

1           **Distribution, diversity and activities of sulfur dioxygenases in**

2                                   **heterotrophic bacteria**

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5   Supplementary Information Contents

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7   1- Supplementary Table

8   Table S1

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10  2- Supplementary Figures

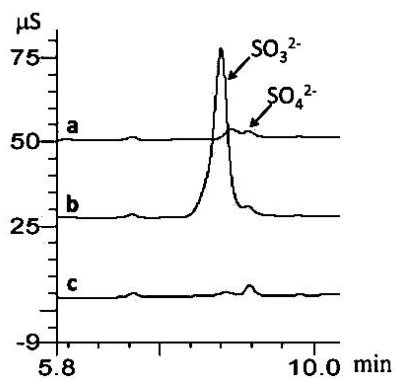
11  Fig.S1

12  Fig.S2

13 **Table S1 Oligonucleotide primers used for plasmid construction**

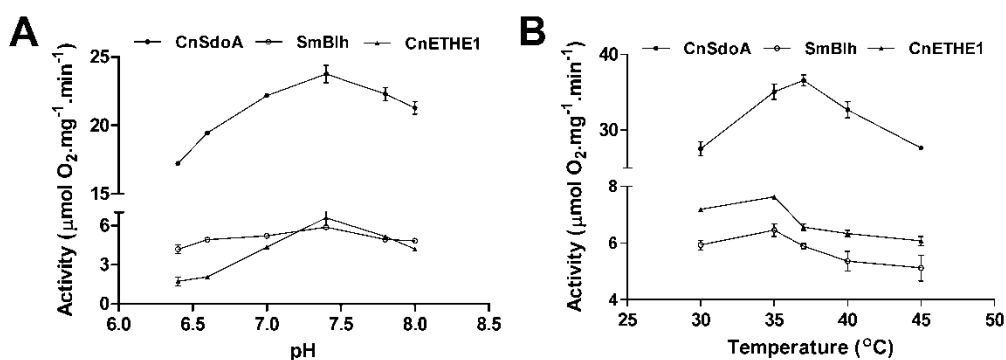
Primers	Nucleotide Sequence <sup>a</sup>	Target protein
AtBlh-f	<u>TAAGAAGGAGATATACAT</u> ATGAAGGCCGTAAGG	AtBlh
AtBlh-r	TGGTGGTGGTGCCTCGAGCCATGTGCTGCCCTCCAGCA	
SmBlh-f	<u>AGAAGGAGATATACAT</u> ATGCCTATTACTGCAATATCC	SmBlh
SmBlh-r	GTGGTGGTGCCTCGAGTTCCCATGCCGCTCCCTG	
CnSdoA-f	<u>TAAGAAGGAGATATACAT</u> ATGACACCGACCATGCCAAGCC	CnSdoA
CnSdoA-r	GTGGTGGTGCCTCGAGGAGGGCGTTGAGGGGAATCT	
BxSdoA-f	<u>TAAGAAGGAGATATACAT</u> ATGACCGCC	BxSdoA
BxSdoA-r	GTGGTGGTGCCTCGAGAAGCGCAT	
PpSdoA-f	<u>TAAGAAGGAGATATACAT</u> ATGATCATCGGCAACAACCTT	PpSdoA
PpSdoA-r	GTGGTGGTGCCTCGAGCAGCTTGTTTCAGCGGGATCT	
PaSdoA-f	<u>TAAGAAGGAGATATACAT</u> ATGTTGAAACCCGACATCAC	PaSdoA
PaSdoA-r	GTGGTGGTGCCTCGAGGAACAGATCCAGCGG	
CnETHE1-f	<u>TAAGAAGGAGATATACAT</u> ATGCAAACCTTCTATCAGCT	CnETHE1
CnETHE1-r	GTGGTGGTGCCTCGAGGGCGCCATGCGGCACGCTTT	
MxETHE1a-f	<u>AGAAGGAGATATACAT</u> ATGCTCTTCCGCCAGCTCT	MxETHE1a
MxETHE1a-r	GTGGTGGTGCCTCGAGATGTGTGAAGCTGCCT	
EcGloB1-f	<u>TAAGAAGGAGATATACAT</u> ATGAATCTTAACAGTATTCCCG	EcGloB1
EcGloB1-r	GGTGGTGGTGCCTCGAGGAACCTATCTTTCTTTGACC	
HiGloB1-f	<u>TAAGAAGGAGATATACAT</u> ATGTTATTTGCTTTACCT	HiGloB1
HiGloB1-r	GTGGTGGTGCCTCGAGGAACATATCTTTTGCTTTGC	
EcGloB2-f	<u>TAAGAAGGAGATATACAT</u> ATGAACTATCGTATTATT	EcGloB2
EcGloB2-r	GTGGTGGTGCCTCGAGCCAGACGGGCATTTCGTCTT	
PaGloB2-f	<u>TAAGAAGGAGATATACAT</u> ATGTCGACATCCCCCGCGCT	PaGloB2
PaGloB2-r	GTGGTGGTGCCTCGAGGCCCTGACGAAGGGGTTC	

14 <sup>a</sup>Underlined sequences indicate overlap sequences with pET30 Ec/Lic.



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16 **Fig. S1 Ion chromatography analysis of the product from GSSH oxidation by**  
 17 **CnSdoA.** The reactions were carried out in typical reaction mixtures with 1 mM  
 18 GSSH and (a) heat-inactivated CnSdoA, (b) CnSdoA, or (c) control without CnSdoA.



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20 **Fig. S2 Optimal of pH and temperature of SDOs.** The SDO activities were  
 21 determined at various pH values within the range of 6.4 to 7.8 in 100 mM KPi buffers  
 22 at 25 °C (A) and at different temperatures range of 25 °C to 45 °C at pH 7.4 (B).