

Supporting Information

A combined bottom-up and top-down strategy to fabricate lanthanide hydrate@2D MOF composite nanosheets for direct white light emission

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Table S1. Crystallographic data and refinement details for HSB-W5d.

Empirical formula	C ₂₂ H ₃₂ N ₄ O ₉ Zn
<i>M</i>	561.88
Crystal system	triclinic
Space group	<i>P</i> -1
<i>a</i> (Å)	9.561(6)
<i>b</i> (Å)	11.067(7)
<i>c</i> (Å)	14.378(9)
α (°)	104.135(2)
β (°)	95.491(9)
γ (°)	115.421(8)
<i>V</i> /Å ³	1297.1(14)
<i>Z</i>	2
<i>D</i> _c /g cm ⁻³	1.439
μ /mm ⁻¹	1.003
2 θ _{range} (°)	4.734 to 54.996
<i>h, k, l, ranges</i>	-12 to 12, -14 to 14, -18 to 18
<i>F</i> (000)	588.0
<i>R</i> ₁ , ^a <i>wR</i> ₂ ^b [<i>I</i> > 2 σ (<i>I</i>)]	0.0692, 0.1695
GOF on <i>F</i> ²	1.162
^a <i>R</i> = $\Sigma(F_o - F_c)/\Sigma F_o $. ^b <i>Rw</i> = $\{\Sigma w[(F_o^2 - F_c^2)^2]/\Sigma w[(F_o^2)^2]\}^{1/2}$.	

Table S2. Selected bond lengths (Å) and angles (°) of HSB-W5d.

Zn1-N1	2.233(3)
Zn1-N2	2.132(4)
Zn1-N4 ^a	2.265(3)
Zn1-O1	1.998(3)
Zn1-O3	2.071(3)
N1-Zn1-N4 ^a	170.20(12)
N2-Zn1-N1	82.08(12)
N2-Zn1-N4 ^a	90.76(12)
O1-Zn1-N1	98.37(12)
O1-Zn1-N2	116.52(13)
O1-Zn1-N4 ^a	90.81(11)
O1-Zn1-O3	101.95(13)
O3-Zn1-N1	94.02(11)
O3-Zn1-N2	141.51(14)
O3-Zn1-N4 ^a	87.41(11)

Symmetry codes: a) -1+X, +Y, +Z.

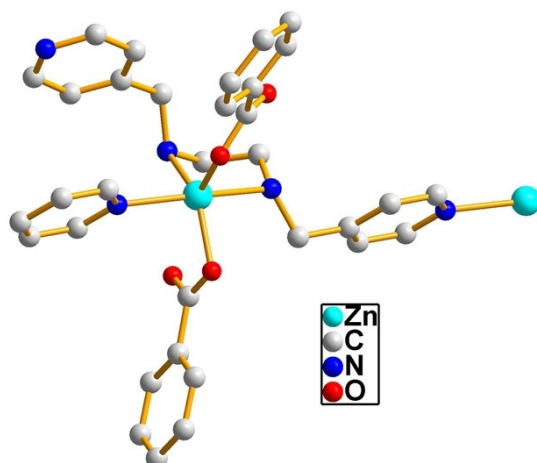
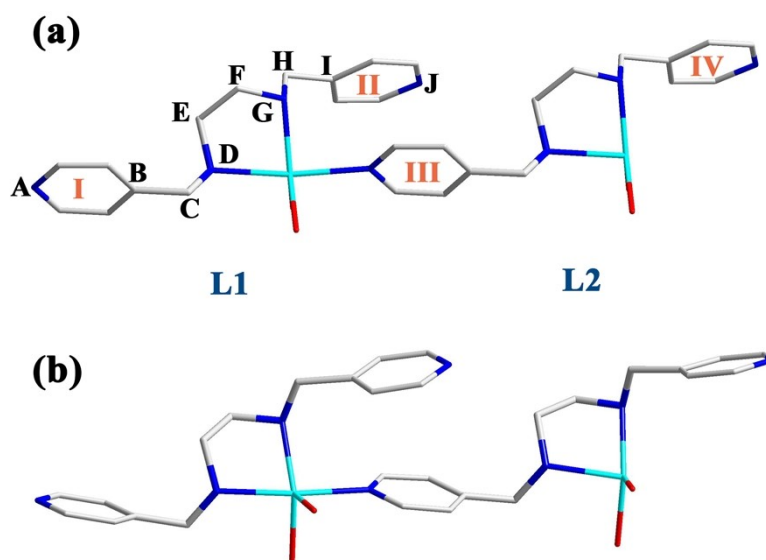


Figure S1. View of the coordination environment of Zn(II) ion in HSB-W5d (hydrogen atoms and free water molecules have been omitted for clarity).



HSB-W5d

Torsion Angles (°)	L1	L2
A-C-H-J	-146.5	-146.5
C-D-G-H	-160.0	-160.0
B-C-D-E	60.1	60.1
C-D-E-F	97.9	97.9
D-E-F-G	57.7	57.7
E-F-G-H	78.2	78.2
F-G-H-I	174.0	174.0

Dihedral Angle (°)	
I-II	16.0
III-IV	16.0
I-III	0
II-IV	0

HSB-W5e

Torsion Angles (°)	L1	L2
A-C-H-J	162.7	131.8
C-D-G-H	158.8	164.6
B-C-D-E	-39.7	-67.2
C-D-E-F	-104.2	-102.7
D-E-F-G	-57.4	-56.0
E-F-G-H	-94.7	-73.9
F-G-H-I	-169.9	-172.2

Dihedral Angle (°)	
I-II	24.0
III-IV	22.2
I-III	7.8
II-IV	14.0

Figure S2. Comparison of the conformation of neighboring hsb-2 ligands in HSB-W5d (a) and HSB-W5e (b).

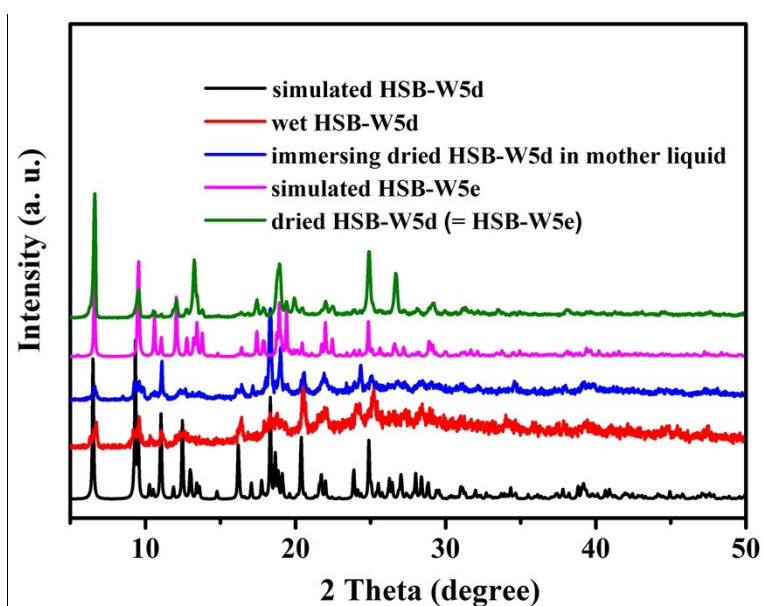


Figure S3. PXRD patterns of simulated HSB-W5d, wet crystal HSB-W5d, as-synthesized HSB-W5d (after drying), as-synthesized HSB-W5d immersing in the mother liquid, and simulated HSB-W5e.

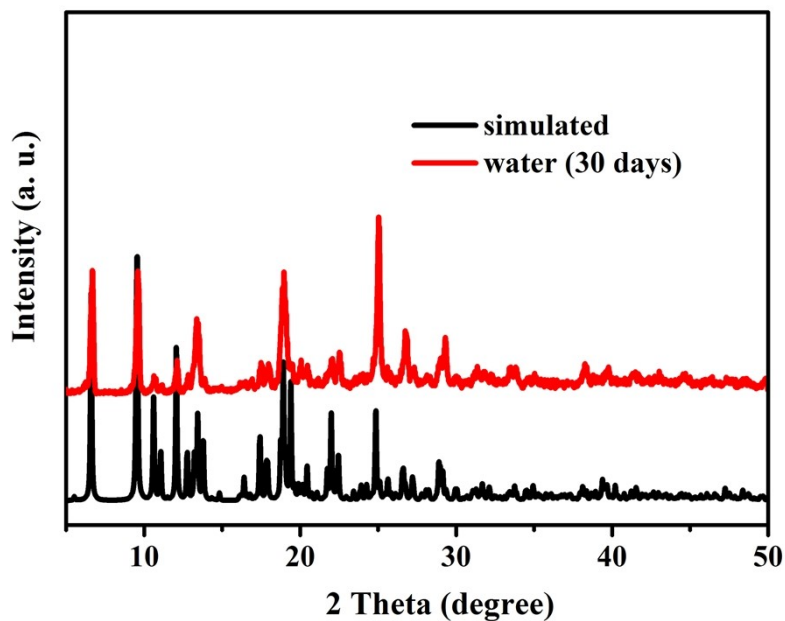


Figure S4. PXRD patterns of HSB-W5 after immersing in water for one month.

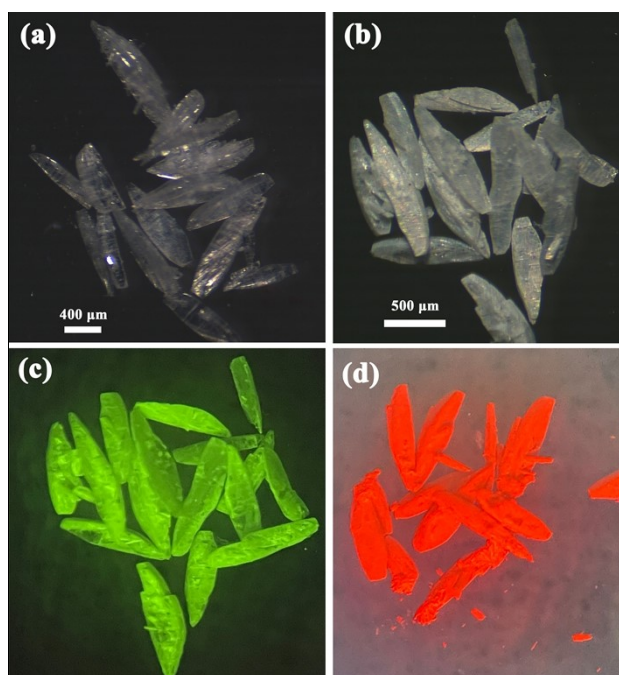


Figure S5. The photographs of as-synthesized HSB-W5 under sunlight (a), Tb(H₂O)₈³⁺@HSB-W5 under sunlight (b), Tb(H₂O)₈³⁺@HSB-W5 irradiated by a standard 254 nm laboratory UV lamp (c), and Eu(H₂O)₈³⁺@HSB-W5 irradiated by a standard 254 nm laboratory UV lamp (d).

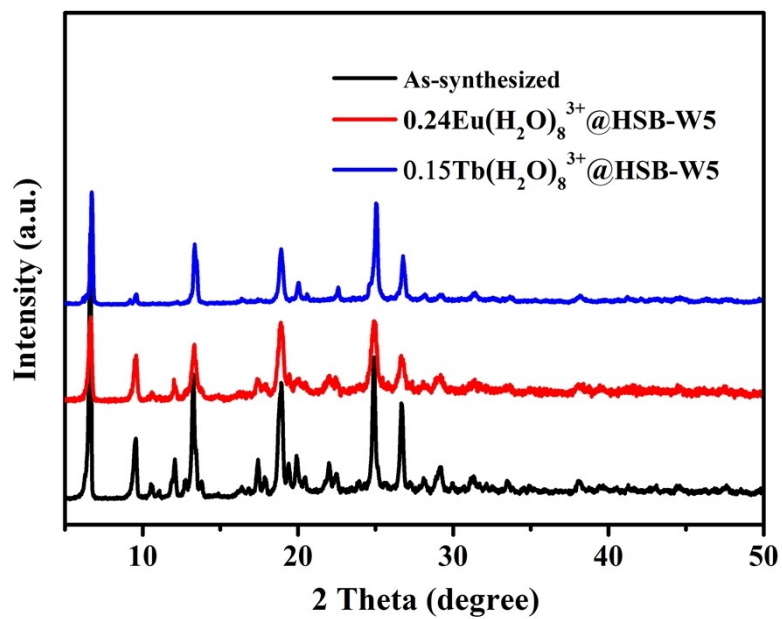


Figure S6. PXRD patterns of $\text{Eu}(\text{H}_2\text{O})_8^{3+}@HSB\text{-W5}$ and $\text{Tb}(\text{H}_2\text{O})_8^{3+}@HSB\text{-W5}$.

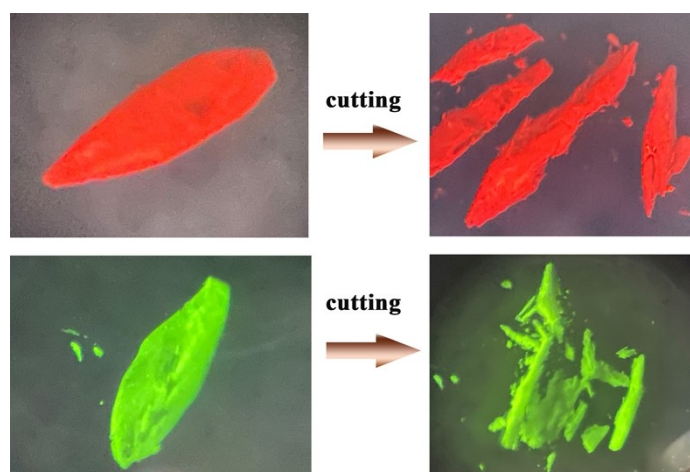


Figure S7. The photographs of crystals $\text{Eu}(\text{H}_2\text{O})_8^{3+}@HSB\text{-W5}$ and $\text{Tb}(\text{H}_2\text{O})_8^{3+}@HSB\text{-W5}$, and their pieces irradiated by a standard 254 nm laboratory UV lamp.

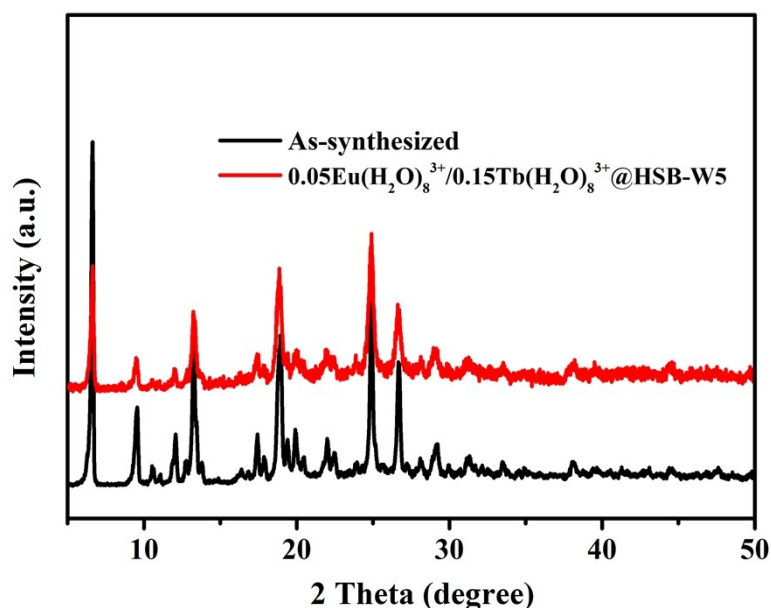


Figure S8. PXRD patterns of $0.03\text{Eu}(\text{H}_2\text{O})_8^{3+}/0.15\text{Tb}(\text{H}_2\text{O})_8^{3+}@HSB-W5$ composite.

Table S3. The amounts of Eu^{3+} in $\text{Eu}(\text{H}_2\text{O})_8^{3+}@HSB-W5$ composites based on ICP measurement.

Composite name	Initial concentration of Eu^{3+} (mol L ⁻¹)	Amount of Zn^{2+} (%)	Amount of Eu^{3+} (%)
$x\text{Eu}(\text{H}_2\text{O})_8^{3+}@HSB-W5$			
$0.08\text{Eu}(\text{H}_2\text{O})_8^{3+}@HSB-W5$	0.009	9.55	1.70
$0.13\text{Eu}(\text{H}_2\text{O})_8^{3+}@HSB-W5$	0.018	9.76	2.91
$0.24\text{Eu}(\text{H}_2\text{O})_8^{3+}@HSB-W5$	0.027	9.41	5.21

Table S4. The amounts of Tb^{3+} in $\text{Tb}(\text{H}_2\text{O})_8^{3+}@HSB-W5$ composites based on ICP measurement.

Composite name	Initial concentration of Tb^{3+} (mol L ⁻¹)	Amount of Zn^{2+} (%)	Amount of Tb^{3+} (%)
$x\text{Tb}(\text{H}_2\text{O})_8^{3+}@HSB-W5$			
$0.04\text{Tb}(\text{H}_2\text{O})_8^{3+}@HSB-W5$	0.002	9.97	1.04
$0.08\text{Tb}(\text{H}_2\text{O})_8^{3+}@HSB-W5$	0.007	9.80	1.87
$0.15\text{Tb}(\text{H}_2\text{O})_8^{3+}@HSB-W5$	0.014	9.71	3.48

Table S5. The amounts of Eu^{3+} and Tb^{3+} in WLE $\text{Eu}(\text{H}_2\text{O})_8^{3+}/\text{Tb}(\text{H}_2\text{O})_8^{3+}@HSB-W5$ composites based on ICP measurement.

Composite name	Amount of Zn^{2+} (%)	Amount of Eu^{3+} (%)	Amount of Tb^{3+} (%)
$x\text{Eu}(\text{H}_2\text{O})_8^{3+}/y\text{Tb}(\text{H}_2\text{O})_8^{3+}@HSB-W5$			
$0.03\text{Eu}(\text{H}_2\text{O})_8^{3+}/0.15\text{Tb}(\text{H}_2\text{O})_8^{3+}@HSB-W5$	9.22	0.72	3.41

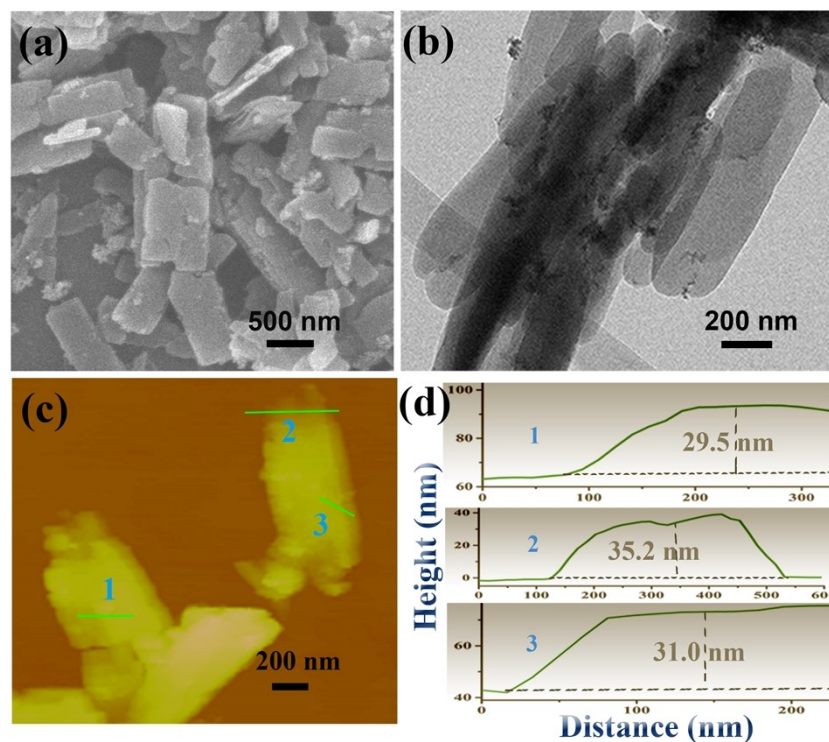


Figure S9. (a) SEM and (b) TEM images of the HSB-W5-NS nanosheets. (c, d) AFM images of the HSB-W5-NS nanosheets and the corresponding height profiles.

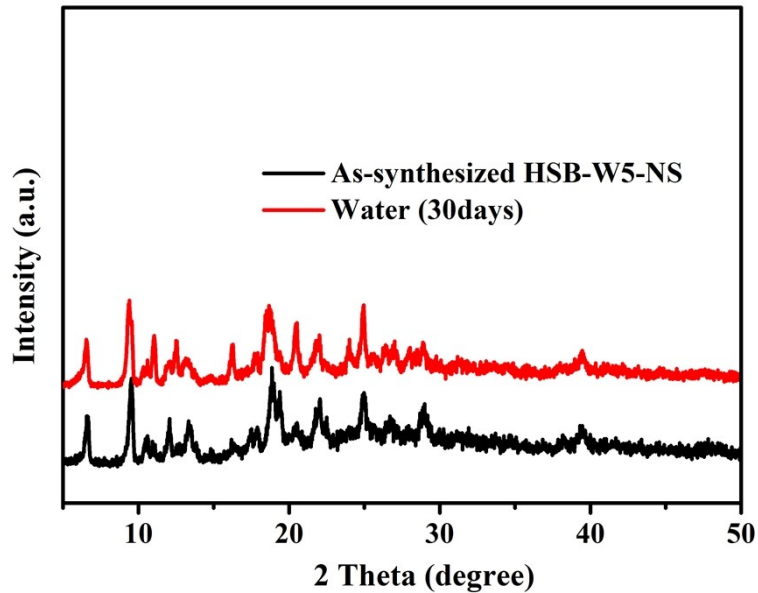


Figure S10. PXRD patterns of HSB-W5-NS after immersing in water for one month.

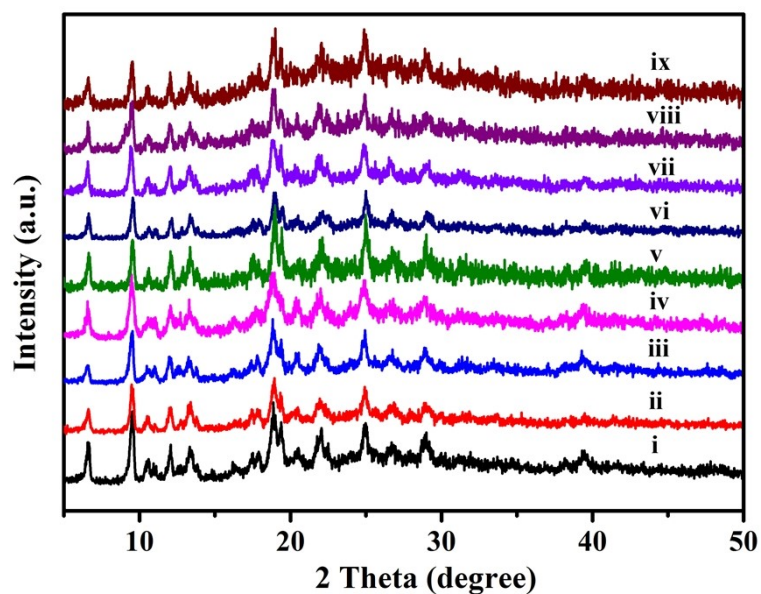


Figure S11. PXRD patterns of as-synthesized HSB-W5-NS (i), $x\text{Eu}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS (ii-v: $x = 0.26-1.60$), and $x\text{Tb}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS (vi-ix: $x = 0.08-0.48$).

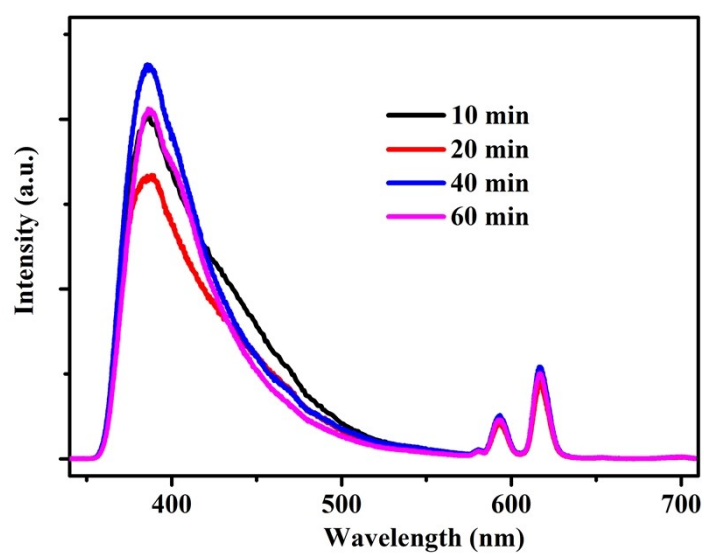


Figure S12. Emission changes with the reaction time.

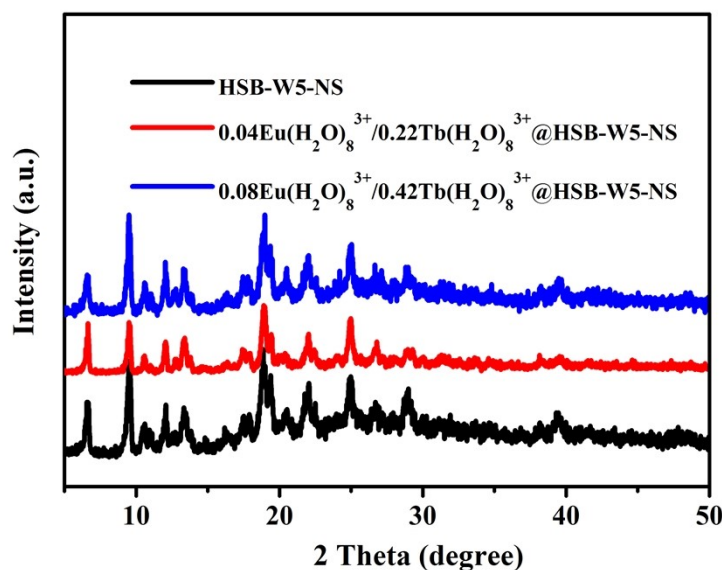


Figure S13. PXRD patterns of $\text{Eu}(\text{H}_2\text{O})_8^{3+}/\text{Tb}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS composite.

Table S6. The amounts of Eu^{3+} in $\text{Eu}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS composites based on ICP measurement.

Composite name	Initial concentration of Eu^{3+} (mol L ⁻¹)	Amount of Zn^{2+} (%)	Amount of Eu^{3+} (%)
$x\text{Eu}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS			
0.26 $\text{Eu}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS	0.001	9.52	5.80
0.55 $\text{Eu}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS	0.002	8.45	10.78
1.24 $\text{Eu}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS	0.003	6.15	17.76
1.60 $\text{Eu}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS	0.004	5.68	21.15

Table S7. The amounts of Tb^{3+} in $\text{Tb}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS composites based on ICP measurement.

Composite name	Initial concentration of Tb^{3+} (mol L ⁻¹)	Amount of Zn^{2+} (%)	Amount of Tb^{3+} (%)
$x\text{Tb}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS			
0.08 $\text{Tb}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS	0.00055	12.80	2.51
0.13 $\text{Tb}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS	0.001	12.23	4.02
0.21 $\text{Tb}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS	0.0013	11.47	5.90
0.48 $\text{Tb}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS	0.0022	9.60	11.24

Table S8. The amounts of Eu^{3+} and Tb^{3+} in WLE $\text{Eu}(\text{H}_2\text{O})_8^{3+}/\text{Tb}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS composites based on ICP measurement.

Composite name	Amount of Zn^{2+} (%)	Amount of Eu^{3+} (%)	Amount of Tb^{3+} (%)
$x\text{Eu}(\text{H}_2\text{O})_8^{3+}/y\text{Tb}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS			
0.04 $\text{Eu}(\text{H}_2\text{O})_8^{3+}/0.22\text{Tb}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS	11.13	1.04	6.03
0.08 $\text{Eu}(\text{H}_2\text{O})_8^{3+}/0.42\text{Tb}(\text{H}_2\text{O})_8^{3+}@$ HSB-W5-NS	9.52	1.76	9.75

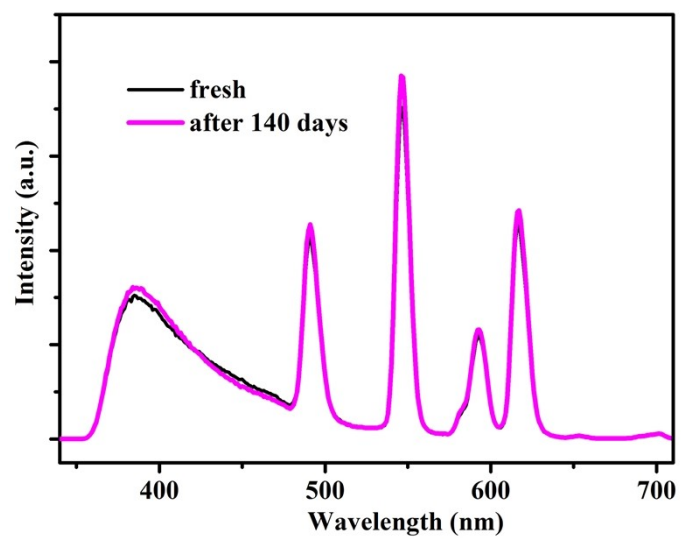


Figure S14. Comparison of the emission spectra of $0.04\text{Eu}(\text{H}_2\text{O})_8^{3+}/0.22\text{Tb}(\text{H}_2\text{O})_8^{3+}@\text{HSB-W5-NS}$ (fresh vs. after 140 days).

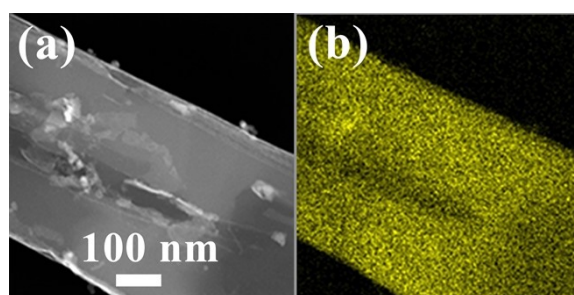


Figure S15. TEM image of $0.04\text{Eu}(\text{H}_2\text{O})_8^{3+}/0.22\text{Tb}(\text{H}_2\text{O})_8^{3+}@\text{HSB-W5-NS}$ (a) and corresponding EDX mapping image for C element (b).