

TABLE S5. Mutations and predictions of protein change on candidate genes to hypersusceptibility of the ancestral Escherich strain isolates

Genes	Gene products	Mutations at the protein level*				Predictions of functional effect			Frequency of the observed allele in Uniprot
		NCTC86	CIP61.11	ATCC4157	DSM301	SIFT <sup>‡</sup>	PolyPhen-2 <sup>§</sup>	PROVEAN <sup>§</sup>	
<i>acrA</i>	Sub-unit of efflux pump AcrAB-ToIC	-	-	-	-				
<i>acrB</i>	Sub-unit of efflux pump AcrAB-ToIC	-	-	K659N	K659N	0.20	Benign (0.001)	Deleterious (-2.666)	1/79
		G679S	G679S	G679S	G679S	0.00	Probably damaging (0.998)	Deleterious (-5.837)	0/79
<i>tolC</i>	Sub-unit of multi druf efflux systems	-	-	-	-				
<i>acrD</i>	Sub-unit of efflux pump AcrAD-ToIC	-	-	-	-				
<i>acrE</i>	Sub-unit of efflux pump AcrEF-ToIC	G191D	G191D	G191D	G191D	0.00	Probably damaging (1.000)	Deleterious (-6.973)	0/46
<i>acrR</i>	DNA binding transcriptional repressor	-	-	-	-				
<i>acrZ</i>	AcrB transmembran domain associated protein	-	-	-	-				
<i>acrS</i>	Regulator/repressor of efflux pump AcrEF-ToIC	W50R	W50R	W50R	W50R	0.00	Probably damaging (0.957)	Deleterious (-13.78)	0/42
<i>marA</i>	DNA binding transcriptional activator	-	-	-	-				
<i>marB</i>	DNA binding transcriptional activator	-	-	-	-				
<i>marR</i>	DNA binding transcriptional repressor	-	-	-	-				
<i>soxS</i>	DNA binding transcriptional regulator	-	-	-	-				
<i>soxR</i>	DNA binding transcriptional dual regulator	-	-	-	-				
<i>evgS</i>	Hybrid sensory histidine kinase in two-component regulatory system	F152L	F152L	F152L	F152L	0.02	Benign (0.006)	Neutral (0.414)	83/83
<i>evgA</i>	DNA binding transcriptional activator	-	-	-	-				
<i>phoP</i>	DNA binding transcriptional regulator	-	K149T	-	-	0.23	Benign (0.002)	Neutral (-1.112)	0/87
<i>phoQ</i>	Sensor protein	-	-	-	-				
<i>sdiA</i>	DNA binding transcriptional activator	-	-	-	-				
<i>rob</i>	DNA binding transcriptional activator	G5C	G5C	G5C	G5C	0.01	Possibly damaging (0.822)	Deleterious (-3.302)	0/86
<i>yajC</i>	Preprotein translocase subunit	-	-	-	-				
		I46M	I46M	I46M	I46M	0.22	Benign (0,000)	Neutral (0.893)	241/253
		V208I	V208I	V208I	V208I	1.00	Benign (0,000)	Neutral (0.899)	222/253
		S365A	S365A	S365A	S365A	1.00	Benign (0,000)	Neutral (2.181)	184/253
<i>mdtC</i>	Multidrug resistance protein	L387I	L387I	L387I	L387I	0.00	Benign (0,394)	Neutral (-1.572)	4/253
		G472V	G472V	G472V	G472V	0.18	Benign (0,000)	Neutral (3.512)	17/253
		L910M	L910M	L910M	L910M	0.18	Benign (0,000)	Neutral (0.013)	152/248
<i>emrA</i>	Sub-unit of efflux pump EmrAB-ToIC	-	-	-	-				
<i>emrB</i>	Sub-unit of efflux pump EmrAB-ToIC	-	-	-	-				
<i>emrY</i>	Sub-unit of efflux pump EmrYK-ToIC	-	-	-	-				
<i>emrK</i>	Sub-unit of efflux pump EmrYK-ToIC	-	-	-	-				
<i>ompA</i>	Outer-membran protein A (porin)	-	-	-	-				
<i>ompF</i>	Outer-membran protein F (porin)	E139G	E139G	E139G	E139G	0.04	Possibly damaging (0.893)	Deleterious (-4.851)	0/101
<i>ompC</i>	Outer-membran porin protein C	-	-	-	-				
<i>ompR</i>	Osmolarity response regulator	T149A	T149A	T149A	T149A	0.28	Benign (0,001)	Deleterious (-2.503)	0/115
<i>envZ</i>	Osmolarity sensor protein	-	-	-	-				

\*Mutations predicted to be deleterious at the protein level at least with two of the three softwares and absent in Uniprot database are highlighted in red

<sup>‡</sup>SIFT software : a score between 0 and 0,05 predicted an alteration of the protein function (altereted) ; a score above 0,05 predicted no alteration (tolerated) : Kumar et al . 2009. Nature protocol. 4:1073-1081

<sup>§</sup>Polyphen-2 software : 3 classes of damaging were predicted : benign, possibly damaged and probably damaged described as follow : Adzhubei et al . 2010. Nature methods. 7(4): 248-249

<sup>§</sup>PROVEAN : a score < -2,5 is predicted as deleterious : Choi et al., 2015.Bioinformatics 31(16): 2745-2747.