of biofilm as the wild-type 1551-2 strain. In conclusion, our data support a role for intimin in LA, for the T3SS translocon in DA, and for T1P in biofilm formation, suggesting that all of these adhesins act in concert to facilitate host intestinal colonization by aEPEC strains.

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