

A ratiometric selective fluorescent probe derived from pyrene for Cu²⁺ detection

Chunwei Yu ^{1,4,5†}, Mei Yang ^{2†}, Shuhua Cui ³, Yuxiang Ji ¹ and Jun Zhang ^{1,4,5*}

¹ *School of Tropical Medicine, Hainan Medical University, Haikou 571199, China; hy0211049@hainmc.edu.cn (C.Y.); hy0211012@hainmc.edu.cn (Y.J.)*

² *School of Public Health and One Health, Hainan Medical University, Haikou 571101, China; yang24364@hainmc.edu.cn (Yang, M.)*

³ *School of Chemical Engineering and Environment, Weifang University of Science and Technology, Shouguang 262700, China; cuishuhua@wfust.edu.cn (Cui, S)*

⁴ *Key Laboratory of Tropical Translational Medicine of Ministry of Education, Hainan Medical University, Haikou 571199, China;*

⁵ *NHC Key Laboratory of Control of Tropical Diseases, School of Tropical Medicine, Hainan Medical University, Haikou 571199, China;*

** Correspondence: hy0211045@hainmc.edu.cn (Zhang, J)*

† These authors contributed equally to this work.

Figure S1 ESI-MS spectra of P

Figure S2 ¹H NMR spectra of P

Figure S3 ¹³C NMR spectra of P

Figure S4 ESI-MS spectra of P₁

Figure S5 Fluorescence response of P(1.0 μM) to 10 μM of Cu²⁺ or the mixture of 50 μM of other metal ions with 10 μM of Cu²⁺.

Figure S6 Job's plot for determining the stoichiometry of P and Cu²⁺. The total concentration was kept 10 μM.

Figure S7 Benesi-Hildebrand plot of P, assuming 1:1 stoichiometry for association between P and Cu²⁺.

Figure S8 ESI-MS spectra of P-Cu²⁺.

Figure S9 Cell viability values (%) estimated by MTT proliferation test versus incubation concentrations of P. HeLa cells were cultured with P in the concentration of 0–10 μM.

Figure S1

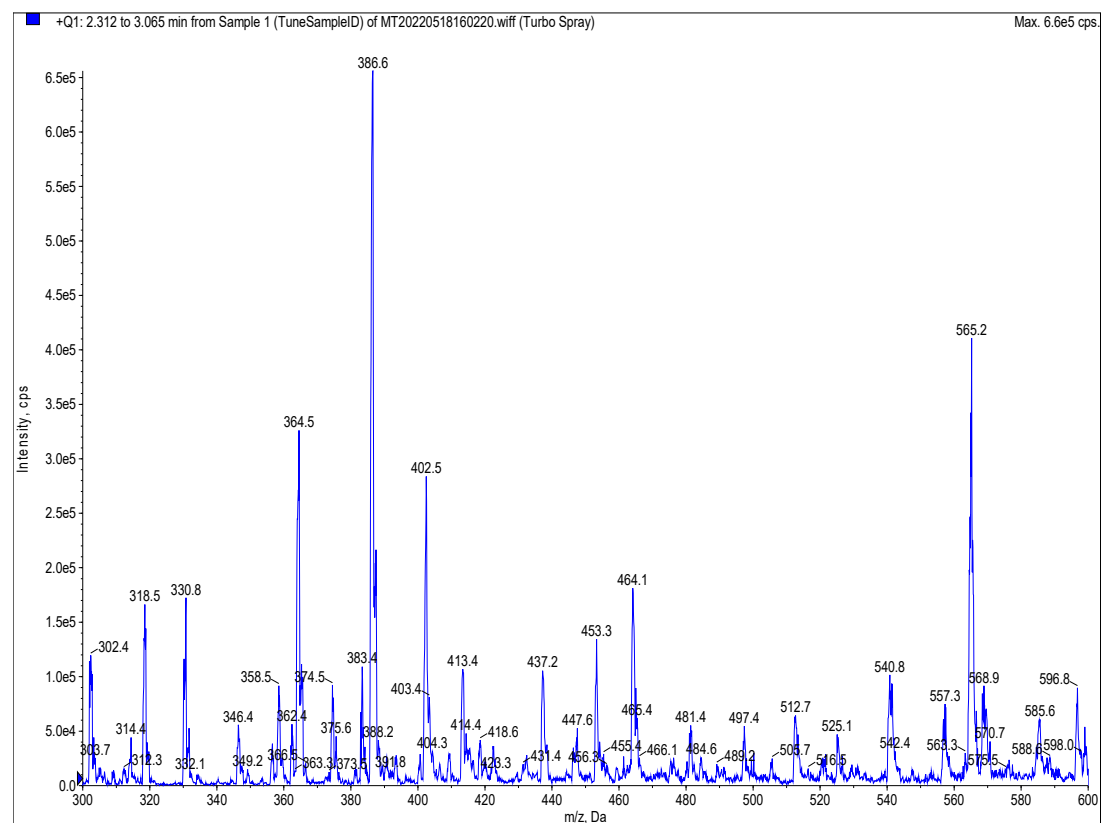


Figure S2

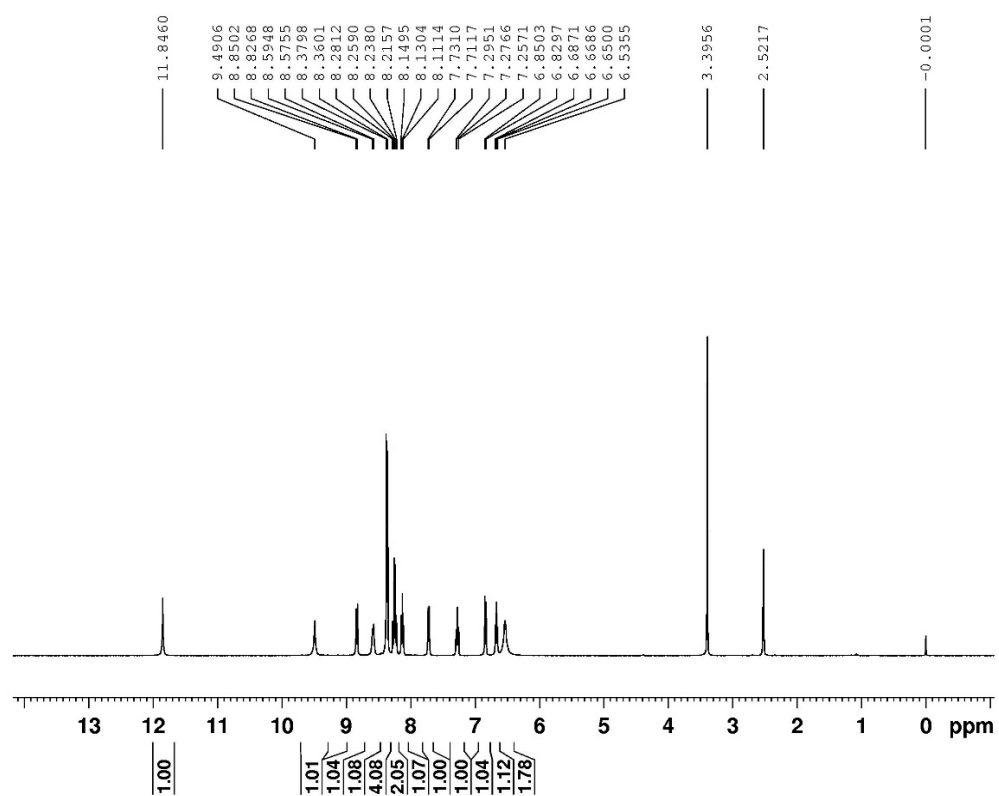


Figure S3

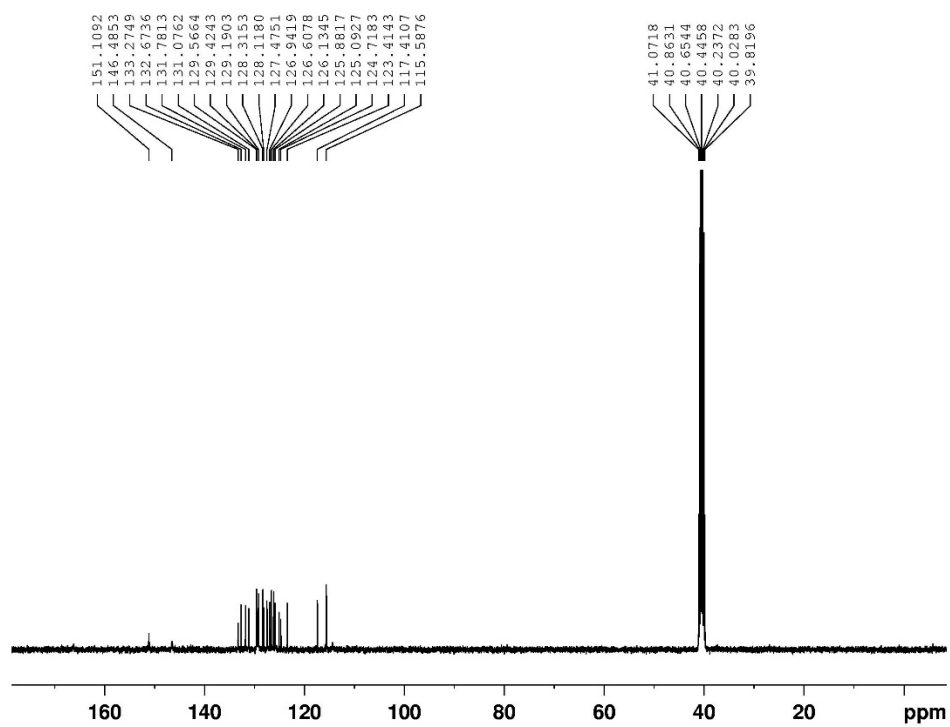


Figure S4

3 #17 RT: 0.27 AV: 1 NL: 5.52E3
F: ITMS + c ESI Full ms [50.00-2000.00]

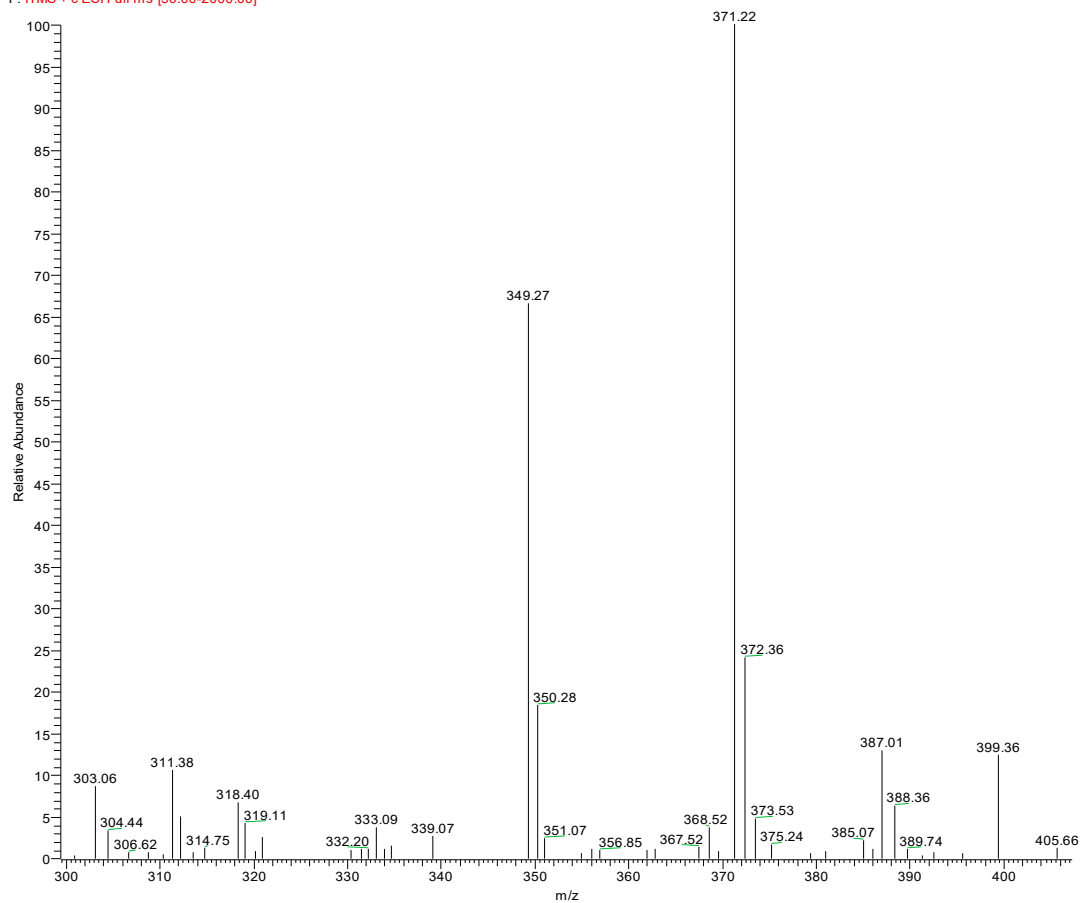


Figure S5

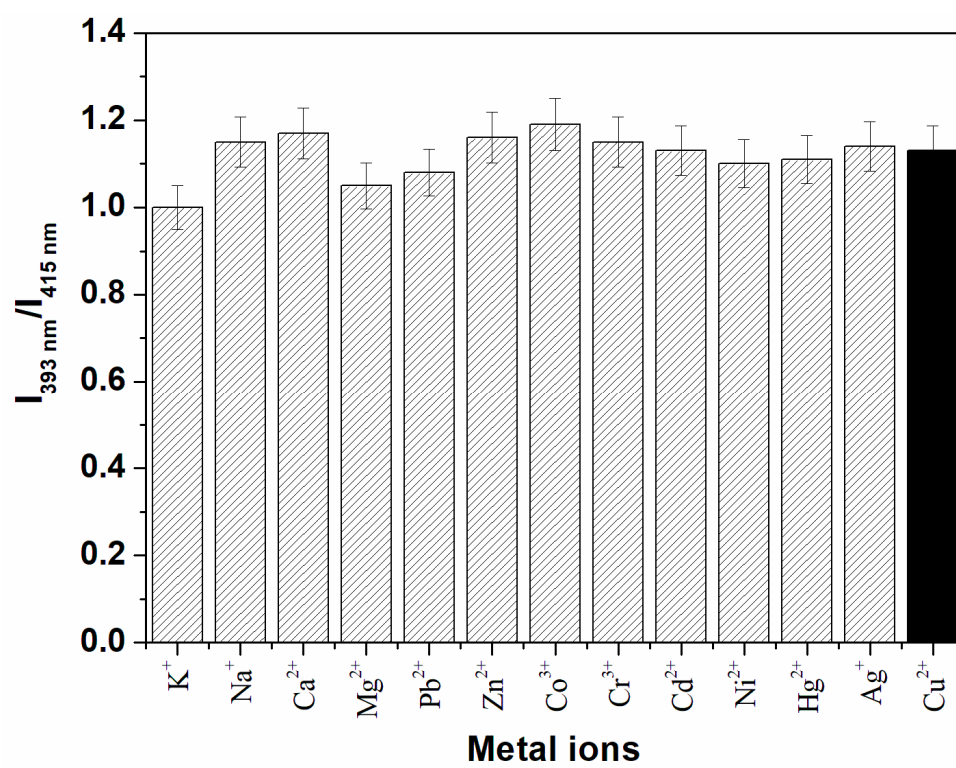


Figure S6

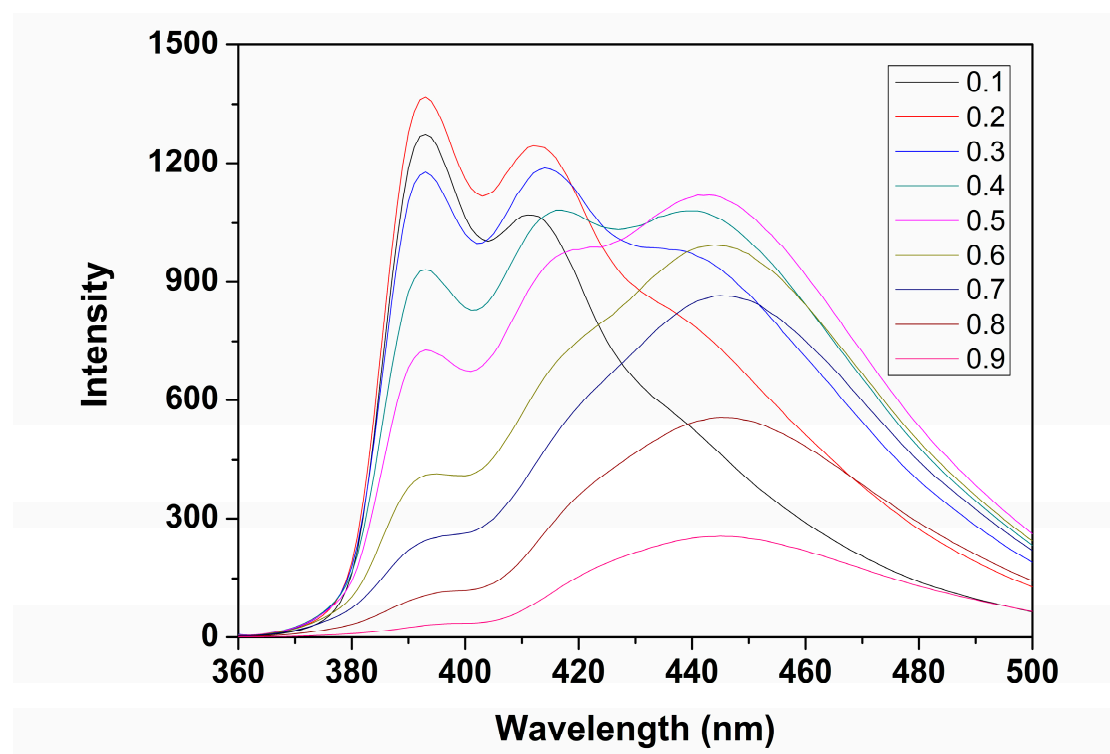


Figure S7

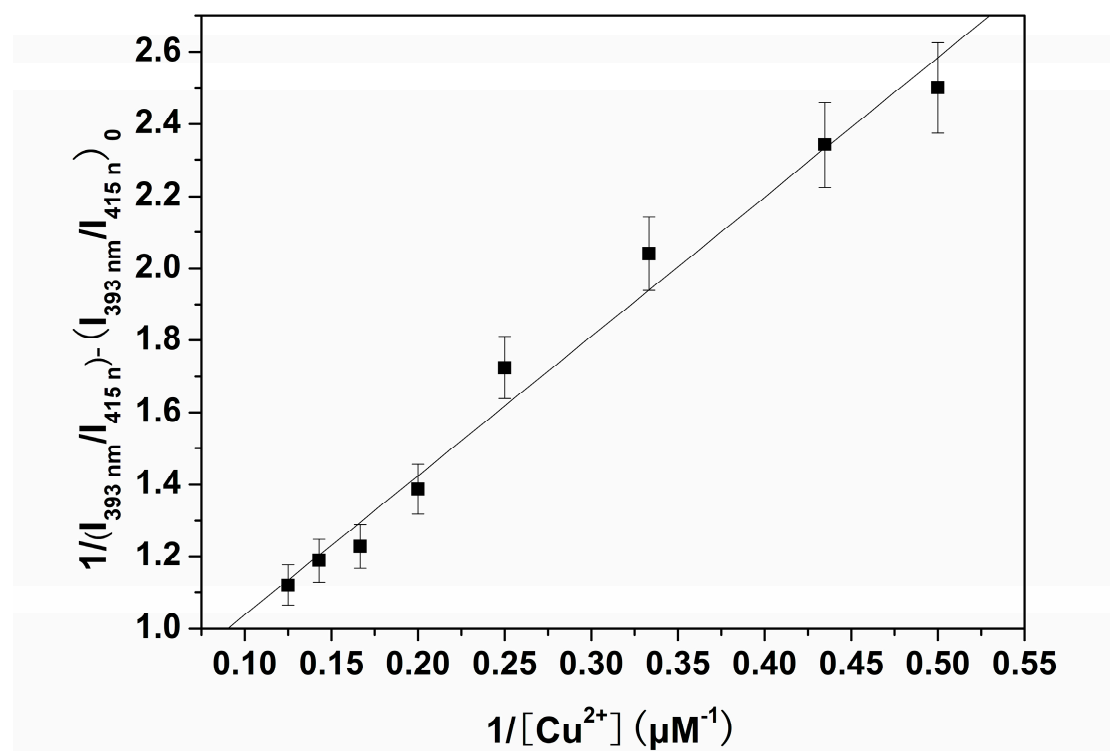


Figure S8

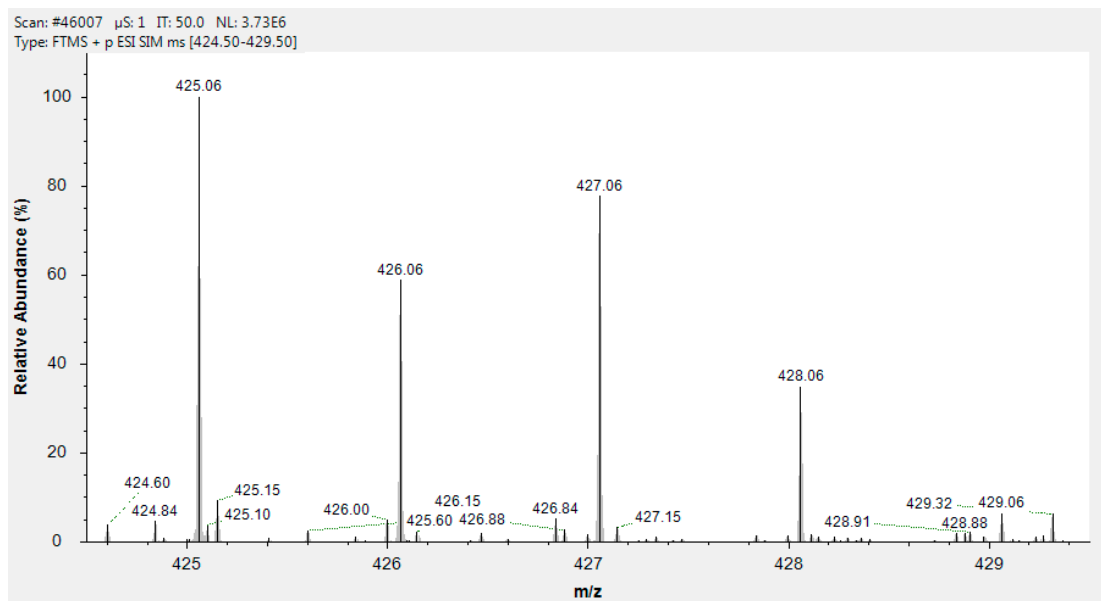


Figure S9

