



Fimbria-Encoding Gene *yadC* Has a Pleiotropic Effect on Several Biological Characteristics and Plays a Role in Avian Pathogenic *Escherichia coli* Pathogenicity

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The extraintestinal pathogen termed avian pathogenic *Escherichia coli* (APEC) is known to cause colibacillosis in chickens. The molecular basis of APEC pathogenesis is not fully elucidated yet. In this work, we deleted a component of the Yad gene cluster (*yadC*) in order to understand the role of Yad in the pathogenicity of the APEC strain SCI-07. *In vitro*, the transcription level of *yadC* was upregulated at 41°C and downregulated at 22°C. The *yadC* expression *in vivo* was more pronounced in lungs than in spleen, suggesting a role in the early steps of the infection. Chicks infected with the wild-type and mutant strains presented, respectively, 80% and 50% mortality rates. The $\Delta yadC$ strain presented a slightly decreased ability to adhere to HeLa cells with or without the D-mannose analog compared with the wild type. Real-time PCR (RT-PCR) assays showed that *fimH* was downregulated ($P < 0.05$) and *csgA* and *ecpA* were slightly upregulated in the mutant strain, showing that *yadC* modulates expression of other fimbriae. Bacterial internalization studies showed that the $\Delta yadC$ strain had a lower number of intracellular bacteria recovered from Hep-2 cells and HD11 cells than the wild-type strain ($P < 0.05$). Motility assays in soft agar demonstrated that the $\Delta yadC$ strain was less motile than the wild type ($P < 0.01$). Curiously, flagellum-associated genes were not dramatically downregulated in the $\Delta yadC$ strain. Taken together, the results show that the fimbrial adhesin Yad contributes to the pathogenicity and modulates different biological characteristics of the APEC strain SCI-07.

Avian pathogenic *Escherichia coli* (APEC) strains cause a variety of extraintestinal infections in poultry collectively known as colibacillosis (1, 2). Although the complete mechanisms of APEC pathogenicity are not fully elucidated, it is believed that colibacillosis starts with colonization of the host's upper respiratory tract, with the bacterium expressing one or more colonization factors known as adhesins (3). The infection can subsequently spread into the lungs and other inner organs, leading to a fatal septicemia (1, 4, 5). Among the known APEC adhesins, those more studied are the type 1 fimbriae, P fimbriae, and curli fimbriae, respectively (6).

Besides their roles in the adhesion and subsequent colonization of environmental surfaces, those fimbriae are assumed to be essential for the establishment of a host-parasite relationship and further disease progression (7). Both type 1 and curli fimbriae play important roles in the initial bacterial colonization of the respiratory epithelium (1, 8), while P fimbriae (9) are important for later stages of infection (1).

In addition to the fimbriae mentioned above, *E. coli* possesses several other fimbrial operons (i.e., Yad, Ycb, Ybg, Yfc, Yra, Sfm, Ygi, and Yeh) that display sequence and organizational homologies to type 1 fimbria and could contribute to *E. coli*'s ability to adhere to and colonize the host epithelia (10, 11).

The adhesion operon Yad is composed of seven genes, *yadN* (major subunit), *ecpD* (usher), *htrE* (chaperone), *yadMLK* (minor subunits), and *yadC* (adhesive tip) (12). The fimbria Yad is more prevalent among uropathogenic *E. coli* (UPEC) than commensal *E. coli* strains and contributes to the adherence to bladder epithelial cells and biofilm formation (10). Moreover, it was demonstrated that *yadK* is important for adhesion of acid-stressed enterohemorrhagic *Escherichia coli* (EHEC) to epithelial cells, increasing bacterial colonization and virulence (12).

In APEC strain O78, the deletion of *yadL* has suggested that the fimbria Yad has a subtle but positive contribution to APEC virulence (13). Furthermore, *in silico* analyses performed by our research group (unpublished data) with 12 *E. coli* genomes, including five APEC genomes, revealed that *yadC* is under positive selection. This suggests the involvement of the fimbria Yad in the host-parasite interactions.

To better understand the role of Yad fimbria in APEC biological characteristics and pathogenicity, we deleted *yadC* in an APEC swollen head syndrome strain (SCI-07) and studied its effects comparing both the wild type and the isogenic mutant through comprehensive *in vitro* and *in vivo* assays.

MATERIALS AND METHODS

Bacterial strains and growth conditions. APEC strain SCI-07 (ONT: H31) was isolated from a laying hen diagnosed with swollen head syn-

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TABLE 1 List of strains and plasmids used in this work

Strain or plasmid	Relevant genotype	Source or reference
Strains		
SCI-07	APEC strain isolated from lesions of a laying hen presenting clinical signs of swollen head syndrome	Our laboratory
DH10β	<i>E. coli</i> K-12 strain	Our laboratory
HB101	<i>E. coli</i> K-12 strain	Our laboratory
DH5α	<i>E. coli</i> K-12 strain	Our laboratory
EIEC strain	Enteroinvasive <i>E. coli</i> O164 strain	Laboratory of UEL, Londrina, Brazil
STM-F98	<i>Salmonella</i> Typhimurium	Laboratory of Ornithopathology, FCAV-UNESP, Jaboticabal, Brazil
Plasmids		
pKD3	<i>cat</i> gene	Datsenko and Wanner (17)
pKD46	Amp ^r -expressing λ <i>red</i> recombination	Datsenko and Wanner (17)
pACYC177	Cloning vector	New England Biolabs

drome. Its genome is available in GenBank (accession no. CP000468.1) (14). The strains were cultured in LB or Dulbecco's modified Eagle's medium (DMEM) at 37°C with appropriate antibiotics when necessary (Table 1). All of the animal experimental protocols were approved by the Ethics Committee of Animal Use-CEUA-UNICAMP (protocol no. 2669-1; Brazilian law no. 11794).

RNA extraction. Total RNA of strain SCI-07 was extracted from LB (*in vitro*) as well as from lungs and spleen of day-old infected chicks. For *in vitro* RNA extraction, culture was grown in LB at 22, 37, or 41°C until reaching an optical density at 600 nm (OD₆₀₀) of 0.5 to 0.6. Afterwards, RNA was extracted using the "RNAeasy minikit" (Qiagen). For *in vivo* assays, an inoculum of 0.1 ml was injected into the right air sac of five 1-day-old chicks. After 24 h postinfection (hpi), the left lung and spleen were removed and homogenized in 1 ml TRIzol solution (Ambion). RNA was extracted using the Purelink RNA minikit (Ambion, Life Technologies).

Relative expression of *yadC* in SCI-07 by qRT-PCR. Real-time PCR (RT-PCR) was performed to quantify the transcription of *yadC* in LB at different temperatures (22, 37, and 41°C). Primers for quantitative RT-PCR (qRT-PCR) were constructed with Primer Express software version 3.0 (Applied Biosystems) (see Table S1 in the supplemental material), using the SCI-07 genome as the template. The qRT-PCR experiments were performed as described previously (15). RNA polymerase subunit A (*rpoA*) was used as the endogenous control (see Table S1). Data were normalized based on the transcription level of *rpoA* in the wild type and then analyzed using the comparative critical threshold cycle (C_T) method described by Walters and Sperandio (16).

Deletion of *yadC*. The $\Delta yadC$ strain was constructed following the methodology described by Datsenko and Wanner (17). To replace gene *yadC* by homologous recombination, a chloramphenicol (Cm) cassette from pKD3 was amplified, and strain SCI-07, harboring the λ *red* plasmid pKD46, was transformed by electroporation as described by de Paiva et al. (15). The $\Delta yadC$ deletion was confirmed by PCR using primers F-*yadC* 1, R-*yadC* 2 (internal), F-*yadC* 3, and R-*yadC* 4 (external) (see Table S1 in the supplemental material).

Construction of the *yadC* complemented strain. For construction of a complemented strain, *yadC* (1,263 bp), along with its promoter, was cloned into the cloning vector pACYC177. Using primers F- $\Delta yadC$ and R- $\Delta yadC$ (see Table S1 in the supplemental material), a fragment containing gene *yadC* and the enzyme *Dra*III and *Hind*III restriction sites was amplified and purified. The fragment was then ligated to plasmid pACYC177 *Dra*III and *Hind*III restriction sites. The recombinant plasmid was electroporated into competent cells of strain DH10β (15). Positive colonies were tested for the presence of *yadC* by PCR using the same complementation primers. The resulting plasmid, pACYC177-*yadC*, was isolated from strain DH10β and then transformed into competent cells of the $\Delta yadC$ strain to make the *C.ΔyadC* strain.

qRT-PCR assays for adhesion- and motility-related genes. Bacterial cultures of wild-type, mutant, and complemented strains were aerobically grown until they reached an OD₆₀₀ of 0.5 to 0.6. RNA from three biological samples was extracted using the RNAeasy minikit (Qiagen). Real-time PCR assays were performed to quantify the transcription of bacterial genes related to adhesion (*fimH*, *csaA*, and *ecpA*) and motility (*fliC*, *flhD*, and *flgE*) and the motility modulator *ompR* in each strain. Primers were designed (see Table S1 in the supplemental material) and qRT-PCR experiments were performed as described previously (6).

Bacterial adhesion to HeLa cells. The SCI-07, $\Delta yadC$, and *C.ΔyadC* strains were evaluated for their ability to adhere to HeLa cells in the presence (1%) and absence of the D-mannose analog. *E. coli* strains O164 (enteroinvasive *E. coli* [EIEC]) and HB101 were used as positive and negative controls, respectively. Monolayers of HeLa cells were grown in DMEM containing 10% fetal bovine serum (FBS) at 37°C in a 5% CO₂ incubator until they reached confluent growth. These incubation conditions were used for all eukaryotic cell lines. HeLa cells were infected with 10 μl of bacterial inocula at a concentration of 10⁷ CFU/ml in triplicate in 24-well plates as described previously (6). The obtained suspension was diluted 1- to 10-fold and plated onto MacConkey agar (MAC) containing 100 μg ml⁻¹ streptomycin (Sm) for colony-forming unit determination.

Bacterial invasion of human epithelial Hep-2 cells. The invasion abilities of the three studied strains were assessed *in vitro* with cultured Hep-2 cells following the methodology described by Scaletsky et al. (18). *E. coli* strains O164 (EIEC) and DH5α were used as positive and negative controls, respectively. A volume of 10 μl of a bacterium-phosphate-buffered saline (PBS) suspension (10⁷ CFU/ml) was inoculated onto monolayers of Hep-2 cells in triplicate. Each well was filled with DMEM and 10% FBS up to 1 ml and incubated for 3 h. The extracellular bacteria were removed by washing the cells twice with PBS. Cells were replenished with fresh DMEM containing 20 mg ml⁻¹ of ampicillin (Amp) for the wild-type and mutant strains and 25 mg ml⁻¹ of kanamycin (Km) for the complementation strain and then incubated for another 3 h. Afterwards, cells were washed with PBS and lysed with subsequent colony-forming unit determination.

Bacterial survival in HD11 macrophage cells. For the macrophage bacterial survival assay, HD11 chicken macrophage cells (19) were infected with the wild-type, $\Delta yadC$, and *C.ΔyadC* strains. *Salmonella enterica* serovar Typhimurium STM-F98 and *E. coli* HB101 were used as positive and negative controls, respectively. A volume of 10 μl of bacterium-PBS suspension (10⁷ CFU/ml) was inoculated onto monolayers of HD11 cells in triplicate. Each well was filled with RPMI 1640, supplemented with 2 mM glutamine and 10% FBS up to 1 ml, and incubated for 1 h. The extracellular bacteria were removed by washing the plate with PBS. After this, 1 ml of the glutamine-FBS-RPMI medium containing 20 mg ml⁻¹ Amp for wild-type and mutant strains and 25 mg ml⁻¹ of Km for the complemented strain was added to each well and incubated for 3 h or

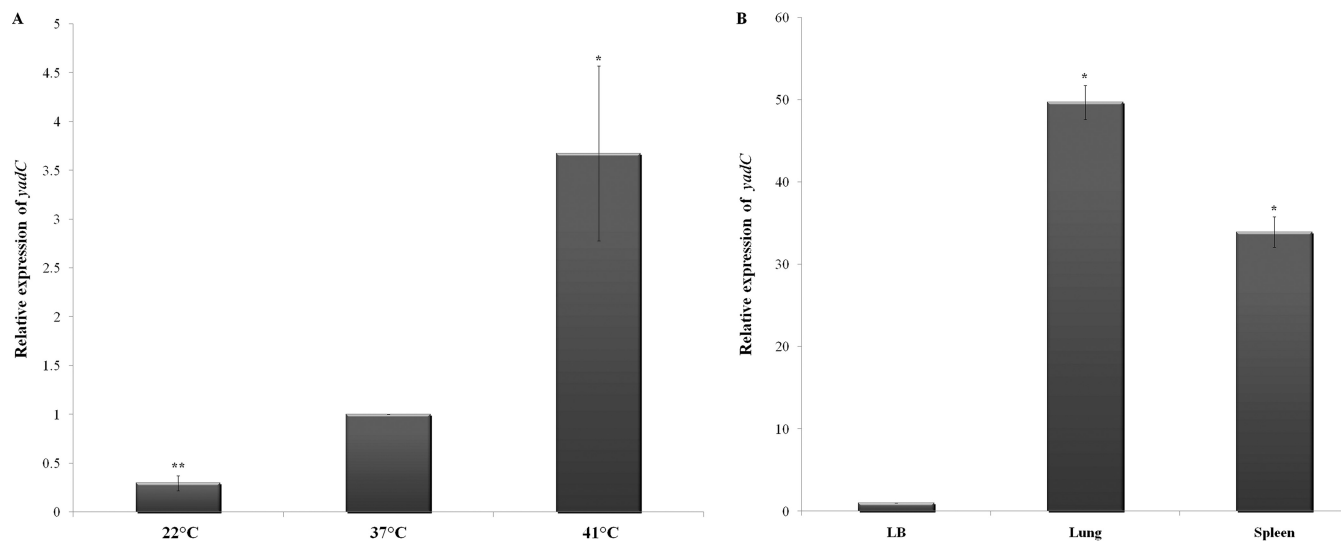


FIG 1 Relative fold expression of *yadC* by strain SCI-07, as determined by qRT-PCR. All levels were based on the comparison with the transcriptions in LB at 37°C. (A) *In vitro* (LB) at 22, 37, and 41°C. (B) *In vivo* (lung and spleen). One asterisk indicates statistical significance at $P < 0.05$, and two asterisks indicate significance at $P < 0.01$.

18 h. Then the plate was washed 3 times with PBS, and cells were lysed with 1% Triton X-100. The lysed cell suspension was used for colony-forming unit determination.

Motility assay. Cultures of each strain grown overnight in LB broth were stabbed into the center of LB–0.3% agar plates. The agar plates were incubated at 37°C and motility (in centimeters) was measured after 6, 8, and 10 h.

Bacterial growth assay. Overnight-incubated cultures (LB broth) of the SCI-07, $\Delta yadC$, and $C.\Delta yadC$ strains were diluted in LB as well as in DMEM (1:100) and grown at 37°C with agitation of 150 rpm. Optical densities of each bacterial culture were monitored spectrophotometrically at 600 nm every 30 min for 6 h.

***In vivo* mortality test.** Cultures from the SCI-07, $\Delta yadC$, and $C.\Delta yadC$ strains were grown (LB medium) at 37°C for 24 h. Briefly, groups of 20 birds for each strain were inoculated in the right thoracic air sac with 0.1 ml of a suspension containing 10^9 CFU or with 0.1 ml of PBS as negative control (15). Birds were observed every 12 h up to 7 days after inoculation, and the numbers of deaths were recorded as described by Antão et al. (20).

Systemic infection. The group of three 1-day-old chicks for each strain (the SCI-07, $\Delta yadC$, and $C.\Delta yadC$ strains) was infected with 0.1 ml of a suspension containing 10^8 CFU, in the right thoracic air sac. At 24 and 48 hpi, chickens were euthanized. Afterwards, samples of the lung, liver, and heart were homogenized, and colony forming unit determination was performed as described previously (15).

Statistical analysis. For statistical analysis, qRT-PCR, motility assays, and agar plate counting experiments were performed in triplicate. Statistical analysis was performed by analysis of variance (ANOVA), with Tukey's test, using ASSITAT software version 7.7.

RESULTS

Transcription level of gene *yadC* changed according to different temperatures. The transcription of gene *yadC* in strain SCI-07 was verified *in vitro* and *in vivo* by qRT-PCR. The transcription level of gene *yadC* was found to be upregulated, by almost 4-fold, when the strain was cultivated at the highest temperature (41°C), which is the usual temperature of chicken bodies, comparing to the incubation at 37°C. On the other hand, it was downregulated, by 0.2-fold, when the strain was cultivated at room temperature

(22°C) (Fig. 1A). *In vivo*, the transcription of gene *yadC* was up-regulated in lungs and spleen compared to its transcription in LB culture (37°C) (Fig. 1B). When comparing lungs and spleen, *yadC* expression was more pronounced in lungs.

The bacterial growth assay performed with the wild-type, mutant, and complemented strains showed the same trend, suggesting that YadC does not play a role in this APEC strain growth (see Fig. S1 in the supplemental material).

Deletion of *yadC* effects in SCI-07 pathogenicity and intrabody survival. To verify if gene *yadC* could play a role in the virulence of strain SCI-07, as suggested by the previous transcription assays, *in vivo* mortality assays were performed. Chicks infected with the wild-type strain (SCI-07) presented an 80% mortality rate ($n = 16$), with 11 animals being killed at the first day of observation. The mutant ($\Delta yadC$) strain presented overall a 50% mortality rate ($n = 10$) during the 7 days of experimentation, with only 8 of the chicks being killed at the first day of observation. The complemented ($C.\Delta yadC$) strain presented a complete restoration and increased virulence levels with a global mortality rate of 90% ($n = 18$) and 14 animals dead on the first day. No mortality was observed for the negative control (data not shown) (Fig. 2).

Moreover, the systemic *in vivo* infection assays demonstrated that less of the $\Delta yadC$ strain was recovered from lungs and heart than the wild-type strain 24 hpi ($P < 0.05$) (Fig. 3A, B, and C). No statistical difference was observed in the number of colony-forming units for any of the analyzed organs 48 hpi.

Role of *yadC* in bacterial adhesion. In order to better understand if *yadC* absence affects the initial infection processes, we performed biological tests, including an adhesion assay with HeLa cells in the presence and absence of the D-mannose analog. The adhesion studies showed that the adhesion levels of the wild-type strain decreased in the presence of the D-mannose analog, indicating that strain SCI-07 expresses type 1 fimbriae as previously described (15). Moreover, strain SCI-07, under these conditions, was still able to express a D-mannose-resistant adhesin, suggesting the expression of another type of fimbria. The $\Delta yadC$ strain, with

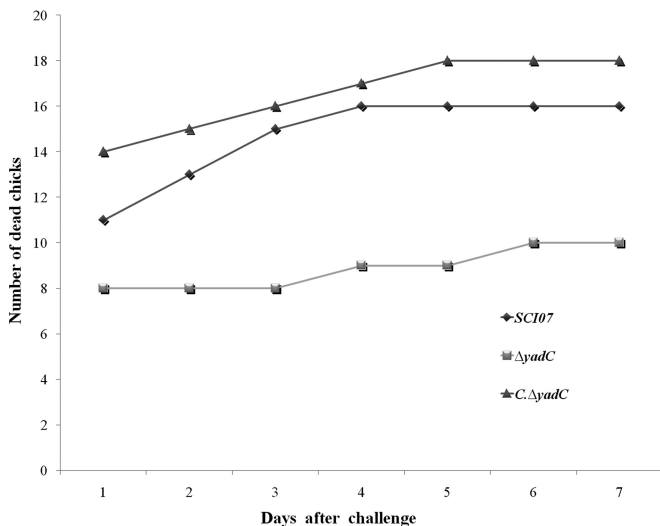


FIG 2 Mortality test in day-old chicks infected with 10^9 CFU of wild-type (SCI-07), mutant ($\Delta yadC$), and complemented (C. $\Delta yadC$) strains.

the same experimental assays, presented a slightly decreased ability to adhere to HeLa cells under both conditions (with and without D-mannose analog) compared with the wild type. This suggests that *yadC* plays a modest direct role in APEC adhesion and may be involved in the regulation of type 1 fimbriae (Fig. 4A). The remaining adhesion capacity of the SCI-07 and $\Delta yadC$ strains either in the presence or in the absence of the D-mannose analog could be due to the possible expression of other adhesins that might exist in strain SCI-07.

To evaluate if the transcription of other fimbriae known to exist in strain SCI-07 could be regulated by gene *yadC*, the qRT-PCR assays for type 1, curli, and *E. coli* common pilus (ECP) were performed *in vitro*. These tests showed that *csgA* and *ecpA* were slightly increased without statistical significance, but *fimH* was downregulated in the $\Delta yadC$ strain (Fig. 5).

Deletion of *yadC* also affects the APEC strain ability to invade Hep-2 cells. The invasion ability of the mutant strain in Hep-2 cells was tested to help us better understand the key role gene *yadC* plays in the $\Delta yadC$ strain's virulence attenuation. This

assay showed that the $\Delta yadC$ strain had a lower number of intracellular bacteria recovered from Hep-2 cells than the wild-type strain ($P < 0.05$) (Fig. 4B).

Replication and survival of strain SCI-07 inside macrophage cells was compromised by *yadC* deletion. The capacity of some pathogenic bacteria to survive and replicate inside macrophages indicates an increased ability to evade host defenses. To test if *yadC* contributes to this process, the ability of the $\Delta yadC$ strain to replicate and survive within the HD11 macrophage cells was tested. The assay showed that the $\Delta yadC$ mutant had lower survivability in this macrophage cell line than the wild-type strain 3 hpi ($P < 0.05$) (Fig. 6).

Deletion of gene *yadC* affects motility. The motility assays showed that the $\Delta yadC$ mutant strain has a decreased motility degree compared with the wild-type strain ($P < 0.01$) (Fig. 7). To better understand how an alteration in expression of fimbriae *Yad* could influence motility or flagellar protein expression in the mutant strain, qRT-PCR analysis was performed to quantify the relative transcription of structural and regulator flagellar genes in all studied strains.

The obtained results showed that the expression levels of *flgE* and *fliC* (structural genes; class II and III, respectively) were slightly decreased in the $\Delta yadC$ strain compared to the wild-type strain. On the other hand, the expression of *flhD* (global regulator; class I) and *ompR* (motility regulator) was slightly increased in the mutant strain (see Fig. S2 in the supplemental material).

DISCUSSION

Escherichia coli's ability to establish colonization in several different niches and to infect the host's surfaces depends first on its ability to adhere, which prevents physical clearance mechanisms and engages the bacteria in a further colonization process. Many putative adhesins have been reported and their roles studied in distinct *E. coli* pathotypes. The *Yad*-encoding gene cluster has been associated with adhesion in an O157:H7 *E. coli* strain (12). In the present work, we showed the role of this gene in an avian pathogenic *E. coli* (APEC) strain.

Because *yadC* was upregulated both at the avian host regular temperature and in *in vivo* assays, it could play an important role in SCI-07 virulence. The high expression of *yadC* in lungs indicates that the fimbria *Yad* plays a role in APEC during the initial

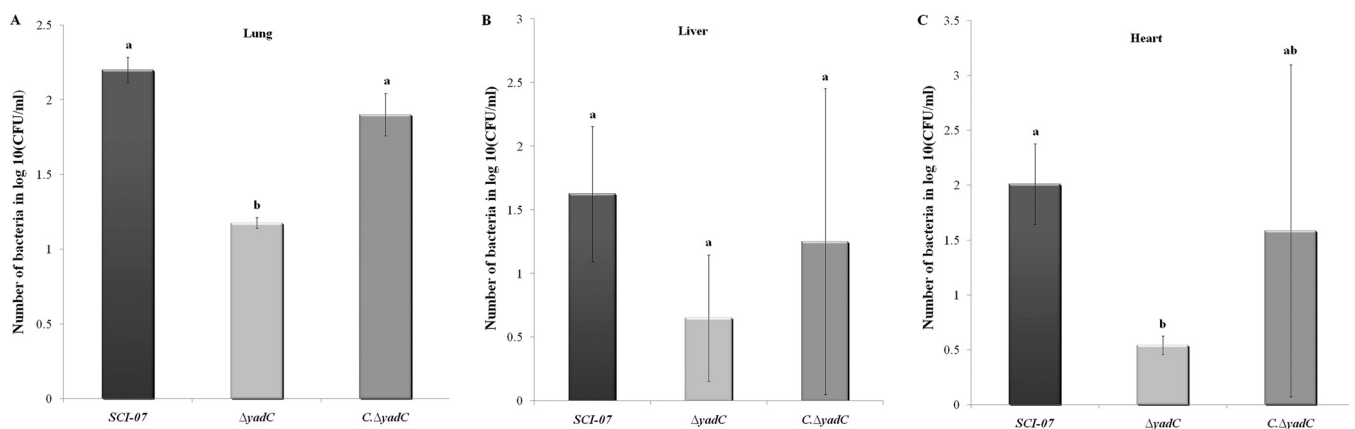


FIG 3 Bacterial recovery of wild-type (SCI-07), mutant ($\Delta yadC$), and complemented (C. $\Delta yadC$) strains from the lungs (A), liver (B), and heart (C) after infection with 10^8 CFU of bacteria. Different letters indicate statistical difference ($P < 0.05$).

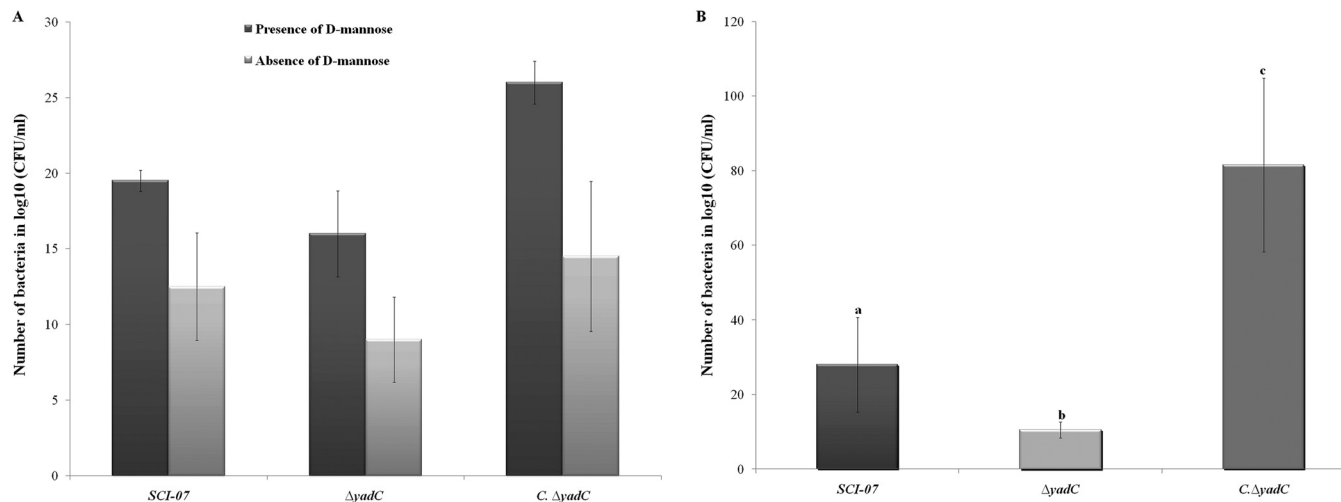


FIG 4 (A) Adhesion of wild-type (SCI-07), mutant ($\Delta yadC$), and complemented ($C.\Delta yadC$) strains to HeLa cells. Although adhesion levels were decreased, no statistical differences were found among the wild-type, mutant, and complemented strains. (B) Invasion of wild-type, mutant, and complemented strains into Hep-2 cells. Different letters indicate statistical difference ($P < 0.05$).

steps of the infection. Moreover, this effect would not be a consequence of an increased number of bacteria once the mutant presented the same bacterial growth as the wild-type and complemented strains.

In a previous work, the biofilm formation capacity associated with the fimbria Yad was more pronounced at lower (30°C) than higher (37°C) temperatures, with an assumed constant level of Yad expression (11). This would indicate that the fimbria Yad works better at lower than higher temperatures, similarly to the temperature-sensitive hemagglutinin (Tsh) (21). The *in vitro* up-regulation at higher temperatures of a *yad* gene cluster component

(*yadC*) found in the present work was also detected in other *yad* components (*ecpD-htrE*) in another work (22). Admitting that biofilm formation is a pathogenicity-related process, the higher transcription of an assumed less effective product could be a way of balancing the ratio of efficiency to quantity in order to keep a constant final function.

In the current proposed model of avian colibacillosis pathogenesis, the respiratory tract is colonized first, with subsequent spread of the pathogenic bacteria to the inner organs (23). The higher *yadC* expression in lungs than in spleen suggests that this adhesin would play a more intense role in the earlier steps of the infection mainly influencing the adhesion process. Moreover, the

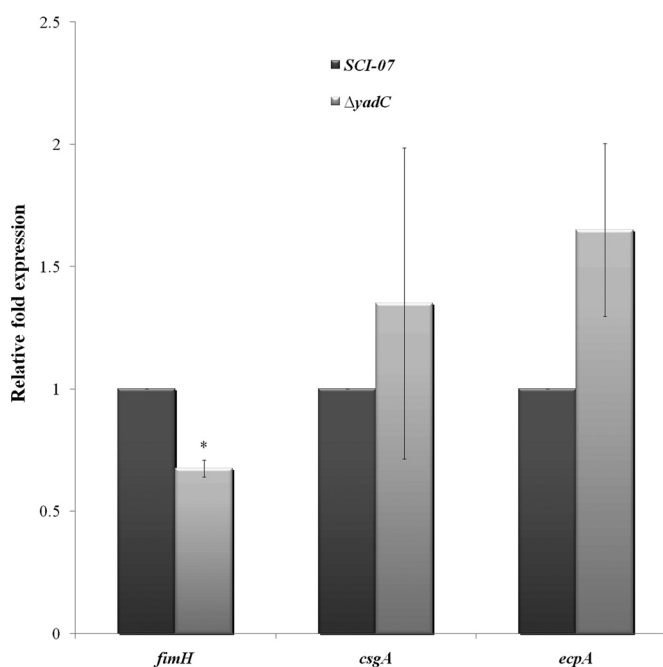


FIG 5 Relative fold expression of fimbrial *fimH*, *csgA*, and *ecpA* genes as determined by qRT-PCR in wild type (SCI-07) and mutant ($\Delta yadC$) strains. An asterisk indicates statistical significance at $P < 0.05$.

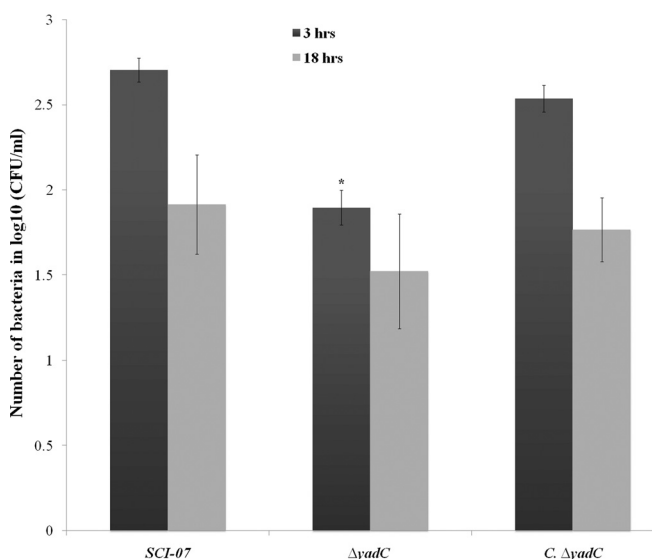


FIG 6 Bacterial survival in avian macrophage cells (HD11) infected with the wild-type (SCI-07), mutant ($\Delta yadC$), and complemented ($C.\Delta yadC$) strains. Infected macrophages were lysed at 3 or 18 h postinfection, and the numbers of bacteria were determined in triplicate. An asterisk indicates statistical significance at $P < 0.05$.

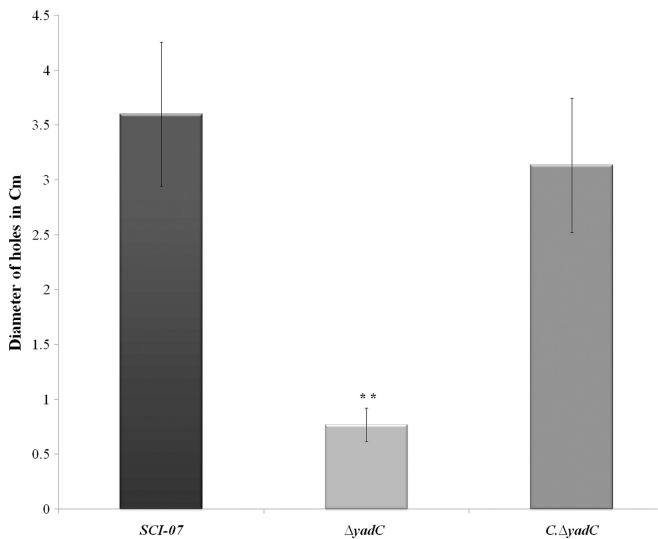


FIG 7 Motility assay in soft agar with the wild type (SCI-07), mutant ($\Delta yadC$), and (C. $\Delta yadC$) complementation strains. The presence of two asterisks indicates statistical significance at $P < 0.01$.

decreased ability of the $\Delta yadC$ mutant strain to survive in macrophages at 3 hpi indicates that the presence of these fimbriae in the wild-type strain would hamper the initial macrophage recognition, facilitating bacterial spreading inside the host body and further septicemic development of the infection.

Although several adhesins are associated with APEC (e.g., type 1 fimbria, curli, and ECP) (23), the complete mechanisms involved in APEC pathogenesis are poorly understood. The roles of Yad-mediated adhesion seem to be distinct between pathogenic and nonpathogenic *E. coli*. In the laboratory strain K-12, Yad does not promote adherence to A549, T24, HeLa, and Henle cells but increases biofilm formation (11). On the other hand, Yad is required for adherence to bladder epithelial cells (UM-UC-3) in uropathogenic *E. coli* (UPEC) (10) and to Caco-2 cells in EHEC under stress conditions (12). Many APEC and UPEC strains share genetic determinants (24) that led to a specialization in causing extraintestinal diseases. The Yad fimbria of the APEC strain tested in this work followed the same trend found with UPEC, contributing to adhesion. Due to the crucial role of adhesion for bacterial colonization and disease progression, it is possible that pathogenic *E. coli* presents further traits absent in commensal strains that mediate Yad-associated adhesion.

The role of *E. coli* adhesins in the internalization into eukaryote cell lines is under debate. Expression of curli fimbriae at high levels was able to mediate the internalization of a uropathogenic *E. coli* strain into bladder cell lines (25). Also, the expression of adhesin AfaD/DraD was suggested to partially mediate the internalization of bacteria into HeLa cells (26). This work showed that the mutant $\Delta yadC$ strain was less invasive in human epithelial cells (Hep-2). It is not possible to confirm if Yad has a crucial role in internalization or if the increased invasion is secondary to increased adhesion.

After physical barriers, bacterial pathogens must overcome the innate host defense in order to produce disease. In this context, the ability to survive in macrophages is an advantage for pathogenic *E. coli*. It was previously demonstrated that the deletion of a gene encoding a DNA-binding protein of APEC (RstA) had a role in diminishing the ability to survive in avian macrophage cell lines

(HD11) (27). The present work showed that Yad increased the survival in macrophages 3 hpi but not 18 hpi. This fact was expected since the battle between APEC and avian macrophages is supposed to take place at the beginning of the infection.

Flagellar motility can be modulated by fimbrial expression. In UPEC, it was shown that a mutant lacking genes encoding type 1 (*fim*) and P (*pap*) fimbriae presented slight decreases in motility compared to the wild-type strain (28). This was similar to the present work, where the $\Delta yadC$ strain was less motile than the wild type. However, the decreased motility of the mutant strain was not accompanied by a significant downregulation of flagellar genes. A similar paradox was observed with an enterotoxigenic *E. coli* (ETEC) strain. In that case, an isogenic $\Delta faeG$ strain (*faeG* encodes the major subunit of F4 fimbriae) did not present a different degree of motility but presented *fliC* (encoding the major flagellin protein) upregulated compared to the wild type (29). This indicates that the level of flagellar gene transcription does not always correspond to the degree of bacterial motility.

Taken together, these results suggest that the fimbrial adhesin YadC plays a role in adhesion, internalization, and motility (all of them basic biological bacterial characteristics) of the APEC strain here studied and contributes to its pathogenicity, particularly in the beginning of the infection process.

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