

Supplementary Information

Luminescent and magnetic [TbEu] 2D Metal-Organic Frameworks

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S1. Infrared spectra of Ln 2D MOFs

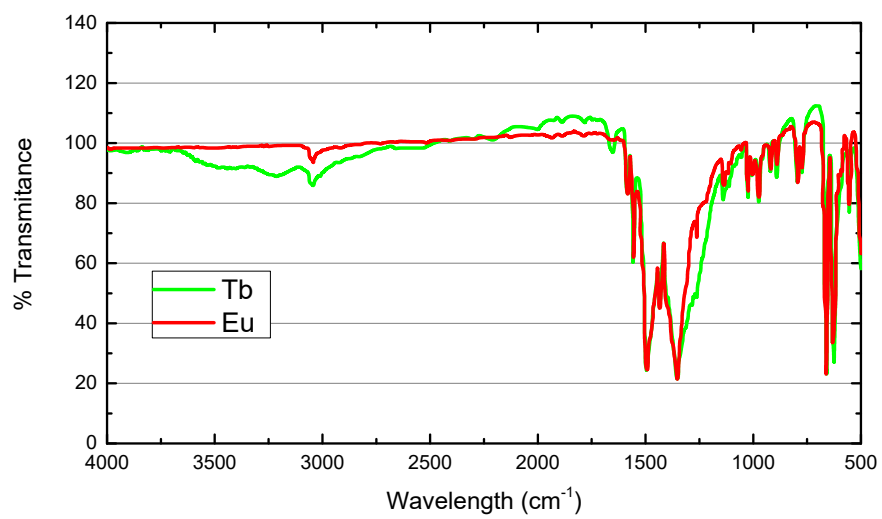
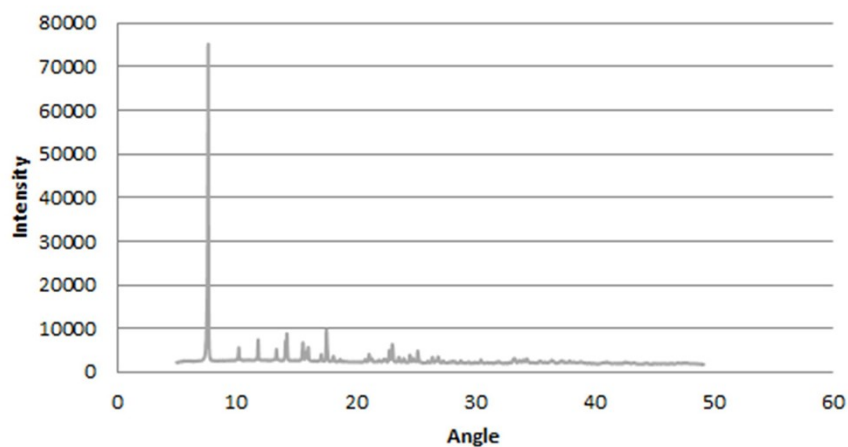


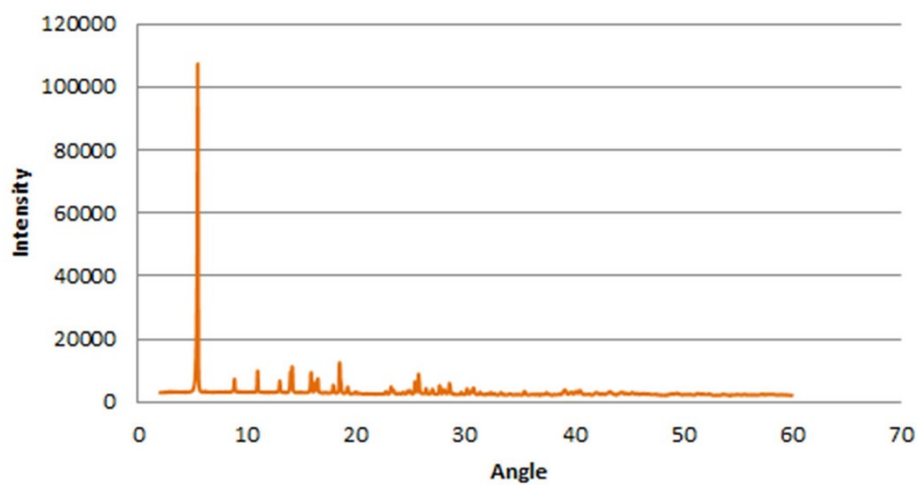
Figure S1. Infrared spectra of homometallic Ln 2DMOFs (Ln= Tb, Eu).

S2. PXRD

Dy



Tb_{0.3}Eu_{0.7}



Tb_{0.2}Eu_{0.8}

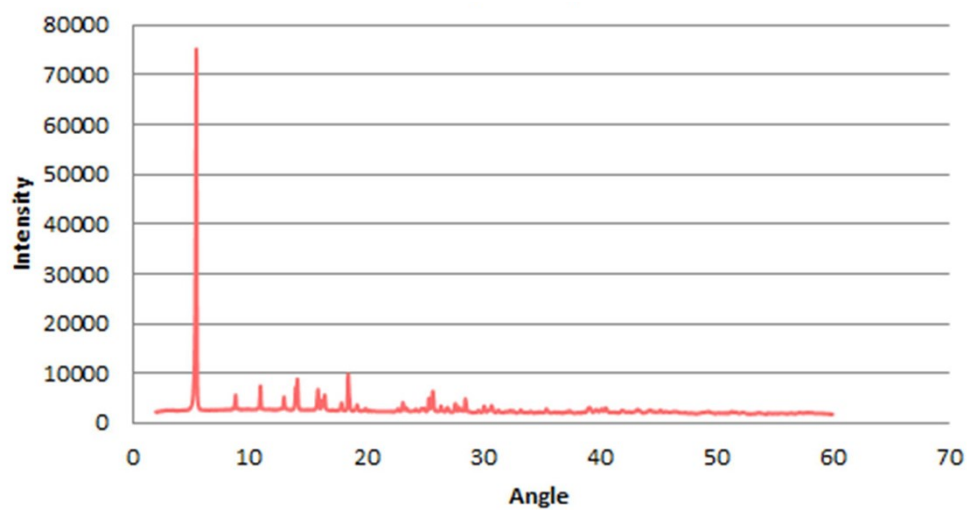


Figure S2. PXRD spectra of the studied mixed compounds (the rest are shown in Fig. 2 of the main paper).

S3. Tb/Eu ratio analysis

| Sample name | ICP | | SEM-EDS | | TEM-EDX | | Fluorescence | |
|--------------------|------|------|---------|------|---------|------|--------------|------|
| | Tb | Eu | Tb | Eu | | | Tb | Eu |
| Tb_xEu_{1-x} | | | | | | | | |
| $Tb_{0.2}Eu_{0.8}$ | 0.17 | 0.83 | 0.25 | 0.75 | | | | |
| $Tb_{0.3}Eu_{0.7}$ | 0.28 | 0.72 | 0.36 | 0.64 | | | | |
| $Tb_{0.4}Eu_{0.6}$ | 0.42 | 0.58 | | | 0.42 | 0.58 | | |
| $Tb_{0.7}Eu_{0.3}$ | 0.71 | 0.29 | | | | | 0.73 | 0.27 |
| $Tb_{0.9}Eu_{0.1}$ | 0.86 | 0.14 | | | | | 0.88 | 0.13 |

Table S3 Tb/Eu ratios determined for the heteronuclear samples by ICP (standard error <2%), SEM-EDS, TEM-EDX and Fluorescence.

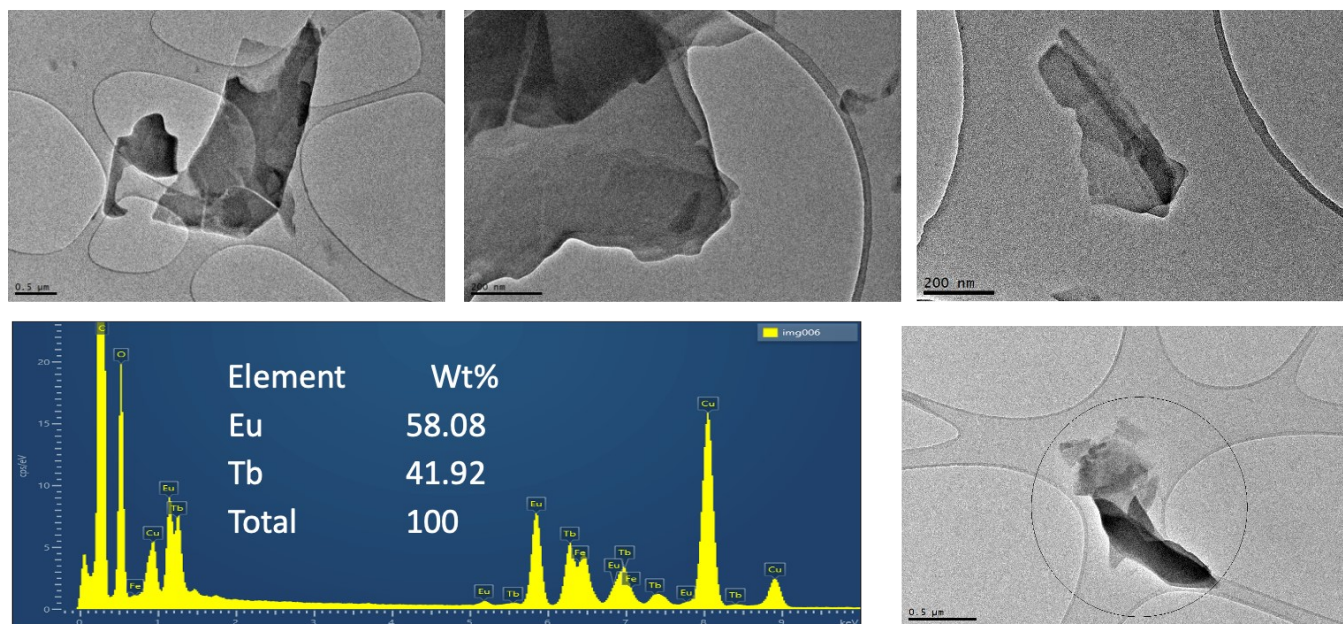


Figure S3. TEM images showing exfoliated flakes of $Tb_{0.4}Eu_{0.6}$ and EDX analysis.

S4. Color emission under UV light

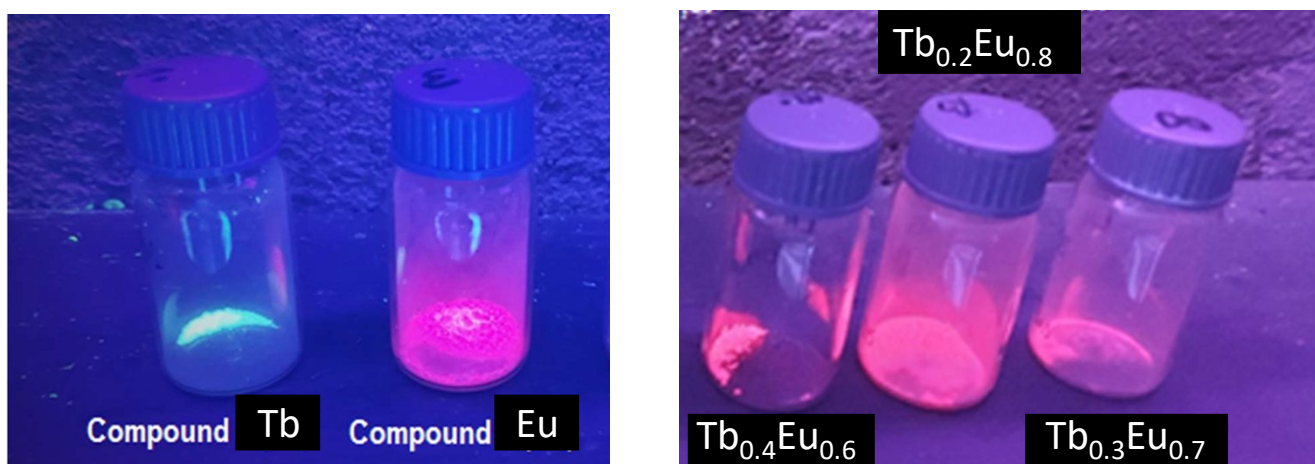


Figure S4. Visible color emission of (left) homonuclear Tb, Eu compounds, and (right) heteronuclear [TbEu] compounds, under UV light.

S5. Luminescence lifetime measurements

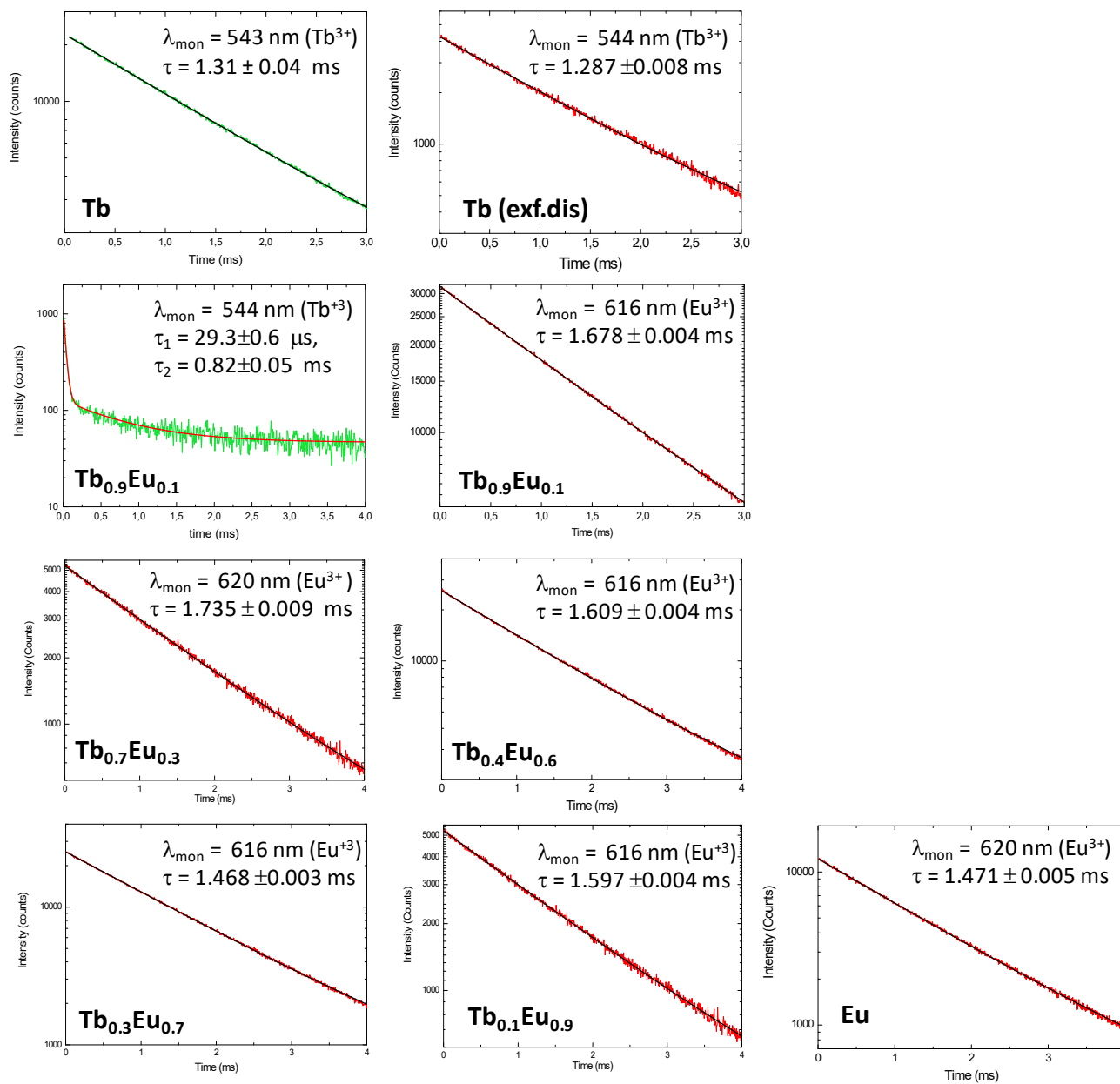


Figure S5. Lifetime measurements for homonuclear Tb and Eu compounds, and heterodinuclear Tb_xEu_{1-x} compounds, excited at $\lambda_{\text{exc}}=280 \text{ nm}$. The decay of either the Eu³⁺ main peak at 620 nm peak, or the Tb³⁺ peak at 544 nm were monitored. For **Tb_{0.9}Eu_{0.1}** the lifetime data were fit to a biexponential law with two time constants (τ_1 and τ_2): $I(t) = A_1 \cdot \exp(-t/\tau_1) + A_2 \cdot \exp(-t/\tau_2) + I_0$, with $A_1=1022 \pm 16$, $\tau_1=29.3 \pm 0.6 \text{ } \mu\text{s}$, $A_2=80 \pm 2$, $\tau_2=0.82 \pm 0.05 \text{ } \mu\text{s}$, $I_0=46.5 \pm 0.8$. For all other compounds the data were fit to an exponential decay law with a single time constant (τ).

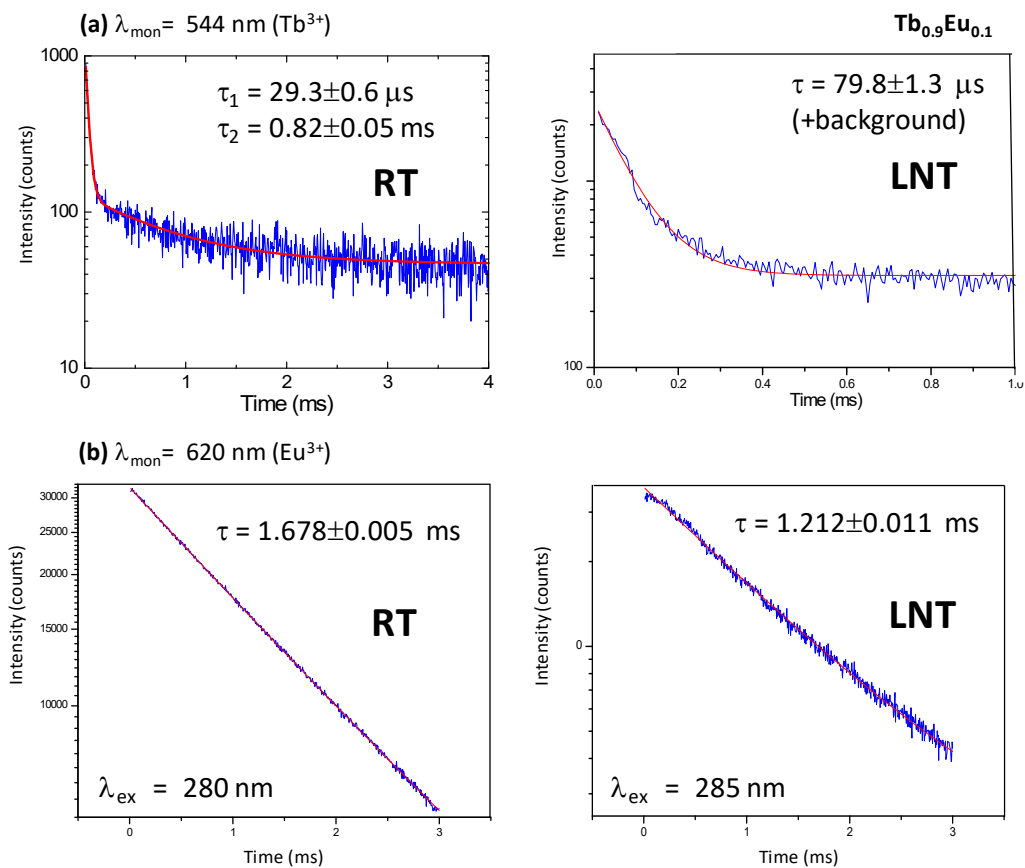


Figure S6. Lifetime measurements for heterodinuclear $\text{Tb}_{0.9}\text{Eu}_{0.1}$ compound, monitored at (a) the Tb^{3+} main peak at 544 nm and (b) the Eu^{3+} main peak at 620 nm peak at, at (left) room temperature (RT), and (right) liquid nitrogen temperature (LNT).

S6. Emission of mixed compound $Tb_{0.9}Eu_{0.1}$

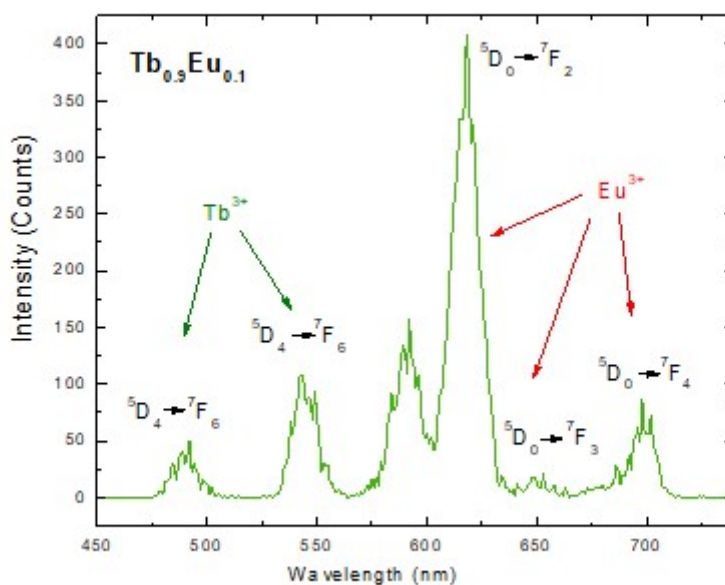


Figure S7. Emission spectra of complex $Tb_{0.9}Eu_{0.1}$ excited at $\lambda_{exc}=280$ nm, measured in. Fluorolog FL-1057, Jobin Yvon HORIBA. The characteristic emission bands for Tb^{3+} and Eu^{3+} are visible.

S7. Ac susceptibility of mixed compound $Tb_{0.9}Eu_{0.1}$

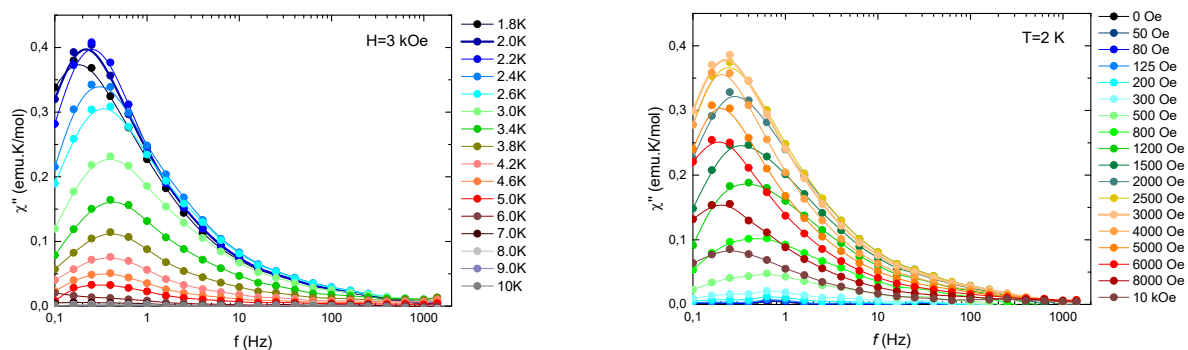


Figure S8. Ac susceptibility results. (Left) $\chi''(f, T)$ at constant magnetic field $H=3$ kOe and (Right) $\chi''(f, H)$ at constant $T=2$ K for mixed compound $Tb_{0.9}Eu_{0.1}$.

S8. Bottleneck effect

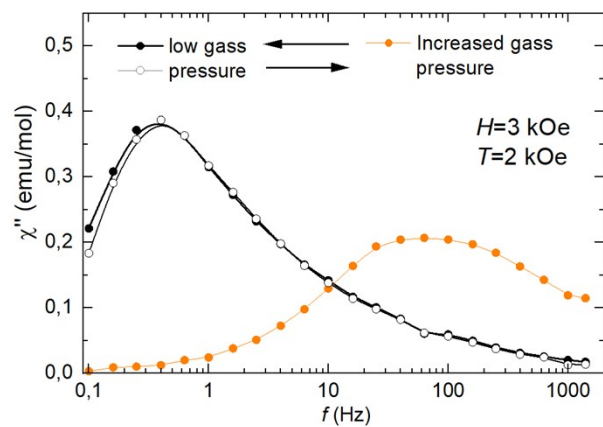


Fig. S9 $\chi''(f)$ measurements on pure **Tb** compound at $H=3$ kOe and $T=2$ K at different experimental SQUID pressure conditions, showing the influence of the bottleneck effect.