

## ***Escherichia coli* type III secretion system 2 (ETT2) is widely distributed in avian pathogenic *Escherichia coli* isolates from Eastern China**

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### **SUMMARY**

Pathogens utilize type III secretion systems to deliver effector proteins, which facilitate bacterial infections. The *Escherichia coli* type III secretion system 2 (ETT2) which plays a crucial role in bacterial virulence, is present in the majority of *E. coli* strains, although ETT2 has undergone widespread mutational attrition. We investigated the distribution and characteristics of ETT2 in avian pathogenic *E. coli* (APEC) isolates and identified five different ETT2 isoforms, including intact ETT2, in 57.6% (141/245) of the isolates. The ETT2 locus was present in the predominant APEC serotypes O78, O2 and O1. All of the ETT2 loci in the serotype O78 isolates were degenerate, whereas an intact ETT2 locus was mostly present in O1 and O2 serotype strains, which belong to phylogenetic groups B2 and D, respectively. Interestingly, a putative second type III secretion-associated locus (*eip* locus) was present only in the isolates with an intact ETT2. Moreover, ETT2 was more widely distributed in APEC isolates and exhibited more isoforms compared to ETT2 in human extraintestinal pathogenic *E. coli*, suggesting that APEC might be a potential risk to human health. However, there was no distinct correlation between ETT2 and other virulence factors in APEC.

**Key words:** Avian pathogenic *Escherichia coli*, distribution, *Escherichia coli* type III secretion system 2.

### **INTRODUCTION**

Many pathogens utilize type III secretion systems (T3SSs) to deliver effectors into eukaryotic cells, which facilitates infections [1, 2]. Most bacteria contain only one T3SS, but some pathogens harbour multiple T3SSs, which function independently in different aspects of pathogenesis. *Salmonella* utilize the *Salmonella* pathogenicity island 1 (SPI-1) T3SS to invade host cells, while the SPI-2 T3SS is required for intracellular survival [3]. Intestinal pathogenic *Escherichia coli*, such as enteropathogenic *E. coli*

(EPEC) and enterohaemorrhagic *E. coli* (EHEC), use the locus of enterocyte effacement (LEE)-encoded T3SS to cause attaching/effacing lesions and diarrhoeal disease [4]. A second T3SS, designated the *E. coli* type III secretion system 2 (ETT2), was identified by sequencing the genome of an EHEC O157 strain [5]; however, the role of ETT2 in *E. coli* infections is less clear.

ETT2 is unable to encode a functional secretion system in most *E. coli* because it has undergone widespread mutational attrition, but it plays a role in regulating bacterial virulence [6, 7]. ETT2 can affect the expression of virulence genes outside the ETT2 cluster, and it is indirectly involved in the virulence of intestinal pathogenic *E. coli* [EHEC and enteroaggregative *E. coli* (EAEC)] [8, 9]. However, ETT2 is

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crucial for the virulence of extraintestinal pathogenic *E. coli* (ExPEC) strains. The ETT2 locus in avian pathogenic *E. coli* (APEC), although degenerate, contributes to virulence and reduced serum survival [10]. Moreover, some evidence suggests that intact ETT2 might deliver effector proteins into human brain microvascular endothelial cells (HBMECs) to enhance bacterial invasion, intracellular survival and virulence of newborn meningitis *E. coli* (NMEC) strains [11].

ETT2 is known to be present, in whole or in part, in the majority of *E. coli* strains; it is most frequent in intestinal pathogenic *E. coli* strains [12–14], but of low prevalence in uropathogenic *E. coli* (UPEC) strains [15]. ExPEC strains (NMEC, UPEC, APEC) share a broad range of similar virulence factors and pathogenic mechanisms and APEC is thought to be a potential reservoir for human ExPEC [16–18]. Thus, this study investigated the prevalence and characteristics of ETT2 in APEC isolates to elucidate the potential zoonotic risk of APEC.

## MATERIALS AND METHODS

### Bacterial strains, growth conditions and DNA preparation

A strain collection containing 245 APEC isolates obtained from chickens, ducks or geese with typical colibacillosis symptoms in Eastern China was screened for the ETT2 locus (Supplementary Table S1). The ETT2-positive EHEC O157 strain ATCC43999 and the ETT2-negative strain DE719 [19] were used as controls in polymerase chain reaction (PCR) amplifications. All *E. coli* isolates were routinely grown in Luria–Bertani broth at 37 °C with aeration. Bacterial genomic DNA was prepared using the TIANamp Bacterial DNA kit (Tiangen, China) according to the manufacturer's guidelines.

### Serotyping

The serotype of each APEC isolate was identified by an allele-specific PCR assay as described previously [20] and confirmed by agglutination with specific hyperimmune rabbit antisera to O1, O2 and O78 (Statens Serum Institut, Denmark) according to the manufacturer's guidelines.

### Phylogenetic grouping

All isolates were classified into phylogenetic groups (A, B1, B2, D) according to the presence of the

*chuA* and *yjaA* genes, and an anonymous DNA fragment, TSPE4.C2 by a triplex PCR as described previously [21].

### Detection of ETT2

Isolates were tested for the presence of ETT2 genes by eight tiling-path PCR (TP-PCR), consisting of several interlocking, PCR-based approaches [6]. A series of primers was designed to amplify long fragments that spanned the entire ETT2 region in EHEC O157:H7 (GenBank accession no. NC002695.1) (Table 1). The structure of ETT2 cluster of EHEC O157:H7, EAEC 042 and NMEC CE10 are displayed in Figure 1a. If the long-fragment PCR was negative, a relevant short overlap PCR with the same primers and/or a deletion-scanning, long-fragment PCR employing primers flanking the lost segments was performed (Fig. 1b). The distribution of the *eip* locus, which encodes homologues of SPI-1 translocators and additional T3SS-related proteins, was determined by PCR with the respective primers (Table 1).

Long-fragment PCRs were performed using LA *Taq* DNA polymerase (TaKaRa, China). For short-fragment PCRs, 1 µl template DNA was added to reaction mixtures (25 µl) containing 2.5 µl of 10× PCR buffer with MgCl<sub>2</sub> (25 mM), 1.5 U *Taq* DNA polymerase (TaKaRa), 2 µl dNTPs (2.5 mM for each dNTP), and 0.5 µl (10 µM) of each primer pair. The PCR mixtures were subjected to the following conditions in an ABI Thermal Cycler (ABI Biosystems, USA): pre-denaturation at 95 °C for 5 min, followed by 30 cycles of 95 °C for 35 s, 50 °C for 30 s and 72 °C for different times, followed by a final extension at 72 °C for 10 min. Following electrophoresis, PCR products were identified by UV illumination, sequenced, and submitted to GenBank.

### Virulence genes

Isolates were screened by a previously described multiplex PCR protocol for the virulence genes *fimC*, *papC*, *tsh*, *mat*, *irp2*, *fyuA*, *iroN*, *iucD*, *iss*, *cvalevi*, *neuC*, *ompA*, *vat*, *ibeA* and *ibeB* [22].

## RESULTS

### Identification and characterization of the ETT2 locus

In total, 57.6% (141/245) APEC isolates were positive for ETT2 gene sequences (Table 2). TP-PCR amplification and sequencing identified five different ETT2

Table 1. Primers used in the present study

Primers	Sequences (5'–3')	Target genes	PCR product size
EC ETT2a-F	ACGCTGATCGTGGGTATCCT	<i>yqeG</i>	3118 bp
EC ETT2a-R	CTCACGATCTGTCAATGATGAG	<i>yqeK</i>	
EC ETT2b-F	GAGTCGAACTCGACATTCATC	<i>yqeK</i>	3584 bp
EC ETT2b-R	CTGGGTTCGAGTTGTAGTGA	<i>ygeH</i>	
EC ETT2c-F	CTGATAACTGGTCTGTCTGGAC	<i>ygeH</i>	3907 bp
EC ETT2c-R	TGCAGATGCACTCGTTTCATC	<i>eprK</i>	
EC ETT2d-F	ACTATCGCCAGTATCAACATCG	<i>eprK</i>	3894 bp
EC ETT2d-R	CTCATACTGTATTGTCTCGGCAG	<i>epaS</i>	
EC ETT2e-F	TGTGAGTAGGGTTGGCAATC	<i>epaS</i>	3310 bp
EC ETT2e-R	GATAGCAGAAGAGAGTGGCAG	<i>epaO</i>	
EC ETT2f-F	GAGCTGGACTCAATACTCCATC	<i>epaO</i>	2872 bp
EC ETT2f-R	GCAGAACTTGACGACGTTTCTG	<i>eivC</i>	
EC ETT2 g-F	CCTGGCTATACGTTTCATTTGC	<i>eivC</i>	6321 bp
EC ETT2 g-R	AGGTCTGGTTGTGACATCGAC	<i>ECs3735</i>	
EC ETT2 h-F	GTCGGATTACGAATCATGAGA	<i>ECs3735</i>	2553 bp
EC ETT2 h-R	GGGATTGATATCTCAGCTCCAC	<i>ECs3738</i>	
EC eicA-F	CAGAGCTGCATGGCATTAC	<i>eicA</i>	369 bp
EC eicA-R	CAAGATAAGCAGTAGCCATCTC	<i>eicA</i>	
EC eipB-F	ACAGCCATTATGGATGCTGA	<i>eipB</i>	610 bp
EC eipB-R	TCGTATCTTGATCGACACCA	<i>eipB</i>	
EC eipX-F	CAAGCAGTTCTACAGAGCAGAC	<i>eipX</i>	738 bp
EC eipX-R	GATCCAGATAGCTACTCATAACTG	<i>eipX</i>	
EC eipD-F	GAGTTCTGCACCTGAAGTGAG	<i>eipD</i>	557 bp
EC eipD-R	CTGAAGACTGACTCATACTGTC	<i>eipD</i>	
EC eilA-F	GTACCTATTGTGATTGTGACCG	<i>eilA</i>	465 bp
EC eilA-R	CATAACAGCAGAATCCAGAGA	<i>eilA</i>	
EC eaeX-F	CGTTACTGTGACGGTTAATGG	<i>eaeX</i>	478 bp
EC eaeX-R	GACGATTGCCAGTACCAATCAG	<i>eaeX</i>	

isoforms among the isolates (GenBank accession nos. KU684467–KU684471). Sequence analysis indicated that the intact ETT2 gene cluster in APEC was similar to that of NMEC CE10, which was designated the type A isoform. Types B, D and E isoforms were characterized by 4.99-, 5.68- and 8.74-kb gene deletions, respectively, in the right end of the ETT2 island. Specifically, there was a 1.3-kb transposase insertion in the transcriptional regulation gene *ygeH*, and a 4.99-kb gene deletion in the ETT2 island of the type C isoform strains (Fig. 1c). Of the five different ETT2 isoforms detected, type E (51.1%, 72/141) was the most widely distributed, followed by type C (14.9%, 21/141) and type B (12.8%, 18/141).

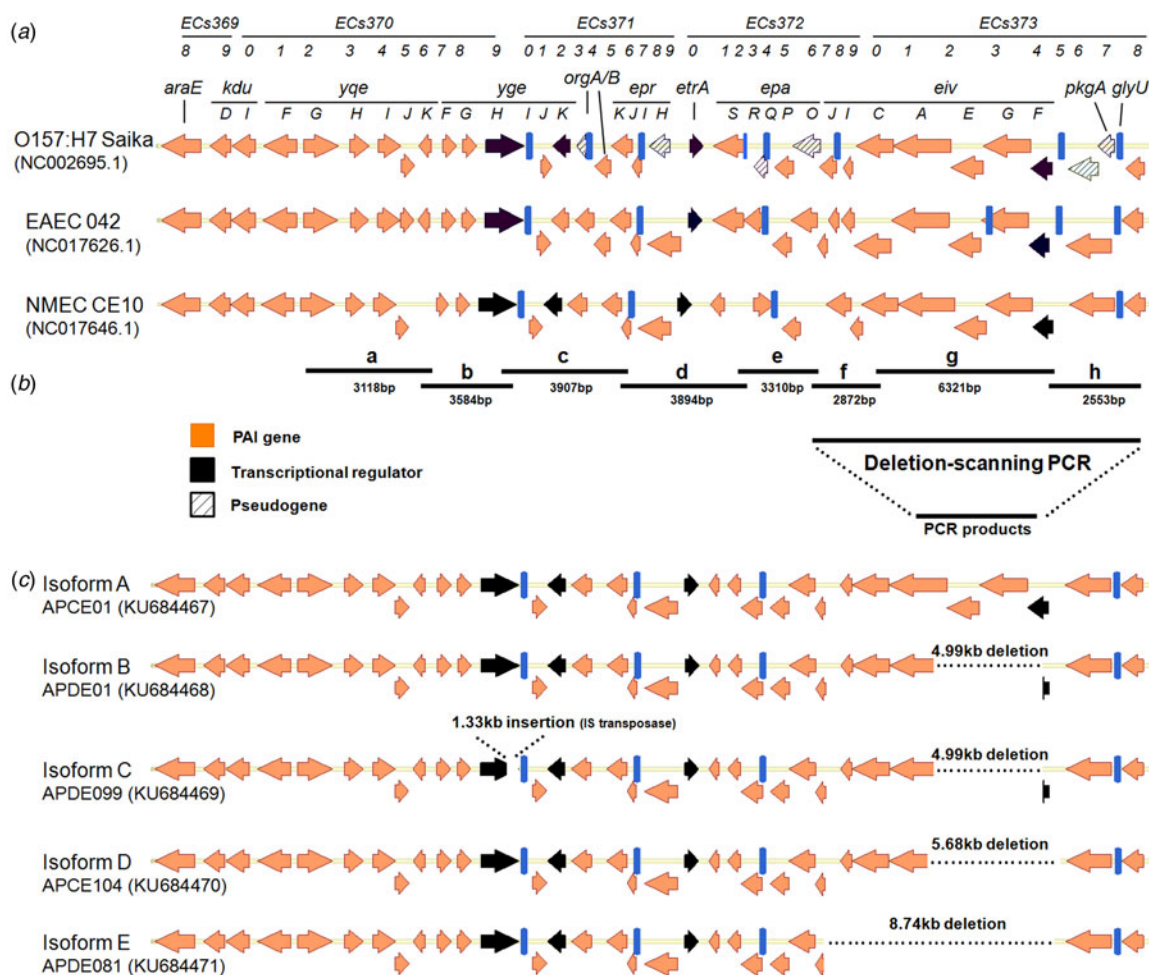
#### ETT2 locus in predominant APEC serotypes

As shown in Table 2, 46.9% (115/245) of the isolates were grouped as serotype O78, 22.5% (55/245) as serotype O2, and 12.7% (31/245) as serotype O1; the remaining 44 isolates (18%), were not typable with these three antisera and were classified as 'Non-O1/

O2/O78' serotypes. The majority (64.4%, 74/115) of the O78 serotype isolates were positive for ETT2, as were 58.2% (32/55) of serotype O2, and 48.4% (15/31) of serotype O1. Correspondingly, of the 141 APEC isolates positive for ETT2, 52.5% (74) were serotype O78, 22.7% (32) serotype O2, and 10.6% (15) serotype O1. The remaining 20 (14.2%) isolates belonged to 'Non-O1/O2/O78' serotypes. An intact ETT2 island (type A) was found only in serotypes O1 and O2 and notably the O78 isolates always contained deleted isoforms of ETT2.

#### Phylogenetic groups D and B2 APEC, and ETT2 cluster

According to the Clermont *E. coli* phylotyping method [21], 78 (31.8%) of the APEC isolates belonged to phylogenetic group B2, while 54 (22%), 52 (21.2%) and 48 (19.6%) of the isolates fell in groups D, A and B1, respectively; 13 (5.8%) of the isolates could not be classified into a phylogenetic group (Table 2). A comparison of the distribution of the



**Fig. 1.** ETT2 structures and lengths of TP-PCR products. (a) Structure of the ETT2 pathogenicity island in EHEC O157:H7, EAEC 042 and NMEC CE10. Homologous genes are vertically aligned with the transcriptional regulator encoding gene shown as black and the pseudogene as a grid. (b) Bold lines indicate the size of the PCR products, and deletion-scanning PCR. (c) Length of gene deletion or insertion of ETT2 isoforms in APEC isolates.

**Table 2.** Distribution of the ETT2 isoforms in APEC isolates

	ETT2 positive ( <i>n</i> = 141), isoforms					ETT2 negative ( <i>n</i> = 104)	Total ( <i>n</i> = 245)
	A ( <i>n</i> = 17)	B ( <i>n</i> = 18)	C ( <i>n</i> = 21)	D ( <i>n</i> = 13)	E ( <i>n</i> = 72)		
<b>Serotypes</b>							
O1	9	2	1	1	2	16	31
O2	5	4	2	9	12	23	55
O78	0	10	16	2	46	41	115
Non-O1/O2/O78	3	2	2	1	12	24	44
<b>Phylogenetic groups</b>							
A	1	8	11	0	11	21	52
B1	1	0	4	2	25	16	48
B2	5	5	2	7	16	43	78
D	9	3	1	4	18	19	54
Unknown	1	2	3	0	2	5	13

ETT2 cluster among phylogenetic groups revealed that 66.7% (32/48), 64.8% (35/54), 59.6% (31/52) and 44.9% (35/78) of the group B1, D, A and B2 isolates, respectively, and 61.5% (8/13) of the unknown group were positive for the ETT2 island. For the 17 isolates that had an intact ETT2 (type A isoform), nine fell in phylogenetic group D, five in B2, whereas types B and C isoforms were predominant in phylogenetic group A isolates, and type E in group B1.

### Type III secretion-associated locus (*eip* locus), virulence genes and ETT2 locus

According to previous studies, the *eip* locus within the EAEC 042 genome is associated with the *sipABCD* homologue cluster of the ETT2 locus [6, 23]. Here, the *eip* locus was only found in APEC strains that possess an intact ETT2 locus and, moreover, was located between the *E. coli* *yicM* and *nlpA* genes (data not shown), which is in the same position as that in EAEC 042 and NMEC CE10.

There was no correlation between the presence or distribution of other virulence genes and the ETT2 locus (Supplementary Table S1).

## DISCUSSION

Pathogenic bacteria utilize several common strategies to overcome host defences and facilitate infections. T3SSs, which deliver effector proteins into host cells, are important virulence mechanisms in Gram-negative bacteria as they interfere with specific cellular and host immune responses to promote bacterial survival and enhance pathogenicity [1, 2]. Two different T3SSs have been identified in *E. coli*. The LEE-encoded T3SS (ETT1) is well-characterized in EHEC and EPEC strains [4], whereas ETT2 is present, in whole or part, in the majority of *E. coli* strains, including many commensal strains [6]. However, until now, no prevalence data for ETT2 in APEC isolates have been published.

The present study shows that over half (57.6%) of 245 APEC isolates harboured the ETT2 locus. We identified five different ETT2 isoforms in these isolates. Sequence analysis showed that there are gene deletions in most APEC strains (types B, C, D, E ETT2 isoforms), indicating that ETT2 has undergone widespread mutational attrition. Similar mutations were also found in other *E. coli* pathovars [12, 13]. A serotyping analysis indicated that ETT2 was present in the predominant APEC serotypes O78, O2 and O1

with over half (52.5%) of the isolates being O78. Further, all the ETT2 elements in isolates of this serotype were variants (not intact), which is consistent with previous studies [10]. Indeed, an intact ETT2 locus was mostly present in serotypes O1 and O2 isolates, which also harboured the *eip* cluster. Further analysis indicated that an intact ETT2 gene cluster was predominant in phylogenetic groups D and B2, as reported previously [6], but by contrast, the distribution of ETT2 mutational isoforms varied.

Our results show that the APEC isolates studied harboured all of the ETT2 isoforms found in human ExPEC (UPEC and NMEC) with a similar frequency as reported for human diarrhoeagenic *E. coli* (65%), Shiga toxin-producing *E. coli* (STEC) (100%), as well as isolates from piglets (85.9%) and cows (47.4%), but was markedly higher than that in UPEC (3%) [7, 12–14]. Published studies suggest that ExPEC (APEC, NMEC, UPEC) strains possess a similar broad range of virulence factors and pathogenic mechanisms [16, 17]. More importantly, the predominant APEC serotypes O1, O2 and O78 have also been implicated in various human diseases, including newborn meningitis and urinary tract infections, indicating that poultry may be a vehicle or even a reservoir for human ExPEC (UPEC and NMEC) [16–18, 24]. Bacteria could acquire a combination of mobile genetic elements, such as the ETT2 locus, through some mechanism of horizontal gene transfer, and become a highly adapted pathogen [25]. The 1.3-kb IS transposase was found in type C isoform ETT2 strains, which might facilitate the transfer of ETT2 among *E. coli* strains.

ETT2 is reported to be associated primarily with the LEE, the high pathogenicity island, the locus of proteolytic activity or Stx2e in intestinal pathogenic *E. coli* (EHEC and STEC) strains [7, 13, 26]. We investigated the association of ETT2 with the distribution of 15 other APEC virulence factors, but failed to find any correlation between them, which possibly indicates that the ETT2 cluster occurs independently of other virulence factors in these strains, perhaps due to different pathogenicity mechanisms and virulence factors.

Characterized T3SS needle complexes are composed of inner and outer membrane rings and a hollow needle. In most *E. coli* strains, ETT2 is highly variable and contains mutations, which leads to an incomplete and non-functional T3SS apparatus [6, 12, 13]. Although degenerate, ETT2 still plays a crucial role in *E. coli* virulence. In EHEC O157:H7,

ETT2 regulates virulence gene expression and protein secretion by the LEE [8]. The EilA regulator of EAEC 042 coordinately activates the expression of effectors and influences bacterial adherence and biofilm formation, which are indirectly involved in virulence [9]. A degenerate ETT2 in APEC contributes to serum survival and virulence. However, it is not involved in the secretion of effectors. The inactivation of ETT2 affects bacterial surface properties, such as pellicle formation, which might decrease bacterial survival during an infection [10].

Intact ETT2 has been shown to be crucial to bacterial virulence and plays a role in invasion, intracellular survival in HBMECs, and virulence of NMEC [11]. Moreover, in the latter study, the ETT2 deletion mutant exhibited the same levels of OmpA and K1 capsules as wild type, which affect the interaction of *E. coli* K1 with HBMECs, indicating that the ETT2 locus associated with NMEC CE10 has novel properties and mechanisms. It has been shown that the ETT2 T3SS not only secretes proteins that are encoded in the T3SS locus, but also those that are encoded outside of the locus [25]. A previous study [6] and our results show that the *eip* locus, which encodes homologues of SPI-1 translocators and T3SS-related proteins, was present in all APEC strains harbouring an intact ETT2 cluster. The *eip* locus might cooperate with ETT2 to construct an active T3SS apparatus. Indeed, *E. coli* DH5 $\alpha$  acquired the ability to secrete the EspB protein when complemented with ETT2 [27]. Additionally, putative T3SS effectors were discovered in EAEC 042, NMEC CE10 and APEC strains [11, 23]. An intact ETT2 may deliver an unidentified effector(s) into host cells to interfere with host defence and facilitate bacterial infection. The exact function of ETT2 remains unknown, and whether and how an intact ETT2 locus encodes a functional secretion system should be investigated in the future to help us prevent poultry colibacillosis and potential human infections [28].

## SUPPLEMENTARY MATERIAL

For supplementary material accompanying this paper visit <http://dx.doi.org/10.1017/S0950268816000820>.

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## DECLARATION OF INTEREST

None.

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