

Genomic Characterization and Transferability of bla_{CTX-M-55}-Carrying Plasmids in *Escherichia coli* from Malawian Broiler Chickens

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Introduction

- •The global spread of extended-spectrum β -lactamase (ESBL)-producing Escherichia coli poses a growing public health concern.
- •While poultry is recognized as a major reservoir of ESBLs, data on the mobility and genetic context of ESBL genes in African settings remain limited.
- •This study investigated the genomic characteristics and transferability of E. coli isolates carrying $bla_{CTX-M-55}$, a commonly detected ESBL genes globally, from Malawian broilers and compared their plasmids with those from other sectors.

Methodology

- •Whole-genome sequencing of 21 bla_{CTX-M-55} -positive *E. coli* isolates in Malawi was performed using short-read sequencing and nine isolates underwent long-read sequencing to resolve complete genome structures, including plasmids.
- •Bacterial conjugation assays were performed to evaluate plasmid transferability using an *E. coli* recipient.
- •Comparative genomic analyses were performed between plasmids in this study and reference plasmids from humans, animals, and environmental sources.

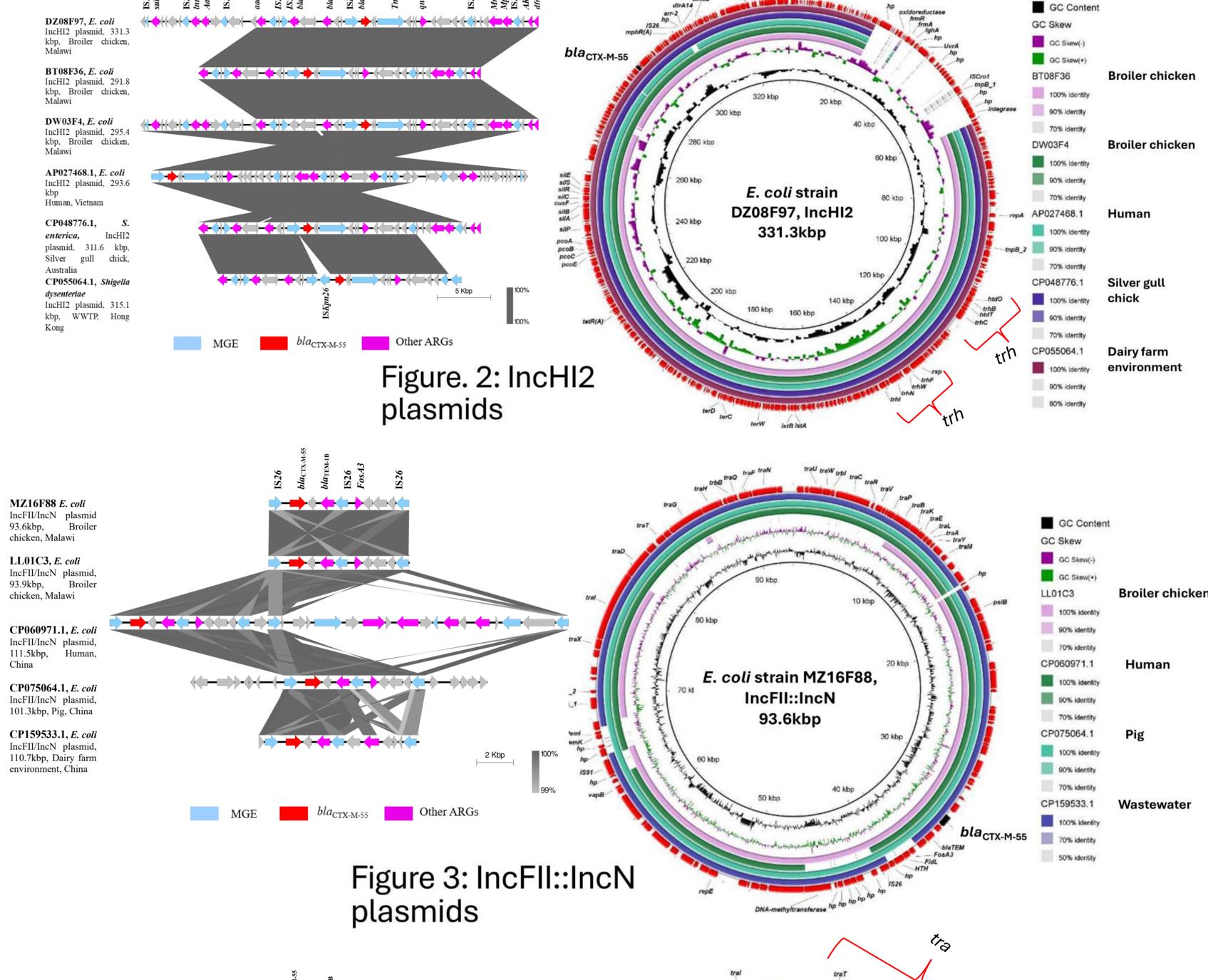
Results

Characteristics of *E. coli* plasmids carrying *bla*_{CTX-M-55} gene

Fifteen isolates were conjugative, including seven harbouring bla_{CTX-M-} 55 on diverse plasmid types: IncFIB (n=1), IncFII (n=1), IncFII::IncN (n=2), IncHI2 (n=3) (Table 1). Several antimicrobial resistance genes (e.g., sul2, tetA, and floR) were frequently identified.

Table 1: Genomic characteristics of *E. coli* plasmids carrying *bla*_{CTX-M-55}

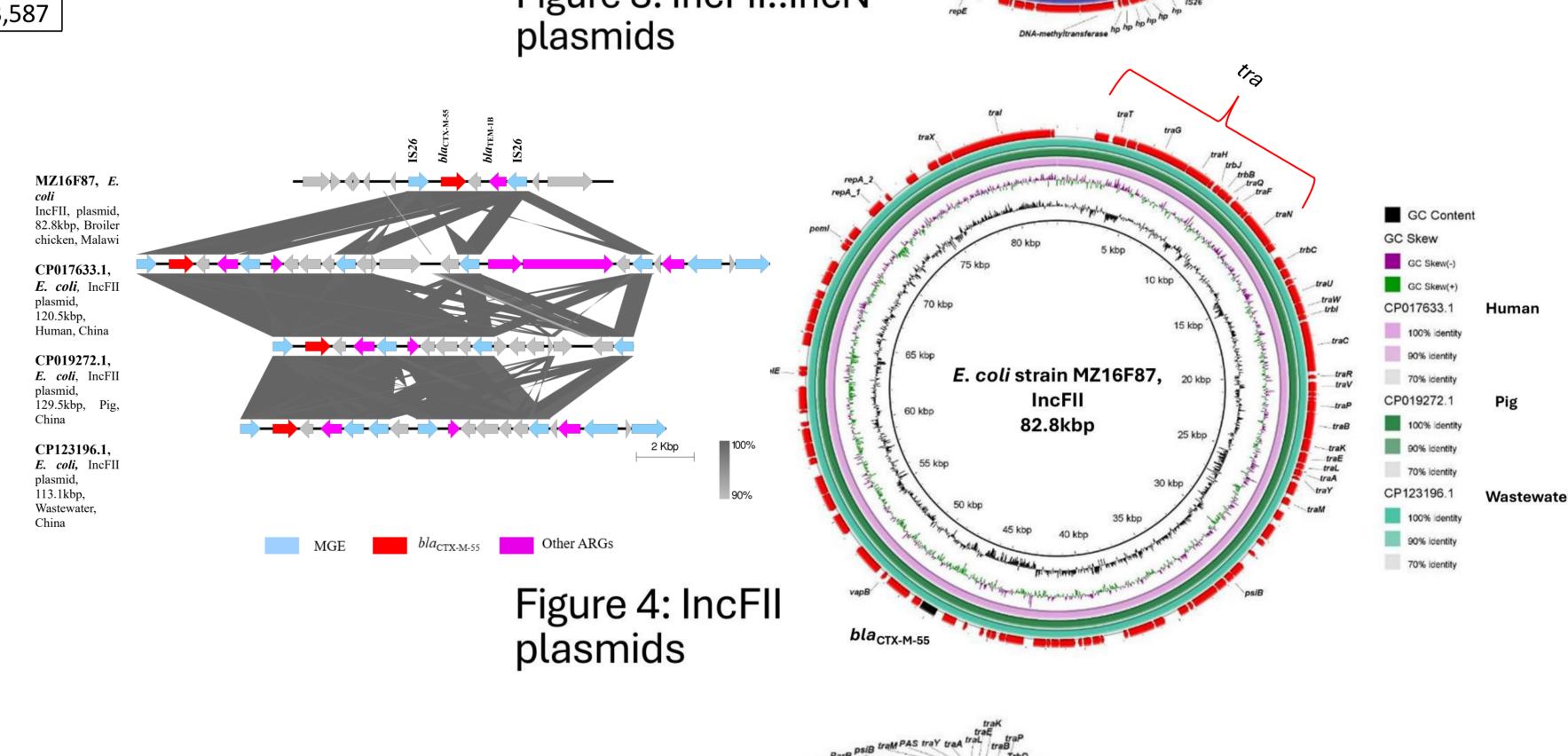
Plasmid	pMLST	Host range	Replicon	AMR	Size (bp)
				aac(3)-IId, aph(3')-Ia, aph(6)-Id, ARR-2, bla CTX-M-	
				55, bla LAP-2, bla TEM-1B, dfrA14, floR, mph(A),	
BT08F36	ST2	IV	IncHI2;IncHI2A	qnrS1, sul3, tet(A)	331,316
				aac(3)-IId, aadA22, aph(3')-Ia, aph(6)-Id, ARR-2,	
				bla CTX-M-55, bla LAP-2, bla TEM-1B, dfrA14, floR,	
DW03F4	ST2	IV	IncHI2;IncHI2A	Inu(F), mph(A), qnrS1, sul3, tet(A)	291,889
			IncFII(pHN7A8);	aph(3'')-Ib, aph(6)-Id, bla CTX-M-55, bla TEM-214,	
LL01C3	F33:A-:B-::repN	-	IncN	fosA3, sul2, tet(A)	93,869
				aac(3)-IId, aadA22, aph(3')-Ia, aph(6)-Id, ARR-2,	
				bla CTX-M-55, bla LAP-2, bla TEM-1B, dfrA14, floR,	
DZ08F97	ST2	IV	IncHI2;IncHI2A	Inu(F), mph(A), qnrS1, sul3, tet(A)	295,410
				aac(3)-IId, aadA2, aph(3'')-Ib, aph(3')-IIa,	
				aph(6)-Id, bla CTX-M-55, bla TEM-1B, bla TEM-216,	
LL12F51	X1	-	IncX1	dfrA12, floR, rmtB, sul1, sul2, tet(A)	46,794
				aac(3)-IId, aadA2, aph(3'')-Ib, aph(3')-Ia, aph(6)-	
			IncFIB(AP00191	Id, bla CTX-M-55, bla TEM-1B, bla TEM-214, dfrA12, floR,	
MZ16F86	F24:A-:B1::repN	Ш	8);IncFII	fosA3, mph(A), sul2, tet(A), tet(A)	170,682
MZ16F87	F33:A-:B-	Ш	IncFII(pHN7A8)	bla CTX-M-55, bla TEM-57, fosA3	82,812
			IncFII(pHN7A8);	bla bla focA2	
MZ16F88	F33:A-:B-	-	IncN	bla CTX-M-55, bla TEM-214, fosA3	93,587

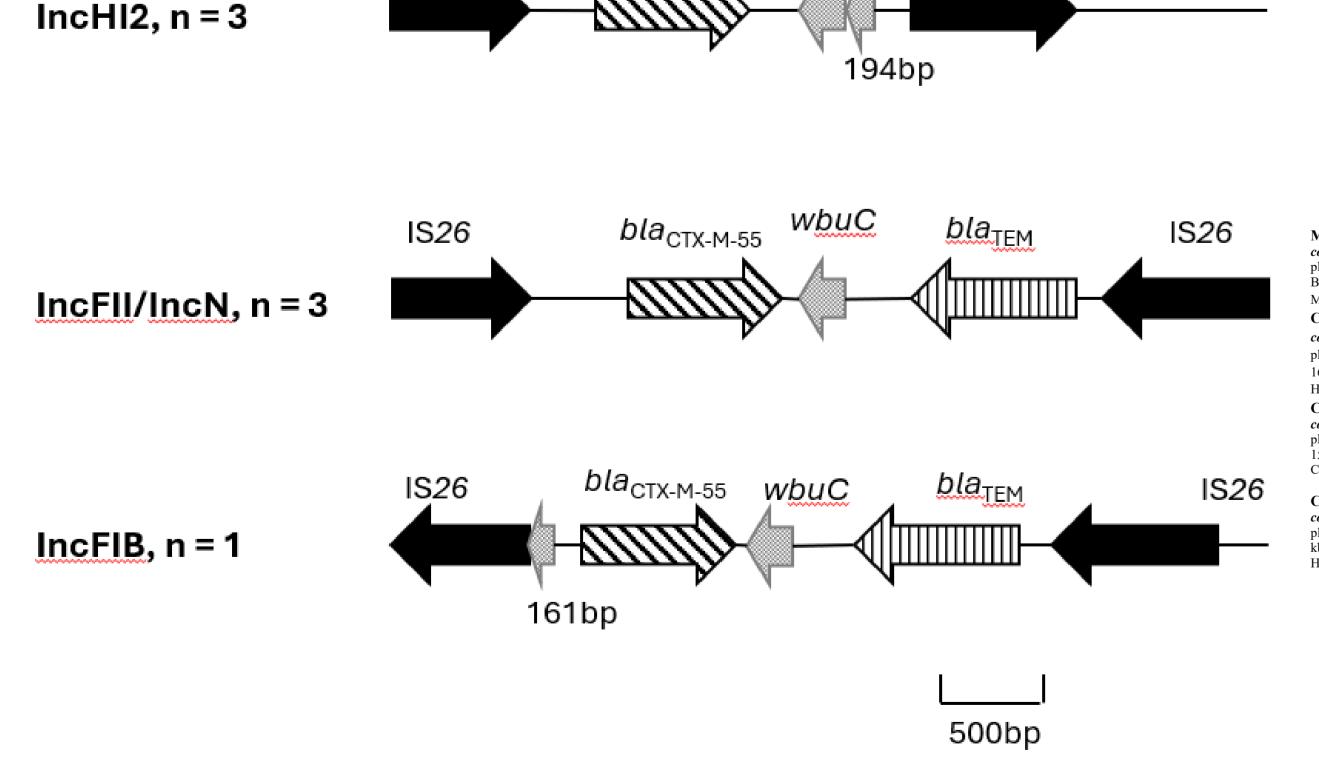


Plasmid structures and the genetic context of $bla_{CTX-M-55}$ gene

Conserved genetic structures flanking bla_{CTX-M-55} were observed across diverse plasmid types and sources (Figures 1 – 5). The aligned regions also included conjugation machinery genes like tra, trh involved in plasmid mobilization through horizontal gene transfer. Virulence genes (iro and sit) were detected only in IncFIB plasmid (Figure 5).

bla_{CTX-M-55} wbuC





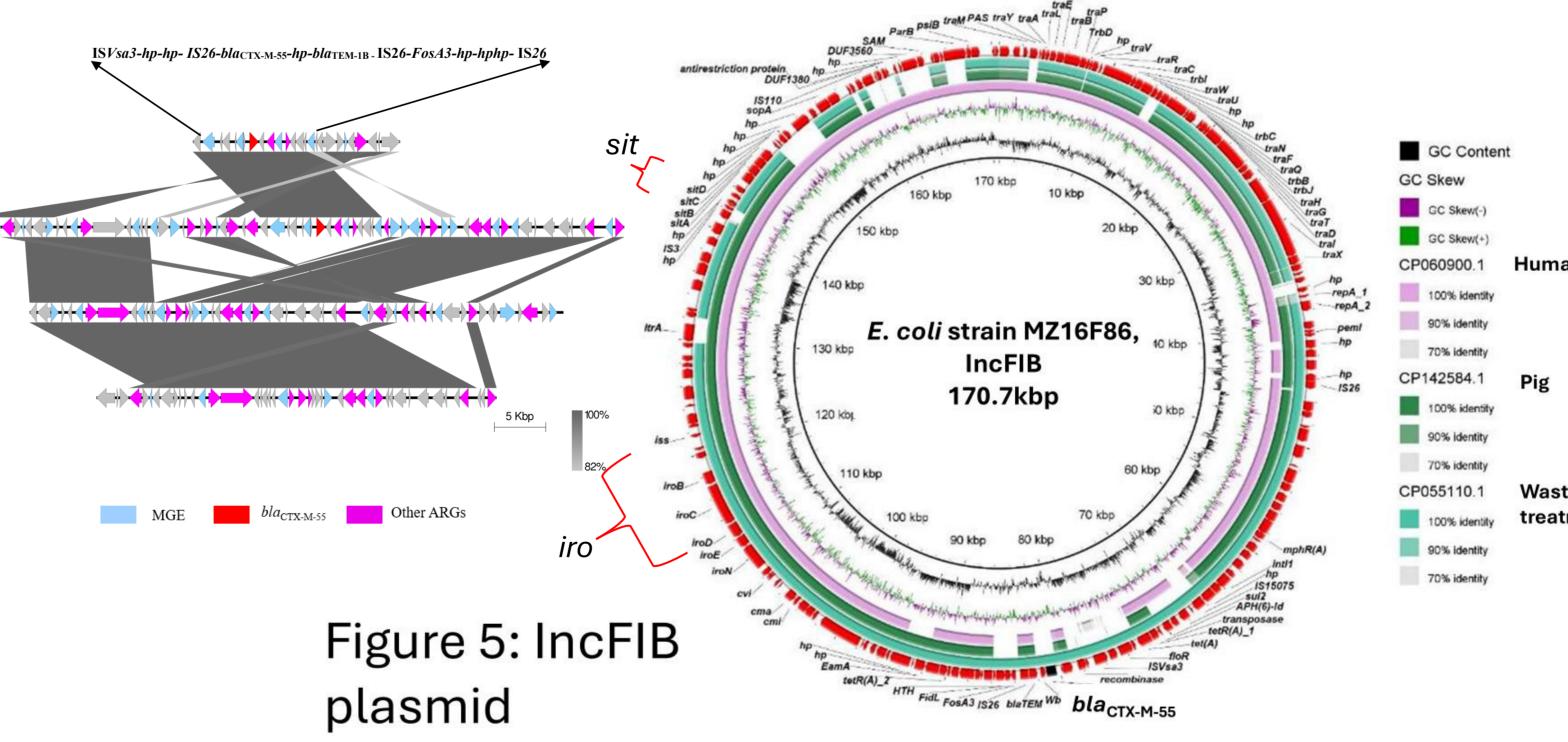


Figure 1: Genetic regions surrounding *bla*_{CTX-M-55}

ISEcp1

Conclusion

- •The presence of multiple incompatibility groups carrying $bla_{CTX-M-55}$ gene shows that it can be hosted by diverse plasmid backbones which increases its dissemination potential.
- •The observed similarities and transferability of poultry-derived plasmids indicate the potential for cross-sectoral transmission of $bla_{CTX-M-55}$ -carrying plasmids through horizontal gene transfer worldwide mediated by mobile genetic elements. These findings support the importance of the One Health approach in antimicrobial resistance surveillance.

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