

## Distinguish cancer cells from normal cells by an organelle-targeted fluorescent marker

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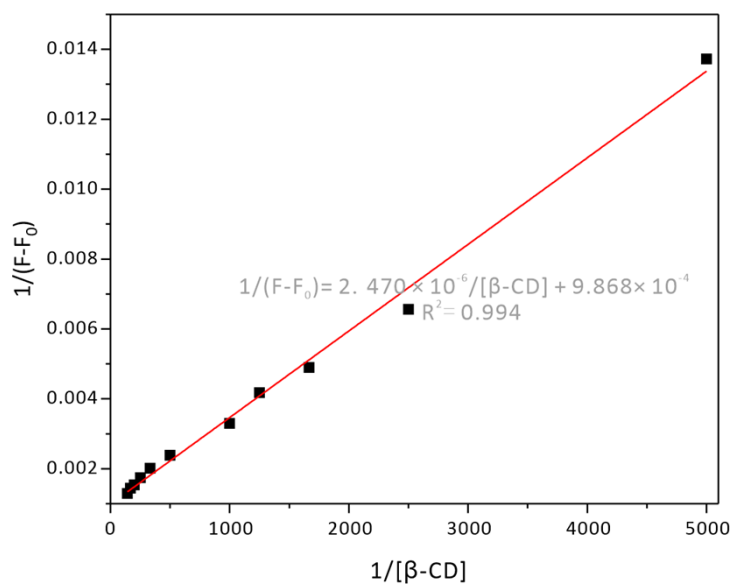
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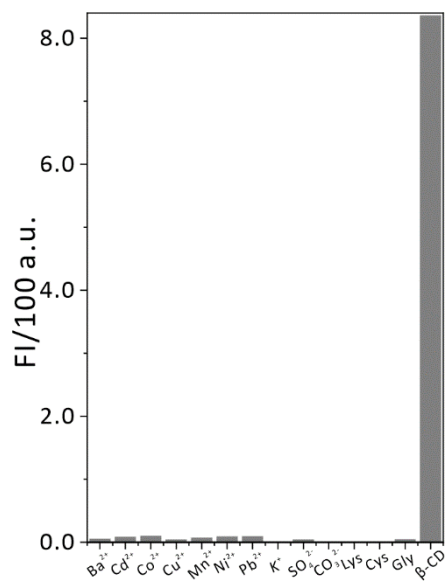
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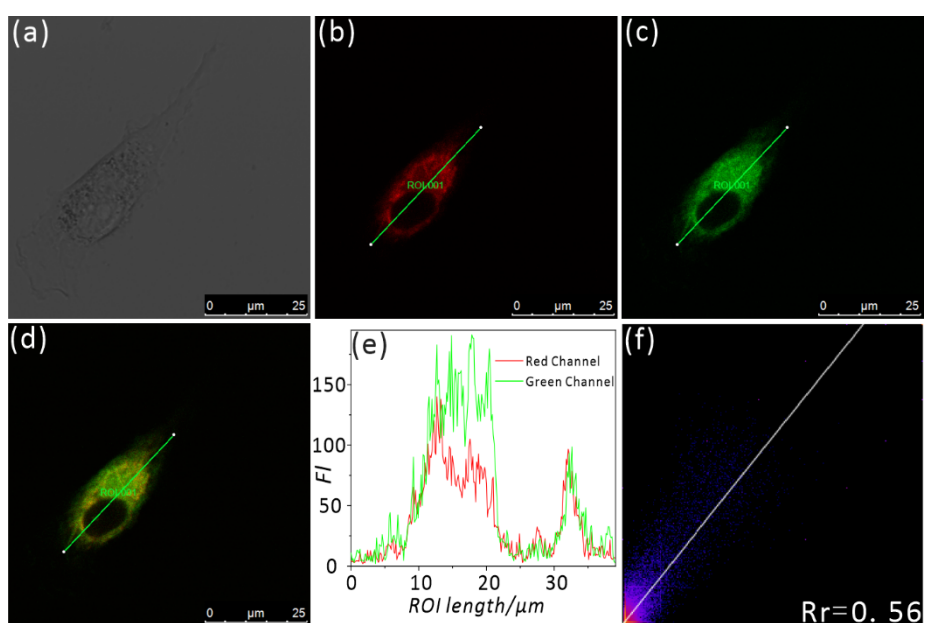
## 1. FIGURES



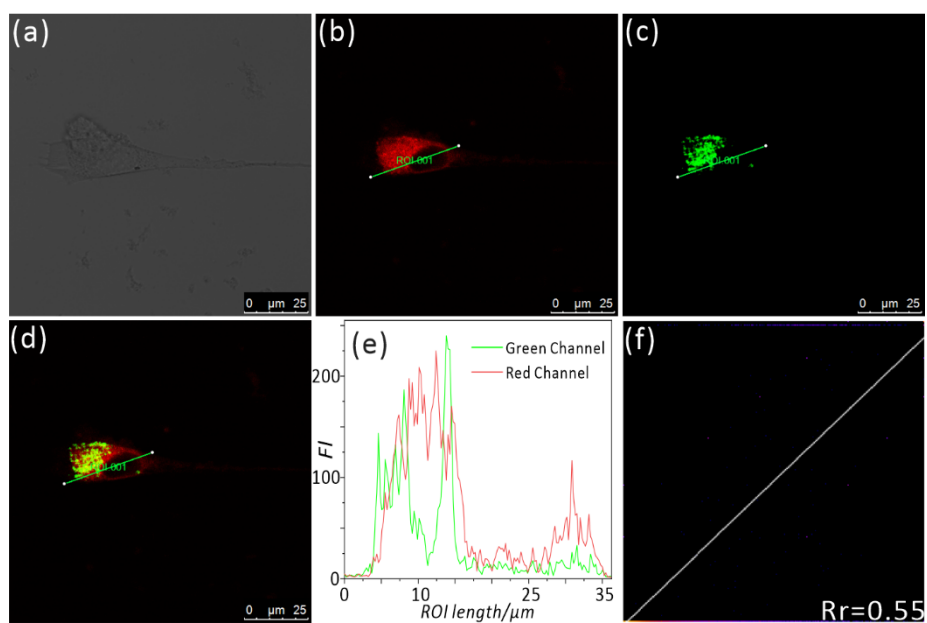
**Fig. S1** Double reciprocal plots:  $1/(F-F_0)$  versus  $1/[\beta\text{-CD}]$ .



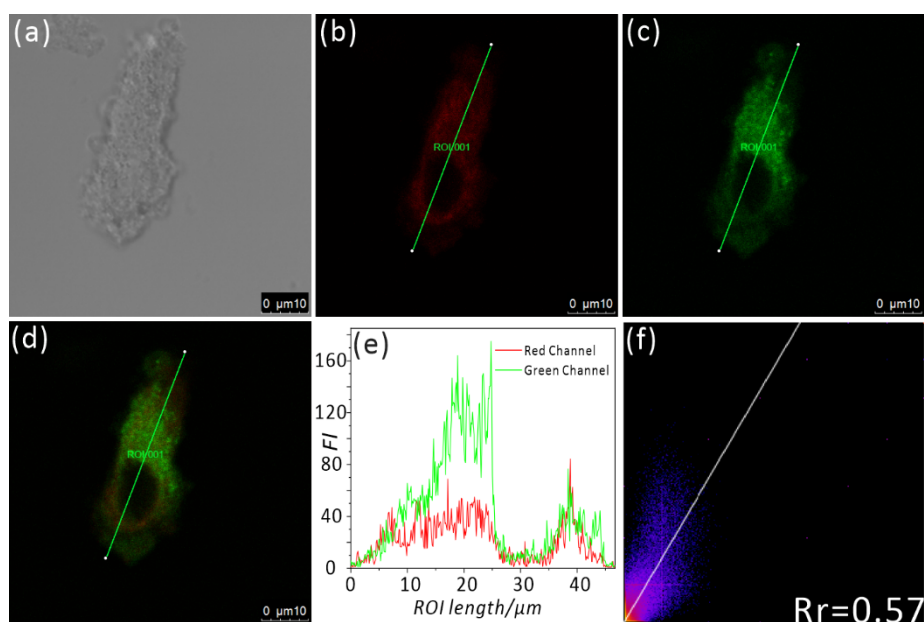
**Fig. S2** The fluorescence intensity of probe **1** (10 μM) towards other interfering ions and biomolecules (1 mM) in double-distilled water containing 10% DMSO. ( $\lambda_{\text{ex}} = 509$  nm, slit widths: 3 nm/5 nm).



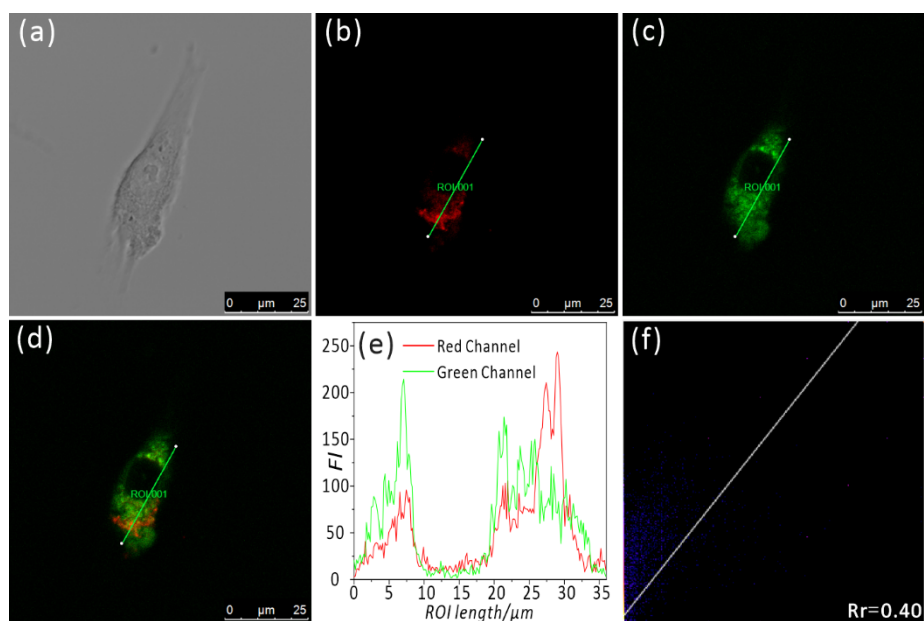
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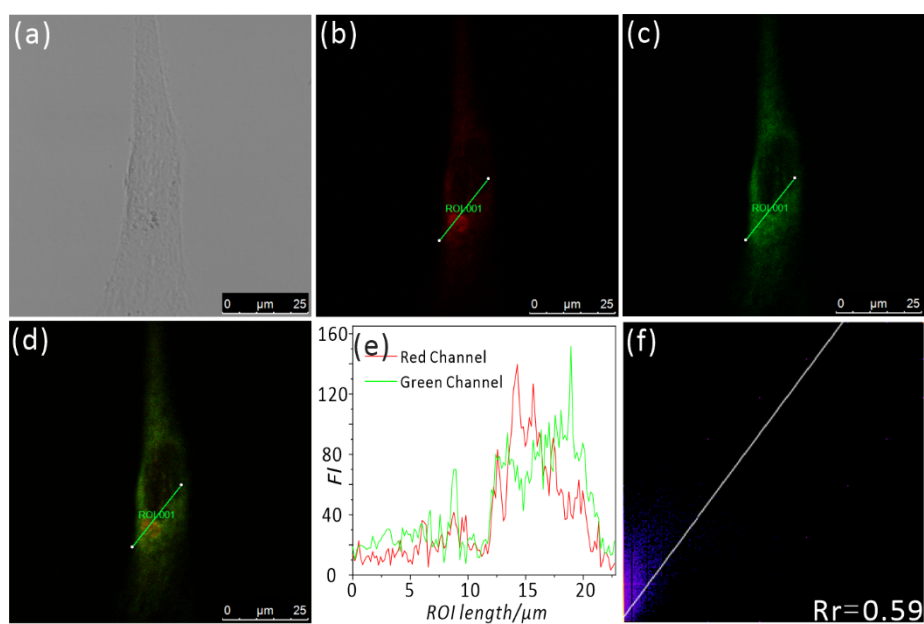
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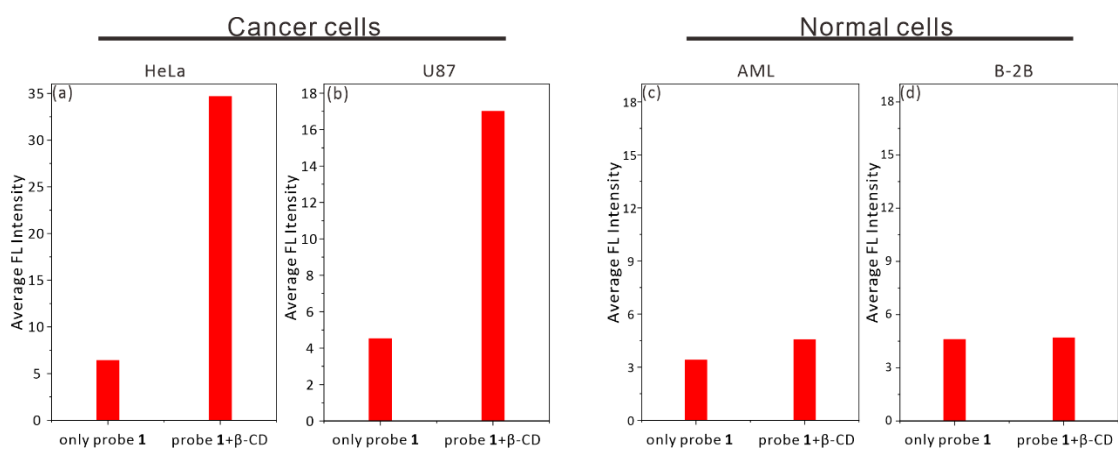
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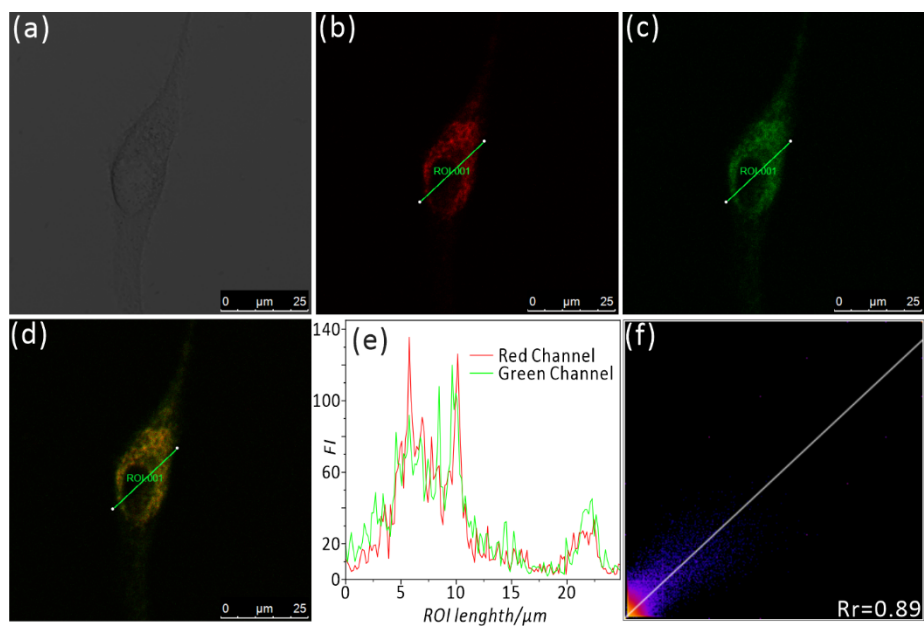
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**Fig. S7** Fluorescence images of B-2B cells costaining with probe **1** (3 μM) and Green ER-Tracker (3 μM). (a) Bright-field image; (b) confocal image (red channel) with probe **1**; (c) confocal image (green channel) of Green ER-Tracker; (d) merged image of (b) and (c); (e) fluorescence intensity of the regions of interest (ROI) across the B-2B cells. (f) fluorescence intensity correlation plot from the red channel and the green channel.



**Fig. S8** Average FL Intensity of cancer cells and normal cells incubated with probe **1** before and after adding β-CD (100 μM).



**Fig. S9** Fluorescence images of HeLa cells costaining with probe **1** (0.3 μM) and Green ER-Tracker (3 μM). (a) Bright-field image; (b) confocal image (red channel) with probe **1** (0.3 μM) after adding β-CD (100 μM); (c) confocal image (green channel) of Green ER-Tracker; (d) merged image of (b) and (c); (e) fluorescence intensity of the regions of interest (ROI) across the HeLa cells. (f) fluorescence intensity correlation plot from the red channel and the green channel.

## 2. APPENDIX

<sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>)

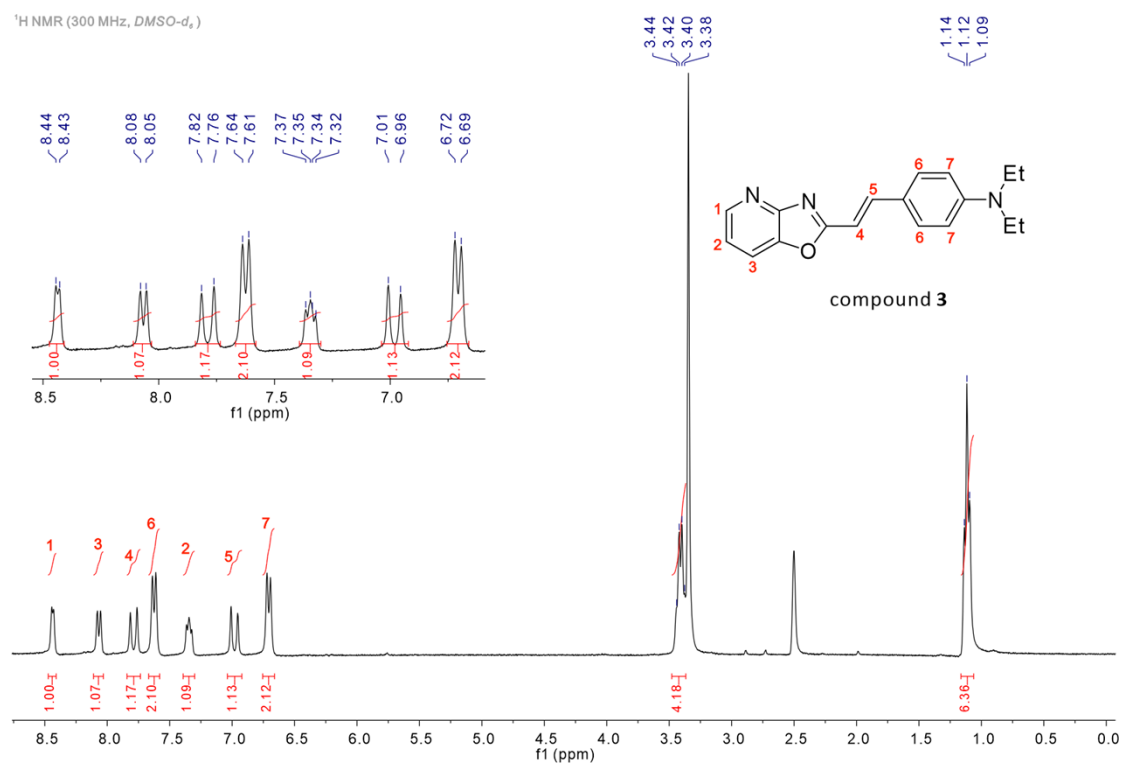


Fig. S10 <sup>1</sup>H NMR spectrum of compound 3 (300 MHz, DMSO-d<sub>6</sub>).

<sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>)

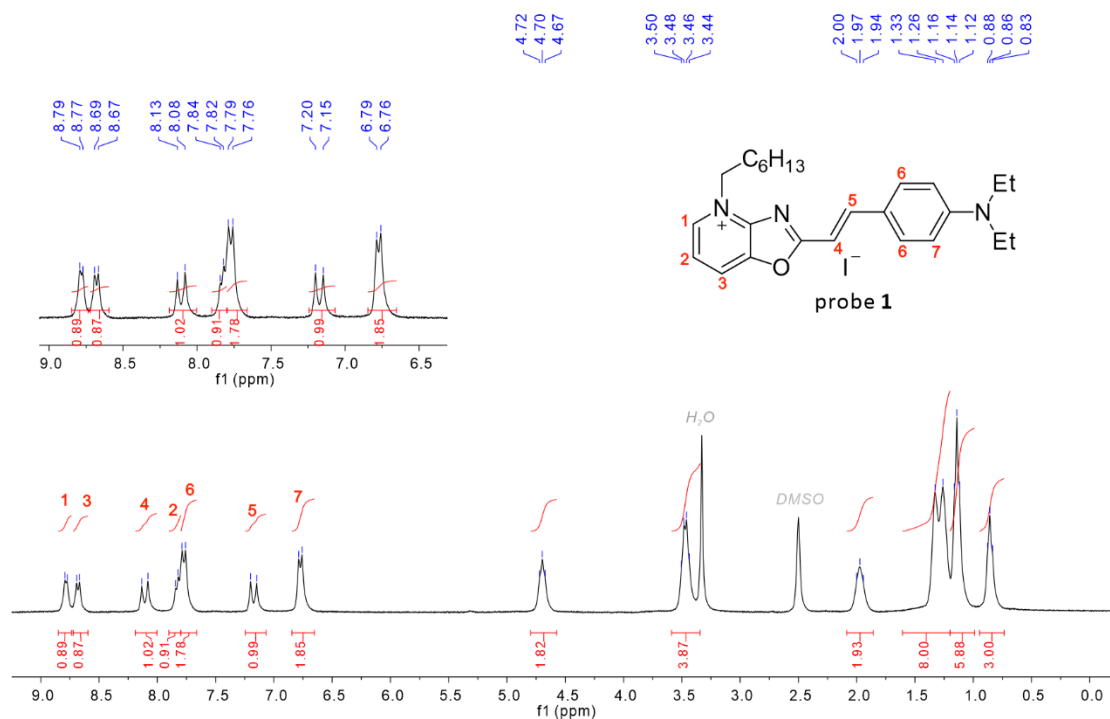


Fig. S11 <sup>1</sup>H NMR spectrum of probe 1 (300MHz, DMSO-d<sub>6</sub>).



**Analysis Info**

Analysis Name D:\Data\wlf\stu-sample\20211019\HHC-13\_BB7\_01\_21474.d  
Method 2011-lcms-ms\_20211202.m  
Sample Name HHC-13  
Comment

Acquisition Date 12/22/2021 10:27:20 PM

Operator bruker  
Instrument micrOTOF-Q III 8228888.20487

**Acquisition Parameter**

Source Type	ESI	Ion Polarity	Positive	Set Nebulizer	0.6 Bar
Focus	Active	Set Capillary	4500 V	Set Dry Heater	180 °C
Scan Begin	50 m/z	Set End Plate Offset	-500 V	Set Dry Gas	4.0 l/min
Scan End	1500 m/z	Set Collision Cell RF	250.0 Vpp	Set Divert Valve	Waste

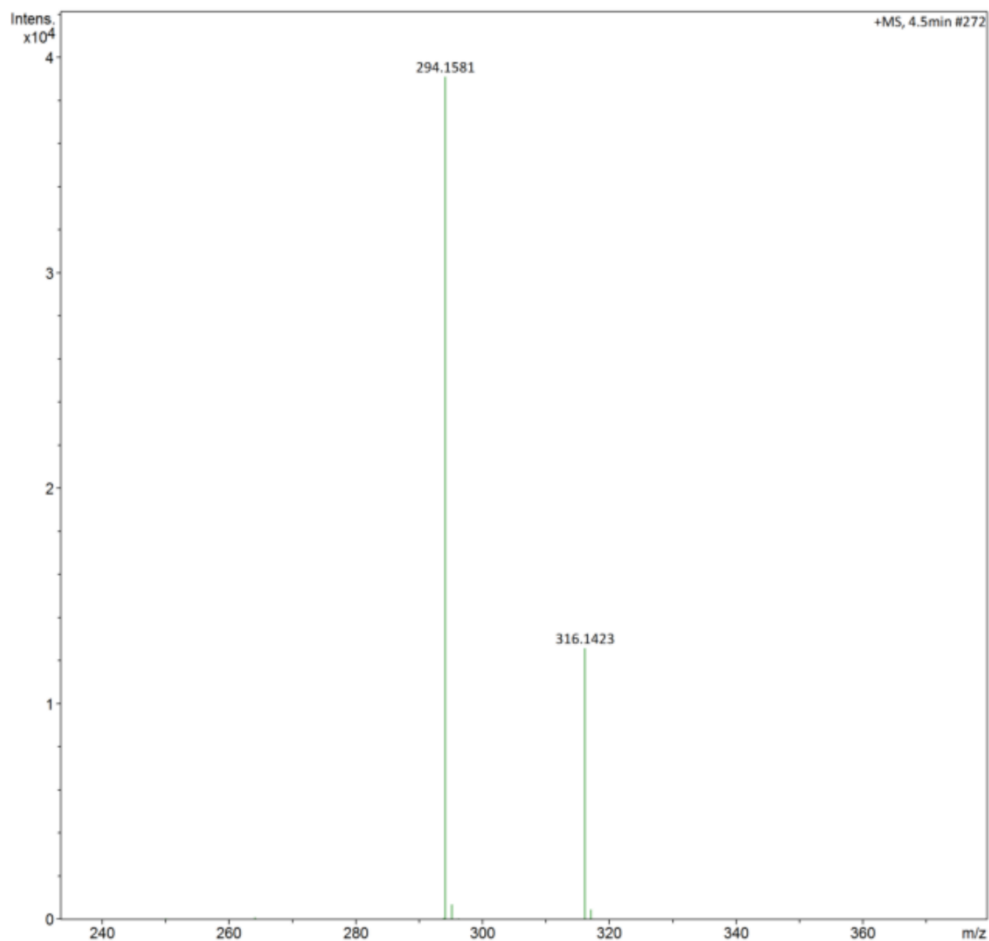


Fig. S12 HRMS (ESI<sup>+</sup>) spectrum of compound 3.

Acquisition Parameter					
Source Type	ESI	Ion Polarity	Positive	Set Nebulizer	0.6 Bar
Focus	Active	Set Capillary	4500 V	Set Dry Heater	180 °C
Scan Begin	50 m/z	Set End Plate Offset	-500 V	Set Dry Gas	4.0 l/min
Scan End	1500 m/z	Set Collision Cell RF	250.0 Vpp	Set Divert Valve	Waste

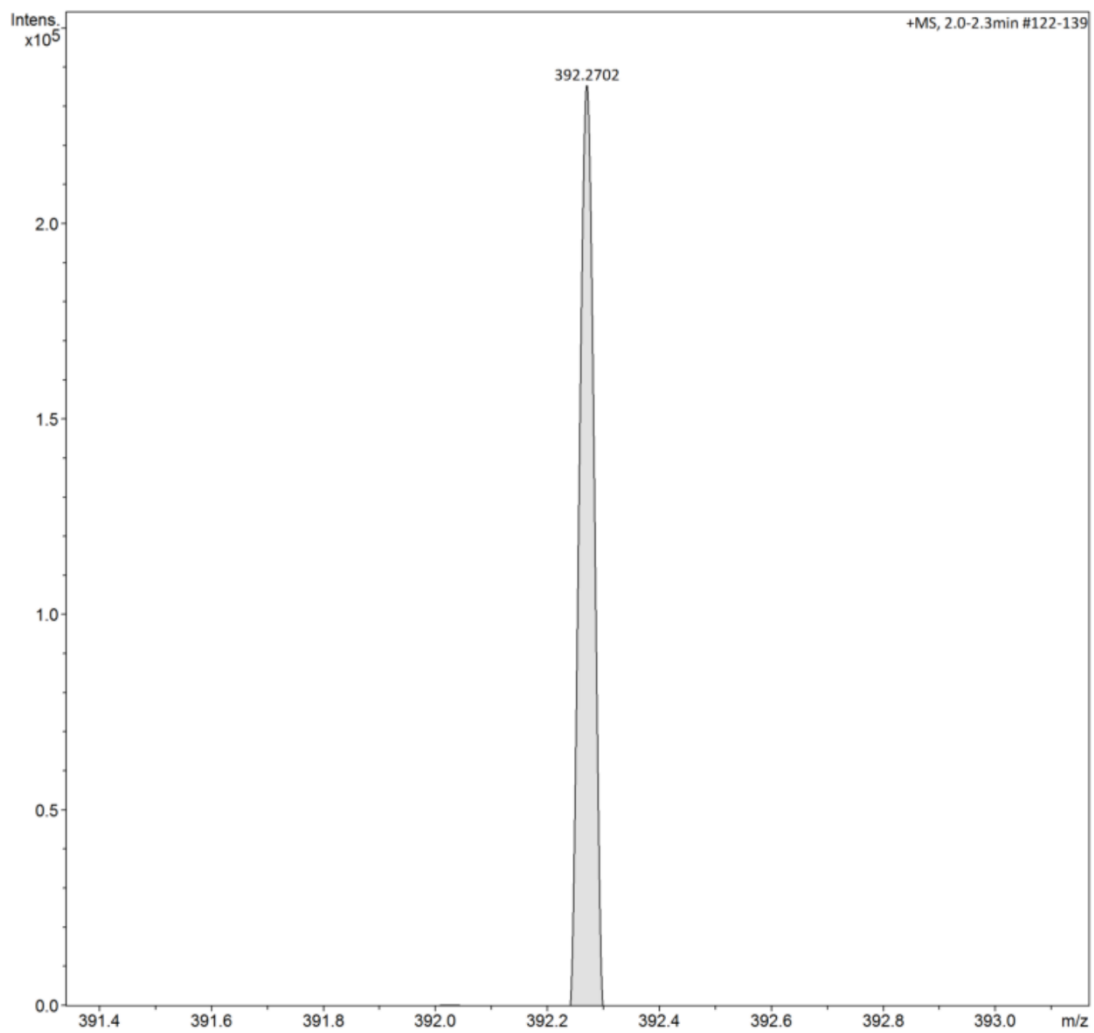


Fig. S13 HRMS (ESI<sup>+</sup>) spectrum of probe 1.

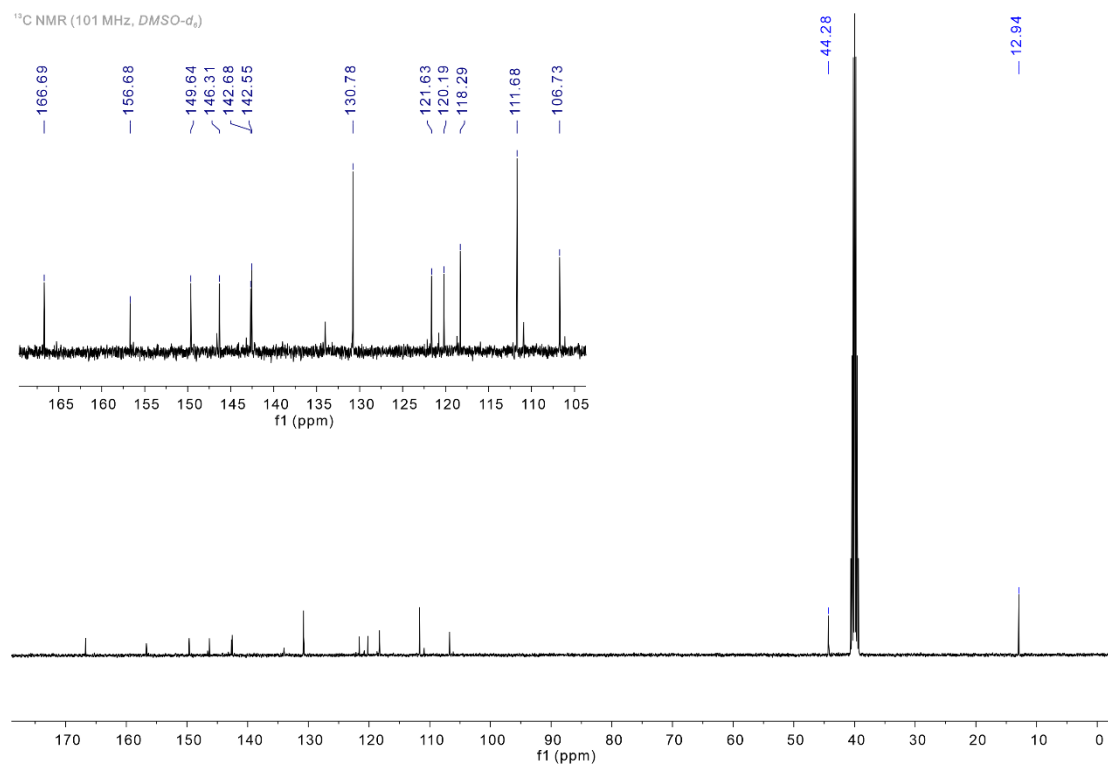


Fig. S14 <sup>13</sup>C NMR spectrum of compound **3** (101MHz, DMSO-d<sub>6</sub>).

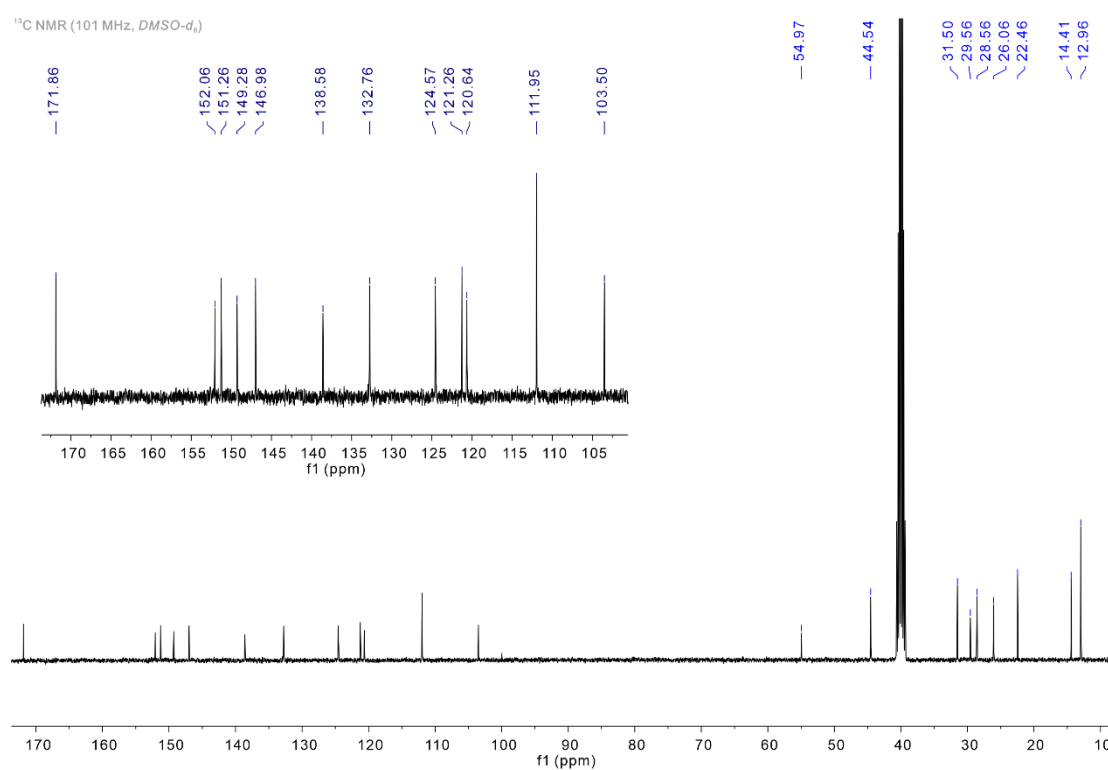


Fig. S15 <sup>13</sup>C NMR spectrum of probe **1** (101 MHz, DMSO-d<sub>6</sub>).