Supporting Information

Detection and discrimination of sulfur dioxide using a

colorimetric sensor array

Chaoqiang Ding¹, Yan Ren¹, Xinyang Liu¹, Jingjing Zeng¹, Xinping Yu¹, Daxiang Zhou¹, Yanjie Li^{1,2*}

¹College of Biology and Food Engineering, Chongqing Three Gorges University, Wanzhou, Chongqing 404100, P.R. China. ²Engineering Technology Research Center for the Development and Utilization of Characteristic Biological Resources in Northeast Chongqing, Chongqing Three Gorges University, Wanzhou, Chongqing 404100, P.R. China.

*Corresponding author: Dr. Yanjie Li

College of biology and food engineering,

Chongqing Three Gorges University,

Wanzhou, Chongqing 404100,

P.R. China.

Tel.: +86 23 5810 2522;

Fax: +86 23 5810 2522;

E-mail address: 20060016@sanxiau.edu.cn (Y.J.Li).

^{*} To whom correspondence should be addressed. E-mail: 20060016@sanxiau.edu.cn

Figure/Table of contents

Composition of the developed 1×6 arrayTabl	e S1
Principle of specific reaction with sulfur dioxide and limits of detection of six prepared dyes	
based on spectrophotometryFigur	e S1
Comparison of LOD of sulfur dioxide by different common methodsTable	e S2
Color difference maps of different interferents with and without sulfur dioxide Figur	e S2
A PCA plot of different interferents with and without sulfur dioxide Figur	re S3
A PCA diagram of sulfur dioxide with different concentrations Figur	e S4
Difference maps of <i>fritillaria</i> samples with different concentrations of sulfur dioxideFigure	e S5

Table S1.	Composition	of the develope	$d 1 \times 6 array$
-----------	-------------	-----------------	----------------------

1	Sulfur dioxide kit(Based on the principle of hydrochloric para-rosaniline)			
2	80µg/mL Brilliant Green in 10mM PBS(pH=7)			
3	46.34µg/mL Malachite Green in 10mM PBS(pH=7)			
4	55µg/mL Basic Fuchsin in 1.47mM Borax buffer solution(pH=9.16)			
5	$1.34 \times 10^{-2} \mu g/mL$ o-Phthalaldehyde + $3.85 \times 10^{-3} \mu g/mL$ ammonium acetate in 4mM potassium dihydrogen phosphate buffer (pH=6.6)			
6	$\frac{3.6 \times 10^{-3} \mu g/mL \ 1,10-Phenanthroline, 3.24 \times 10^{-3} \mu g/mL \ Iron(III) \ chloride + CTAB(0.5\%)}{in \ 0.5Mm \ Sodium \ acetate \ anhydrous \ solution(pH=5.5)}$			

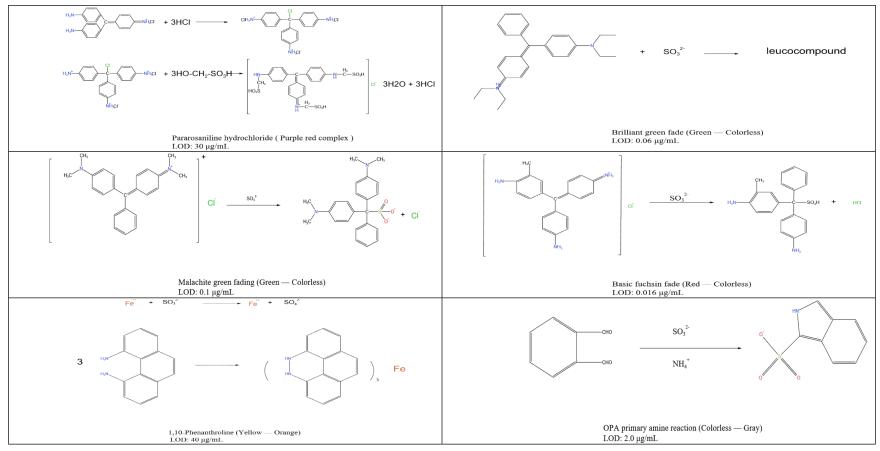


Figure S1.Principle of specific reaction with sulfur dioxide and limits of detection of six prepared dyes based on spectrophotometry.

Note: The above six reactions are based on the specific color reaction with sulfite.

Methods	LOD (µg/mL)	References
Spectrophotometry	4.0	Ref [15]
Acid-base titration	1.50	Ref [16]
Fluorescence method	6.29×10 ⁻³	Ref [17, 18]
Gas chromatography-mass spectrometry	1.5×10 ⁻³	Ref [19]
High-performance liquid chromatography	0.30	Ref [20]
Electrochemical analysis	0.40	Ref [21]
Enzyme-linked immunosorbent assays	0.46	Ref [22]
Fourier-transform infrared spectroscopy	0.80	Ref [23]
Colorimetric sensor array	0.406	

Table S2. Comparison of detection limits of sulfur dioxide by different common methods

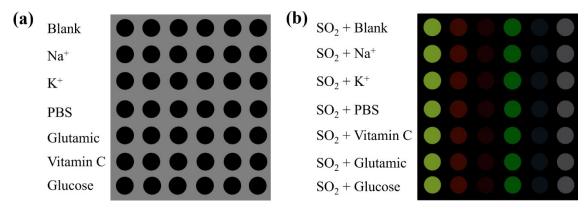


Figure S2. Color difference maps of different interferents with and without sulfur dioxide. (a) Difference maps of different interferents with final concentration of 100 μ g/mL. (b) Difference maps of different interferents with final concentration of 100 μ g/mL and sulfur dioxide with final concentration of 10 μ g/mL.

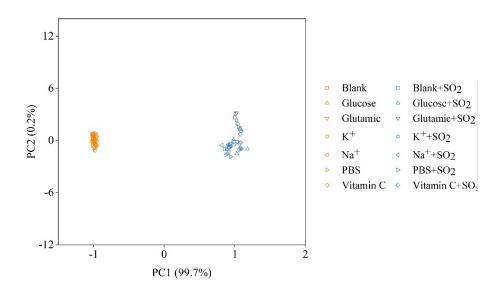


Figure S3. A PCA plot of different interferents with and without sulfur dioxide.

Note: interferents (e.g., Blank, Na⁺, K⁺, PBS, vitamin C, glutamic acid, glucose, final concentration 100 μ g /mL), sulfur dioxide (final concentration 100 μ g/mL). All of the experiments were performed in quintuplicate.

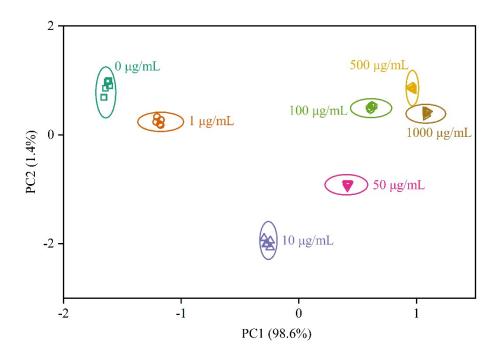


Figure S4. A PCA diagram of sulfur dioxide with different concentrations. All of the experiments were performed in quintuplicate.

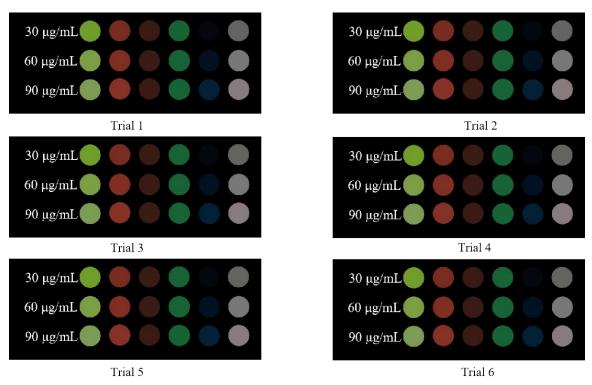


Figure S5. Difference maps of *fritillaria* samples with different concentrations of sulfur dioxide. All of the experiments were performed in sextuplicate.