

Correction to “Cyanine-Based Fluorescent Probe for Highly Selective Detection of Glutathione in Cell Cultures and Live Mouse Tissues”

Jun Yin, Younghee Kwon, Dabin Kim, Dayoung Lee, Gyoungmi Kim, Ying Hu, Ji-Hwan Ryu,* and Juyoung Yoon*

J. Am. Chem. Soc. 2014, 136, 5351–5358. DOI: 10.1021/ja412628z

Supporting Information

Pages 5352–5253. The conditions for detection in vitro should be corrected owing to the instrument failure.¹

All measurements of probes (5 μ M) were performed with amino acids (500 μ M) in PBS buffer solution (10 mM, pH 7.36) containing 0.5% DMSO ($\lambda_{\text{ex}} = 635$ nm, $\lambda_{\text{em}} = 785$ nm, slit: 10/10 nm). Figure 2, shown here, reflects those corrections, and the Supporting Information has also been

corrected. Other related descriptions of fluorescence spectra followed the above conditions.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/jacs.6b05099.

UV/vis absorption and fluorescence spectra of probes 1 and 2, mass spectra after treatment of probes 1 and 2 with thiols, fluorescence images of mouse tissues, frontier molecular orbital profiles, and ¹H NMR, ¹³C NMR, and MS spectra of 1, 2, and 3 (corrected) (PDF)

ACKNOWLEDGMENTS

Our study of reaction of probe 2 with glutathione has been supported by NUS (National University of Singapore) in Singapore. We also thank them for using our probe in their research.

REFERENCES

(1) Li, L.; Wijaya, H.; Samanta, S.; Lam, Y.; Yao, S. Q. *Sci. Rep.* 2015, 5, 11522.

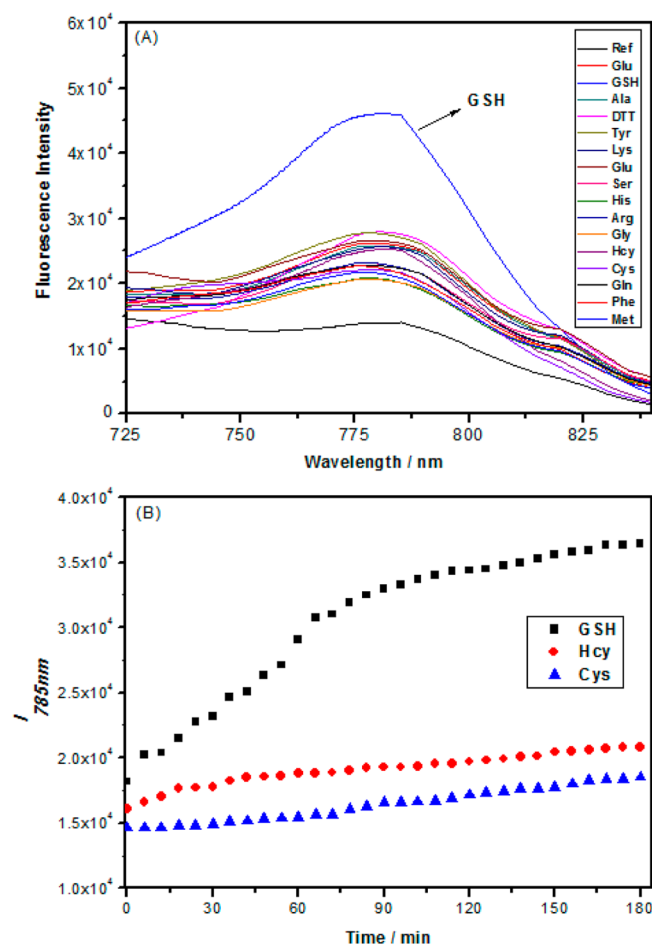


Figure 2. (A) Fluorescence response of probe 2 (5 μ M) to various amino acids (500 μ M). Each spectrum was recorded 180 min following addition of the amino acid. (B) Time-dependent fluorescence changes of probe 2 (5 μ M) upon addition of GSH, Cys, and Hcy (500 μ M) in PBS (10 mM, pH 7.36) containing 0.5% DMSO. $\lambda_{\text{ex}} = 635$ nm, $\lambda_{\text{em}} = 785$ nm, slit: 10/10 nm.

Published: June 1, 2016