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Supporting Information

A Naphthalimide Based Novel "Turn-On" Fluorescence Approach for Uric Acid Determination and Monitoring of Xanthine Oxidase Activity

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Fig. S1: Uv-Vis absorption spectra of S3.



Fig. S2: Fluorescence stability study of S3 after exposing to UV light at different time interval.



Fig. S3: Effect of pH over fluorescence emission of S3 in the presence of uric acid



Fig. S4: (A) Change in the fluorescence emission of probe S3 when different concentration of xanthine was added to probe solution. The concentration of enzyme remained fixed 15 U/mL with incubation time is 25 min. (B) Change in fluorescence emission of probe at different concentration of XO while the concentration of xanthine remained 300 μ M constant.



Fig. S5: (A) pH dependent fluorescence emission response of probe S3 in the presence of xanthine and XO. (B) Temperature dependent fluorescence emission response of probe S3 in the presence of Xanthine and XO (15 U/mL).



Fig. S6: Time dependent change in fluorescence emission of probe **S3** in response to enzymatic action. Where black, red and blue colour represents the addition of 2, 5 and 10 U/mL of XO.



Fig. S7: Molecular docking study of xanthine and probe S3 (A) Docking pose probe S3 with XO (B) Docking pose of xanthine with XO.





Fig. S11: ¹HNMR of S3



Fig. S13: HRMS data of S3