

Novel luminescent EuBTC@HA composite with dual sensing properties

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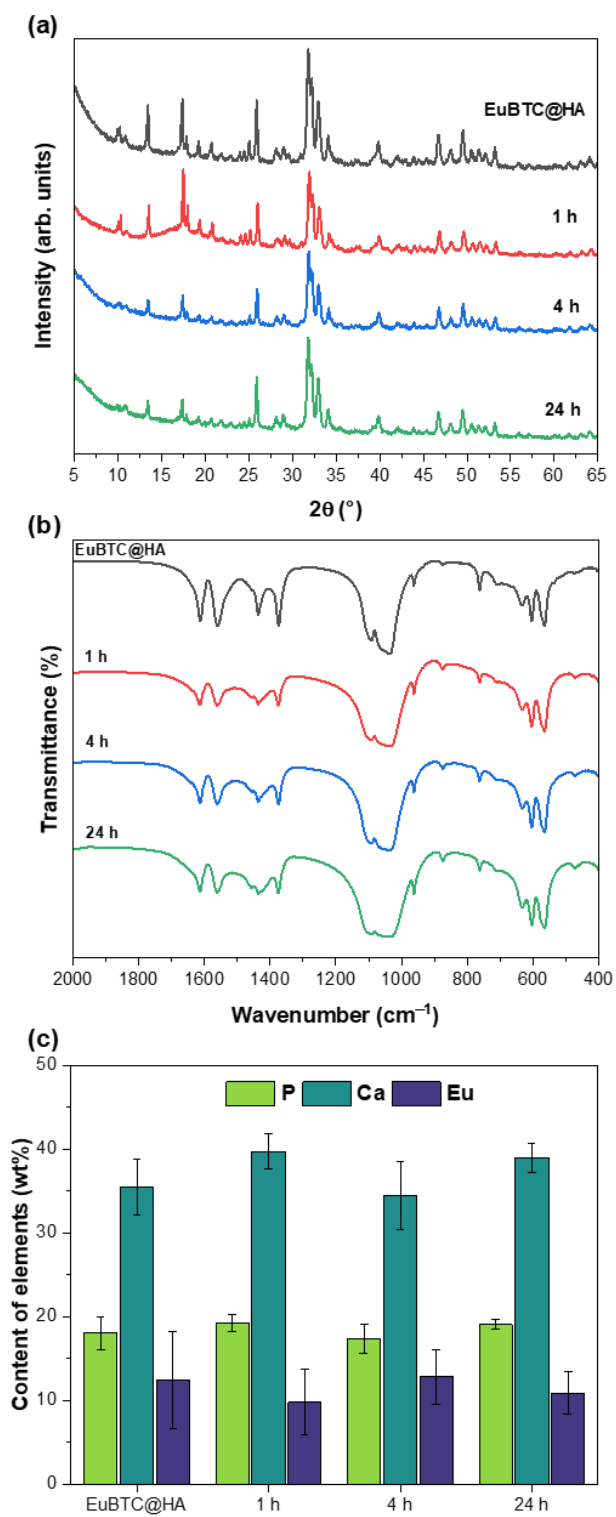


Figure S1. Stability of EuBTC@HA in water determined by PXRD (a), FT-IR spectroscopy (b), and analysis of composition monitored by EDS (c).

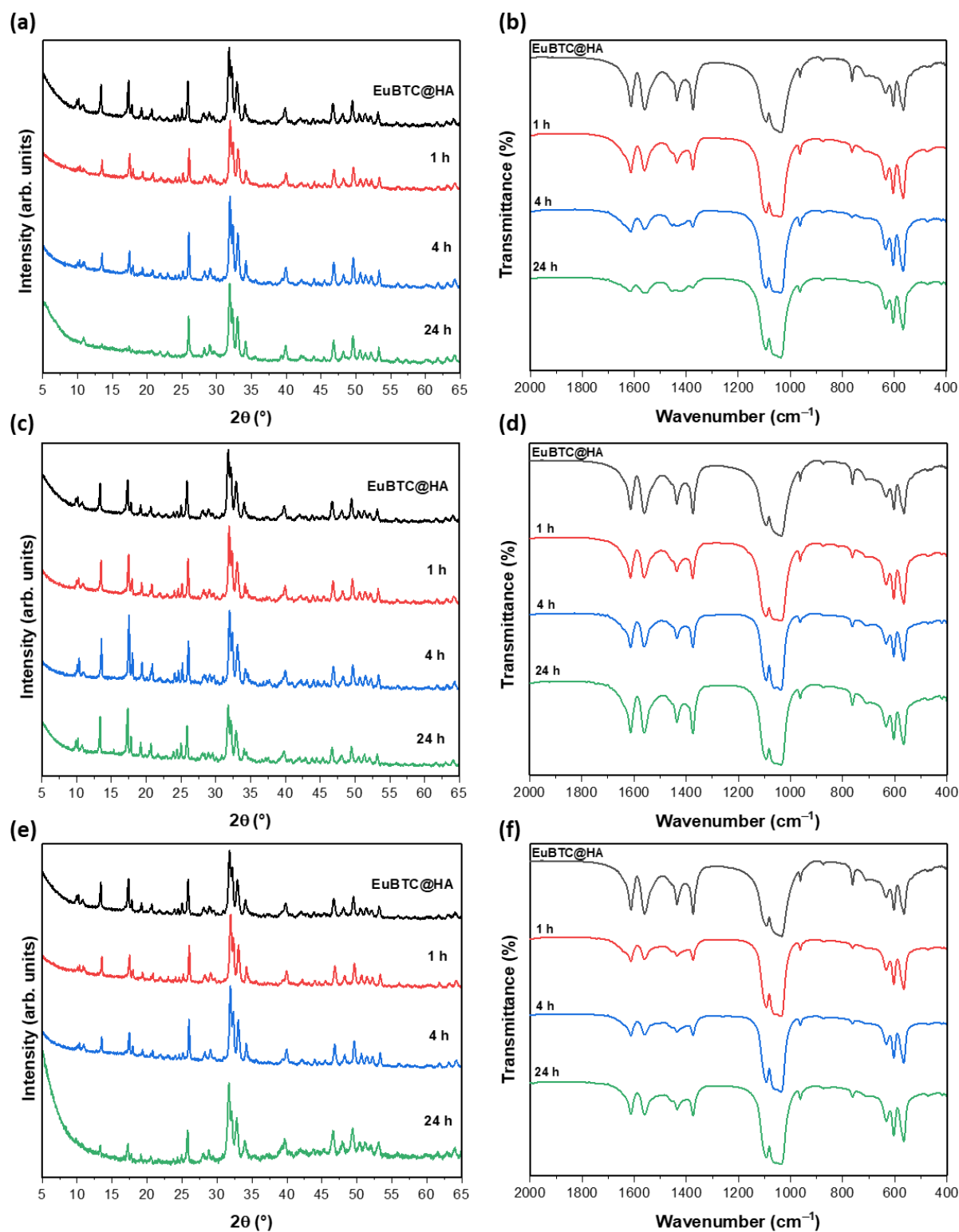


Figure S2. Stability of EuBTC@HA determined by PXRD (a,c,e) and FT-IR spectroscopy (b,d,f) in buffers of pH equal to 4.5 (a,b), 6.8 (c,d), and 8.8 (e,f), respectively.

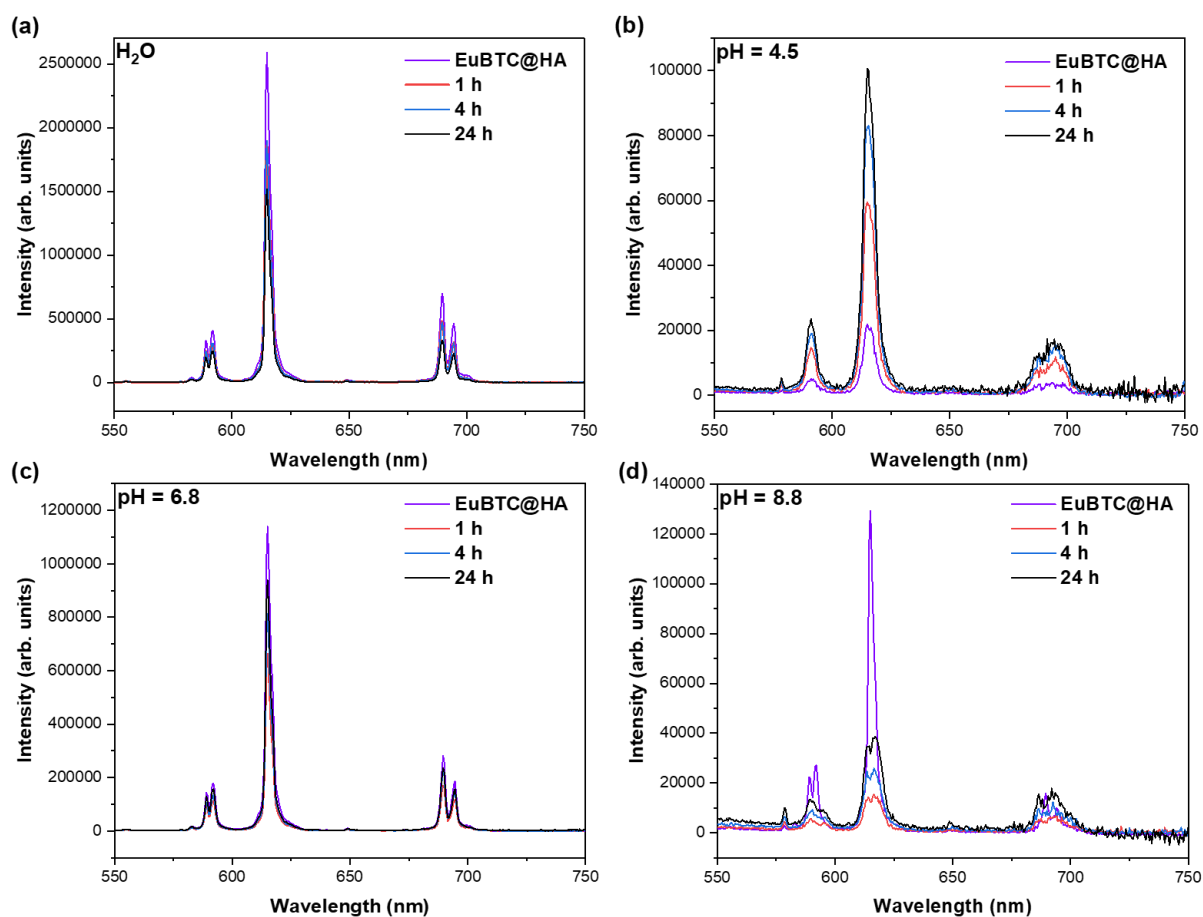


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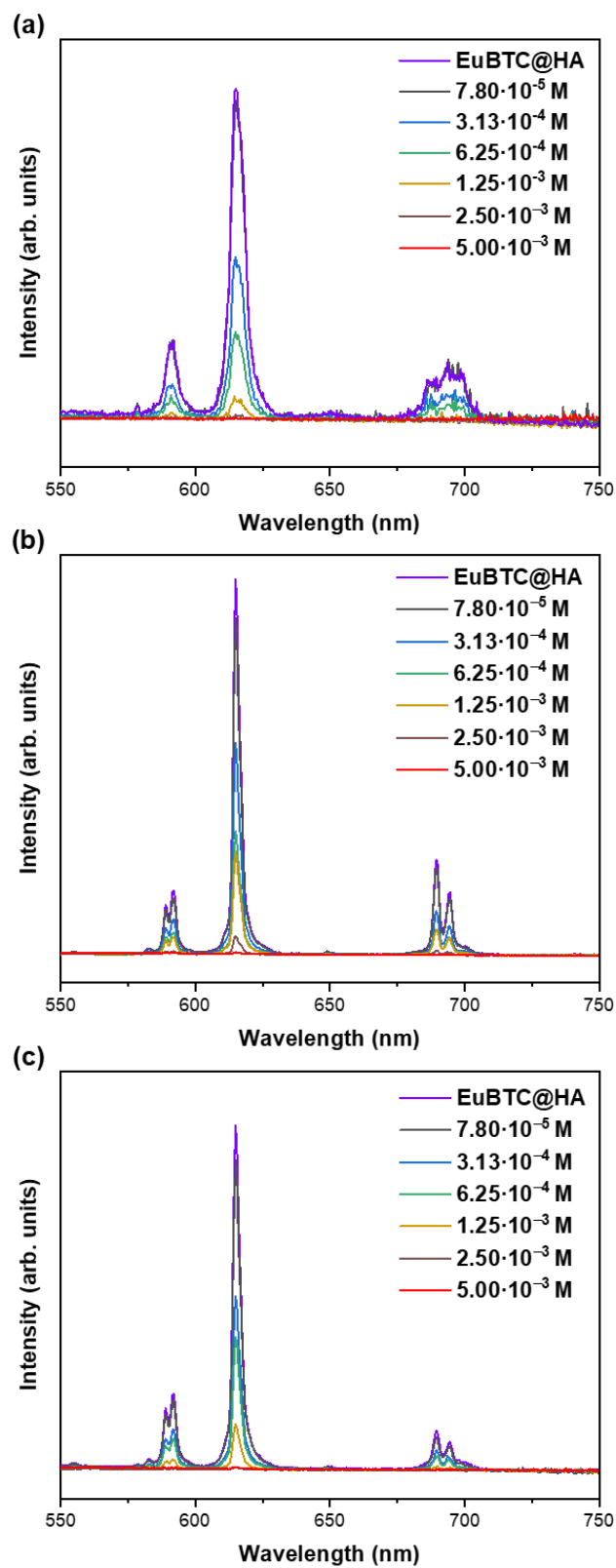


Figure S4. Emission spectra of EuBTC@HA for various concentrations of Fe^{3+} ions at pH 4.5 (a), 6.8 (b), and 8.8 (c).

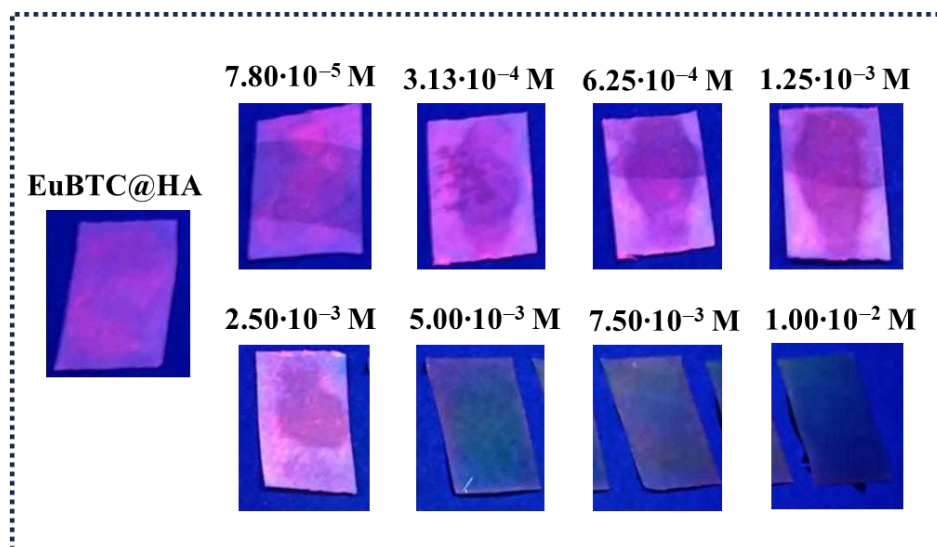


Figure S5. Photographs of the filter paper coated with EuBTC@HA used for the detection of Fe³⁺ ions in the concentration range of $7.80 \cdot 10^{-5}$ M to $1.00 \cdot 10^{-2}$ M under UV light ($\lambda = 254$ nm).

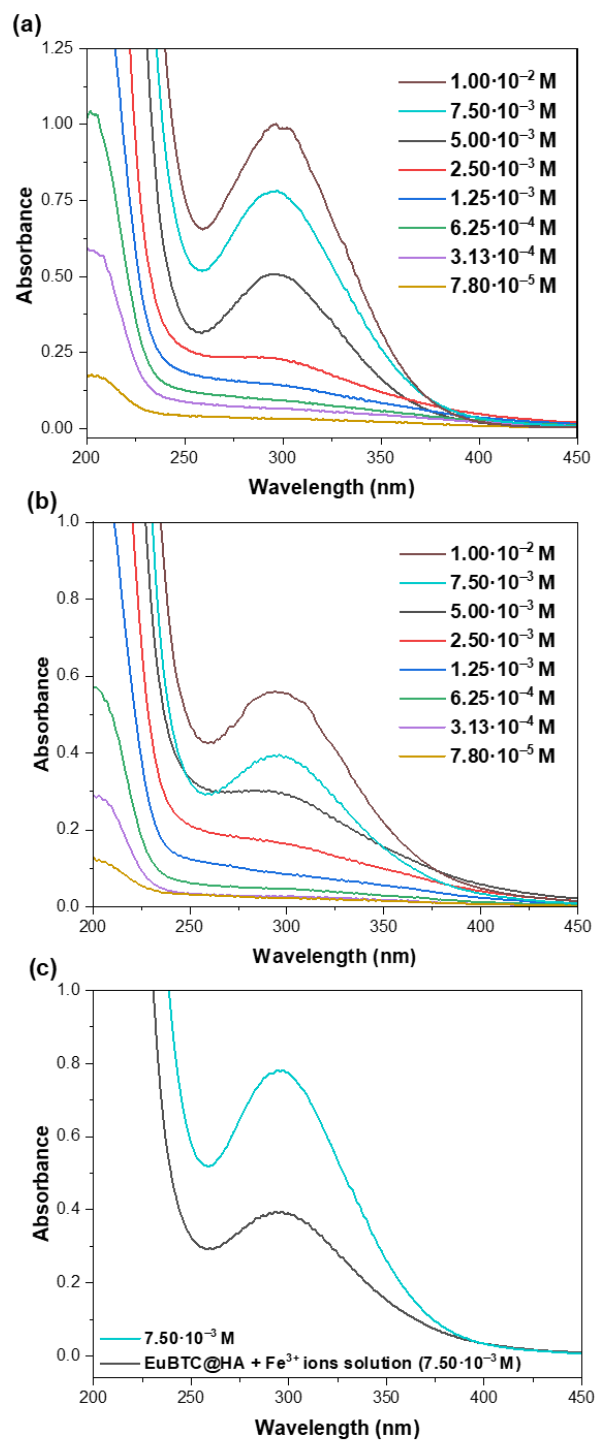


Figure S6. UV-Vis spectra of aqueous solution of iron(III) nitrate before (a) and after detection using EuBTC@HA (b), along with a comparison of the spectra for a selected concentration of $7.50 \cdot 10^{-3}$ M (c).

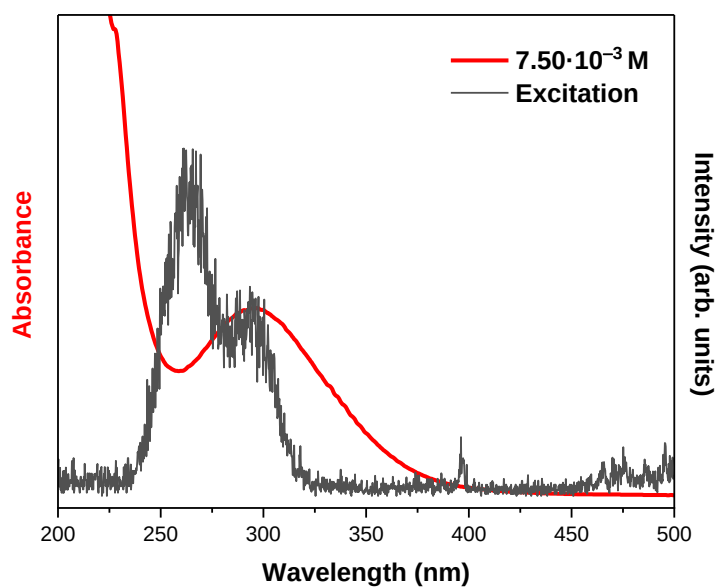


Figure S7. Overlay of the absorption spectrum of an aqueous solution of iron(III) nitrate (with a concentration of $7.50 \cdot 10^{-3} \text{ M}$) and the excitation spectrum of EuBTC@HA.

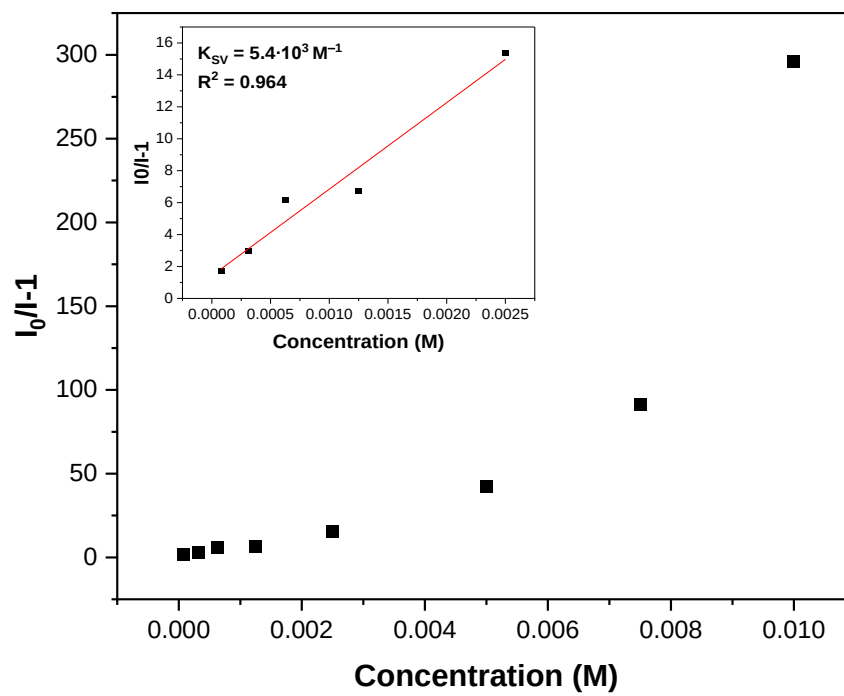


Figure S8. Stern–Volmer plot against the concentrations of iron(III) nitrate aqueous solutions with an inserted linear range.

Table S1. Average luminescence lifetimes of EuBTC@HA before and after detection of Fe³⁺ in the concentration range of 7.80·10⁻⁵ M to 1.00·10⁻² M.

Concentration (M)	Average lifetime (ms)
EuBTC@HA	0.18
7.80·10 ⁻⁵	0.18
3.13·10 ⁻⁴	0.18
6.25·10 ⁻⁴	0.18
1.25·10 ⁻³	0.18
2.50·10 ⁻³	0.18
5.00·10 ⁻³	0.18
7.50·10 ⁻³	0.012
1.00·10 ⁻²	0.0075

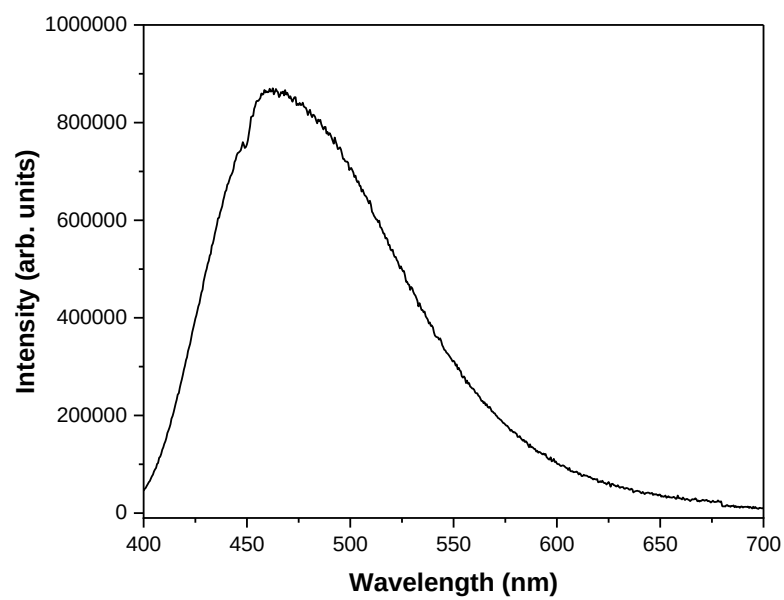


Figure S9. Emission spectrum of OFX at a concentration of $9.75 \cdot 10^{-6}$ M ($\lambda_{\text{exc}} = 265$ nm).

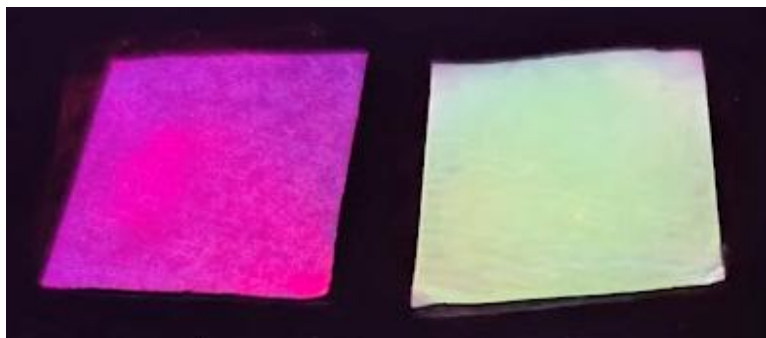


Figure S10. Photograph of the filter paper under UV light ($\lambda = 254$ nm) coated with EuBTC@HA (left) and after the addition of a few drops of OFX at a concentration of $2.43 \cdot 10^{-6}$ M (right).

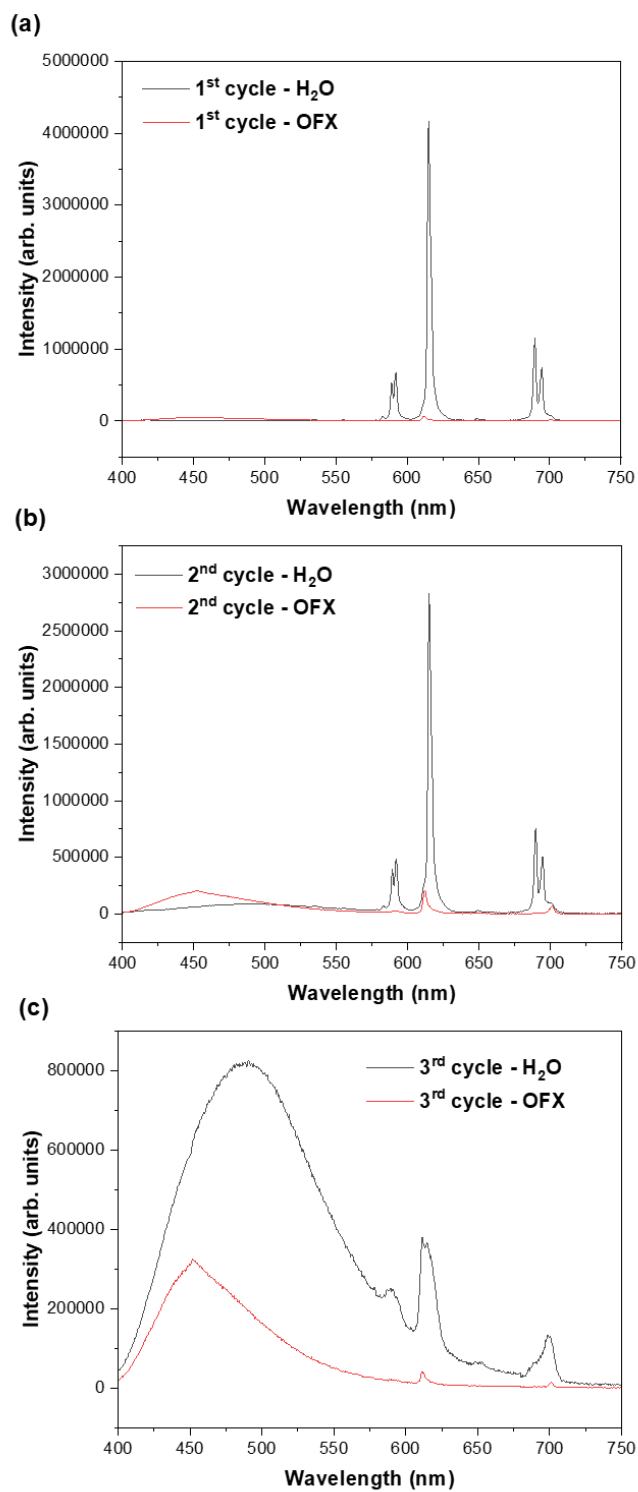


Figure S11. Emission spectra of EuBTC@HA after 1st (a), 2nd (b), and 3rd (c) cycle of OFX detection at a concentration of $5.00 \cdot 10^{-3}$ M.

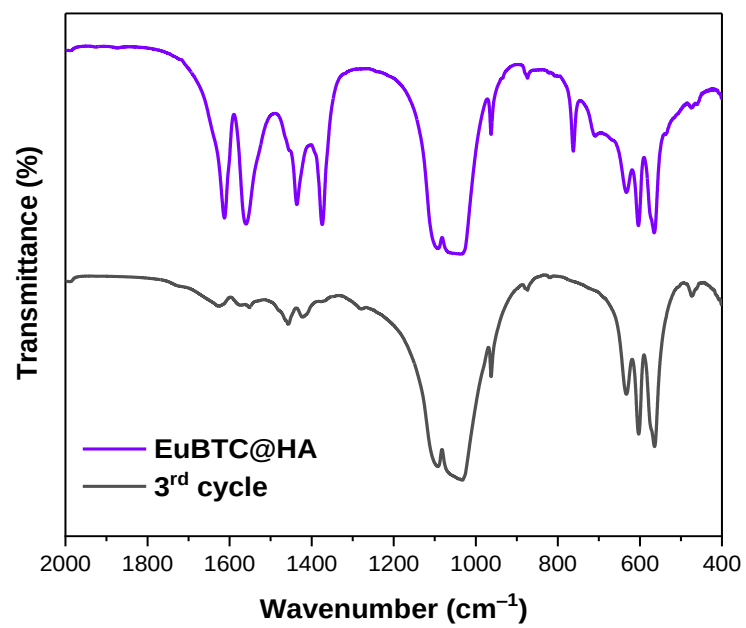


Figure S12. FT-IR spectra of EuBTC@HA before and after the 3rd cycle of OFX detection.

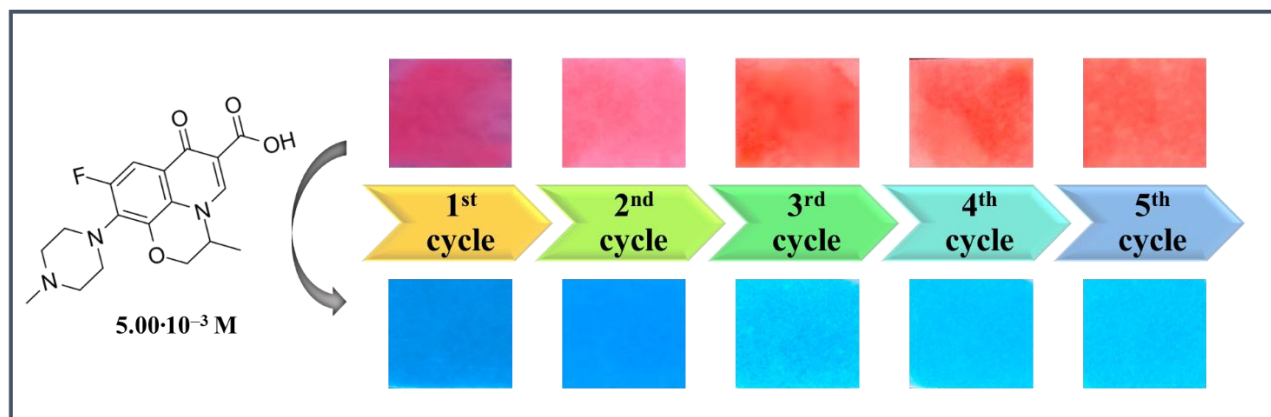


Figure S13. Photographs of the filter paper coated with EuBTC@HA used for the five cycles of OFX detection at a concentration of $5.00 \cdot 10^{-3} \text{ M}$ under UV light ($\lambda = 254 \text{ nm}$).

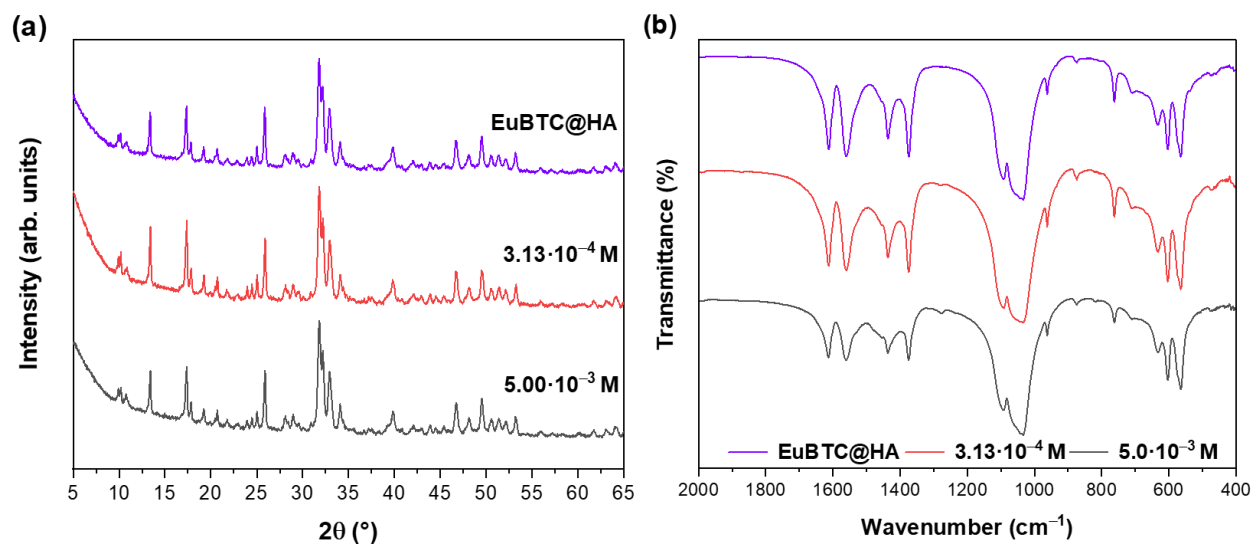


Figure S14. PXRD patterns (a) and FT-IR spectra (b) of EuBTC@HA before and after detection of OFX at concentrations of $3.13 \cdot 10^{-4}$ M and $5.00 \cdot 10^{-3}$ M.

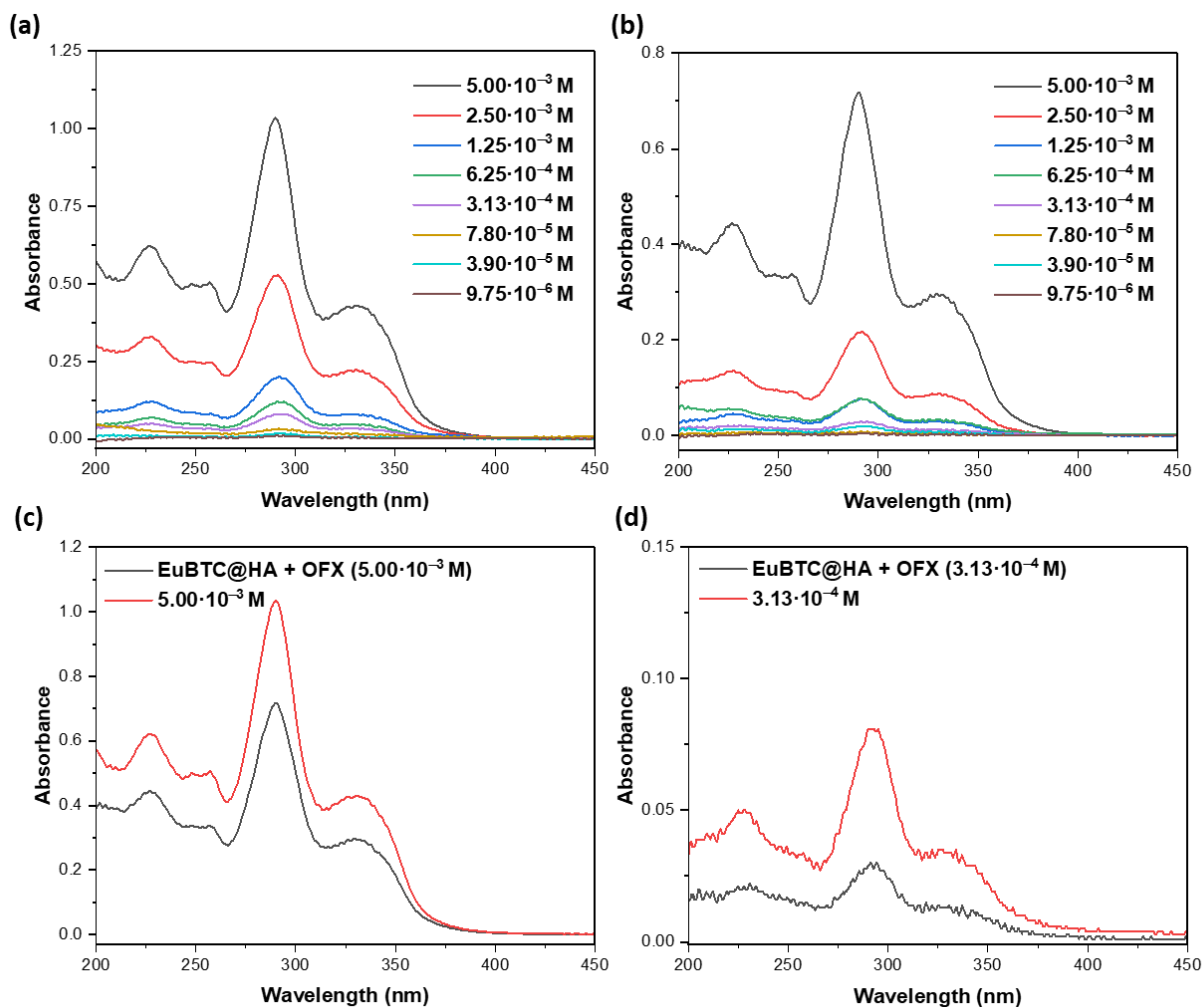


Figure S15. UV-Vis spectra of OFX solution at different concentrations before (a) and after detection using EuBTC@HA (b), along with a comparison of spectra for selected concentrations of $5.00 \cdot 10^{-3} \text{ M}$ (c) and $3.13 \cdot 10^{-4} \text{ M}$ (d).

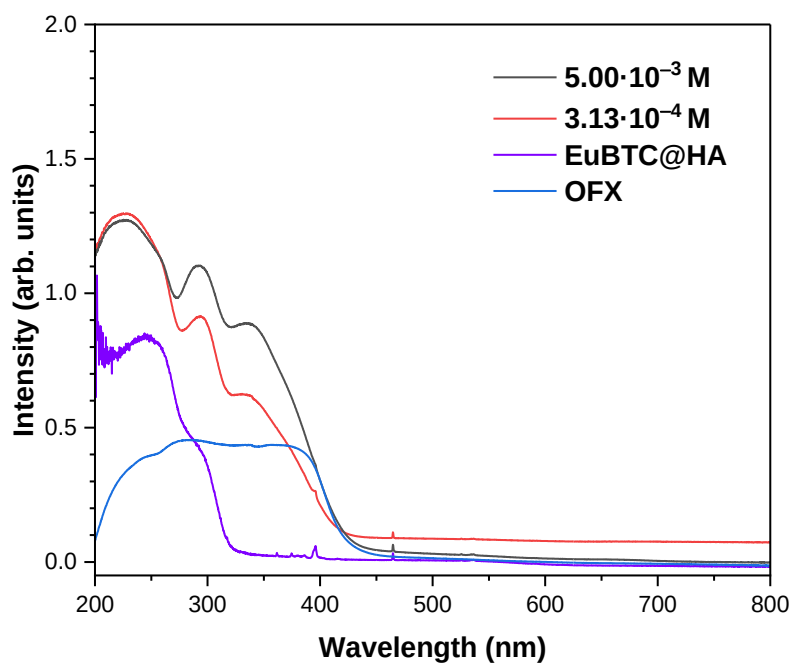


Figure S16. Characterization of OFX and EuBTC@HA, before and after detection of OFX at concentrations of $3.13 \cdot 10^{-4} \text{ M}$ and $5.00 \cdot 10^{-3} \text{ M}$ using UV-Vis diffuse reflectance spectroscopy.

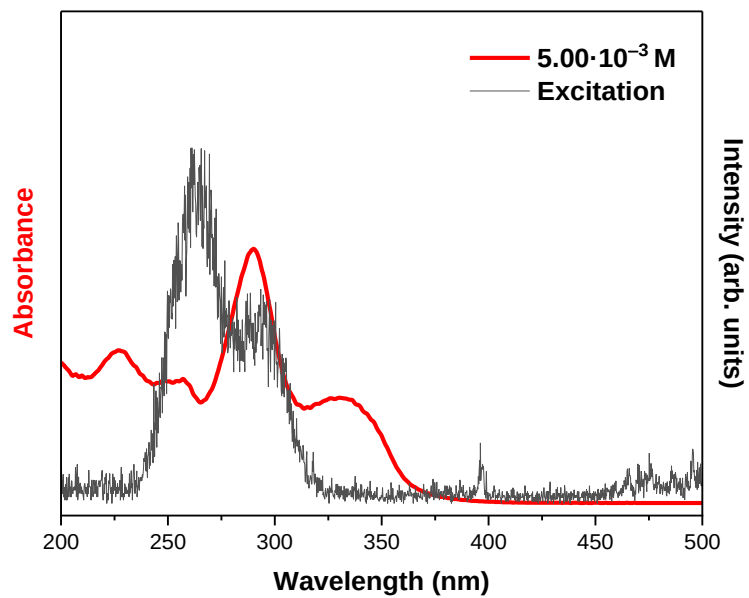


Figure S17. Overlay of the absorption spectrum of OFX (at a concentration of $5.00 \cdot 10^{-3}$ M) and the excitation spectrum of EuBTC@HA.

Table S2 Calculated HOMO and LUMO energies for H₃BTC and OFX.

	E_{HOMO} (eV)	E_{LUMO} (eV)	$\Delta E(E_{LUMO} - E_{HOMO})$ (eV)
H₃BTC	-8.0501	-2.361	5.689
OFX	-6.151	-1.948	4.203