

Table S1. Plasmids and bacterial strains used in this study

Strains and plasmids	Genotype and relevant characteristics	Source or references
E. coli strains		
DH5 α	F- ϕ 80d <i>lacZ</i> Δ M15 Δ (<i>lacZYA-argF</i>) U169 <i>recA1 end A1 hsdR17</i> (rk-, mk+) <i>phoA</i> , <i>supE44</i> λ - <i>thi-1 gyrA96 relA1 phoA</i>	TransGen Biotech
β 2163	F-, RP4-2-Tc::Mu Δ dapA::(erm-pir)	Demarre et al., 2005
S17-1 λ pir	Tpr Smr <i>recA thi pro hsdR</i> -M+RP4:2-Tc: Mu:Km Tn7 λ pir	Simon et al., 1983
S17-1-pBBR1-MCS1- <i>hcp2</i>	S17-1 containing plasmid of pBBR1-MCS1- <i>hcp2</i> , Cmr	This study
<i>Vibrio aginolyticus</i>		
HY9901	Wild-type, isolated from the diseased fish	Cai et al., 2007
Δ <i>hcp2</i>	HY9901 carrying an in-frame deletion of <i>hcp2</i>	This study
HY9901-pBBR1-MCS1	HY9901 containing plasmid of pBBR1-MCS1, Cmr (HY9901-ev)	This study
Δ <i>hcp2</i> -pBBR1-MCS1	HY9901 Δ <i>hcp2</i> containing plasmid of pBBR1-MCS1, Cmr (Δ <i>hcp2</i> -ev)	This study
Δ <i>hcp2</i> -pBBR1-MCS1- <i>hcp2</i>	HY9901 Δ <i>hcp2</i> containing plasmid of pBBR1-MCS1- <i>hcp2</i> , Cmr (Δ <i>hcp2</i> -com)	This study
Plasmids		
pMD19-T	Cloning vector; Amp ^r	TaKaRa
pBBR1-MCS1	Expression vector; pBBR1 Rep, pBBR1 oriV; Cmr	Kovach et al., 1995
pLP12	oriTRP4 oriVR6Kvmi480 PBAD, Cmr	Luo et al., 2015
pLP12- <i>hcp2</i>	pLP12 derivative containing homologous arms of <i>hcp2</i>	This study