

## Supplementary Information

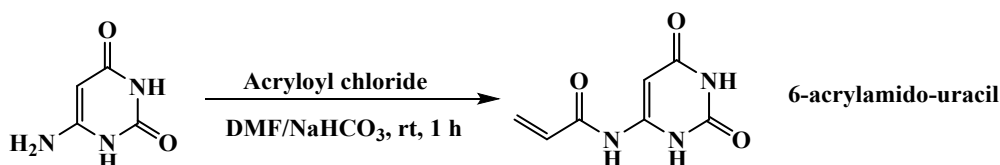
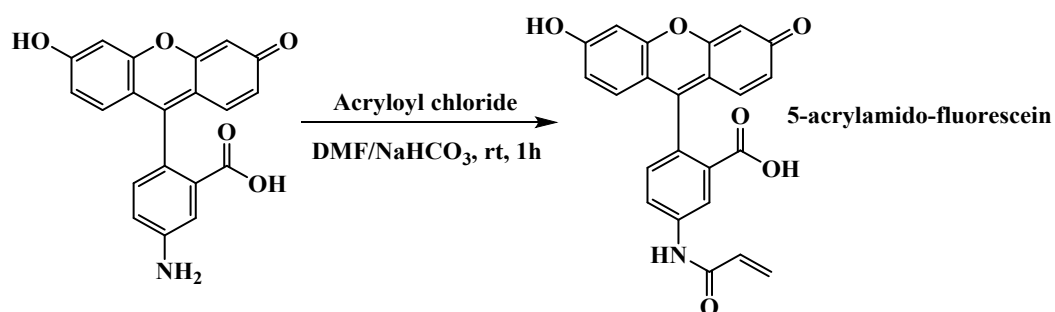
### Specific Visual-volumetric Sensor for Mercury Ion Based on Smart Hydrogel

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#### Step1



#### Step2

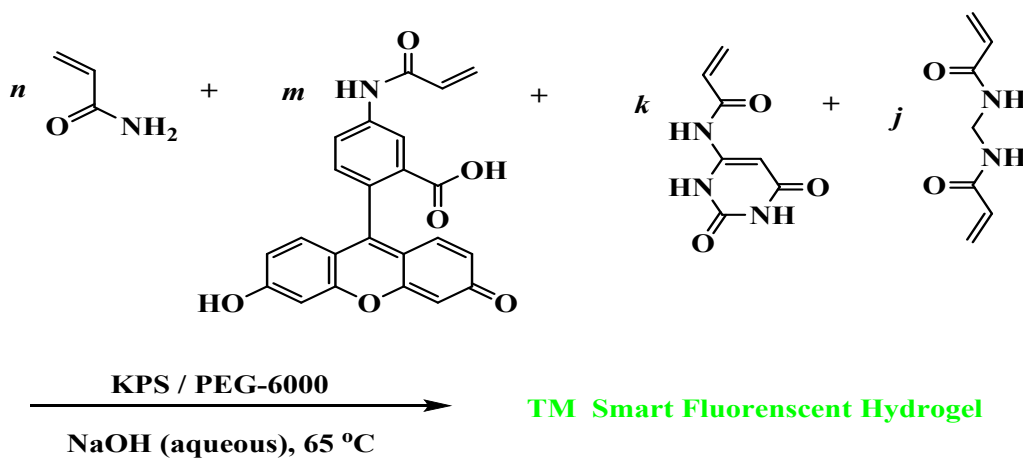


Fig. S1 Synthetic route of the smart fluorescent hydrogel

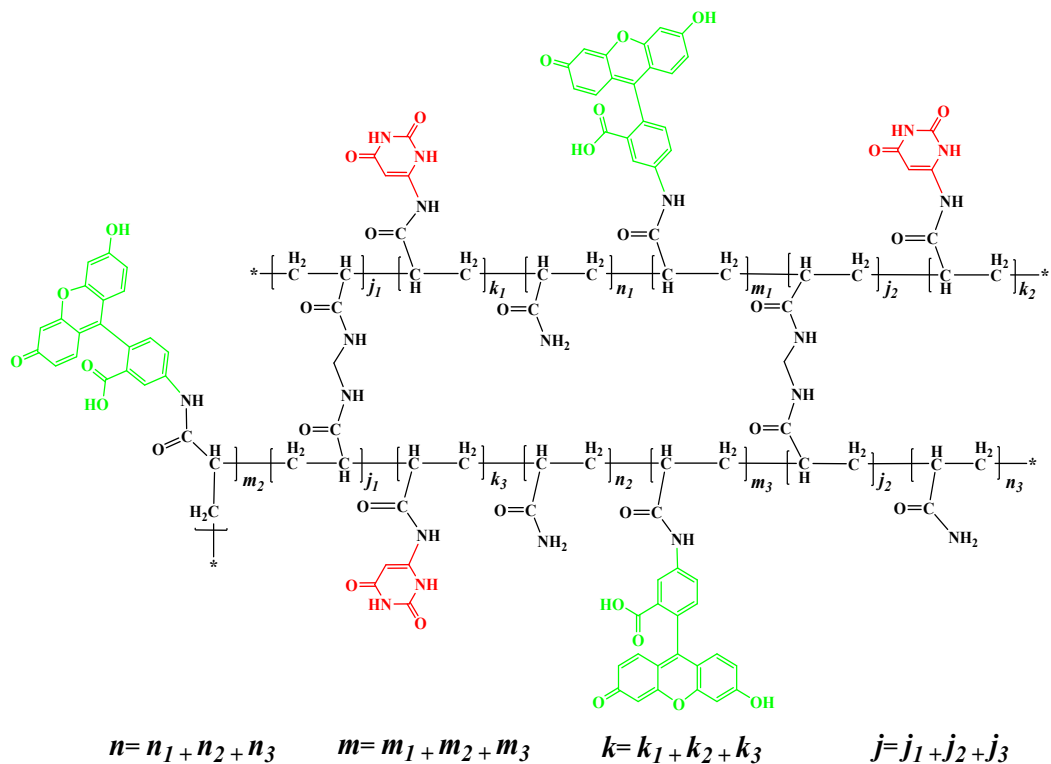


Fig. S2 The structure of the smart fluorescent hydrogel

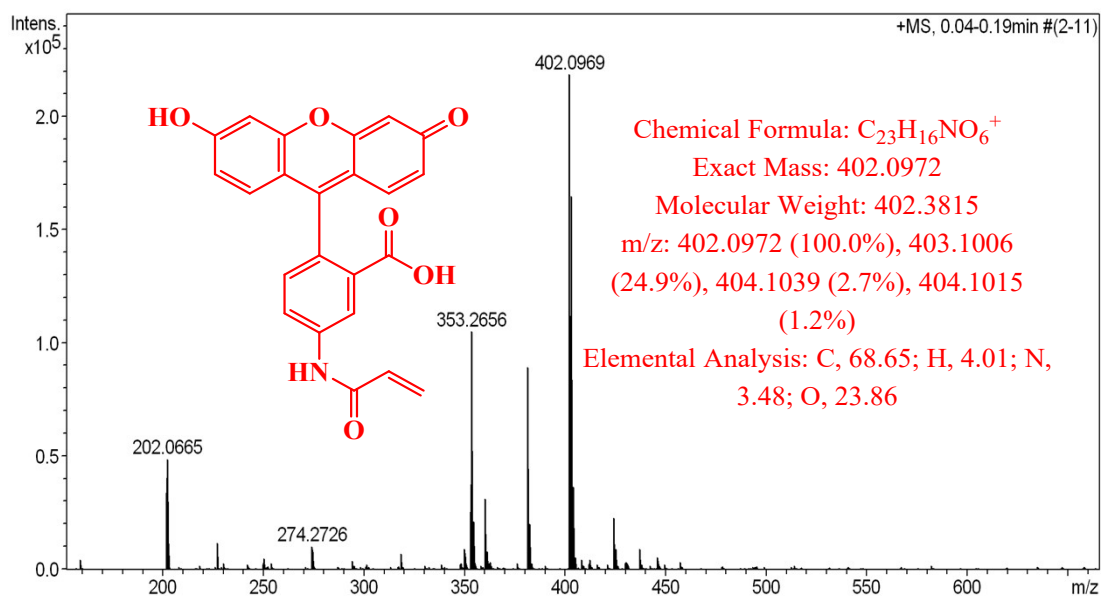


Fig. S3 Mass spectrum of the prepared 5-acrylamido-fluorescein

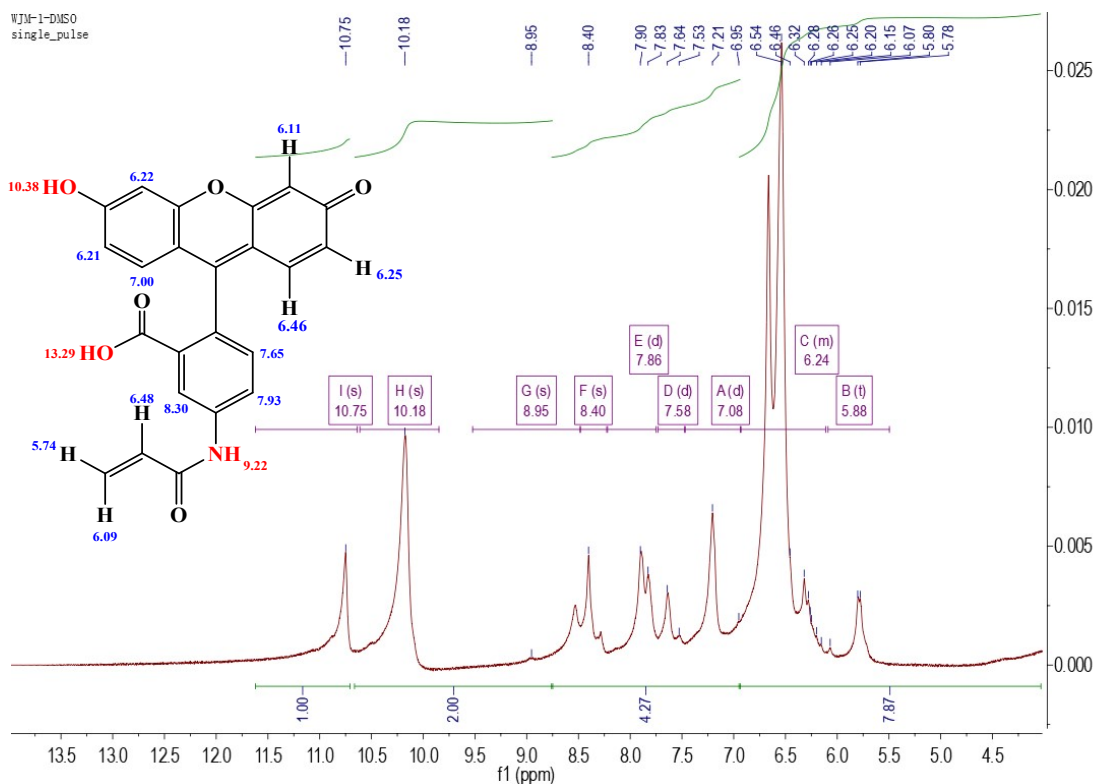


Fig. S4  $^1\text{H}$  NMR spectrum of 5-acrylamido-fluorescein

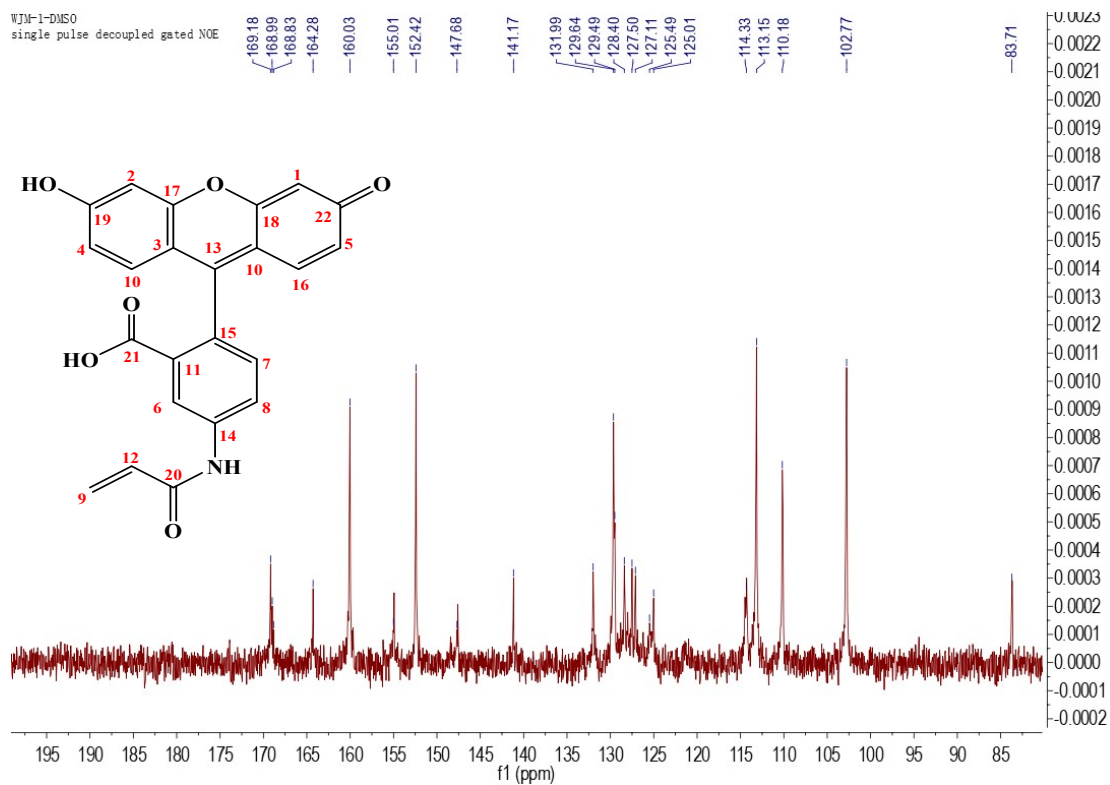
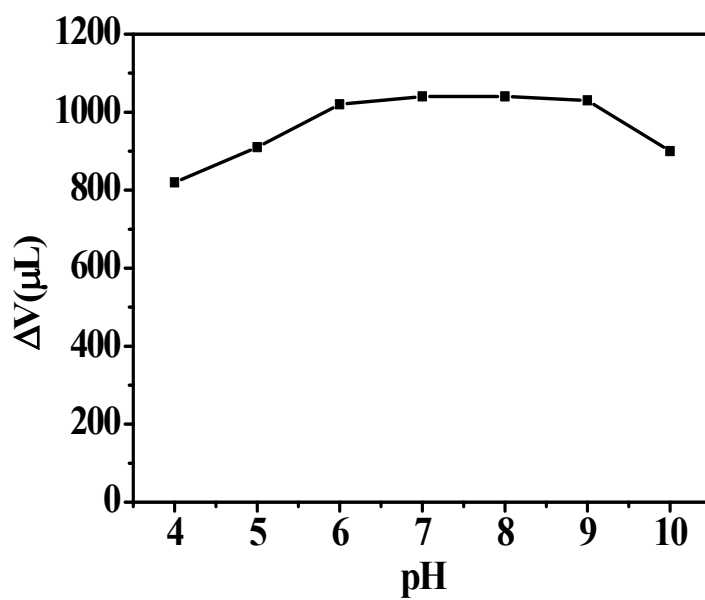


Fig. S5  $^{13}\text{C}$  NMR spectrum of 5-acrylamido-fluorescein



**Fig. S6** Response characteristics of fluorescent hydrogels to  $Hg^{2+}$ . Volume changes of the fully swelled fluorescent hydrogel after soaking in  $3.0 \times 10^{-4} \text{ mol} \cdot L^{-1}$  NaCl solution (left) and  $1.0 \times 10^{-4} \text{ mol} \cdot L^{-1}$   $Hg(NO_3)_2$  solution (right) for 30 min.



**Fig. S7** Effect of pH on the  $Hg^{2+}$ -induced volume shrinkage of the obtained hydrogel.  $Hg^{2+}$ ,  $5.0 \times 10^{-5} \text{ mol} \cdot L^{-1}$ ; pH 4~10, by adding 0.2 M NaOH to B-R buffer (0.04 M  $H_3PO_4$ , HAc and  $H_3BO_3$ ).

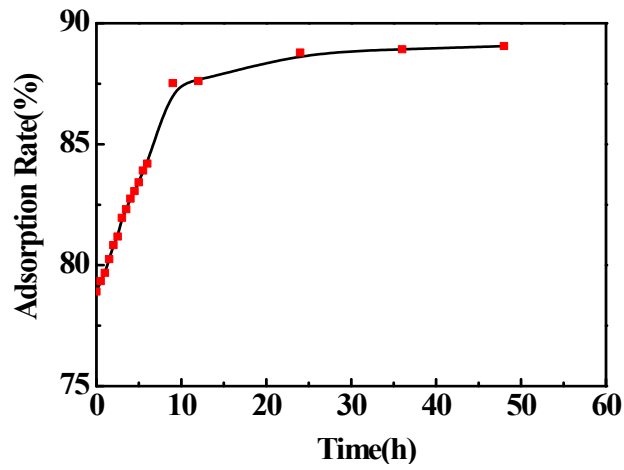


Fig. S8 Effect of time on the adsorption rate for  $Hg^{2+}$

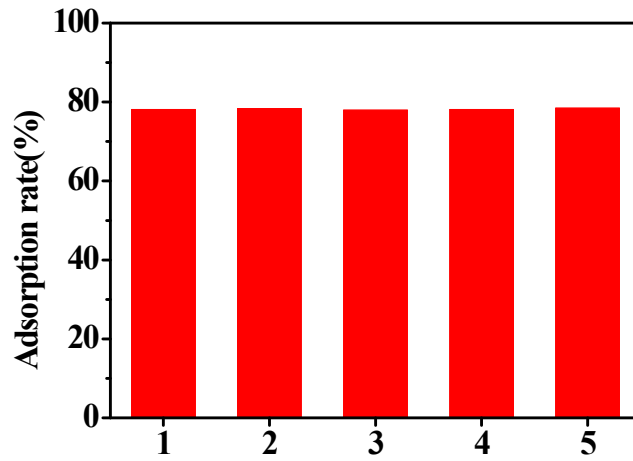


Fig. S9 The adsorption rate of 30 minutes for  $Hg^{2+}$

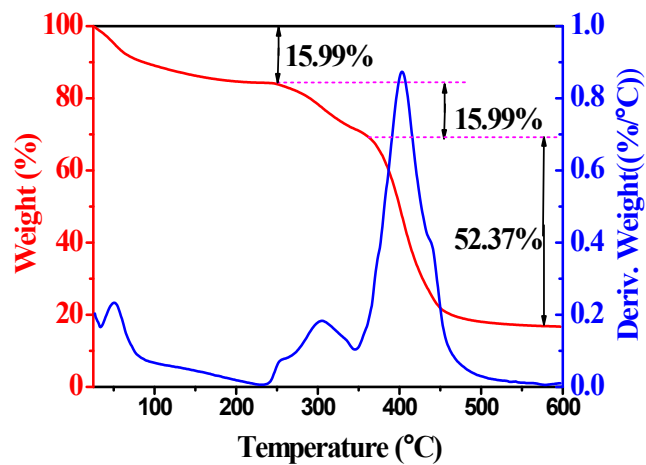


Fig. S10 TGA result of the obtained hydrogel. Rate of the temperature rise, 5.0 °C/min; Temperature range, 27~600 °C.

Table S1 Comparison of the reported and the present nitrogen base-based sensors for  $Hg^{2+}$  determination

Sensor type	Advantages	Disadvantages	Ref.
Electrochemical sensor	High sensitivity, fast response, and can be used in opaque system	Complicated in sensor preparation, determination results affected by non-specific adsorption, and electrochemistry workstation is required	[1-20]
Colorimetric sensor	Low-cost, naked-eye readout or a simple instrument is requirement, and can be used for field detection	Color vision difference limits the sensitivity, which usually makes it is difficult to accurately quantify and generally be used for qualitative or semi-quantitative analysis	[21-33]
Fluorescence sensor	High sensitivity, fast response speed, wide linear range and short time consuming	Environmental factors greatly influenced the determination results, and there may exist the problems such as photobleaching and low fluorescent intensity, etc; Fluorescence spectrophotometer is required and rarely available for field detection	[34-56]
The present sensor	Easy operate, ultra - low cost and suitable for on-site inspection; Naked-eye signal readout in line with human's perception nature of "seeing is believing" and power-free of instrument requirement	Sensitivity needs to be further improved that limited by the currently obtained sensing materials	—

## References

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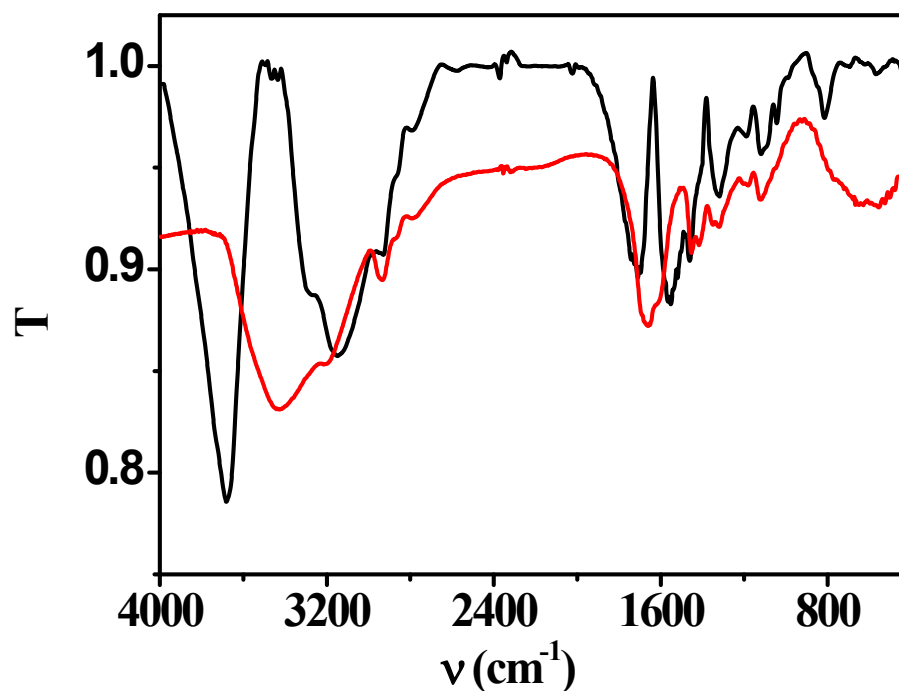


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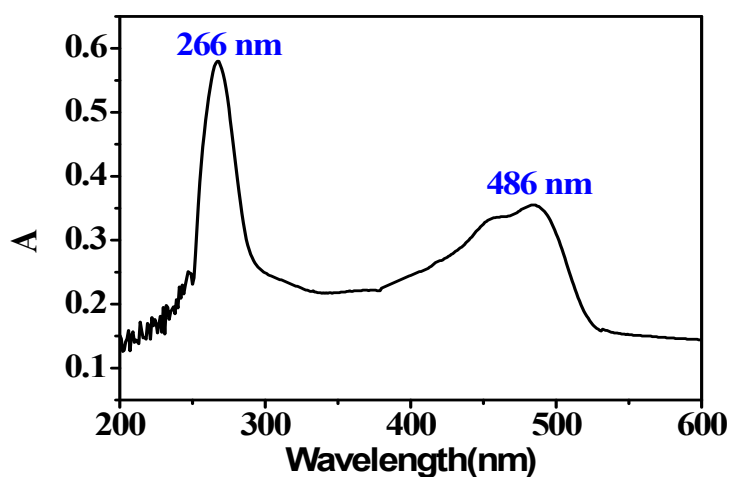


**Fig.S11** IR spectra of polyacrylamide and the fluorescent hydrogel

In IR spectra, the red curve is the IR absorption spectrum of polyacrylamide. Peaks  $3199.37\text{ cm}^{-1}$  and  $3416.61\text{ cm}^{-1}$  representative the symmetric and asymmetric stretching vibration of amide N-H,  $1667.40\text{ cm}^{-1}$  stands for the symmetric stretching vibration of C=O,  $2936.99\text{ cm}^{-1}$  and  $2776.17\text{ cm}^{-1}$  for asymmetric and symmetric stretching vibration of  $-\text{CH}_2-$ . Peak  $1458.62\text{ cm}^{-1}$  belongs to the bending vibration of  $-\text{CH}_2-$  in the polymer chains; The black curve is the IR absorption spectrum of the obtained fluorescent hydrogel. Compared with polyacrylamide, the most obvious changes take place at the absorption peak of  $3416.61\text{ cm}^{-1}$ ,  $3199.37\text{ cm}^{-1}$ , and  $1667.40\text{ cm}^{-1}$ . The position of  $3416.61\text{ cm}^{-1}$  and  $3199.37\text{ cm}^{-1}$  shifts towards high wave number and gain the strongest absorption peak in  $3680.40\text{ cm}^{-1}$ , which should be attributed to the asymmetric stretching vibration of O-H that comes from the

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fluorescein monomer; the position of  $1667.40\text{ cm}^{-1}$  also shifts towards high wave number and gain a strong absorption peak in  $1696.33\text{ cm}^{-1}$  and stands for the symmetric stretching vibration of C=O; meanwhile, a new strong absorption peak appears in  $1541.73\text{ cm}^{-1}$ , which should be attributed to the stretching vibration of the C=C from the uracil monomer and the skeleton vibration of the benzene ring from the fluorescein. The above changes in the position of the key absorption peaks indicate that the obtained hydrogel is indeed a uracil and fluorescein functionalized polymer.



**Fig.S12** UV-visible absorption spectra of the fully swollen fluorescent hydrogel