

Table S2: Strains, plasmids, and oligonucleotides

Name	Description	Source
Bacterial Strains		
<i>Vibrio cholerae</i> ¹		
E7946	Wildtype <i>Vibrio cholerae</i> El Tor Ogawa derivative, SmR	(1)
E7946 $\Delta hapR$	E7946 derivative lacking <i>hapR</i>	This work
E7946 Δcrp	E7946 derivative lacking <i>crp</i>	This work
E7946 $\Delta hapR\Delta crp$	E7946 derivative lacking <i>hapR</i> and <i>crp</i>	This work
E7946 $\Delta luxO$	E7946 derivative lacking <i>luxO</i>	This work
E7946 $\Delta hapR$	E7946 derivative <i>hapR::Spec^R</i>	This work
E7946 $\Delta luxO\Delta hapR$	E7946 derivative lacking <i>luxO</i> , <i>hapR::Spec^R</i>	This work
E7946 $\Delta murP$	E7946 derivative lacking <i>murP</i>	(2)
<i>Escherichia coli</i>		
DH5 α	<i>fhuA2</i> $\Delta(argF-lacZ)$ U169 <i>phoA glnV44</i> $\Phi 80$ $\Delta(lacZ)$ M15 <i>gyrA96 recA1 relA1 endA1 thi-1 hsdR17</i>	NEB
T7 Express	<i>fhuA2 lacZ::T7 gene1 [lon] ompT gal sulA11 R(mcr-73::mini</i> <i>Tn10--Tet^S)2 [dcm] R(zgb-210::Tn10--Tet^S) endA1 $\Delta(mcrC-$</i> <i>mrr)114::IS10</i>	NEB
JCB387	$\Delta nirB$, Δlac	(3)
S17 λpir	<i>lacU169 (lacZM15), recA1, endA1, hsdR17, thi-1, gyrA96,</i> <i>relA1, pir</i>	(4)
Plasmids		
pSR	pBR322-derived plasmid. Features cloning site upstream λ oop transcription terminator. AmpR	(5)
pRW50T	A broad-host range <i>lacZ</i> expression vector encoding <i>oriT</i> of pRK, TetR, Tra+	(6)
pAMNF	Plasmid for basal expression of N-terminal 3xFLAG-tagged proteins, KanR	(7)
pAMCF	Plasmid for basal expression of C-terminal 3xFLAG-tagged proteins, KanR	(7)
pRK2013	Helper plasmid for conjugation, KanR, oriColE1, RK2-, Mob+, Tra+	(8)
pKAS32	Suicide plasmid for mutant strain construction, AmpR	(9)
Oligonucleotides ^{2,3}		
Oligonucleotides for cloning <i>hapR</i> or <i>luxO</i> (5' to 3')		
<i>hapR</i> pAMNF fwd	GGCTGCGGTACCATGGACGCATCAATCGAAAAACGC	This work
<i>hapR</i> pAMNF rev	GCCCGAAGCTTCTAGTTCTTATAGATACACAGCATAT TGAGG	This work

<i>luxO</i> pAMCF fwd	GGCTGCGGTACCATGGTAGAAGACACGGCGTCGGTG GCGGCGCTGTATCGTTCTTACCTCACACCGCTGGATA TTGATATCAATATCGTGGGT <u>ACGGGAC</u>	This work
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<i>luxO</i> pAMCF rev.1	ATCGCGTCCCCTACCCACGATATTG	This work
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<i>luxO</i> pAMCF rev.2	GCCCGAAGCTTCCGTTCTTCTTTTTCTTTTAC	This work
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Oligonucleotides for cloning regulatory regions in pRW50T and pSR (5' to 3')

<i>PmurQ</i> fwd	AAAAGAATTCCACCAATCTGGCGGCCACTC	This work
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<i>PmurQ</i> rev	TTTTAAGCTTCATAAGGCTTCTCGGCAAAT	This work
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<i>PhapR</i> fwd	AAAAGAATTCCATACCATTCTCGTTGTGTT	This work
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<i>PhapR</i> rev	TTTTAAGCTTCATAGGGGTATATCCTTGCC	This work
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<i>PVC0585</i> fwd	AAAAGAATTCCATAGGGGTATATCCTTGCC	This work
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<i>PVC0585</i> rev	TTTTAAGCTTCATACCATTCTCGTTGTGTT	This work
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<i>PVC0502</i> fwd	AAAAGAATTGACAAAAGTTTGTGGCCGC	This work
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<i>PVC0502</i> rev	TTTTAAGCTTCATAGTATTGGCTTTGGCAT	This work
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<i>PmutH</i> fwd	AAAAGAATTCACCAGACCACATGAAGATCC	This work
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<i>PmutH</i> rev	TTTTAAGCTTCATAAAGGCTTTCGGTTTGG	This work
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<i>PleuO</i> fwd	AAAAGAATTCTAATGAACTGACTAACTCA	This work
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<i>PleuO</i> rev	TTTTAAGCTTCATTGCGTCTTTTTTATCT	This work
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<i>PVC0433</i> fwd	AAAACAATTGTACCTGCAACTTCAAGTAGT	This work
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<i>PVC0433</i> rev	TTTTAAGCTTCATTATTTTTTAATCACCAA	This work
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<i>PVC0241</i> fwd	AAAAGAATTCCTATAGATGCGAACAGTTGC	This work
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<i>PVC0241</i> rev	TTTTAAGCTTCATAGTATGCTGACTACTGC	This work
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<i>PrfaD</i> fwd	AAAACAATTGCATAGTATGCTGACTACTGC	This work
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<i>PrfaD</i> rev	TTTTAAGCTTCATTATGAATTCCTATAGAT	This work
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<i>PVC0620</i> fwd	AAAAGAATTCAAACCGTACCCGTTTTGCGAG	This work
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<i>PVC0620</i> rev	TTTTAAGCTTACTAGTAAGGAACAGCTATG	This work
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<i>PVC0688</i> fwd	AAAAGAATTCATAGCGTTTGTCTTTTGT	This work
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<i>PVC0688</i> rev	TTTTAAGCTTCATTACTATCCATTTTTTCA	This work
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<i>PVC1298</i> fwd	AAAAGAATTCATAAACTCTTTGTATAATT	This work
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<i>PVC1298</i> rev	TTTTAAGCTTCATGATAGTTTTGTAAATTAT	This work
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<i>PVC1375</i> fwd	AAAACAATTGCATCCACTTCTTCCTTATTA	This work
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<i>PVC1375</i> rev	TTTTAAGCTTCATATCAAAGTGGTTGGGA	This work
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<i>PVC1376</i> fwd	AAAACAATTGCATATCAAAGTGGTTGGGA	This work
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<i>PVC1376</i> rev	TTTTAAGCTTCATCCACTTCTTCCTTATTA	This work
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<i>PVC1403</i> fwd	AAAAGAATTCATTTTTTGGGTAAATCGATA	This work
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<i>PVC1403</i> rev	TTTTGGATCCCATGGAAAACCTCGTTGTTT	This work
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<i>PVC1405</i> fwd	AAAAGAATTCATGGAAAACCTCGTTGTTT	This work
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PVC1405 rev	TTTTGGATCCCATTTTTTGGGTAATCGATA	This work
PVC1436 fwd	AAAAGAATTCCCAATTCGTAAAGAAGCTAA	This work
PVC1436 rev	TTTTGGATCCCATATCTGCTGAGGCTTTAG	This work
PVC2352 fwd	AAAAGAATTCTCAGCACAATATCTCGCGCC	This work
PVC2352 rev	TTTTAAGCTTCATGCCGATGAGGCTCATAA	This work
PVCA0218 fwd	AAAAGAATTCCATATAAACCTCACTGACTC	This work
PVCA0218 rev	TTTTAAGCTTCATCGTTTACTTCTTATCAT	This work
PVCA0219 fwd	AAAAGAATTCCATCGTTTACTTCTTATCAT	This work
PVCA0219 rev	TTTTAAGCTTCATATAAACCTCACTGACTC	This work
PVCA0662 fwd	AAAAGAATTCTCAGTTTGTAATTGAGGAA	This work
PVCA0662 rev	TTTTAAGCTTCATATCAAATATGAATCCTT	This work
PVCA0663 fwd	AAAAGAATTCCATATCAAATATGAATCCTT	This work
PVCA0663 rev	TTTTAAGCTTCATGGTGTTACCTACTTGTT	This work
PVCA0906 fwd	AAAAGAATTCATTTTTATCCTGACCCCATATA	This work
PVCA0906 rev	TTTTGGATCCCATGCCTACGCCTATCGCCG	This work
PVCA0960 fwd	AAAAGAATTCCATCATGAGTTATATTTACA	This work
PVCA0960 rev	TTTTAAGCTTCATCCCTCAATCCTCAGTTT	This work
PVCA0961 fwd	AAAAGAATTCCATCCCTCAATCCTCAGTTT	This work
PVCA0961 rev	TTTTAAGCTTCATCATGAGTTATATTTACA	This work

Oligonucleotides for sequencing cloned DNA in plasmid constructs (5' to 3')

pRW50 seq fwd	GTTCTCGCAAGGACGAGAATTC	This work
pRW50 seq rev	GTCGTTGAACTGAGCCTGAAATTCAGG	This work
pSR seq fwd	GTGCCACCTGACGTCTAAGAAACC	This work
pSR seq rev	GCAACCGAGCGTTCTGAACAAATCC	This work
pAM seq fwd	ATTTATTCCAATGTCACACACTTTTCGC	This work
pAM seq rev	GAAACGCCGTAGCGCCGATGGTAGT	This work

Oligonucleotides for amplifying DNA for radio labelling (5' to 3')

pSR footprinting fwd	[BIOTIN]-CACGAGGCCCTTTCGTCTTCTC	This work
pSR footprinting rev	GGAGTTCTGAGGTCATTACTGGAG	This work
pSR footprinting rev	CACGAGGCCCTTTCGTCTTCTC	This work
pSR footprinting rev	[BIOTIN]-GGAGTTCTGAGGTCATTACTGGAG	This work

Oligonucleotides for cloning DNA fragments in pKAS32 (5' to 3')

pKAS32 fwd	GCAGGCACAAGCGGCCGCCTGCAGCTGGCGCCA TCGATACGCGTACGTCCG	This work
pKAS32 rev	CACGGTTTCATTAACAACCGGTACCTCTAGAACT	This work

	ATAGCTAGCATGCGCAAATTTAAAGCGCTG	This work
<i>crp</i> arm up fwd	CGGTTGTTAATGAAACCGTGGATATTAATGC	This work
<i>crp</i> arm up rev	ATCGGGGCACCTAGCCGATTTTTCCGGTTTC	This work
<i>crp</i> arm down fwd	AATCGGCTAGGTGCCCGATAACCCGTC	This work
<i>crp</i> arm down rev	AGGCGGCCGCTTGTGCCTGCGCAGCCAA	This work
<i>hapR</i> arm1 fwd	CTAGAGGTACCGTTGTTAATCCCAACCCCGATT GGTAATC	This work
<i>hapR</i> arm1 rev	TGTGTTTCATTTTCTTGGGCAGCACAAAG	This work
<i>hapR</i> arm2 fwd	GCCCAAGAAAATGAAACACACAGTTGAAGTC	This work
<i>hapR</i> arm2 rev	CGCCAGCTGCAGGCGGCCGCTATGCGGTCGATGT GCTGA	This work

Oligonucleotides for mutagenesis of *crp* or *hapR* (5' to 3')

hapR R123E fwd	TGCTTCAACCGAGGACGAAGTTTGGC	This work
hapR R123A fwd	TGCTTCAACCGCTGACGAAGTTTGGCC	This work
hapR R123 rev	CTCCACTCAAACCAGACTTTG	This work
<i>crp</i> E55R fwd	GATCAAAGATCGTGAAGGTAAAGAGATGATTCTC	This work
<i>cpr</i> E55A fwd	GATCAAAGATGCGGAAGGTAAAGAGATGATTC	This work
<i>crp</i> E55 rev	AGTACCGCAACTGAACCT	This work

¹E7946 $\Delta luxO$, $\Delta hapR$, $\Delta luxO\Delta hapR$ and $\Delta murP$ derivatives are synonymous with strains SAD764, SAD793, TND3765 and SAD268.

²sequences in italic are sites for restriction endonucleases for use during cloning

³underlined sequences encode amino acid substitutions or remove unwanted sites for restriction endonucleases

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