

Supplementary Materials

Table S1: Primers used in this study; Table S2: Bacterial strains and plasmids used in this study;

Table S1. Strains and plasmids used in this study

Strains and vectors	Characteristic	Source
Strains		
<i>Acidovorax citrulli</i>		
Aac5	Wild type strain, Amp ^r	This lab
$\Delta hrpX$	<i>hrpX</i> markerless knock-out mutant of Aac5, Amp ^r	This lab
$\Delta hrpG$	<i>hrpG</i> markerless knock-out mutation of Aac5, Amp ^r	This lab
$\Delta hrcJ$	<i>hrcJ</i> markerless knock-out mutation of Aac5, Amp ^r	This lab
WT- <i>aopV</i> - <i>cyaA</i>	Wild-type strain Aac5 containing pBBRNolac- <i>cyaA</i> carrying <i>aopV</i> with its native promoter; Amp ^r ; Km ^r	This study
$\Delta hrcJ$ - <i>aopV</i> - <i>cyaA</i>	$\Delta hrcJ$ containing pBBRNolac- <i>cyaA</i> carrying <i>aopV</i> with its native promoter; Amp ^r ; Km ^r	This study
<i>Escherichia coli</i>		
DH5 α	F-, <i>supE44</i> $\Delta lacU169$ ($\Phi 80/lacZ\Delta M15$), <i>hsd R17</i> , <i>recA1</i> , <i>endA1</i> , <i>gyrA96thi-1 relA1</i>	TIANGEN, Beijing, China
<i>Agrobacterium tumefaciens</i>		
GV3101	The wild-type strain, Rif ^r	BIOMED, Beijing, China
Vectors		
pBBR1MCS-2	Broad host range expression vector with <i>lac</i> promoter, Km ^r	This lab
pBBRNolac- <i>cyaA</i>	pBBRNolac-4FLAG containing <i>cyaA</i> gene without promoter	This lab
pBBRNolac- <i>aopV</i> - <i>cyaA</i>	pBBRNolac- <i>cyaA</i> containing <i>Aave_3085</i> gene with its native promoter	This study
pBI121	Plant expression vector containing 35S promoter, Km ^r	This study
pBI121-3FLAG	pBI121 containing 3FLAG tag, Km ^r	This lab
pBI121-eGFP	pBI121 containing eGFP tag, Km ^r	This lab
pBI121-3FLAG-AopV	pBI121-3FLAG containing AopV	This study
pBI121-GFP-AopV	pBI121-GFP containing AopV	This study
pCAMBIA1300-nLUC	Plant expression vector containing 35S promoter, carrying nLUC tag, Km ^r	Lab collection
pCAMBIA1300-cLUC	Plant expression vector containing 35S promoter, carrying cLUC tag, Km ^r	Lab collection
pCAMBIA1300-nLUC-AopV	pCAMBIA 1300-cLUC fused with AopV for LCI assay, Km ^r	This study

pCAMBIA1300-cLU C-ADT6	pCAMBIA1300-cLUC fused with ADT6 for LCI assay, Km ^r	This study
pSPYNE®173	Plant expression vector containing 35S promoter, carrying nYFP tag, Km ^r	Lab collection
pSPYCE(M)	Plant expression vector containing 35S promoter, carrying cYFP tag, Km ^r	Lab collection
pSPYNE®173-AopV	pSPYNE®173 containing AopV for BiFC assay, Km ^r	This study
pSPYCE(M)-ADT6	pSPYCE(M) containing ADT6 for BiFC assay, Km ^r	This study
pGBKT7-53	Positive control bait	Lab collection
pGADT7-T	Positive control prey	Lab collection
pGBKT7-Lam	Negative control bait	Lab collection
pGBKT7-AopV	pGBKT7 containing AopV for yeast two hybrid assay, Km ^r	This study
pGADT7-ADT6	pGADT7 containing ADT6 for yeast two hybrid assay, Amp ^r	This study

The Km^r, Amp^r and Rif^r indicate resistance to kanamycin, ampicillin, and rifampicin, respectively.

Table S2. Primers used in this study

Primer name	Primer sequence (5'-3')	Description
HB- <i>aopV</i> -F	CTAGTTCTAGAGCGGCCGCCA	For cloning the <i>aopV</i> full-length open reading frame (ORF) with its native promoter in the pBBRNolac- <i>aopV</i> - <i>cyaA</i> vector
HB- <i>aopV</i> -R	GGTACCGGGCCCCCTCGAG	
RT- <i>aopV</i> -F	CCGTCAGGCGAGTCATTTCT	For detecting <i>aopV</i> mRNA in RT-qPCR assay
RT- <i>aopV</i> -R	TTAGGGCTATGTGTTCCGGC	
121GFP- <i>aopV</i> -F	GAGCTGGACAAGGGTCTAGACGA AGATGCGGCTGTGGAA	For cloning the <i>aopV</i> full-length ORF into pBI121-eGFP vector
121GFP- <i>aopV</i> -R	TCTATCGATCAATCAGGATCCCCC CGGAGCGCGCGGCGCC	
<i>Ac</i> - <i>cyaA</i> -F	TCCCCCGGGCTGCAGGAATTC GATGGGCAGCAATCGCATCA	For cloning the <i>cyaA</i> into pBBRNolac vector to construct the pBBRNolac- <i>cyaA</i> vector
<i>Ac</i> - <i>cyaA</i> -R	ATCCTTGTAATCGGTAAGCTT GCTGTCATAGCCGGAATCCT	
121FLAG- <i>aopV</i> -F	GAGAACACGGGGGACTCTAGCG AAGATGCGGCTGTGGAA	For cloning the <i>aopV</i> full-length ORF in pBI121-3FLAG vector
121FLAG- <i>aopV</i> -R	GTCATCCTTGTAATCCCCGGCCCC GGAGCGCGCGGCGCC	
121FLAG-F	GGACTCTAGAGGATCCCCGGGGA TTACAAGGATGAC	For inserting the 3FLAG tag into pBI121 vector to construct the pBI121-3FLAG vector
121-FLAG-R	CGATCGGGGAAATTCGAGCTCTC ACTTGTCATCGTCAT	
121GFP-F	GGACTCTAGAGGATCCCCGGGAT GGTGAGCAAGGGCGAG	For inserting the eGFP tag into pBI121 vector to construct the pBI121-eGFP vector
121GFP-R	CGATCGGGGAAATTCGAGCTCTC ACTTGACAGCTCGTCCATG	
RT- <i>rpoB</i> -F	GCGACAGCGTGCTCAAAGTG	Reference gene of <i>A. citrulli</i>

RT- <i>rpoB</i> -F	GCCTTCGTTGGTGCGTTTCT	strain Aac5 in RT-qPCR assay
<i>NbAcre31</i> -F	AATTCGGCCATCGTGATCTTGGTC	For detecting <i>NbAcre31</i> mRNA in RT-qPCR assay
<i>NbAcre31</i> -R	GAGAAACTGGGATTGCCTGAAGG A	
<i>NbEF1α</i> -F	AAGGTCCAGTATGCCTGGGTGCTT GAC	Reference gene of <i>N. benthamiana</i> in RT-qPCR assay
<i>NbEF1α</i> -R	AAGAATTCACAGGGACAGTTCCA ATACCA	
<i>NbPti5</i> -F	CCTCCAAGTTTGAGCTCGGATAGT	For detecting <i>NbPti5</i> mRNA in RT-qPCR assay
<i>NbPti5</i> -R	CCAAGAAATTCTCCATGCACTCT GTC	
AD-ADT6-F	GCCATGGAGGCCAGTGAATTCAT GGACGCTCAGAACTTCACG	For cloning the <i>ADT6</i> into pGADT7-T vector
AD-ADT6-R	CAGCTCGAGCTCGATGGATCCTC AAGCACTTTTGGGGCCAA	
nLUC- <i>aopV</i> -F	ACGGGGGACGAGCTCGGTACCGA AGATGCGGCTGTGGAA	For cloning the <i>aopV</i> into pCAMBIA1300-nLUC vector
nLUC- <i>aopV</i> -R	CGCGTACGAGATCTGGTTCGACCC CGGAGCGCGCGGCGCC	
cLUC-ADT6-F	TACGCGTCCCGGGGCGGTACATG GACGCTCAGAACTTC	For cloning the <i>ADT6</i> full-length ORF into pCAMBIA1300-cLUC vector
cLUC-ADT6-R	ACGAAAGCTCTGCAGGTCTGAAGC ACTTTTGGGGCCAATG	
173-AopV-F	CCCAGGCCTACTAGTGGATCCGA AGATGCGGCTGTGGAA	For cloning the <i>aopV</i> into pSPYNE [®] 173 vector
173-AopV-R	CCCGGGAGCGGTACCCTCGACCC CGGAGCGCGCGGCGCC	
M-ADT6-F	TGGCGCGCCACTAGTGGATCATG GACGCTCAGAACTTC	For cloning the <i>ADT6</i> into pSPYCE(M) vector
M-ADT6-R	CCCGGGAGCGGTACCCTCGAGAG CACTTTTGGGGCCAATG	