Table S1. Plasmids used in this study.

Plasmid	Relevant characteristics ^a	Reference or source
pGEM-T Easy	Ap ^r , cloning vector	Promega
pPS858-Eco	Ap ^r , Gm ^r , source of Gm ^r -GFP-FRT cassette	[67]
pEX18Ap	Ap ^r , gene replacement vector	[67]
pEX18Ap.ABC58-ko.Gm-GFP	$\mbox{Ap}^{\prime},\mbox{Gm}^{\prime},\mbox{contains a 2.8-kb fusion fragment of PA14_58420,}\mbox{Gm}^{\prime}-\mbox{GFP-}\emph{FRT}\mbox{ cassette, and PA14_58500}$	This study
pEX18Gm	Gm ^r , gene replacement vector	[67]
pEX18Gm.SBP58350-ko	Gm ^r , contains a 1.1-kb fusion fragment of PA14_58350 and PA14_58360	This study
pEX18Gm.SBP58390-ko	Gm ^r , contains a 1.5-kb fusion fragment of PA14_58390	This study
pEX18Gm.SBP58420-ko	Gm ^r , contains a 1.2-kb fusion fragment of PA14_58420	This study
pEX18Gm.SBP70200-ko	Gm ^r , contains a 1.4-kb fusion fragment of PA14_70200	This study
pBBR1MCS-5	Gm ^r , broad-host-range cloning vector	[69]
pBBR5.DppA1	${\rm Gm}^{\rm r}$, contains a 2.0-kb fragment carrying the SBP PA14_58350 including upstream promoter region in opposite orientation with respect to the $\it lac$ promoter	This study
pBBR5.DppA2	${\rm Gm}^{\rm f}$, contains a 1.7-kb fragment carrying the SBP PA14_58360 including upstream promoter region in opposite orientation with respect to the $\it lac$ promoter	This study
pBBR5.DppA3	${\rm Gm}^{\rm r}$, contains a 1.9-kb fragment carrying the SBP PA14_58390 including upstream promoter region in opposite orientation with respect to the $\it lac$ promoter	This study
pBBR5.DppA4	Gm^r , contains a 1.8-kb fusion fragment carrying the SBP PA14_58420 including upstream promoter region in opposite orientation with respect to the lac promoter	This study
pBBR5.DppA5	${\rm Gm}^{\rm r}$, contains a 2.0-kb fragment carrying the SBP PA14_70200 including upstream promoter region in opposite orientation with respect to the {\it lac} promoter	This study
pFLP2	Ap ^r , source of FLP recombinase	[67]

^a Antibiotic resistance: Ap^r, ampicillin resistance; Gm^r, gentamicin resistance