Supplementary Material

A novel coumarin derivatives-modified cellulose fluorescent probe for selective and sensitive detection of CN⁻ in food sample

Jiali Kou, Zhiyuan Meng, Xiaoyuan Wang, Zhonglong Wang*, Yiqin Yang* Co-Innovation Center of Efficient Processing and Utilization of Forest Resources, International Innovation Center for Forest Chemicals and Materials, College of Light Industry and Food, College of Chemical Engineering, Nanjing Forestry University, Nanjing, 210037, China

*Corresponding author: Dr. Zhonglong Wang

Phone: +86-25-85428369, Fax: +86-25-85428369

Email: wang_zhonglong@njfu.edu.cn

*Corresponding author: Dr. and Prof. Yiqin Yang

Phone: +86-25-85427118, Fax: +86-25-85427118

Email: wsfyyq@njfu.edu.cn

1. Experimental instruments

Fourier transform-infrared (FT-IR) spectra were recorded on Nicolet spectrometer using powder-pressed KBr. UV-*vis* absorption spectra were obtained on Shimadzu UV-2450 spectrophotometer, and fluorescence spectra were acquired on PerkinElmer LS55 spectrophotometer. ¹H NMR spectra were gained on Bruker AV 600 spectrometer in DMSO- d_6 , and ESI-MS spectra were measured on LTQ Orbitrap XL mass spectrometer. XPS spectra data were obtained on Shimadzu AXIS UltraDLD Xray photoelectron spectrometer with monochromatic Al Ka radiation (hv = 1486.6 eV, 600 W). The element compositions of chemicals were measured on PE2400II organic elemental analyzer (EA). The microscopic morphologies of samples were obtained using Quanta 200 scanning electron microscope (SEM). X-ray diffraction (XRD) patterns were determined with Rigaku Ultima IV horizontal X-ray diffractometer. Thermogravimetric data were collected on Netzsch TGA-209 F1 thermogravimetric analyzer at a heating rate of 20 °C/min under nitrogen atmosphere over a range of 25-800 °C.

2. Figures



Scheme S1. The synthesis route of ADC and FBC.

- Fig. S1. ¹H NMR spectra of compound ADC.
- Fig. S2. ¹H NMR spectra of compound FBC.

Fig. S3. Fluorescence emission intensity of ADC and DCB excited at 365 nm.

Fig. S4. photostability of DCB-CA (1×10^{-4} g/mL) and DCB-CA + CN⁻(1×10^{-4}

g/mL +100.0 μM CN⁻) were measured in DMF solution.

Fig. S5. The HRMS spectra of DCB

Fig. S6. The HRMS spectra of DCB after reaction with CN⁻.

Fig. S7. Stress-strain curve of DCB-CA film.

Fig. S8. The relationship between the absorption intensity at 450 nm and immersion time of **DCB-CA** film in water.

Fig. S9. Photographs of the film being placed in the water for 48 h under day light (a) and 365 nm UV lamp (b).



Fig. S1 ¹H NMR spectra of compound ADC.



Fig. S2 ¹H NMR spectra of compound FBC.



Fig. S3 Fluorescence emission intensity of ADC and DCB excited at 365 nm.



Fig. S4 photostability of DCB-CA (1×10^{-4} g/mL) and DCB-CA + CN⁻ (1×10^{-4}

 $g/mL + 100.0 \ \mu M \ CN^{-}$) were measured in DMF solution.



Fig. S5 The HRMS spectra of DCB



Fig. S6 The HRMS spectra of DCB-CN.



Fig. S7 Stress-strain curve of DCB-CA film.



Fig. S8 Photographs of the film being placed in the water for 48 h under day light (a) and 365 nm UV lamp (b).



Fig. S9 The relationship between the absorption intensity at 450 nm and immersion time of DCB-CA film in water.

Table S1 Comparison of DCB-CA with other CN⁻ probes.

Sensors	Structure of probe	Methods	LOD (μM)	Food samples application	Materials	References
R		Fluorometry	1	No	Strips	1

P1	HOOC HOOC HOC HOC HN N HN HN HN HN CrizH2 HN NH HN CrizH2 HN NH	Fluorometry	8	Yes	Strips	2
L1	ОН N-N НО - CH HC CH HC СН НС	Fluorometry	0.95	Yes	Strips	3
DSS		Colormetric	5.81	No	No	4
AIN	O-N HO	Colormetric	12.3	No	No	5
NBF	F∑B√F OB O I	Fluorometry	2.23	Yes	No	6
ML'CT+L' L'CT		Colorimetric	6.0	No	No	7
DCB-CA		Fluorometry	0.58	Yes	Film	This work

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