

Table 1. Bacterial strains and plasmids used in this study

Strain or plasmid	Genotype or relevant characteristic	Ref. or source
<i>Escherichia coli</i>		
XL1-Blue	<i>RecA1</i> , <i>endA1</i> , <i>gyrA96</i> , <i>thi-1</i> , <i>hsdR17</i> , <i>supE44</i> , <i>relA1</i> , <i>lac</i> , [<i>F'</i> , <i>proAB</i> , <i>lacI^RZΔM15</i> , <i>Tn10</i> (<i>tet^R</i>)]	1, Stratagene
B ER2566	<i>F</i> ⁻ , λ^- , <i>fhuA2</i> [<i>lon</i>], <i>ompT</i> , <i>lacZ</i> : <i>T7</i> , <i>geneI</i> , <i>gal</i> , <i>sulA11</i> , Δ (<i>mcrC-mrr</i>) <i>114</i> :: <i>IS10</i> , <i>R</i> (<i>mcr-73</i> :: <i>miniTn10</i>) <i>2</i> , <i>R</i> (<i>zgb-210</i> :: <i>Tn10</i>) <i>1</i> , (<i>TetS</i>), <i>endA1</i> [<i>dcm</i>]	2, NEB
M15(pREP4)	<i>Lac</i> , <i>ara</i> , <i>gal</i> , <i>mtl</i> , <i>recA</i> ⁺ , <i>uvr</i> ⁺ , [pREP4, <i>lacI</i> , <i>kan</i> ^r]	3
SK6600	<i>F</i> ⁻ , <i>ara</i> ^r , Δ (<i>lac-proAB</i>), <i>rpsL</i> , <i>F80</i> , <i>lacZΔM15(r⁺, m⁺)</i>	4
SK6600ispG::neo ^R	<i>F</i> ⁻ , <i>ara</i> ^r , Δ (<i>lac-proAB</i>), <i>rpsL</i> , <i>F80</i> , <i>lacZΔM15(r⁺, m⁺)</i> , <i>ispG</i> :: <i>neo</i> ^R (<i>neo</i> ^R)	This study
Plasmids		
pMAL-C2	High-copy maltose binding protein expression vector	NEB
pMALispG	Expression of <i>ispG</i> (<i>gcpE</i>) from <i>Escherichia coli</i> as maltose binding fusion protein	This study
pBluescript SKII ⁻	High-copy cloning vector	Stratagene
pACYC184	Low-copy cloning vector	5, NEB
pMAK705	Low-copy vector with temperature sensitive origin for DNA replication (<i>Cm</i> ^R)	4
pMAKispGneo ^R	pMAK705 with a 1.7-kb fragment containing the <i>ispG</i> gene and adjacent regions interrupted by a neomycin resistance cassette	This study
pBSipp	Expression of <i>mk</i> , <i>pmk</i> , and <i>dpmid</i>	This study
pACYCipp	Expression of <i>mk</i> , <i>pmk</i> , and <i>dpmid</i>	This study
pNCO113	High-copy expression vector	ATCC, PTA-852
pNCOidi	Expression of <i>idi</i> from <i>E. coli</i>	This study

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