

## Appendix A. Strains and plasmids used in this study

Strain or plasmid	Description <sup>a</sup>	Source or reference
<b>Burkholderia</b>		
<b>glumae (B. glumae)</b>		
LMG 2196 <sup>T</sup>	Wild type strain of <i>B. glumae</i>	Kindly provided by Prof. Guanlin Xie from Zhejiang University
LMG 2196 <sup>Δsyp</sup>	LMG 2196 deletion mutation defective in <i>syp</i>	This study
<b>Escherichia coli</b>		
DH5α	Bacterial host for gene cloning	Laboratory collection
S17-1(λ pir)	Bacterial host for gene cloning	Laboratory collection
BL21(DE3)	Bacterial host for protein expression	Novagen
<b>Plasmids</b>		
pGEM-T	Amp <sup>R</sup> ; cloning vector	Promega
pUFR034	IncW Nm <sup>f</sup> Mob <sup>+</sup> mob (P) lacZ alpha Par <sup>+</sup> cos	Laboratory collection
pKMS1	Km <sup>R</sup> ; R6K-based suicide vector; requires the <i>pir</i> -encoded π protein for replication	Laboratory collection
LMG 2196- pUFR034	Empty vector control	This study
pKMS1- <i>syp</i>	Km <sup>R</sup> ; Used to create knockout mutant of LMG 2196 <sup>syp</sup>	This study
pUFR034- <i>syp</i>	Km <sup>R</sup> ; Used to create complement of LMG 2196 <sup>syp</sup>	This study

<sup>a</sup>Km<sup>R</sup> and Amp<sup>R</sup> indicate kanamycin and ampicillin resistant

## Appendix B Schematic diagram about gene knockout of *syp*.



## Appendix C List of oligonucleotide polymerase chain reaction (PCR) primers used in this study

Primer name	Nucleotide sequence (5'-3') <sup>a</sup>	Target PCR product of function
Pr1-F	TGAGGATCCTTCTGCAGGCTGATGAAATC (B)	1009 bp
Pr1-R	CCGGAATTCGGCTTGC GGTAAGGTGTG (E)	
Pr2-F	CCGGAATTCGAGGTGTACATCACGTAGGC (E)	1220 bp
Pr2-R	TGTCAGTCGAGAGCTGCAGGAACTGATCG (S)	
Pr3-F	CCGGAATTC GCAGATAGACCTGCGTGTTG (E)	948 bp
Pr3-R	TGTCAGTCGA AGGTCCGGCTCAGCTACC (S)	
Pr4-F	CCGGAATTC AAGCGCTGGTAGCGATAGTC (E)	1014 bp
Pr4-R	CCGGAATTC GCACCCGTCTACAACATCG (S)	
Pr5-F	CCGGAATTC CTATGCGCTCGTATCGAGGT (E)	1072 bp
Pr5-R	CCGGAATTC GATATCGGCTCGCTGGTCT (S)	
<i>syp</i> -F	TGAGGATCCATGAATAACAATGTGCCGAA (B)	23,421 bp including <i>syp</i>
<i>syp</i> -R	TGTCAGTCGATCACGGCGCGTCCGGCGCGT (S)	

<sup>a</sup> Underlined nucleotides in some of the PCR primers represent restriction sites of enzymes indicated in parentheses (E=*Eco*RI; B=*Bam*HI; S=*Sal*I). The corresponding enzymes were used to excise the obtained PCR products.