

Supplementary Table S3 All plasmids used in this study

Plasmid	Description	Origin
pBARGPE1 series of overexpression plasmids		
pBARGPE1	pBARGPE1 is a broad host shuttle plasmid and fungal expression plasmid. The strong promoter <i>gpdA</i> from <i>Aspergillus nidulans</i> can be used to initiate the expression of exogenous genes. Bacteria can be screened with <i>AmpR</i> , and glufosinate-ammonium (<i>Bar</i>) can be screened when transformed into filamentous fungi.	J. Yang, W. W. Yang, J. Feng, J. Chen, M. Jiang and X. Zou, <i>J. Biotechnol.</i> , 2018, 275, 24-30.
pBARGPE1- <i>gpdA-adh-trapC</i>	<i>adh</i> overexpression plasmid, used to amplify <i>adh</i> expression cassette	This study
pBARGPE1- <i>gpdA-acs-trapC</i>	<i>acs</i> overexpression plasmid, used to amplify <i>acs</i> expression cassette	This study
pBARGPE1- <i>gpdA-adh-acs-trapC</i>	<i>adh-acs</i> fusion overexpression plasmid, used to amplify <i>adh-acs</i> fusion protein expression cassette	This study
pBARGPE1- <i>gpdA-eGFP-icl1-trapC</i>	<i>eGFP-icl1</i> fusion overexpression plasmid, used to amplify <i>eGFP-icl1</i> fusion protein expression cassette	This study
pBARGPE1- <i>gpdA-eGFP-icl2-trapC</i>	<i>eGFP-icl2</i> fusion overexpression plasmid, used to amplify <i>eGFP-icl2</i> fusion protein expression cassette	This study
pBARGPE1- <i>gpdA-eGFP-mls-trapC</i>	<i>eGFP-mls</i> fusion overexpression plasmid, used to amplify <i>eGFP-mls</i> fusion protein expression cassette	This study
pBARGPE1- <i>gpdA-icl1-mls-trapC</i>	<i>icl1-mls</i> fusion overexpression plasmid, used to amplify <i>icl1-mls</i> fusion protein expression cassette	This study
pBARGPE1- <i>gpdA-cat8-trapC</i>	<i>cat8</i> overexpression plasmid, used to amplify <i>cat8</i> expression cassette	This study
pBARGPE1- <i>gpdA-cat8-myc¹³-trapC</i>	<i>cat8-myc¹³</i> fusion overexpression plasmid, used to amplify <i>cat8-myc¹³</i> fusion protein expression cassette	This study
pBARGPE1- <i>gpdA-pck-trapC</i>	<i>pck</i> overexpression plasmid, used to amplify <i>pck</i> expression cassette	This study
pBARGPE1- <i>gpdA-Asp. PGK-hyg-trapC</i>	<i>Asp. PGK-hyg</i> fusion overexpression plasmid, used to amplify <i>Asp. PGK-hyg</i> fusion protein expression cassette	This study

pBARGPE1- <i>gpdA-Asp. FBA-hyg-trapC</i>	<i>Asp. FBA-hyg</i> fusion overexpression plasmid, used to amplify <i>Asp. FBA-hyg</i> fusion protein expression cassette	This study
pBARGPE1- <i>gpdA-Asp. PGI-hyg-trapC</i>	<i>Asp. PGI-hyg</i> fusion overexpression plasmid, used to amplify <i>Asp. PGI-hyg</i> fusion protein expression cassette	This study
pBARGPE1- <i>gpdA-Rt. FBA-hyg-trapC</i>	<i>Rt. FBA-hyg</i> fusion overexpression plasmid, used to amplify <i>Rt. FBA-hygA</i> fusion protein expression cassette	This study
pBARGPE1- <i>gpdA-Rt. PGI-hyg-trapC</i>	<i>Rt. PGI-hyg</i> fusion overexpression plasmid, used to amplify <i>Rt. PGI-hyg</i> fusion protein expression cassette	This study
pK2 series shuttle plasmids		
pK2- <i>ptrpc-hyg-trpCt</i>	It has the replication function of pK2 plasmid and can replicate in <i>E. coli</i> and <i>Agrobacterium tumefaciens</i> . With Ti plasmid on the left and right sides of the marginal region sequence, so that the DNA can be integrated into the genome of the infected cell. <i>Kan</i> is used as a screening label for bacterial transformants, and <i>hyg</i> is used as a screening label for infecting cell transformants.	J. Yang, W. W. Yang, J. Feng, J. Chen, M. Jiang and X. Zou, <i>J. Biotechnol.</i> , 2018, 275, 24-30.
pK2- <i>ptrpc-neoR-trpCt</i>	It has the replication function of pK2 plasmid and can replicate in <i>E. coli</i> and <i>Agrobacterium tumefaciens</i> . With Ti plasmid on the left and right sides of the marginal region sequence, so that the DNA can be integrated into the genome of the infected cell. <i>Kan</i> is used as a screening label for bacterial transformants, and <i>neoR</i> is used as a screening label for infecting cell transformants.	This study
pK2- <i>ptrpc-nat-trpCt</i>	It has the replication function of pK2 plasmid and can replicate in <i>E. coli</i> and <i>Agrobacterium tumefaciens</i> . With Ti plasmid on the left and right sides of the marginal region sequence, so that the DNA can be integrated into the genome of the infected cell. <i>Kan</i> is used as a screening label for bacterial transformants, and <i>nat</i> is used as a screening label for infecting cell transformants.	This study
pK2- <i>ptrpc-bar-gus-trpCt</i>	It has the replication function of pK2 plasmid and can replicate in <i>E. coli</i> and <i>Agrobacterium tumefaciens</i> . With Ti plasmid on the left and right sides of the marginal region sequence, so that the DNA can be integrated into the genome of the infected cell. <i>Kan</i> is used as a screening label for bacterial transformants, and <i>bar</i> is used as a screening label for infecting cell transformants.	This study
pK2- <i>gpdA-adh-acs-trapC-ptrpc-hyg-trpCt</i>	The plasmid inserted into the <i>adh-acs</i> fusion expression cassette for <i>Agrobacterium</i> -mediated	This study

	transformation	
pK2- <i>gpdA-cat8-trapC-ptRPC-neoR-trpCt</i>	The plasmid inserted into the <i>cat8</i> expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>gpdA-icl1-mls-trapC-ptRPC-bar-gus-trpCt</i>	The plasmid inserted into the <i>icl1-mls</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>gpdA-pck-trapC-ptRPC-nat-trpCt</i>	The plasmid inserted into the <i>pck</i> expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>gpdA-eGFP-icl1-trapC-ptRPC-hyg-trpCt</i>	The plasmid inserted into the <i>eGFP-icl1</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>gpdA-eGFP-icl2-trapC-ptRPC-hyg-trpCt</i>	The plasmid inserted into the <i>eGFP-icl2</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>gpdA-eGFP-mls-trapC-ptRPC-hyg-trpCt</i>	The plasmid inserted into the <i>eGFP-mls</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>gpdA-cat8-myc¹³-trapC-ptRPC-hyg-trpCt</i>	The plasmid inserted into the <i>cat8-myc¹³</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>gpdA-Asp. PGK-hyg-trapC-ptRPC-hyg-trpCt</i>	The plasmid inserted into the <i>Asp. PGK-hyg</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>gpdA-Asp. FBA-hyg-trapC-ptRPC-hyg-trpCt</i>	The plasmid inserted into the <i>Asp. FBA-hyg</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>gpdA-Asp. PGI-hyg-trapC-ptRPC-hyg-trpCt</i>	The plasmid inserted into the <i>Asp. PGI-hyg</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>gpdA-Rt. FBA-hyg-trapC-ptRPC-hyg-trpCt</i>	The plasmid inserted into the <i>Rt. FBA-hyg</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>gpdA-Rt. PGI-hyg-trapC-ptRPC-hyg-trpCt</i>	The plasmid inserted into the <i>Rt. PGI-hyg</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
Knockout plasmids		
p426-loxp- <i>gRNA</i> -Cas9-2Sac II	Contains Cas protein and <i>gRNA</i> amplification template. <i>AmpR</i> serves as a screening tag for bacterial transformants.	Y. Zhang, J. Feng, P. Wang, J. Xia, X. R. Li and X. Zou, <i>Gene</i> , 2019, 709, 8-

		16.
PCB004	Bacterial expression plasmid, <i>AmpR</i> as a selection label	This study
p426-loxp-gRNA(<i>cat8</i>)-Cas9-2Sac II	<i>cat8</i> knockout plasmid	This study
PCB004- <i>cat8</i> -up-donor	<i>cat8</i> knockout split marker upstream donor amplification template	This study
PCB004- <i>cat8</i> -down-donor	<i>cat8</i> knockout split marker downstream donor amplification template	This study
Promoter engineering adaptation plasmids		
pUC19- <i>Rt. FBA</i>	<i>Rhodospiridium toruloides</i> promoter <i>Rt. FBA</i> amplification template	This study
pUC19- <i>Rt. PGI</i>	<i>Rhodospiridium toruloides</i> promoter <i>Rt. PGI</i> amplification template	This study
PCB004- <i>Asp. PGK-CYC1</i>	<i>Asp. PGK</i> -driven template plasmid for the construction of expression cassettes, <i>AmpR</i> as a screening label for bacterial transformants	This study
PCB004- <i>Asp. FBA-CYC1</i>	<i>Asp. FBA</i> -driven template plasmid for the construction of expression cassettes, <i>AmpR</i> as a screening label for bacterial transformants	This study
PCB004- <i>Asp. PGI-CYC1</i>	<i>Asp. PGI</i> -driven template plasmid for the construction of expression cassettes, <i>AmpR</i> as a screening label for bacterial transformants	This study
PCB004- <i>Rt. FBA-CYC1</i>	<i>Rt. FBA</i> -driven template plasmid for the construction of expression cassettes, <i>AmpR</i> as a screening label for bacterial transformants	This study
PCB004- <i>Rt. PGI-CYC1</i>	<i>Rt. PGI</i> -driven template plasmid for the construction of expression cassettes, <i>AmpR</i> as a screening label for bacterial transformants	This study
PCB004- <i>Asp. PGK-adh-acs-CYC1</i>	<i>Asp. PGK</i> -driven <i>adh-acs</i> fusion expression cassette amplification template	This study
PCB004- <i>Asp. FBA-adh-acs-CYC1</i>	<i>Asp. FBA</i> -driven <i>adh-acs</i> fusion expression cassette amplification template	This study
PCB004- <i>Asp. PGI-adh-acs-CYC1</i>	<i>Asp. PGI</i> -driven <i>adh-acs</i> fusion expression cassette amplification template	This study
PCB004- <i>Rt. FBA-adh-acs-CYC1</i>	<i>Rt. FBA</i> -driven <i>adh-acs</i> fusion expression cassette amplification template	This study
PCB004- <i>Rt. PGI-adh-acs-CYC1</i>	<i>Rt. PGI</i> -driven <i>adh-acs</i> fusion expression cassette amplification template	This study
PCB004- <i>Asp. PGK-cat8-CYC1</i>	<i>Asp. PGK</i> -driven <i>cat8</i> expression cassette amplification template	This study
PCB004- <i>Asp. FBA-cat8-CYC1</i>	<i>Asp. FBA</i> -driven <i>cat8</i> expression cassette amplification template	This study
PCB004- <i>Asp. PGI-cat8-CYC1</i>	<i>Asp. PGI</i> -driven <i>cat8</i> expression cassette amplification template	This study

PCB004- <i>Rt. FBA-cat8-CYCI</i>	<i>Rt. FBA</i> -driven <i>cat8</i> expression cassette amplification template	This study
PCB004- <i>Rt. PGI-cat8-CYCI</i>	<i>Rt. PGI</i> -driven <i>cat8</i> expression cassette amplification template	This study
PCB004- <i>Asp. PGK-pck-CYCI</i>	<i>Asp. PGK</i> -driven <i>pck</i> expression cassette amplification template	This study
PCB004- <i>Asp. FBA-pck-CYCI</i>	<i>Asp. FBA</i> -driven <i>pck</i> expression cassette amplification template	This study
PCB004- <i>Asp. PGI-pck-CYCI</i>	<i>Asp. PGI</i> -driven <i>pck</i> expression cassette amplification template	This study
PCB004- <i>Rt. FBA-pck-CYCI</i>	<i>Rt. FBA</i> -driven <i>pck</i> expression cassette amplification template	This study
PCB004- <i>Rt. PGI-pck-CYCI</i>	<i>Rt. PGI</i> -driven <i>pck</i> expression cassette amplification template	This study
pK2- <i>Asp. PGK-adh-acs-trapC-ptipc-hyg-trpCt</i>	The plasmid inserted into the <i>Asp. PGK</i> -driven <i>adh-acs</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>Asp. FBA-adh-acs-trapC-ptipc-hyg-trpCt</i>	The plasmid inserted into the <i>Asp. FBA</i> -driven <i>adh-acs</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>Asp. PGI-adh-acs-trapC-ptipc-hyg-trpCt</i>	The plasmid inserted into the <i>Asp. PGI</i> -driven <i>adh-acs</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>Rt. FBA-adh-acs-trapC-ptipc-hyg-trpCt</i>	The plasmid inserted into the <i>Rt. FBA</i> -driven <i>adh-acs</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>Rt. PGI-adh-acs-trapC-ptipc-hyg-trpCt</i>	The plasmid inserted into the <i>Rt. PGI</i> -driven <i>adh-acs</i> fusion expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>Asp. PGK-cat8-trapC-ptipc-neoR-trpCt</i>	The plasmid inserted into the <i>Asp. PGK</i> -driven <i>cat8</i> expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>Asp. FBA-cat8-trapC-ptipc-neoR-trpCt</i>	The plasmid inserted into the <i>Asp. FBA</i> -driven <i>cat8</i> expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>Asp. PGI-cat8-trapC-ptipc-neoR-trpCt</i>	The plasmid inserted into the <i>Asp. PGI</i> -driven <i>cat8</i> expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>Rt. FBA-cat8-trapC-ptipc-neoR-trpCt</i>	The plasmid inserted into the <i>Rt. FBA</i> -driven <i>cat8</i> expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>Rt. PGI-cat8-trapC-ptipc-neoR-trpCt</i>	The plasmid inserted into the <i>Rt. PGI</i> -driven <i>cat8</i> expression cassette for Agrobacterium-mediated	This study

	transformation	
pK2- <i>Asp. PGK-pck-trapC-ptrpc-nat-trpCt</i>	The plasmid inserted into the <i>Asp. PGK</i> -driven <i>pck</i> expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>Asp. FBA-pck-trapC-ptrpc-nat-trpCt</i>	The plasmid inserted into the <i>Asp. FBA</i> -driven <i>pck</i> expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>Asp. PGI-pck-trapC-ptrpc-nat-trpCt</i>	The plasmid inserted into the <i>Asp. PGI</i> -driven <i>pck</i> expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>Rt. FBA-pck-trapC-ptrpc-nat-trpCt</i>	The plasmid inserted into the <i>Rt. FBA</i> -driven <i>pck</i> expression cassette for Agrobacterium-mediated transformation	This study
pK2- <i>Rt. PGI-pck-trapC-ptrpc-nat-trpCt</i>	The plasmid inserted into the <i>Rt. PGI</i> -driven <i>pck</i> expression cassette for Agrobacterium-mediated transformation	This study
