

**Supplementary Table S1: Strains and plasmids used in this study.**

	<b>Description</b>	<b>Reference</b>
<b>Strain</b>		
E2348/69	EPEC serotype O127:H6	Levine et al 1978
E2348/69 $\Delta gspD$	E2348/69 $\Delta gspD$	Baldi et al 2012
E2348/69 $\Delta sslE$	E2348/69 $\Delta sslE$	Baldi et al 2012
E2348/69 $\Delta yacC$	E2348/69 $\Delta yacC$	This Study
E2348/69 $\Delta aspS$	E2348/69 $\Delta aspS$	This Study
E2348/69 $\Delta gspD \Delta yacC$	E2348/69 $\Delta gspD \Delta yacC$	This Study
E2348/69 $\Delta gspD \Delta aspS$	E2348/69 $\Delta gspD::Cm^r, \Delta aspS$	This Study
BL21 (DE3)	<i>E. coli</i> strain: F- <i>ompT hsdSB</i> (rB- mB-) <i>dcm gal</i> (DE3)	Invitrogen
BL21 (DE3) $\Delta gspD \Delta aspS$	BL21 (DE3) $\Delta gspD::Cm^r, \Delta aspS$	This Study
Rosetta (DE3)	<i>E. coli</i> strain: F- <i>ompT hsdSB</i> (rB- mB-) <i>dcm gal</i> (DE3)	Novagen
<b>Plasmid</b>		
pBAD24	<i>ori</i> pMB1, Amp <sup>r</sup>	Guzman et al 1995
pET DUET-1	<i>ori ColE1, lacI</i> gene, Amp <sup>r</sup> , used for assembly assays in BL21(DE3)	Novagen
pET DUET-1 GspD-C <sub>4</sub>	E2348/69 <i>gspD-C<sub>4</sub></i> was cloned into NcoI/HindIII sites of pET DUET-1	This Study
pET DUET-1 GspD-C <sub>4</sub> YacC	E2348/69 <i>yacC</i> was cloned into NdeI/XhoI sites of pET DUET-1 GspD-C <sub>4</sub>	This Study
pET DUET-1 GspD-C <sub>4</sub> AspS	E2348/69 <i>aspS</i> was cloned into NdeI/XhoI sites of pET DUET-1 GspD-C <sub>4</sub>	This Study
pFT-A	<i>ori</i> R101, <i>repA101ts, flp</i> , Amp <sup>r</sup>	Posfai et al 1997
pGEM-T-Easy	<i>ori</i> pMB1, Amp <sup>r</sup>	Promega
pKD3	FRT flanked Cm <sup>r</sup> gene, Cm <sup>r</sup> , Amp <sup>r</sup>	Datsenko and Wanner, 2000

pKD4	FRT flanked Kan <sup>r</sup> gene, Kan <sup>r</sup> , Amp <sup>r</sup>	Datsenko and Wanner, 2000
pKD46	λ Red recombinase, <i>ori</i> R101, <i>repA101ts</i> , Amp <sup>r</sup>	Datsenko and Wanner, 2000
pJP117	E2348/69 <i>gspD-C<sub>4</sub></i> is cloned into the <i>NcoI/XbaI</i> sites of pNM12 (pBAD24 modified to include <i>MscI</i> site). Used for assembly assays in E2348/69	This Study
pJP168	P <sub>BAD</sub> promoter and <i>araC</i> replaced by P <sub>tetA</sub> and <i>tetR</i>	Baldi et al 2012
pJP181	E2348/69 <i>gspD-C<sub>4</sub></i> is cloned into the <i>NcoI/HindIII</i> sites of pJP168. Used for sucrose fractionation in E2348/69.	This Study
pKV1066	H10407 <i>yghG</i> was cloned into a modified pET-22b(+) vector (Novagen) to encode <i>pelB</i> periplasmic signal sequence, His <sub>6</sub> tag and tobacco etch virus protease cleavage site; <i>ori ColE1</i> , <i>lacI</i> gene, Amp <sup>r</sup> ; used for binding assay with the secretin S-domain	This study
pKV1111	Sequence corresponding to 60 C-terminal residues of H10407 <i>gspD</i> (S-domain) was cloned into a pET-22b(+) vector to encode a fusion with <i>malE</i> gene; <i>ori ColE1</i> , <i>lacI</i> gene, Amp <sup>r</sup> ; used for binding assay with the pilotin	This study
pKV1112	VC395_1831 from <i>V. cholerae</i> 569B (corresponds to VC1703 from <i>V. cholerae</i> O1 biovar El Tor str. N16961) was cloned into a modified pET-22b(+) vector to encode <i>pelB</i> periplasmic signal sequence, His <sub>6</sub> tag and tobacco etch virus protease cleavage site; <i>ori ColE1</i> , <i>lacI</i> gene, Amp <sup>r</sup>	This study

## REFERENCES

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