

Supplementary Information for

Multi-stimulated far-UVC luminescence for solar-blind imaging

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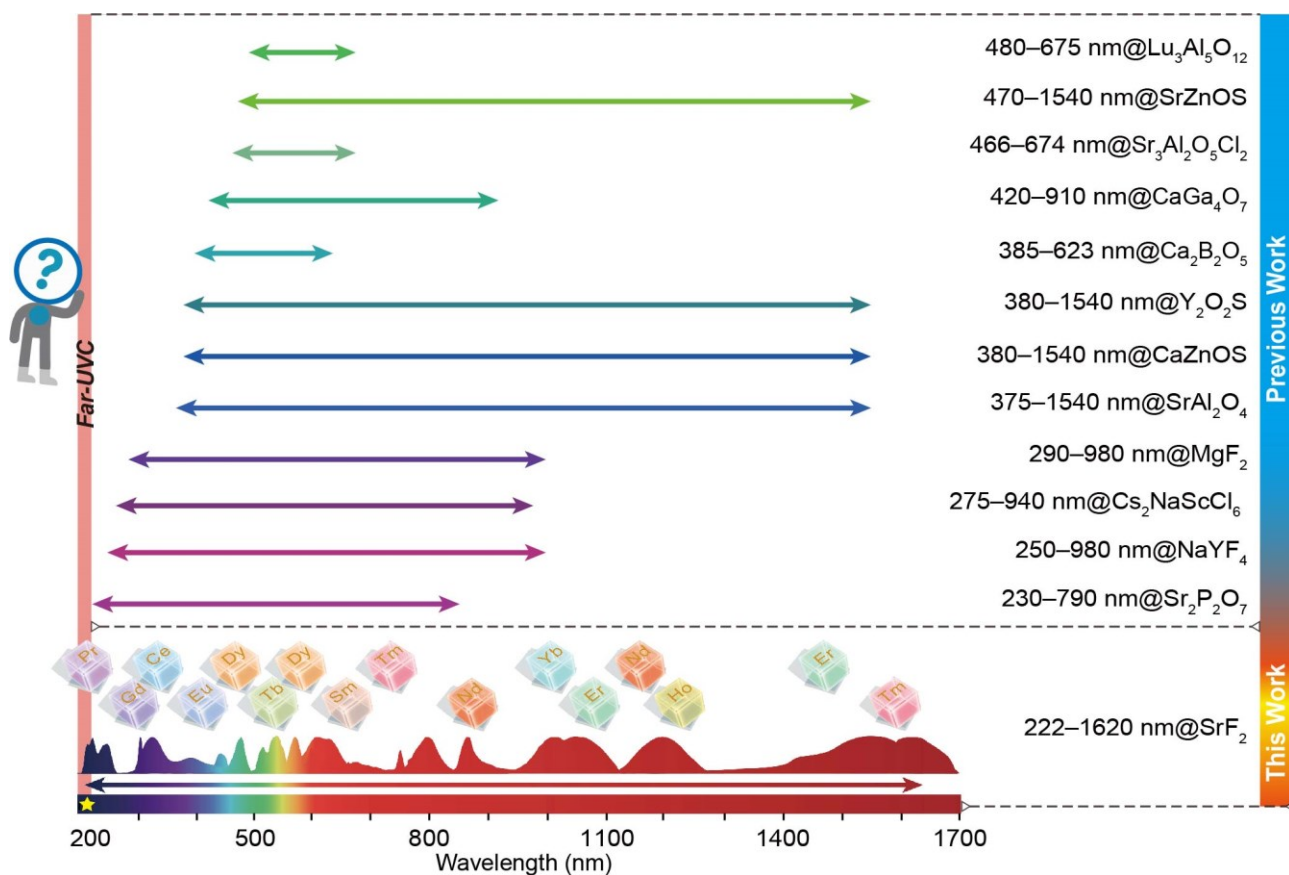
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Supplementary information file includes:

- **Supplementary Figs. 1 to 51;**
- **Supplementary Tables 1 to 4;**
- **Supplementary Notes 1 and 2;**
- **Supplementary references.**



Supplementary Fig. 1 | Comparison of mechanoluminescence spectral coverage of different systems.

Supplementary Table 1 | Comparison of mechanoluminescence spectral coverage of SrF₂ doped with different lanthanides to previously reported systems.

Host matrix	Doped ions	Emission band (nm)	References
CaZnOS	Mn ²⁺ , Cu ⁺ , Bi ³⁺ , Sb ³⁺ , Ce ³⁺ , Pr ³⁺ , Nd ³⁺ , Sm ³⁺ , Eu ³⁺ , Tb ³⁺ , Dy ³⁺ , Ho ³⁺ , Er ³⁺ , Tm ³⁺ , Yb ³⁺	380–1540	1, 2, 3, 4, 5, 6
SrZnOS	Mn ²⁺ , Pr ³⁺ , Nd ³⁺ , Sm ³⁺ , Tb ³⁺ , Dy ³⁺ , Ho ³⁺ , Er ³⁺ , Tm ³⁺ , Yb ³⁺	420–1540	7
Y ₂ O ₂ S	Pr ³⁺ , Nd ³⁺ , Sm ³⁺ , Tb ³⁺ , Dy ³⁺ , Ho ³⁺ , Er ³⁺ , Tm ³⁺ , Yb ³⁺	480–1540	8, 9
Lu ₂ O ₂ S	Eu ³⁺	580–705	8
La ₂ O ₂ S	Eu ³⁺	580–705	8
Gd ₂ O ₂ S	Eu ³⁺	615–705	8
(Zn,Cd)S	Mn ²⁺ , Ag ⁺ , Au ⁺ , Cu ²⁺	430–620	10
BaZnOS	Mn ²⁺ , Bi ³⁺	480–630	11, 12
ZnS	Cu ²⁺ , Mn ²⁺ , Mn ²⁺ /Te ²⁺	456–648	13, 14, 15, 16
MgF ₂	Mn ²⁺ , Tm ³⁺ , Yb ³⁺ /Yb ²⁺ , Dy ³⁺ , Tb ³⁺ , Eu ³⁺ /Eu ²⁺ , Sm ³⁺ , Er ³⁺ , Ho ³⁺	280–1000	17, 18, 19, 20
KMgF ₃	Tb ³⁺	494–621	21
CaF ₂	Tb ³⁺ , Sm ³⁺ , Tm ³⁺ , Eu ^{2+/3+}	425–825	22, 23
NaYF ₄	Pr ³⁺ , Sm ³⁺ , Tb ³⁺ , Dy ³⁺ , Ho ³⁺ , Er ³⁺ , Tm ³⁺	250–980	24, 25, 26
NaLuF ₄	Tb ³⁺ , Ho ³⁺	486–647	27, 28
ZnF ₂ /ZnO	Mn ²⁺	585	29
NaCa ₂ GeO ₄ F	Mn ²⁺	570	30
Sr ₃ (Al,Ga)O ₄ F	Eu ³⁺ , Tb ³⁺	488–704	31
Cs ₂ ZrCl ₆	Te ⁴⁺	576	32
CaCl ₂	Eu ²⁺	432	33
(Na,K)Cl	crystal	Not mentioned	34
Sr ₅ (PO ₄) ₃ Cl	Eu ²⁺	455	35
Sr ₃ Al ₂ O ₅ Cl ₂	Dy ³⁺ , Eu ²⁺ , Tb ³⁺ , Ce ³⁺ , Tm ³⁺ /Eu ³⁺	466–674	36, 37, 38
Sr ₅ (PO ₄) ₃ (F,Cl,Br)	Eu ³⁺ /Eu ²⁺	446–700	39
Sr ₅ (PO ₄) ₃ Cl	Eu ²⁺ , Eu ²⁺ /Mn ²⁺	400–700	40
Cs ₂ NaScCl ₆	Sb ³⁺ , Pr ³⁺ , Nd ³⁺ , Sm ³⁺ , Tb ³⁺ , Dy ³⁺ , Ho ³⁺ , Er ³⁺ , Tm ³⁺ , Yb ³⁺ , Gd ³⁺ , Ce ³⁺ , Ag ⁺ /Bi ³⁺ , Mn ²⁺	275–940	41
Cs ₂ HfCl ₆	Te ⁴⁺	557	42
Ca ₆ BaP ₄ O ₁₇	Ce ³⁺	488	43
Ca ₉ Bi(PO ₄) ₇	Ce ³⁺ , Bi ²⁺	356–682	44
SrMg ₂ (PO ₄) ₂	Eu ²⁺	412	45, 46, 47
CaZr(PO ₄) ₂	Eu ²⁺	474	48, 49, 50

Sr ₈ MgCe(PO ₄) ₇	Ce ³⁺ , Tb ³⁺ , Mn ²⁺	373–620	51
Ca ₆ BaP ₄ O ₁₇	Eu ²⁺	540	52
Ca ₉ Al(PO ₄) ₇	Tb ³⁺ , Mn ²⁺	480–648	53
Sr ₂ P ₂ O ₇	Pr ³⁺ , Tb ³⁺ , Dy ³⁺ , Sm ³⁺ , Eu ²⁺ , Ce ³⁺ , Sn ²⁺ , Mn ²⁺	230–800	54, 55
Ca ₉ Al(PO ₄) ₇	Ce ³⁺	344–700	56
Ca ₉ NaMg(PO ₄) ₇	Tb ³⁺	370–624	57
SrAl ₂ O ₄	Eu ²⁺ /Dy ³⁺ , Ce ³⁺ /Ho ³⁺ , Eu ²⁺ /Er ³⁺ , Eu ²⁺ /Cr ³⁺ /Nd ³⁺ , Ce ³⁺ , Eu ²⁺ , Nd ³⁺ /Eu ²⁺ , Er ³⁺ /Eu ²⁺ , Ce ³⁺ /Ho ³⁺	375–1540	58
LaAlO ₃	Cr ³⁺	685–750	59
Lu ₃ Al ₅ O ₁₂	Ce ³⁺ , Pr ³⁺ , Eu ³⁺ , Tb ³⁺ , Sm ³⁺ , Ho ³⁺ , Dy ³⁺ , Er ³⁺ , Nd ³⁺ , Tm ³⁺	480–725	60
Y ₃ Al ₅ O ₁₂	Ce ³⁺ , Cr ³⁺	560–688	61, 62
CaYAl ₃ O ₇	Eu ²⁺ , Ce ³⁺	421–445	63, 64
ZnAl ₂ O ₄	Mn ²⁺	512	65
SrMgAl ₁₀ O ₁₇	Ce ³⁺	400–480	66
SrMgAl ₆ O ₁₁	Eu ²⁺	512	67
BaMgAl ₁₂ O ₁₇	Eu ²⁺	450	68
CaLaAl ₃ O ₇	Tb ³⁺	413–622	69
BaLaAl ₃ O ₇	Eu ³⁺	595–615	70
CaGa ₄ O ₇	Pr ³⁺ , Tb ³⁺ , Er ³⁺ , Dy ³⁺ , Mn ²⁺ , Sm ³⁺ , Eu ³⁺ , Tm ³⁺ , Nd ³⁺	420–910	71, 72
Ca ₅ Ga ₆ O ₁₄	Tb ³⁺	484–625	73
Y ₃ Ga ₅ O ₁₂	Tb ³⁺	380–690	74
MgGa ₂ O ₄	Mn ²⁺ , Cr ³⁺ , Bi ³⁺	480–710	75, 76, 77,
CaGa ₂ O ₄	Pr ³⁺ /Li ⁺	485–651	78
Ca ₃ Ga ₄ O ₉	Tb ³⁺ , Eu ³⁺	380–701	79, 80
SrGa ₁₂ O ₁₉	Cr ³⁺ , Cr ³⁺ /Pr ³⁺	710–750	81, 82, 83
Y ₃ Al ₂ Ga ₃ O ₁₂	Gd ³⁺	310	84
Lu ₃ Al ₂ Ga ₃ O ₁₂	Cr ³⁺ , Tb ³⁺	380–720	85, 86
Y ₃ Al ₃ Ga ₂ O ₁₂	Ce ³⁺	528	87
CaSrGa ₄ O ₈	Tb ³⁺	480–630	88
Y ₃ GaO ₆	Tb ³⁺	484–625	89
LiGa ₅ O ₈	Matrix, Cr ³⁺ , Pr ³⁺	525–716	81, 90, 91
LiGaO ₂	Fe ³⁺ , Mn ²⁺	660–746	92, 93
ZnGa ₂ O ₄	Mn ²⁺ , Cr ³⁺	505–692	75, 94
Lu ₃ Al ₂ Ga ₃ O ₁₂	Tb ³⁺ , Eu ³⁺	383–709	95
(Li,Na) ₂ (Zn,Mg)GeO ₄	Mn ²⁺	515–535	96
Zn ₃ Ga ₂ GeO ₈	Cr ³⁺	700	81

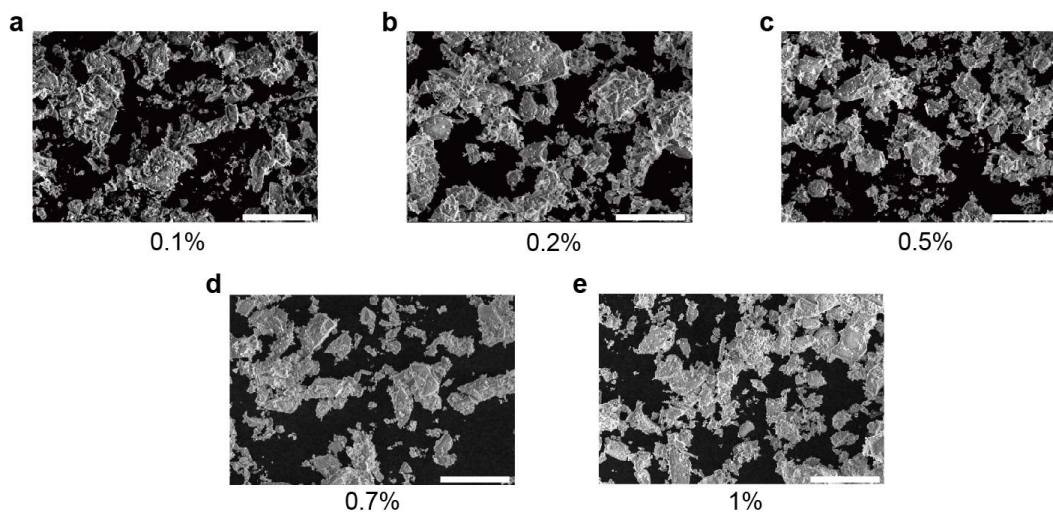
CaZnGe ₂ O ₆	Mn ²⁺	676	97, 98
Ca ₂ YGa ₃ Ge ₂ O ₁₂	Cr ³⁺ , Cr ³⁺ /Nd ³⁺	767–1340	99
Mg ₃ Ga ₂ GeO ₈	Cr ³⁺	740–801	100
MgGeO ₃	Mn ²⁺	691	101
CaMgGe ₂ O ₆	Mn ²⁺	670	102
Zn ₂ (Ge _{0.9} Si _{0.1})O ₄	Mn ²⁺	535	103
Na ₂ ZnSiO ₄	Mn ²⁺	514	96
Sr ₂ MgSi ₂ O ₇	Eu ²⁺ /Dy ³⁺ , Eu ²⁺	470–580	104, 105
Ca ₂ MgSi ₂ O ₇	Eu ²⁺	545	106
Ca ₂ Al ₂ SiO ₇	Ce ³⁺	400	107
Sr(Ba,Ca,Sr)MgSi ₂ O ₇	Eu ²⁺	450–490	108
(Ca,Sr)Al ₂ Si ₂ O ₈	Eu ²⁺	403–428	109
(Ca,Sr) ₈ Mg ₃ Al ₂ Si ₇ O ₂₈	Ce ³⁺	395–405	110
Ca ₂ Ga ₂ SiO ₇	Tb ³⁺	495–620	111
Y ₃ Ga ₃ MgSiO ₁₂	Cr ³⁺	731–780	112
CaNb ₂ O ₆	Pr ³⁺	613	113
Ca ₂ Nb ₂ O ₇	Pr ³⁺	613	113
Ca ₃ Nb ₂ O ₈	Pr ³⁺	613	113
Sr ₂ Nb ₂ O ₇	Pr ³⁺	red	114
NaNbO ₃	Pr ³⁺ /Er ³⁺	613	115
LiNbO ₃	Pr ³⁺ , Nd ³⁺	619–895	116, 117
(Na,Mg)NbO ₃	Pr ³⁺	605	118
LiTaO ₃	Tb ³⁺ , Pr ³⁺ /Gd ³⁺	480–625	119, 120
(Ba,Ca)TiO ₃	Pr ³⁺	620	121, 122, 123
Sr ₃ Sn ₂ O ₇	Sm ³⁺ , Nd ³⁺ , Nd ³⁺ /Yb ³⁺	570–1350	124, 125, 126, 127, 128, 129, 130
Sr ₂ SnO ₄	Sm ³⁺	574–745	131
SrZn ₂ S ₂ O	Mn ²⁺	610	132
ZrO ₂	Sm ³⁺ /Nd ³⁺ , Ti ⁴⁺	478–660	133, 134
Ga ₂ O ₃	Cr ³⁺ /In ³⁺	715–830	135
BaSi ₂ O ₂ N ₂	Eu ²⁺	500	8
Ba _{0.5} Sr _{0.5} Si ₂ O ₂ N ₂	Eu ²⁺	505	61
B ₂ O ₃ -SiO ₂ -ZnO	Tb ³⁺ , Tm ³⁺ , Eu ³⁺	455–710	136
Ca ₂ B ₂ O ₅	Tb ³⁺ , Dy ³⁺ , Sm ³⁺ , Eu ³⁺ , Ce ³⁺	350–800	137
(Ca,Sr,Ba) ₂ (Ga,Ln,Sc, Y)SbO ₆	Cr ³⁺	825–1010	138
BaSi ₂ O ₂ N ₂	Eu ²⁺	498	139
ZnB ₂ O ₄	Mn ²⁺	550	140

Supplementary Table 2 | Comparison of radius of different lanthanides and Sr^{2+} in an octahedral coordination environment.

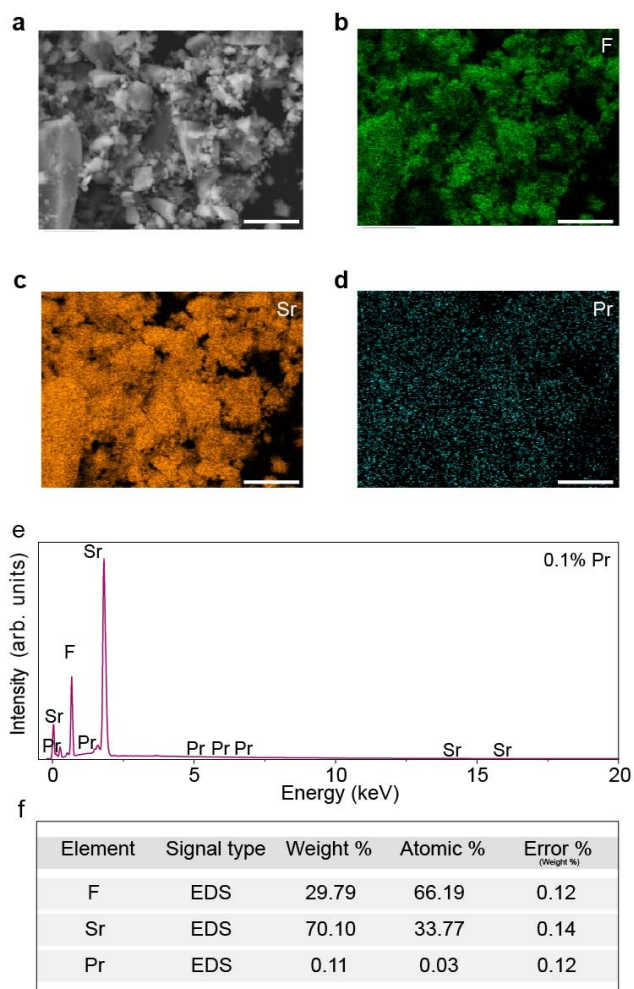
Ion	Ion radius (pm)
Sr^{2+}	126
Ce^{3+}	114.3
Pr^{3+}	112.6
Nd^{3+}	110.9
Sm^{3+}	107.9
Eu^{2+}	125
Eu^{3+}	106.6
Gd^{3+}	105.3
Tb^{3+}	104.0
Dy^{3+}	102.7
Ho^{3+}	101.5
Er^{3+}	100.4
Tm^{3+}	99.4
Yb^{3+}	98.5

Supplementary Table 3 | Structural information of SrF₂:Pr³⁺ determined by the refinement results.

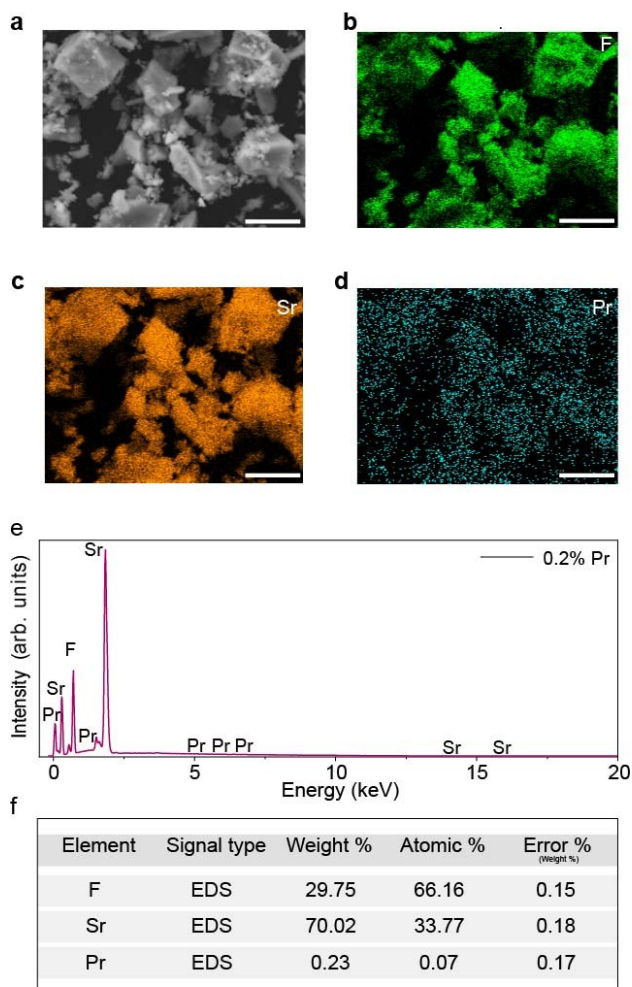
SrF₂:Pr³⁺	
Space group	<i>Fm-3m</i>
R_p	9.96
R_{wp}	13.72
χ^2	2.336
a (Å)	5.7986
b (Å)	5.7986
c (Å)	5.7986
$\alpha=\beta=\gamma$	90.000°



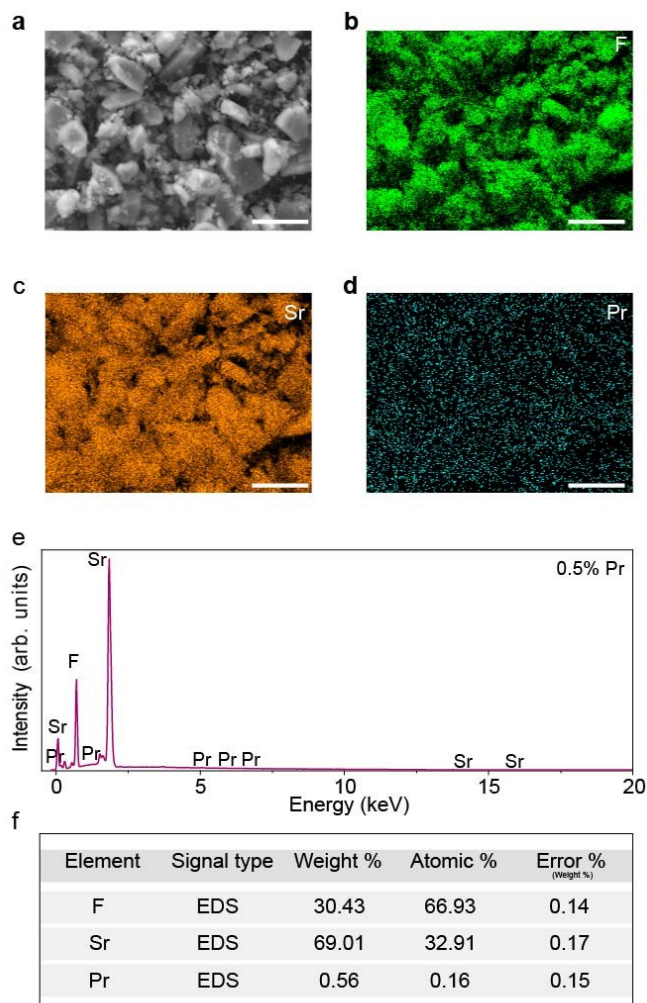
Supplementary Fig. 2 | Scanning electron microscopy images of SrF₂ doped with different concentrations. a SrF₂:0.1%Pr³⁺, b SrF₂:0.2%Pr³⁺, c SrF₂:0.5%Pr³⁺, d SrF₂:0.7%Pr³⁺, and e SrF₂:1%Pr³⁺. Scale bar: 50 μ m.



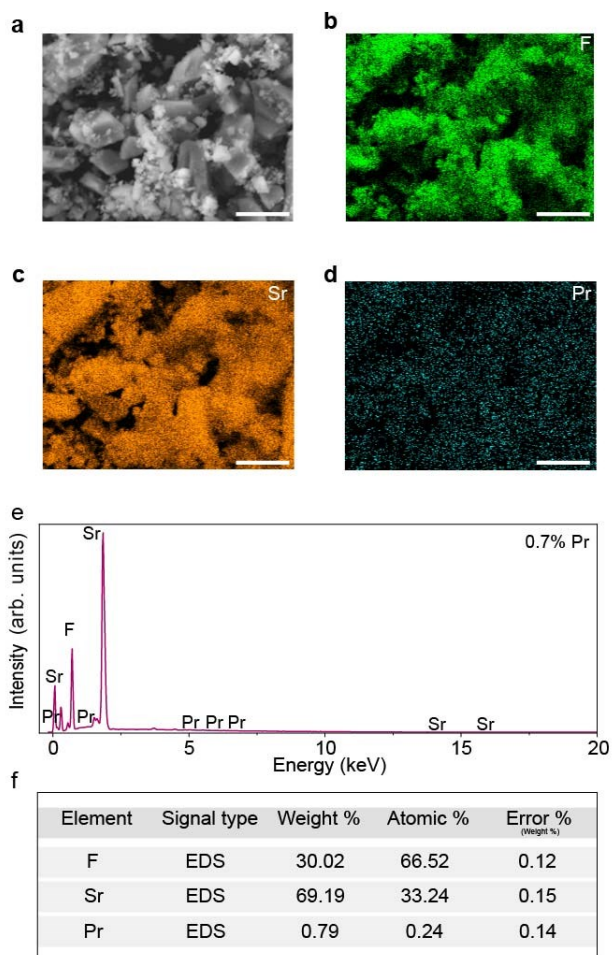
Supplementary Fig. 3 | Elemental analysis of $\text{SrF}_2:0.1\%\text{Pr}^{3+}$. **a** Scanning electron microscopy image, **b** F, **c** Sr, and **d** Pr elemental mappings, **e** Energy dispersive spectroscopy spectrum and **f** Elemental analysis data of $\text{SrF}_2:0.1\%\text{Pr}^{3+}$. Scale bar: 50 μm .



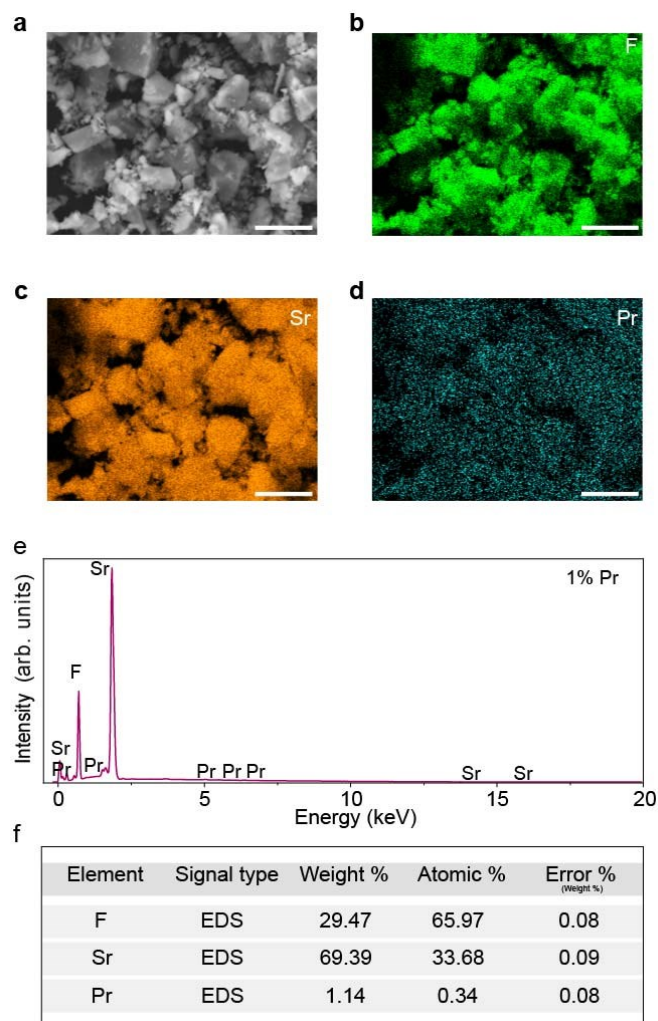
Supplementary Fig. 4 | Elemental analysis of $\text{SrF}_2:0.2\%\text{Pr}^{3+}$. **a** Scanning electron microscopy image, **b** F, **c** Sr, and **d** Pr elemental mappings, **e** Energy dispersive spectroscopy spectrum and **f** Elemental analysis data of $\text{SrF}_2:0.2\%\text{Pr}^{3+}$. Scale bar: 50 μm .



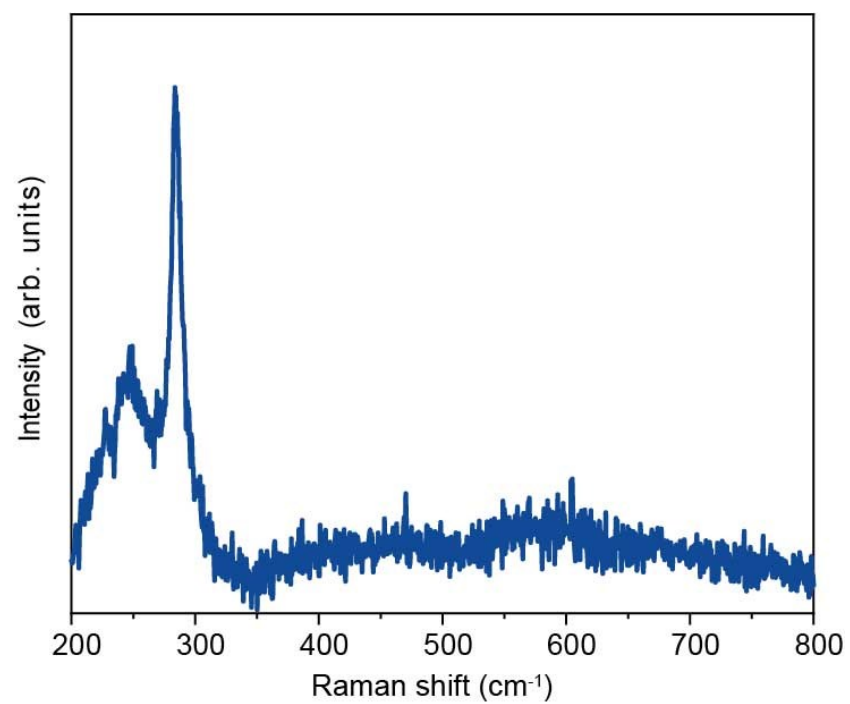
Supplementary Fig. 5 | Elemental analysis of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$. **a** Scanning electron microscopy image, **b** F, **c** Sr, and **d** Pr elemental mappings, **e** Energy dispersive spectroscopy spectrum and **f** Elemental analysis data of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$. Scale bar: 50 μm .



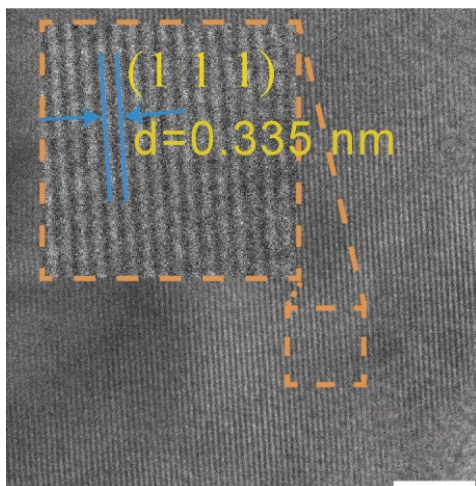
Supplementary Fig. 6 | Elemental analysis of $\text{SrF}_2:0.7\%\text{Pr}^{3+}$. **a** Scanning electron microscopy image, **b** F, **c** Sr, and **d** Pr elemental mappings, **e** Energy dispersive spectroscopy spectrum and **f** Elemental analysis data of $\text{SrF}_2:0.7\%\text{Pr}^{3+}$. Scale bar: 50 μm .



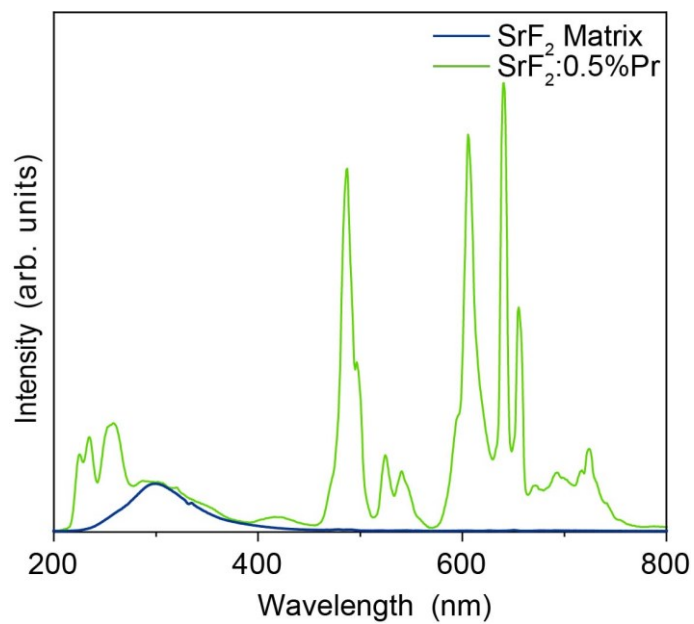
Supplementary Fig. 7 | Elemental analysis of $\text{SrF}_2:1\%\text{Pr}^{3+}$. **a** Scanning electron microscopy image, **b** F, **c** Sr, and **d** Pr elemental mappings, **e** Energy dispersive spectroscopy spectrum and **f** Elemental analysis data of $\text{SrF}_2:1\%\text{Pr}^{3+}$. Scale bar: 50 μm .



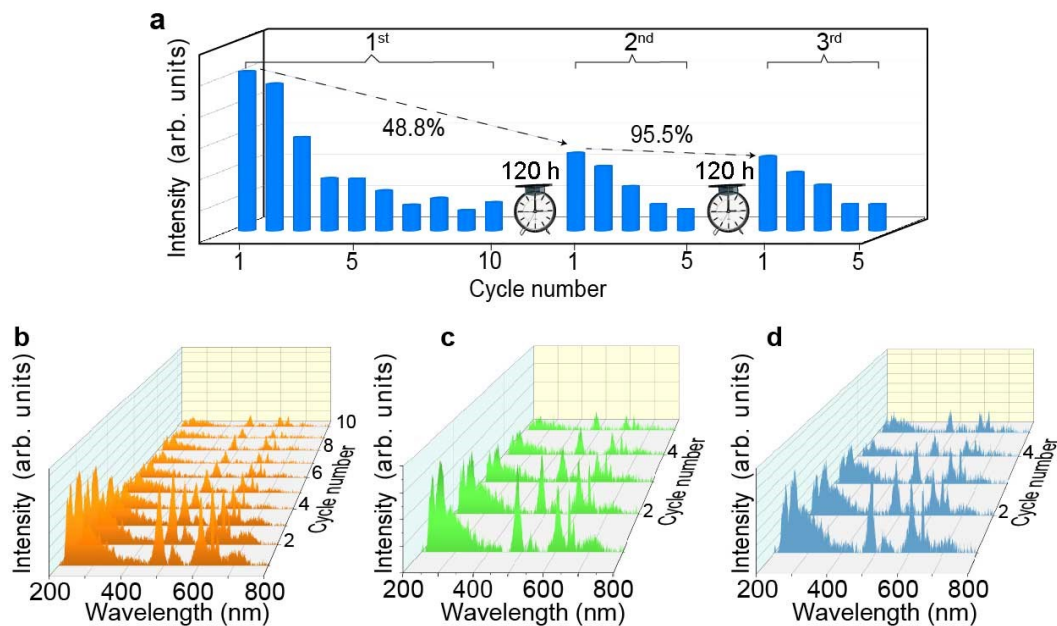
Supplementary Fig. 8 | Raman spectrum of SrF₂:0.5%Pr³⁺.



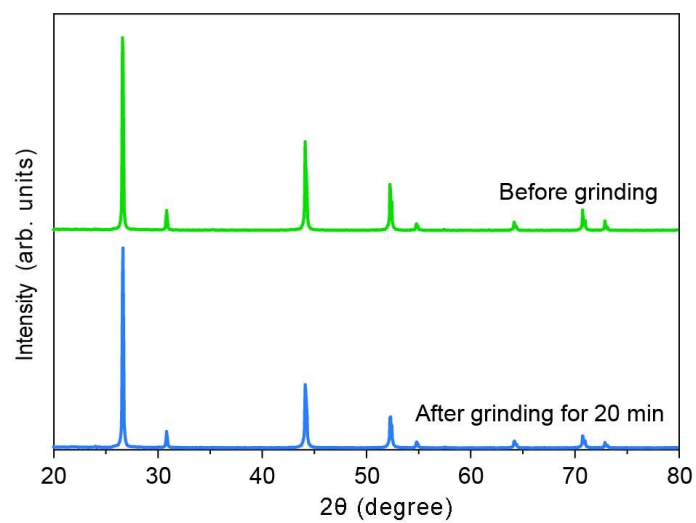
Supplementary Fig. 9 | High-resolution transmission electron microscopy image of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$ (Scale bar: 5 nm).



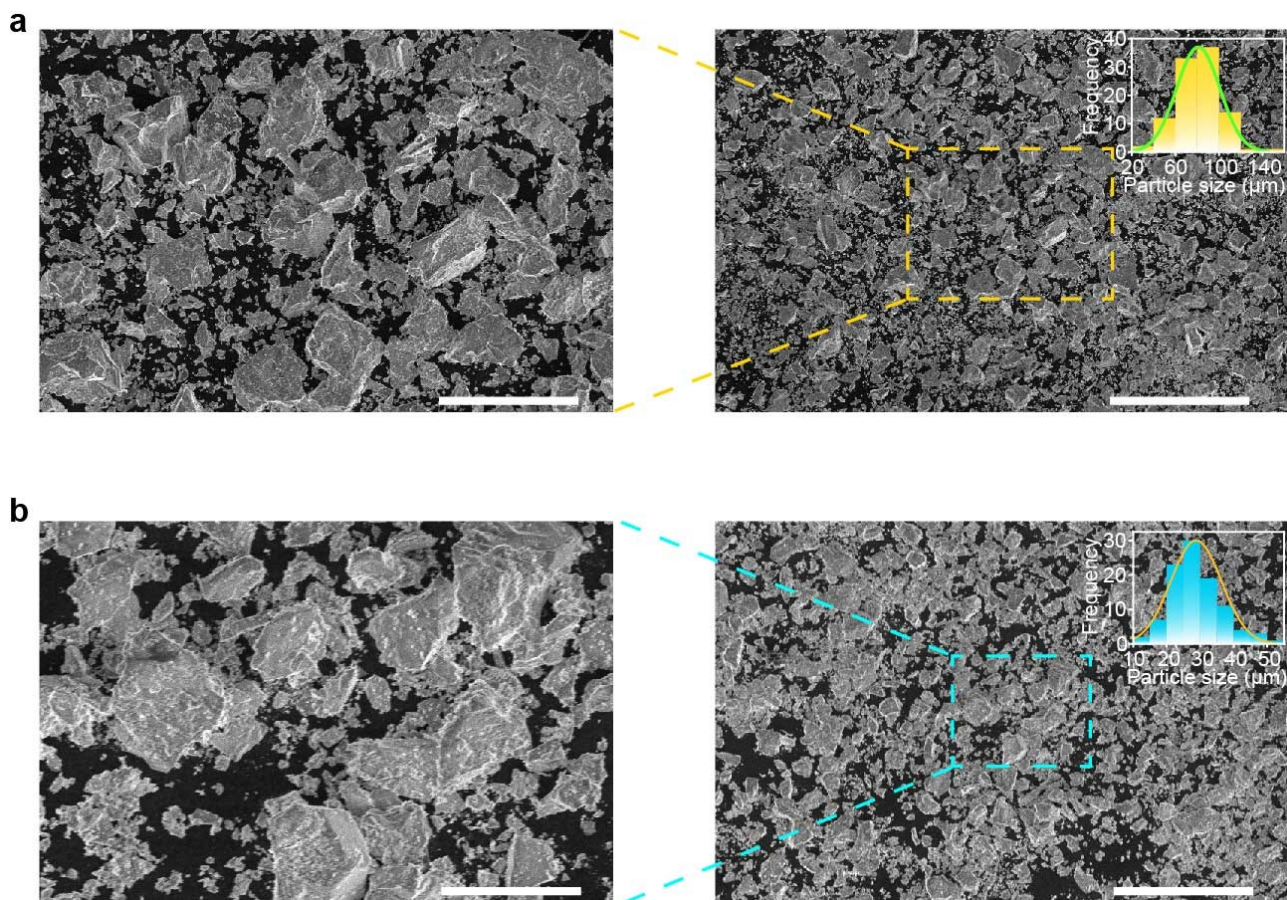
Supplementary Fig. 10 | Comparison of photoluminescence spectra between SrF₂:0.5%Pr³⁺ and SrF₂ matrix. The broadband peak at 300 nm is attributed to the luminescence from the self-trapped exciton of the host.



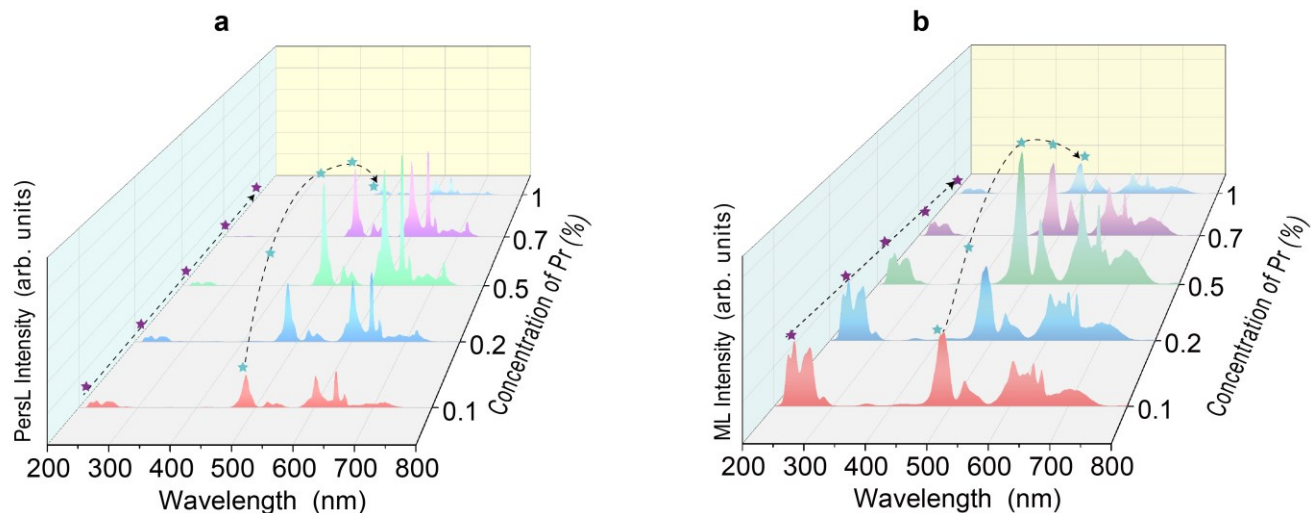
Supplementary Fig. 11 | Repeatability test of self-recoverable mechanoluminescence of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$ before X-ray charging. **a** Repeatability test results of self-recoverable mechanoluminescence intensity. Original ML spectra collected during the **b** First, **c** Second and **d** Third cycles.



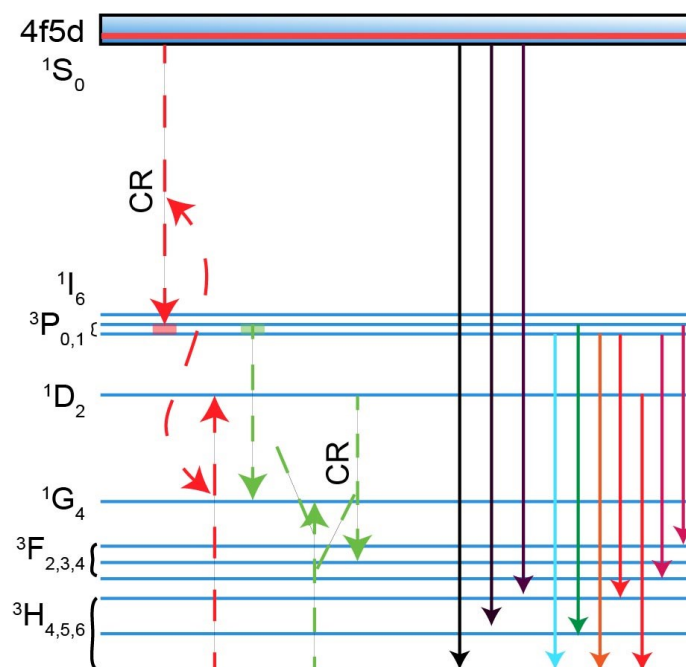
Supplementary Fig. 12 | X-ray diffraction patterns of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$ before and after grinding for 20 min.



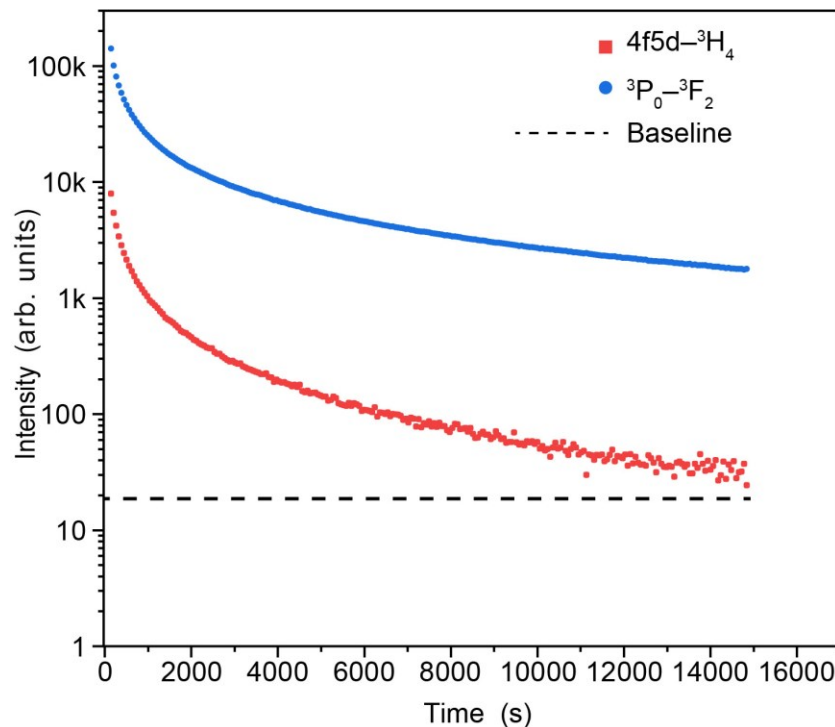
Supplementary Fig. 13 | Comparison of particle size of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$ before and after grinding. Scanning electron microscopy images of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$: **a** Before (Scale bar: 500 μm) and **b** After continuous grinding for 20 min (Scale bar: 200 μm).



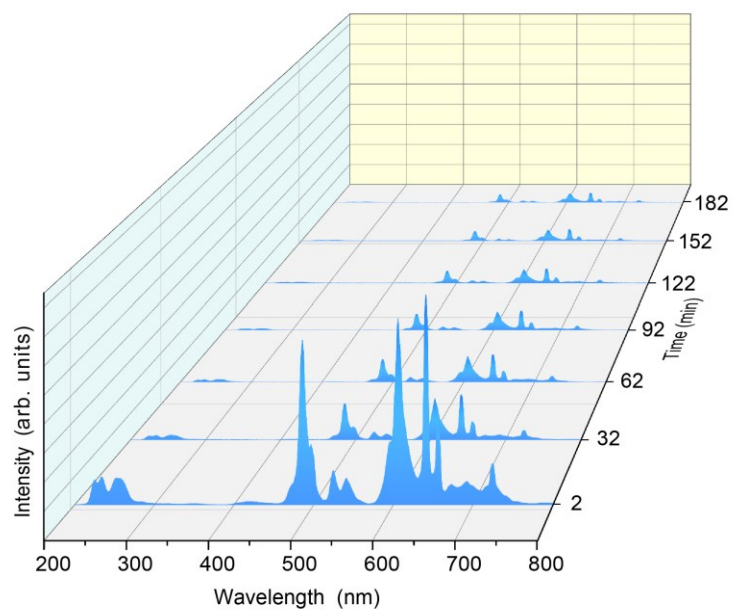
Supplementary Fig. 14 | Persistent luminescence (PersL) and mechanoluminescence (ML) spectra of SrF₂ doped with different concentration of Pr³⁺. a PersL spectra and b ML spectra of SrF₂ doped with $x\%$ mol Pr³⁺ ($x = 0.1, 0.2, 0.5, 0.7$ and 1).



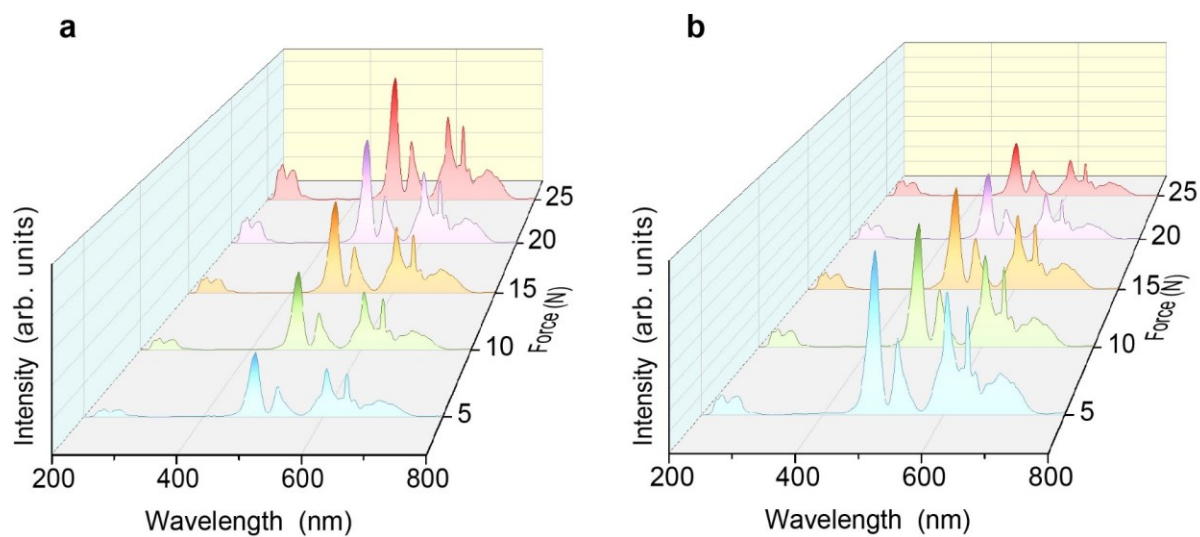
Supplementary Fig. 15 | Energy levels diagram of Pr^{3+} and possible cross-relaxation (CR) processes.



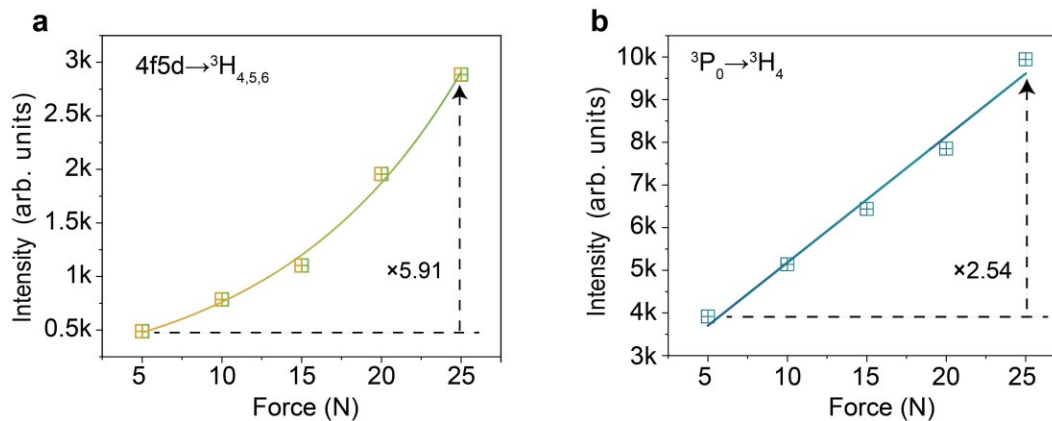
Supplementary Fig. 16 | Comparison of visible and UVC persistent luminescence (PersL) decay curves of SrF₂:0.5%Pr³⁺. For the visible PersL decay curve, the $^3P_0 \rightarrow ^3F_2$ transition of Pr³⁺ was monitored. For the UVC PersL decay curve, the $4f5d \rightarrow ^3H_4$ transition of Pr³⁺ was monitored. Before test, the sample of SrF₂:0.5%Pr³⁺ was subjected to X-ray irradiation for 5 min.



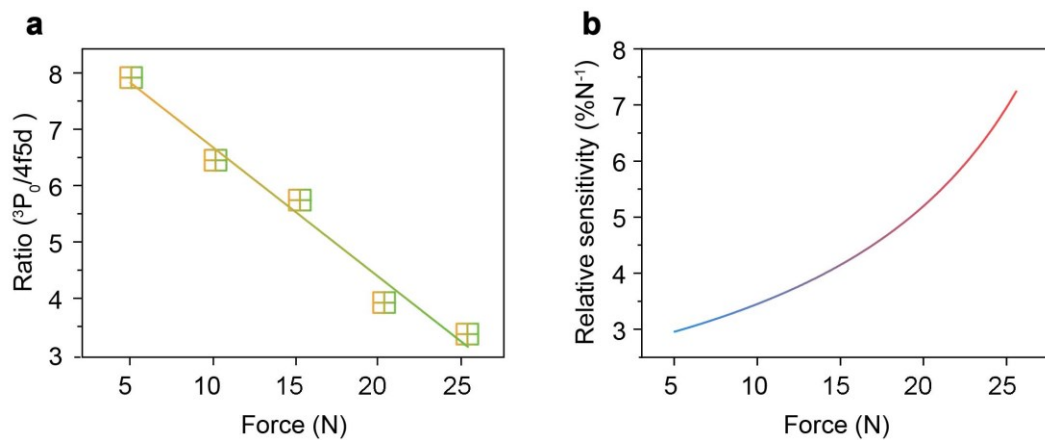
Supplementary Fig. 17 | Persistent luminescence spectra of SrF₂:0.5%Pr³⁺ following 5 min of X-ray irradiation.



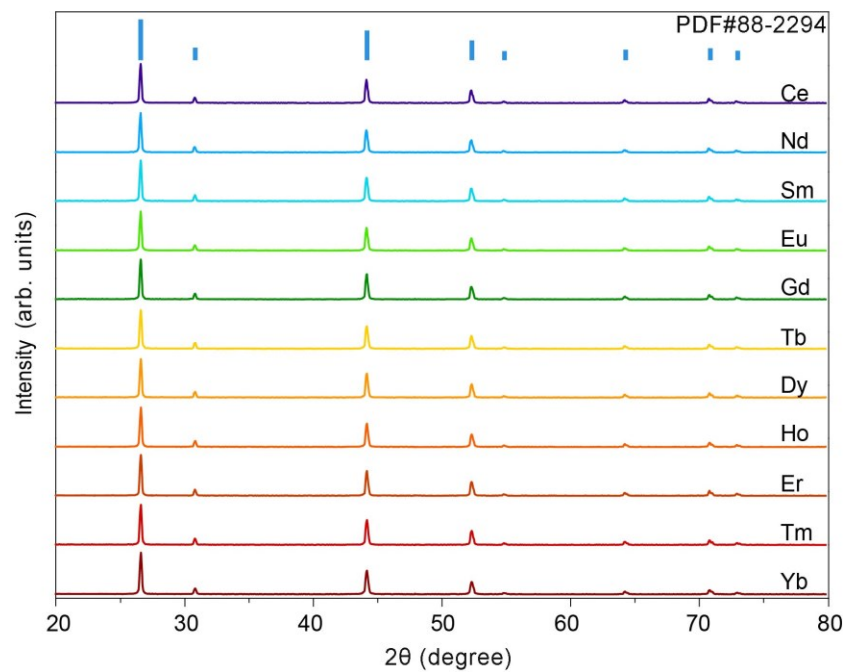
Supplementary Fig. 18 | Mechanoluminescence spectra under different forces. a Original and **b** Normalized mechanoluminescence spectra of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$.



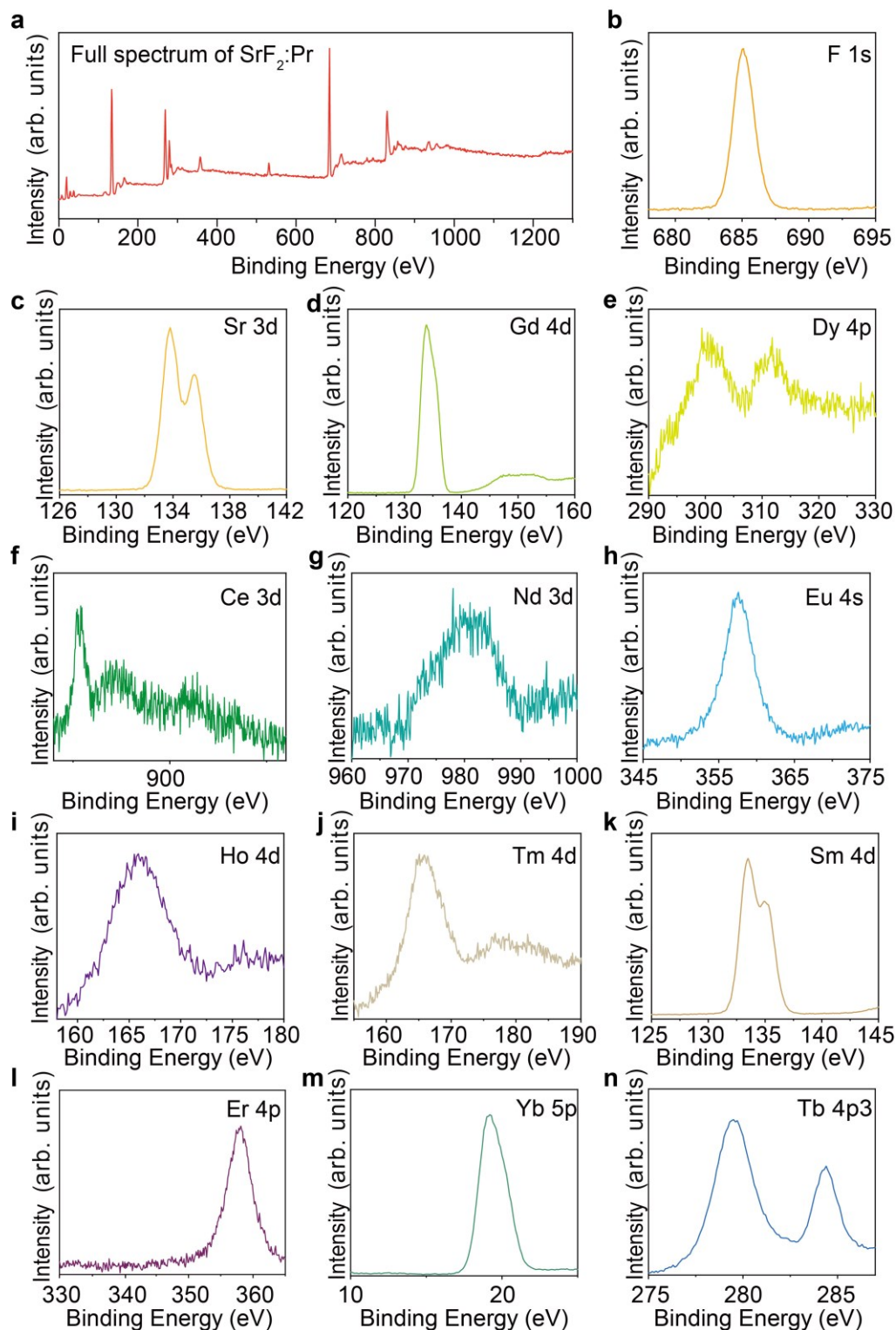
Supplementary Fig. 19 | Effect of force on mechanoluminescence intensity. Mechanoluminescence intensity of **a** $4f5d \rightarrow {}^3\text{H}_{4,5,6}$ and **b** ${}^3\text{P}_0 \rightarrow {}^3\text{H}_4$ transitions of Pr^{3+} of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$ as a function of external force in the range of 5–25 N.



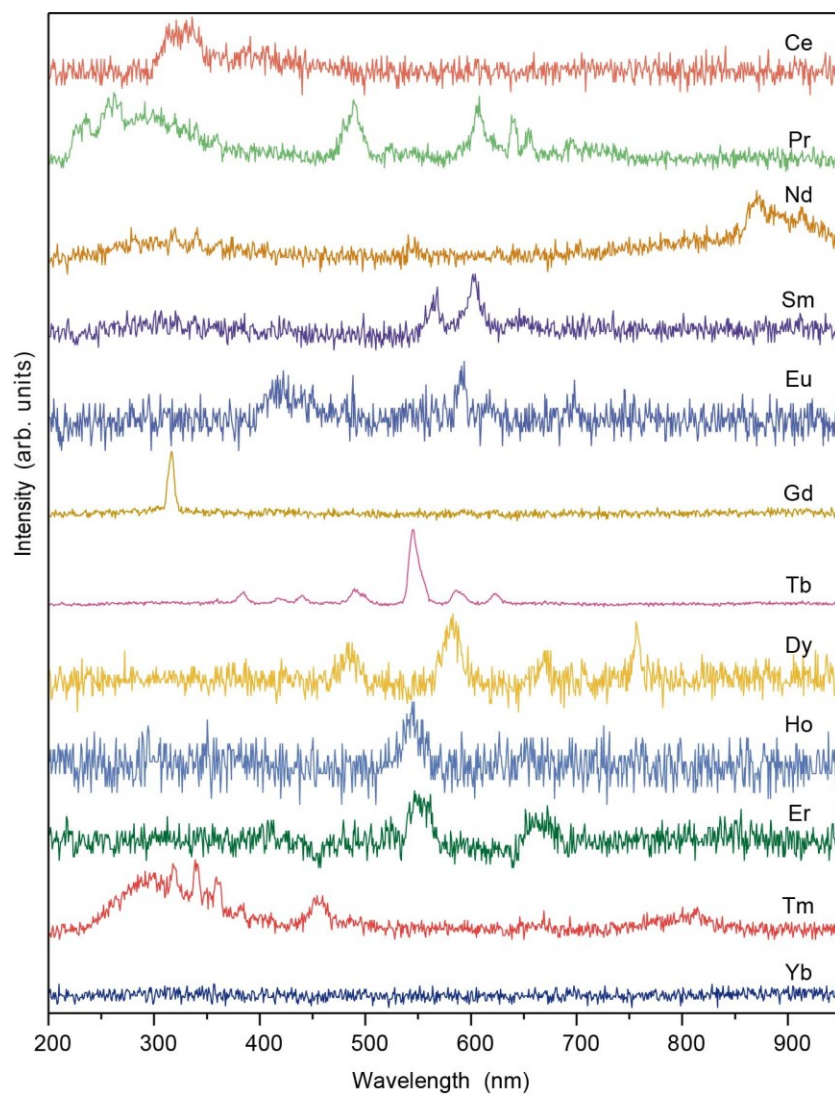
Supplementary Fig. 20 | Ratiometric stress sensing. **a** Mechanoluminescence intensity ratio between the $^3P_0 \rightarrow ^3H_4$ and $4f5d \rightarrow ^3H_{4,5,6}$ transitions of Pr^{3+} of $SrF_2:0.5\%Pr^{3+}$, and **b** Relative sensitivity of proposed ratiometric stress sensing method as a function of external force in the range of 5–25 N.



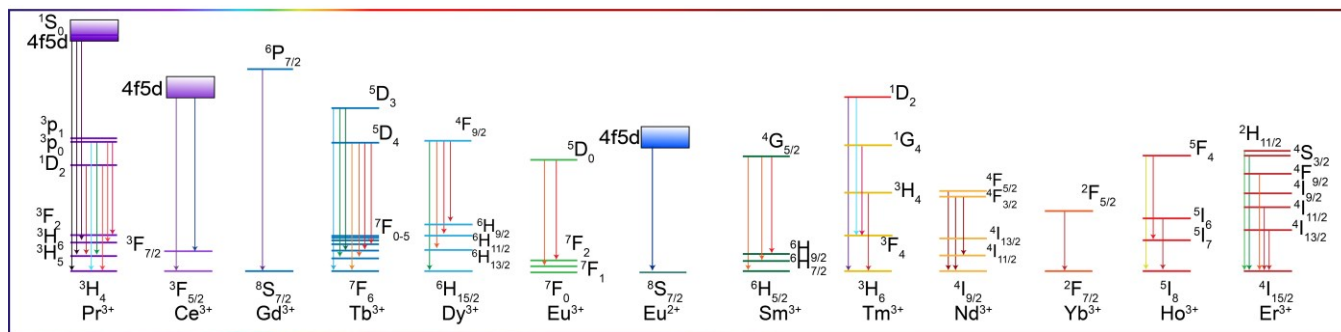
Supplementary Fig. 21 | X-ray diffraction patterns of SrF₂ doped with different lanthanides (0.5%mol).



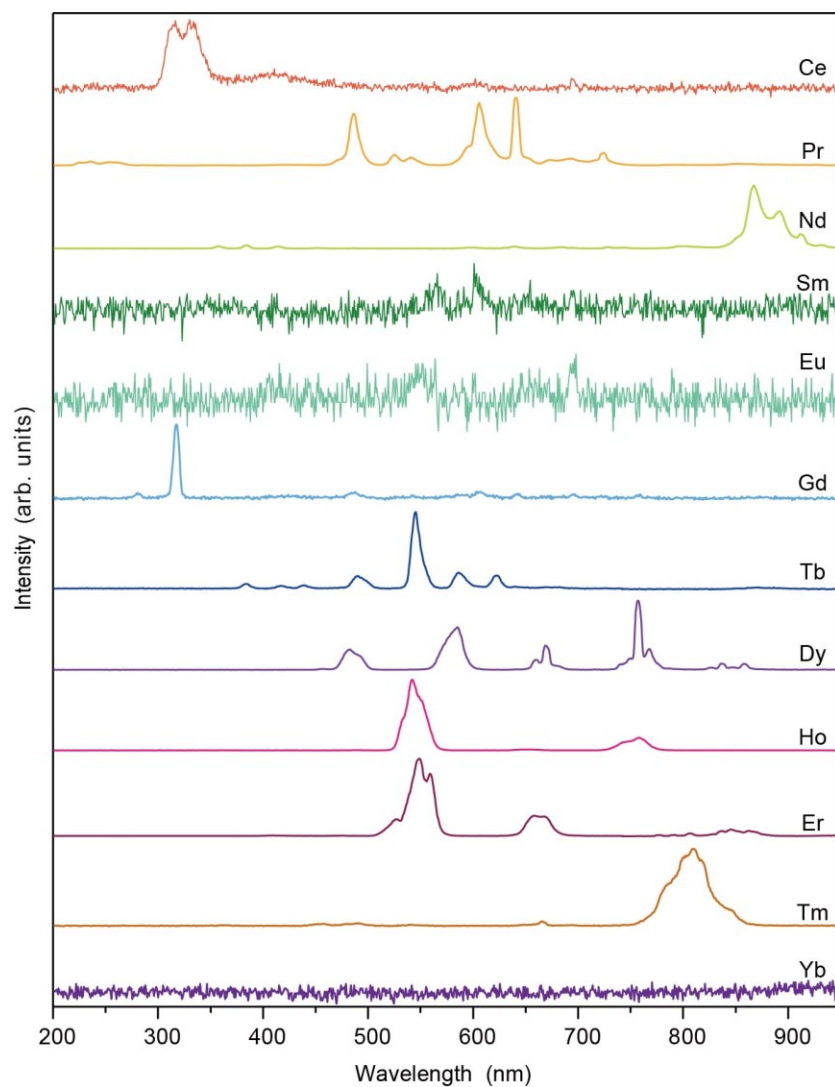
Supplementary Fig. 22 | X-ray photoelectron spectroscopy results of SrF_2 doped with different lanthanides (0.5%mol). a Full spectrum of $\text{SrF}_2:\text{Pr}$. **b–n** specific lanthanides: **b** F, **c** Sr, **d** Gd, **e** Dy, **f** Ce, **g** Nd, **h** Eu, **i** Ho, **j** Tm, **k** Sm, **l** Er, **m** Yb and **n** Tb.



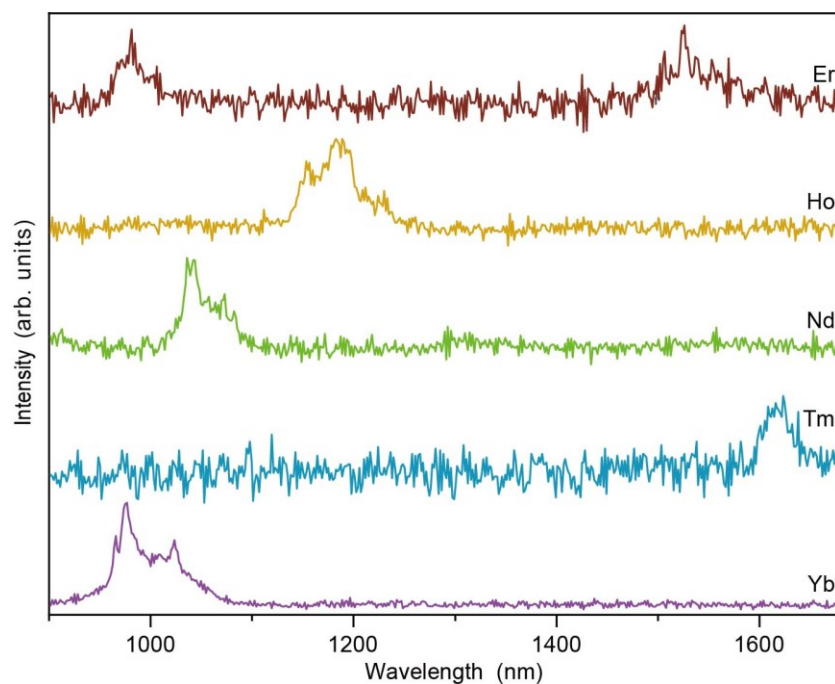
Supplementary Fig. 23 | Mechanoluminescence spectra of SrF_2 doped with different lanthanides (0.5%mol) before X-ray charging.



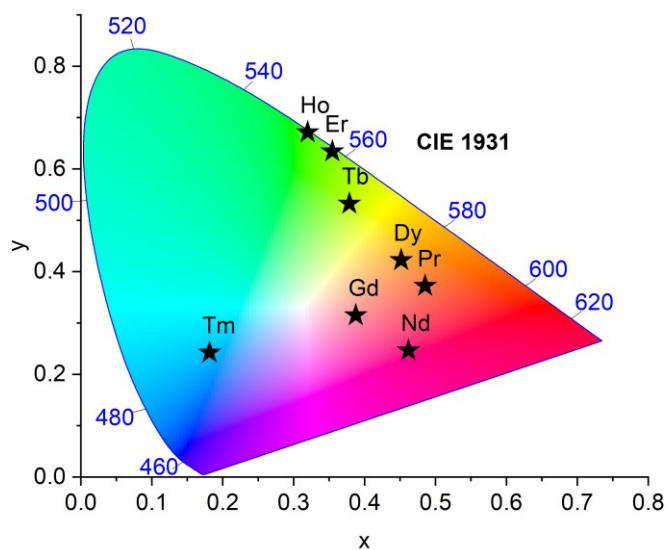
Supplementary Fig. 24 | Energy level diagrams of Pr^{3+} , Ce^{3+} , Gd^{3+} , Tb^{3+} , Dy^{3+} , $\text{Eu}^{2+,3+}$, Sm^{3+} , Tm^{3+} , Nd^{3+} , Yb^{3+} , Ho^{3+} and Er^{3+} .



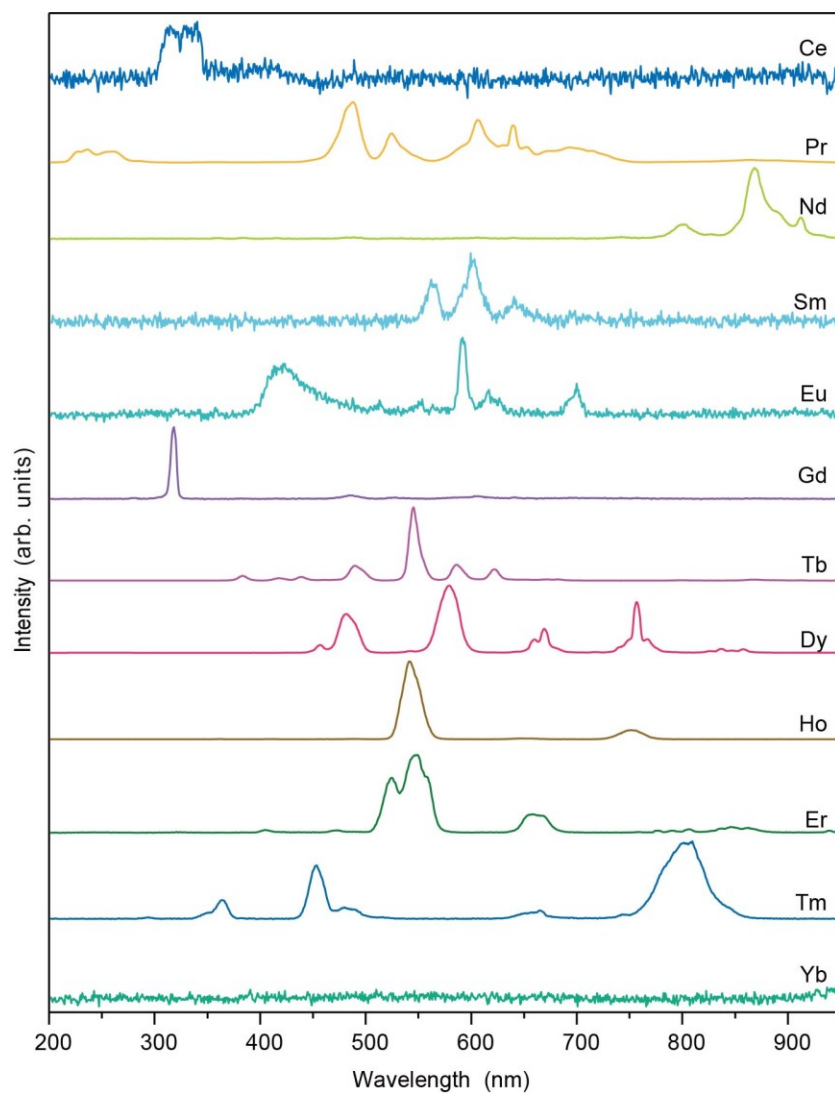
Supplementary Fig. 25 | Persistent luminescence spectra of SrF_2 doped with different lanthanides (0.5%mol) after X-ray irradiation over the 200–950 nm range.



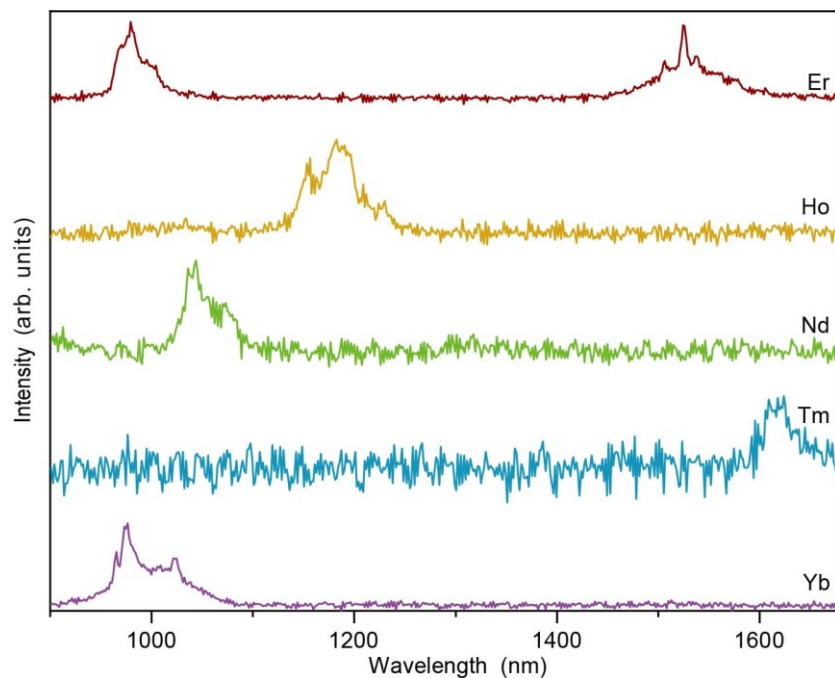
Supplementary Fig. 26 | Persistent luminescence spectra of SrF₂ doped with different lanthanides (0.5%mol) after X-ray irradiation over the 900–1680 nm range.



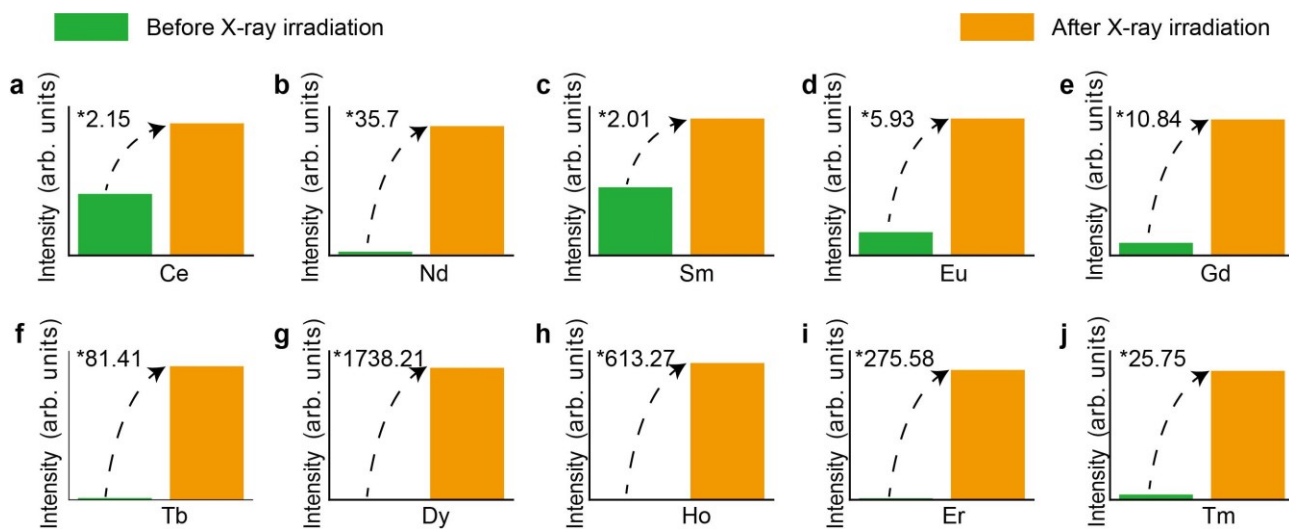
Supplementary Fig. 27 | Commission Internationale de l'Éclairage (CIE) chromatic coordinates of persistent luminescence from SrF₂ doped with different lanthanides (0.5%mol).



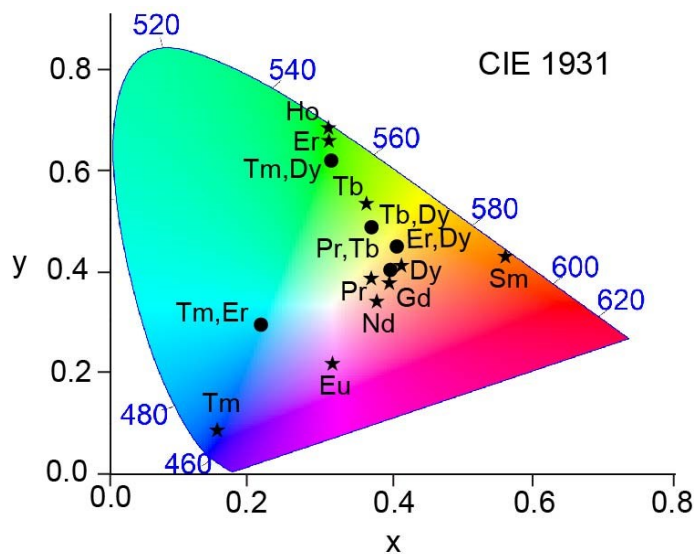
Supplementary Fig. 28 | Mechanoluminescence spectra of SrF_2 doped with different lanthanides (0.5%mol) after X-ray irradiation over the 200–950 nm range.



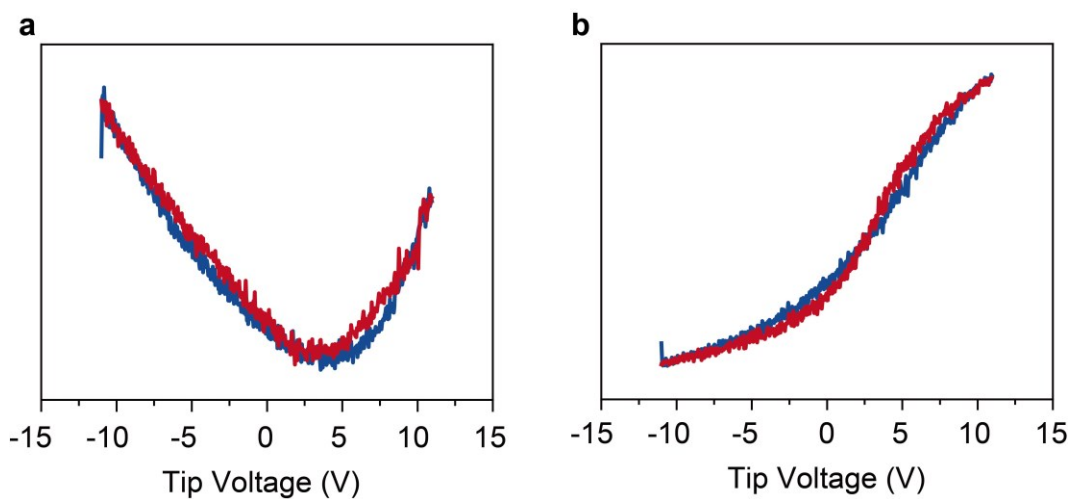
Supplementary Fig. 29 | Mechanoluminescence spectra of SrF₂ doped with different lanthanides (0.5%mol) after X-ray irradiation over the 900–1680 nm range.



Supplementary Fig. 30 | Comparison of mechanoluminescence intensity of SrF₂ doped with different lanthanides (0.5%mol) before and after X-ray irradiation. a Ce, b Nd, c Sm, d Eu, e Gd, f Tb, g Dy, h Ho, i Er and j Tm.

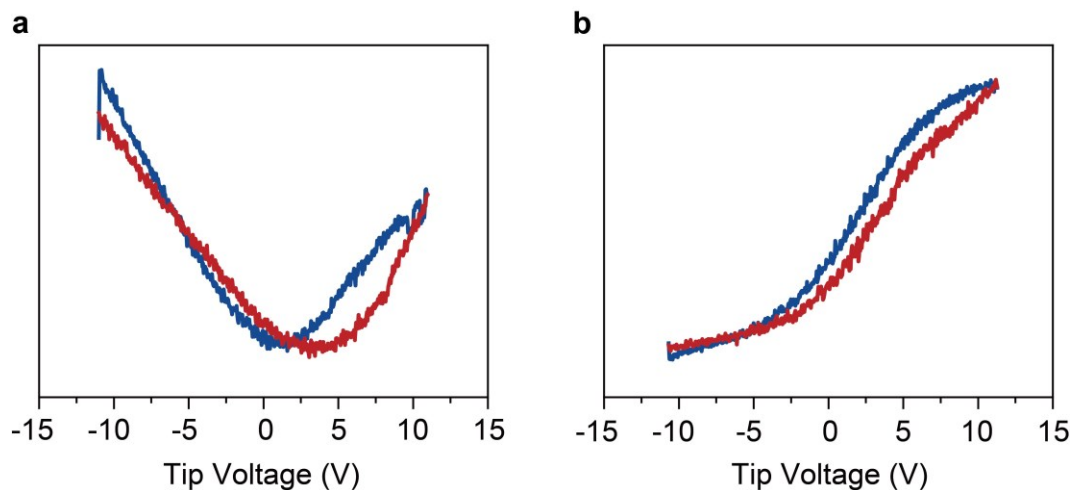


Supplementary Fig. 31 | Commission Internationale de l'Éclairage (CIE) chromatic coordinates of mechanoluminescence from SrF₂ doped with different lanthanides (0.5%mol). Circle symbols represent doubly doped samples with both dopants in 0.5%mol concentration.



Supplementary Fig. 32 | Amplitude curves (bi-directional bias) and ferroelectric phase curve of SrF₂.

a Amplitude curve. The red and blue curves correspond to signal variations for the bias voltage changing from -10 to +10 V and from +10 to -10 V, respectively. **b** Standard ferroelectric phase curve.



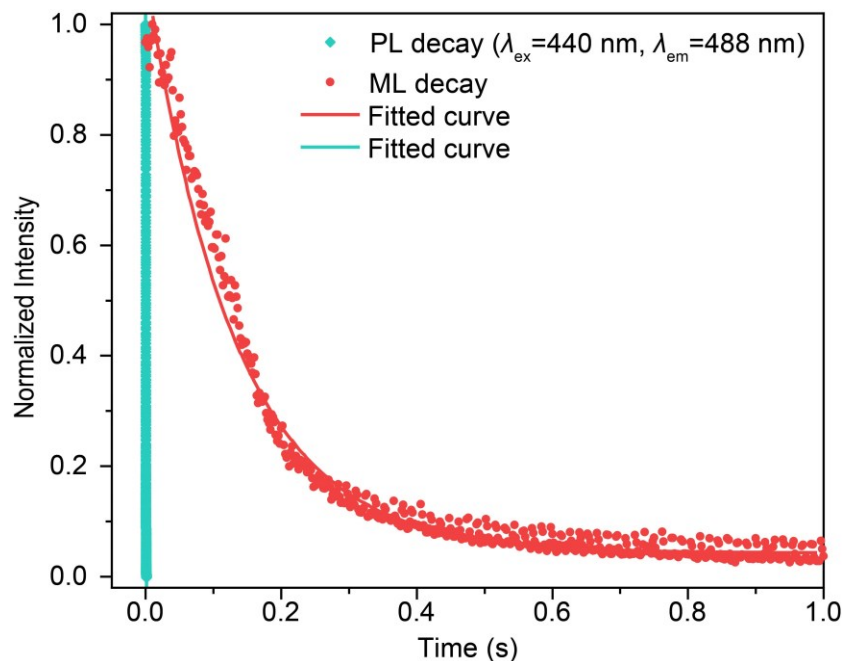
Supplementary Fig. 33 | Amplitude curves (bi-directional bias) and ferroelectric phase curve of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$. **a** Amplitude curve. The red and blue curves correspond to signal variations for the bias voltage changing from -10 to +10 V and from +10 to -10 V, respectively. **b** Standard ferroelectric phase curve.

Supplementary Table 4 | Criteria for evaluating the mechanism of mechanoluminescence (ML), summarized based on the literature.

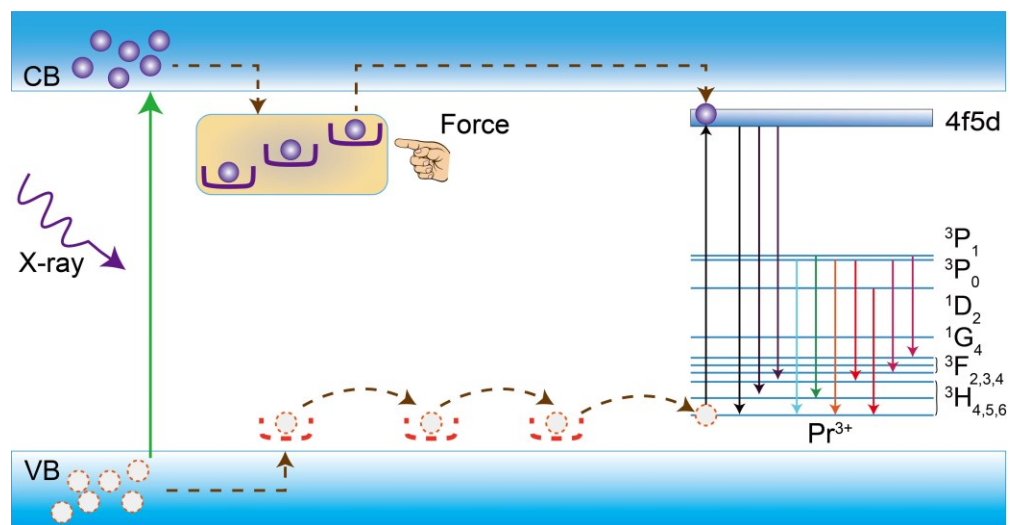
Number	Materials	PDMS	SG/SR	PU	ER	Powder	PFM	ML mechanism	References
1	Ca ₆ BaP ₄ O ₁₇ :Ce	√	○	×	×	×	○	Triboelectrification	43
2	B ₃ O ₃ -SiO ₂ - ZnO:Tb/Tm/Eu	√	√	×	×	×	×	Triboelectrification	136
3	Sr ₃ Al ₂ O ₅ Cl ₂ :Dy	√	√	×	×	×	○	Triboelectrification	36
4	Ca ₉ Bi(PO ₄) ₇ :Ce/Bi	√	√	×	×	×	○	Triboelectrification	44
5	SrGa ₁₂ O ₁₉ :Cr	√	√	×	×	×	○	Triboelectrification	83
6	CaF ₂ :Tb	√	√	×	×	×	×	Triboelectrification	22,23
7	Sr ₅ (PO ₄) ₃ Cl:Eu	√	√	×	×	×	○	Triboelectrification	35
8	Sr ₃ Al ₂ O ₅ Cl ₂ :Eu/Tb/Ce	√	√	×	×	×	○	Triboelectrification	37
9	ZnB ₂ O ₄ :Mn	√	○	○	○	×	○	Triboelectrification	140
10	CaZnGe ₂ O ₆ :Mn	√	√	√	×	×	○	Triboelectrification	98
11	BaMgAl ₁₂ O ₁₇ :Eu	√	○	○	○	○	×	Triboelectrification	68
12	Lu ₃ Al ₅ O ₁₂ :Ce	√	√	√	×	×	√	Triboelectrification	60
13	LiNbO ₃ :Pr	○	○	○	√	○	○	Piezoelectricity	116
14	Ca ₂ MgSi ₂ O ₇ :Eu	×	○	○	√	○	○	Piezoelectricity	106
15	Ga ₂ O ₃ :Cr	√	○	○	○	○	√	Piezoelectricity	135
16	MgF ₂ :Yb	√	√	√	√	√	√	Piezoelectricity	19
17	SrMg ₂ (PO ₄) ₂ :Eu	√	√	√	√	√	√	Piezoelectricity	141
18	This work	√	√	√	√	√	√	Piezoelectricity and Triboelectrification	

√ Luminous × Non-luminous ○ Lacking

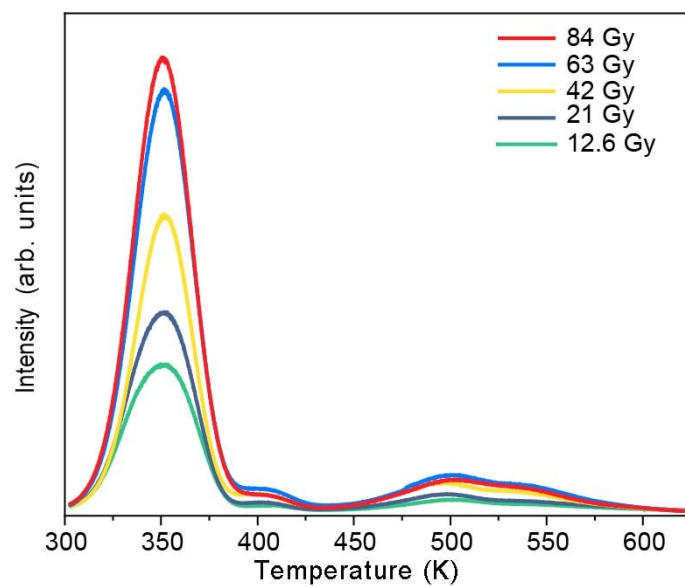
PDMS: Polydimethylsiloxane; SG: Silicon gel; SR: Silicon resin; PU: Polyurethane; ER: Epoxy resin;
PFM: Piezoresponse force microscopy.



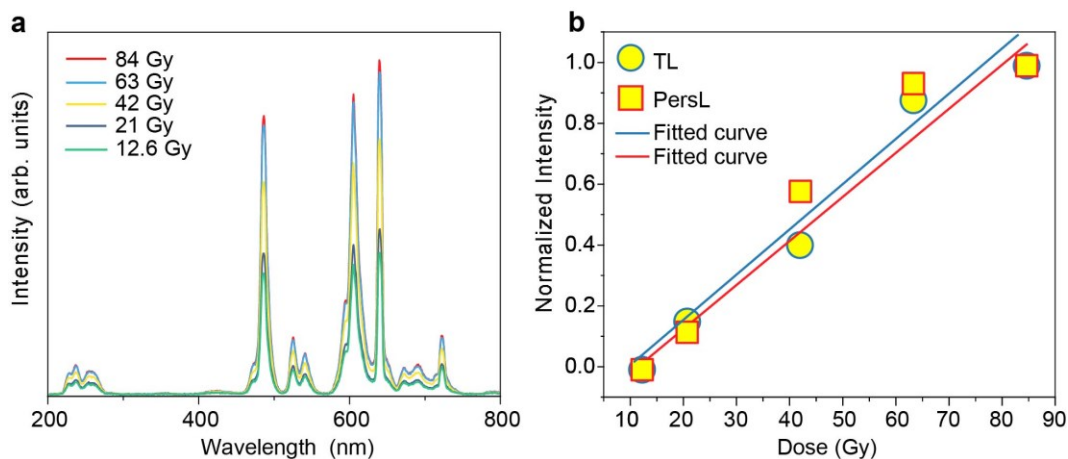
Supplementary Fig. 34 | Decay curves of photoluminescence (PL, $\lambda_{\text{ex}}=440$ nm, $\lambda_{\text{em}}=488$ nm) and trap-controlled mechanoluminescence (ML) of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$. By using the single-exponential function, the intrinsic lifetime of PL was fitted to 239 μs , a typical value of the lifetime of $4f \rightarrow 4f$ transition of Pr^{3+} . When the dynamic force was no longer applied to the sample, the trap-controlled ML continued for over 137 ms, three orders of magnitude longer than that of PL. It suggests that the processes of PL and trap-controlled ML are totally different for $\text{SrF}_2:0.5\%\text{Pr}^{3+}$. As trap-controlled ML involves the release and migration of charge carriers, it lasts significantly longer than the PL process. Therefore, such fact could also be regarded as indirect evidence that the defects indeed participated in the trap-controlled ML.



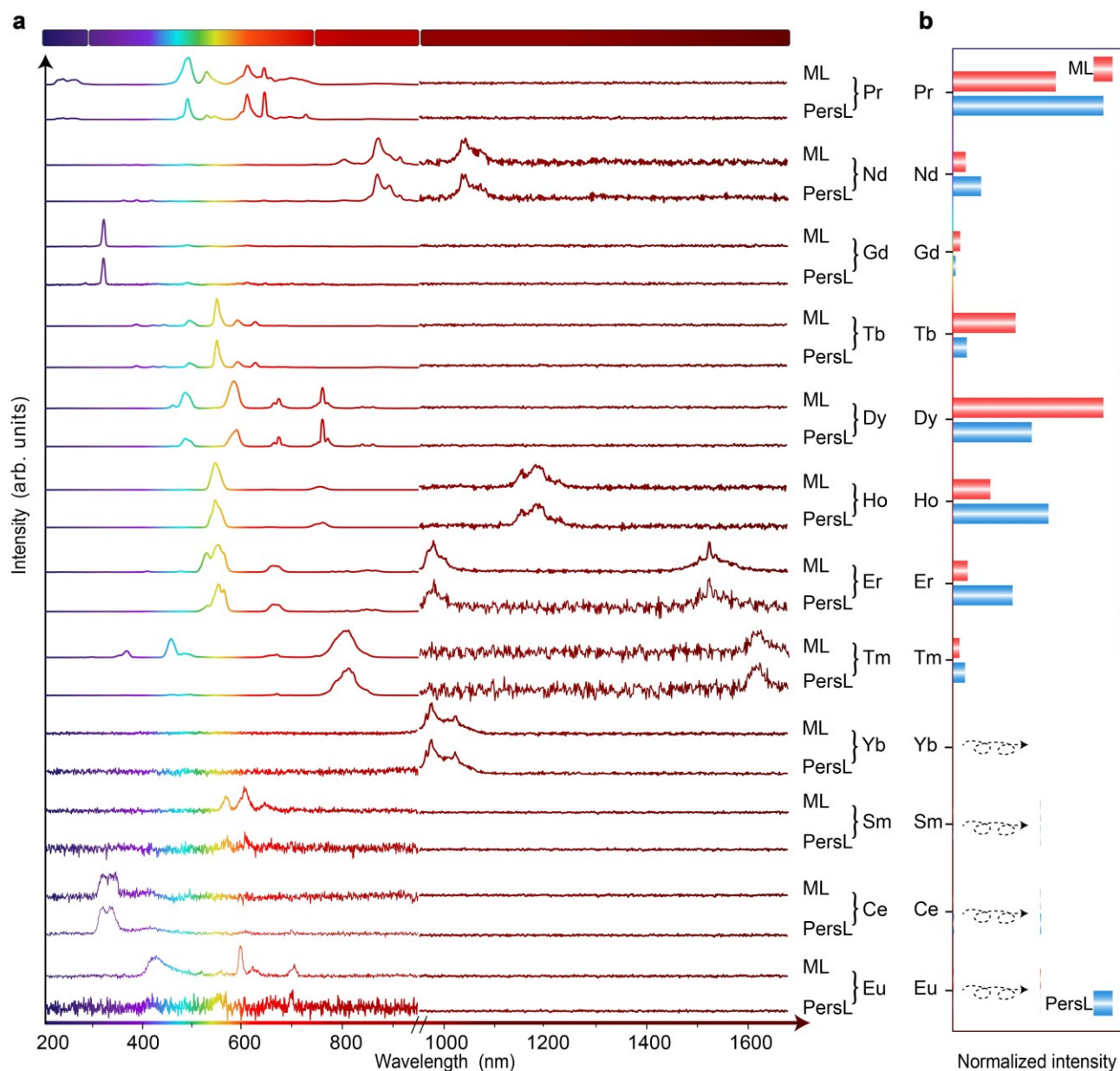
Supplementary Fig. 35 | Possible mechanism of trap-controlled mechanoluminescence behavior in Pr^{3+} -doped SrF_2 . CB: conduction band; VB: valence band.



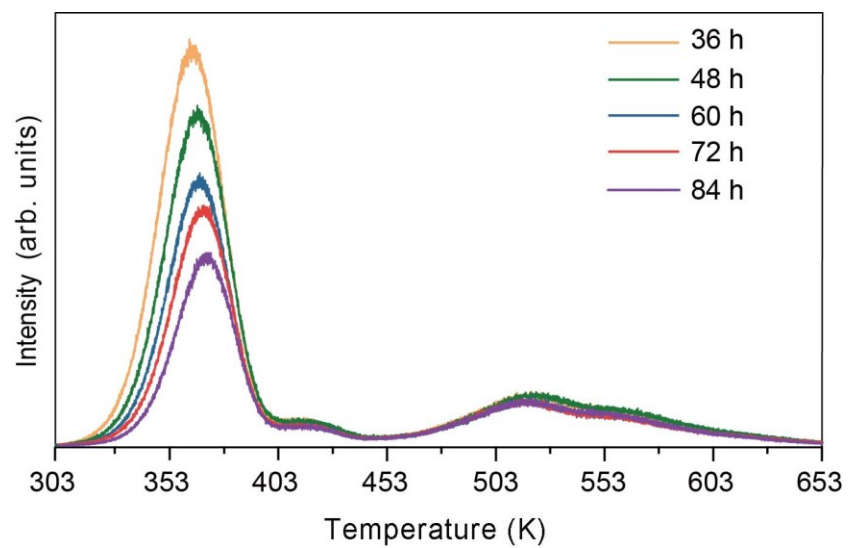
Supplementary Fig. 36 | Thermoluminescence curves of SrF₂:0.5%Pr³⁺ after being charged by X-ray of different doses (heating rate: 5 K/min).



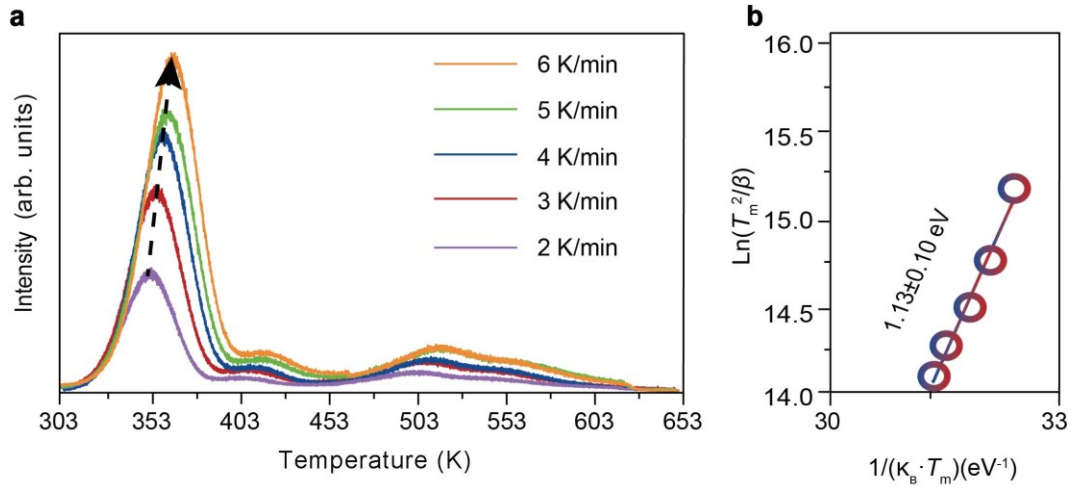
Supplementary Fig. 37 | Correlation between persistent luminescence (PersL) and thermoluminescence (TL). **a** PersL spectra of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$ after being charged by X-ray of different doses, collected at 2 min after the stoppage of X-ray. **b** Comparison of normalized PersL intensity of the spectra shown in **a** and normalized TL intensity of the curves shown in Supplementary Fig. 36.



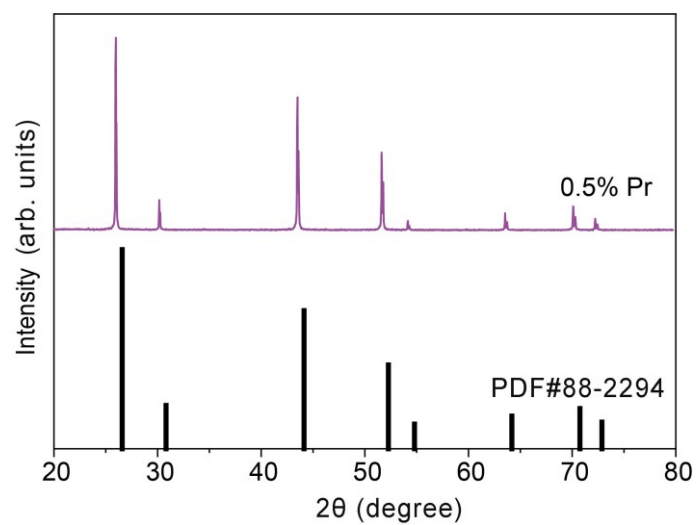
Supplementary Fig. 38 | Correlation between mechanoluminescence (ML) and persistent luminescence (PersL). **a** Comparison of ML and PersL spectra of SrF₂ doped with different lanthanides (0.5%mol) after 5 min of X-ray irradiation. **b** Comparison of ML and PersL intensities of SrF₂ doped with different lanthanides (0.5%mol).



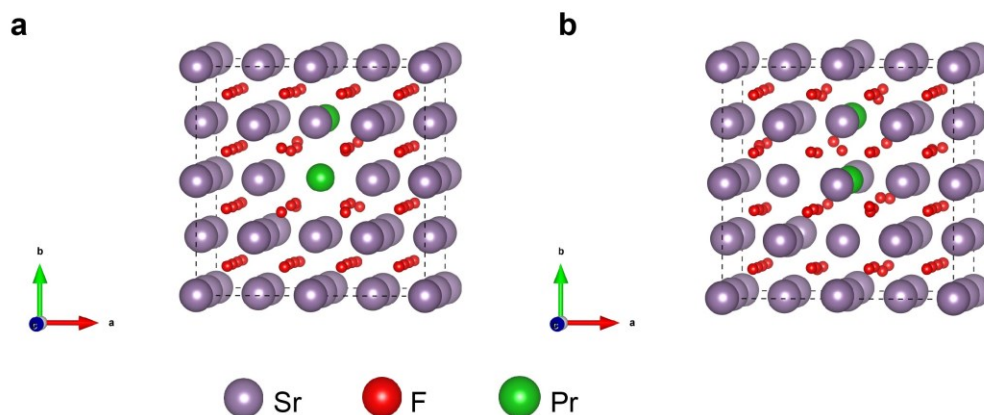
Supplementary Fig. 39 | Thermoluminescence curves of SrF₂:0.5%Pr³⁺ with various waiting times post-charging after 5 min of X-ray irradiation (heating rate: 5 K/min).



Supplementary Fig. 40 | Analysis of trap distribution. **a** Thermoluminescence curves of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$ measured after 5 min of X-ray irradiation with different heating rates. **b** Fitted trap depth derived from **a**. By using the heating rate method, we estimated the depth of the main trap to be 1.13 eV. T_m : The temperature (in K) at the maximum of the thermoluminescence glow peak. β : Heating rate (in K/s). k : Boltzmann constant.



Supplementary Fig. 41 | X-ray diffraction patterns of SrF₂:0.5%Pr³⁺ after 10 min of X-ray irradiation.



Supplementary Fig. 42 | Charge compensation mechanisms. Two specific kinds of defects in SrF_2 crystal, including Tb^{3+} replacing Sr^{2+} $[\text{Tb}_{\text{Sr}}]^+$, Sr vacancy $[\text{V}_{\text{Sr}}]^{2-}$ and F vacancy $[\text{V}_{\text{F}}]^+$, taking 6.25% Pr^{3+} doping as an example. **a** Two Tb^{3+} ions substituting for two Sr^{2+} ions, with charge neutrality maintained by the creation of one Sr vacancy, described as $2[\text{Tb}_{\text{Sr}}]^+ + [\text{V}_{\text{Sr}}]^{2-}$. **b** Two Tb^{3+} ions substituting for two Sr^{2+} ions, with charge neutrality maintained by the simultaneous formation of two F vacancies and two Sr vacancies, expressed as $2[\text{Tb}_{\text{Sr}}]^+ + 2[\text{V}_{\text{F}}]^+ + 2[\text{V}_{\text{Sr}}]^{2-}$. The atomic coordinates of the optimized computational models are provided in the Source Data file as data.

Supplementary Table 5 | Density functional theory calculated cohesive energies (E_{coh}) for the different defects in the SrF₂ crystal, including $2[\text{Pr}_{\text{Sr}}]^+ + [\text{V}_{\text{Sr}}]^{2-}$ and $2[\text{Tb}_{\text{Sr}}]^+ + 2[\text{V}_{\text{F}}]^+ + 2[\text{V}_{\text{Sr}}]^{2-}$.

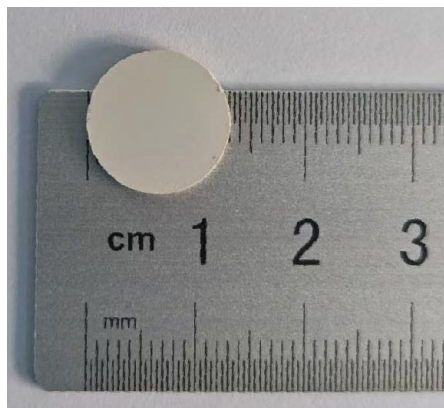
Defects	Number of atoms (N)	Supercell E_s (eV)	Free state energy (eV)			Cohesive energies E_{coh} (eV)
			E_{Sr}	E_{F}	E_{Pr}	
$2[\text{Pr}_{\text{Sr}}]^+ + [\text{V}_{\text{Sr}}]^{2-}$	95	-546.93	-0.02	-0.45	-0.19	-5.44
$2[\text{Tb}_{\text{Sr}}]^+ + 2[\text{V}_{\text{F}}]^+ + 2[\text{V}_{\text{Sr}}]^{2-}$	92	-526.90	-0.02	-0.45	-0.19	-5.41

Supplementary Note 1 | Charge compensation mechanisms.

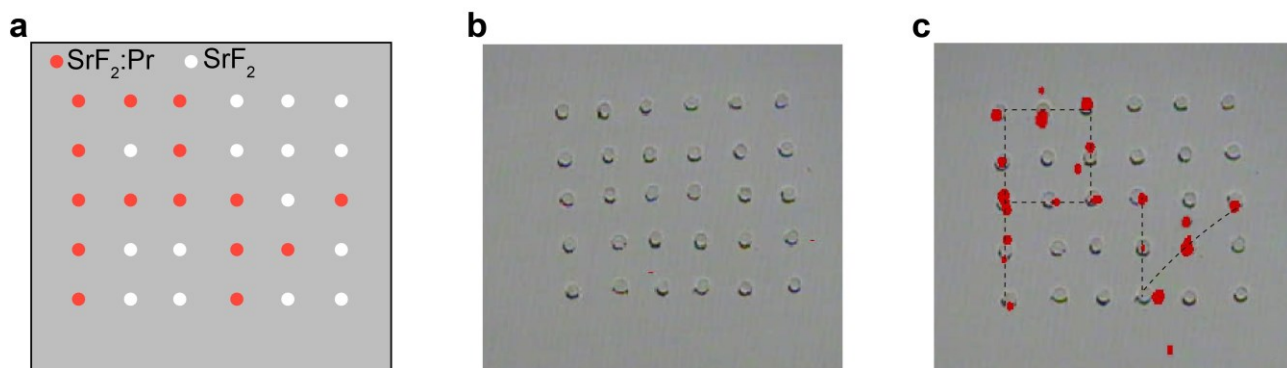
Supplementary Table 5 presents the calculated cohesive energies of the cell (E_{coh}) corresponding to different defects within the SrF₂ crystal. The definition of E_{coh} is as follows: $E_{\text{coh}} = (E_s - a \times E_{\text{Sr}} - b \times E_{\text{F}} - c \times E_{\text{Pr}})/N$, where E_s represents the total energy of the supercell, E_{Sr} , E_{F} and E_{Pr} are the free state energies of the Sr, F and Pr atoms, respectively; a, b and c denote the numbers of the corresponding atomic species in the supercell; and N is the total number of atoms in the supercell. The results indicate that the atomic E_{coh} (-5.44 eV) for the $2[\text{Pr}_{\text{Sr}}]^+ + [\text{V}_{\text{Sr}}]^{2-}$ defect cluster is the smallest. It means that two Pr^{3+} dopants would replace two Sr^{2+} cations in the SrF₂ crystal, resulting in one Sr^{2+} vacancy to keep the charge balance.



Supplementary Fig. 43 | A quick response (QR) code made of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$ for inspection at customs. Upon exposure to X-ray irradiation, the QR code is charged. After the release of room temperature persistent luminescence, force stimulation can be applied to the QR code to achieve mechanoluminescence for secondary inspection.



Supplementary Fig. 44 | Photograph of the SrF₂:0.5%Pr³⁺ disk with a diameter of 13 mm.



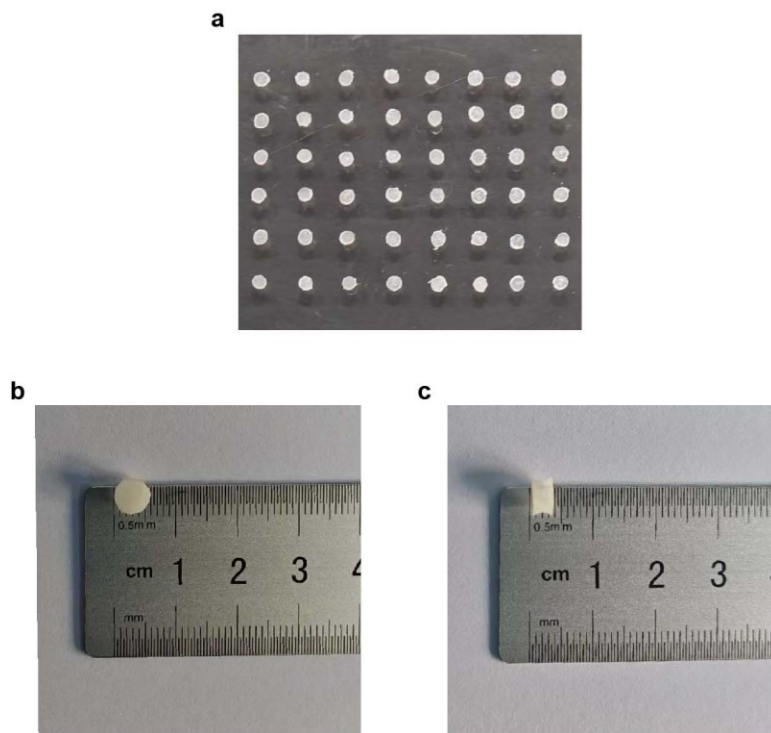
Supplementary Fig. 45 | Information storage and reading depending on UVC luminescence. **a** Schematic diagram of 5*6 array. The white and red dots represent SrF_2 and $\text{SrF}_2:\text{Pr}^{3+}$, respectively. **b** Photograph of the array before X-ray irradiation. **c** Photograph of the array after being charged by X-ray for 5 min. The hidden patterns 'P' and 'r' can be captured by a solar blind camera (dotted line).



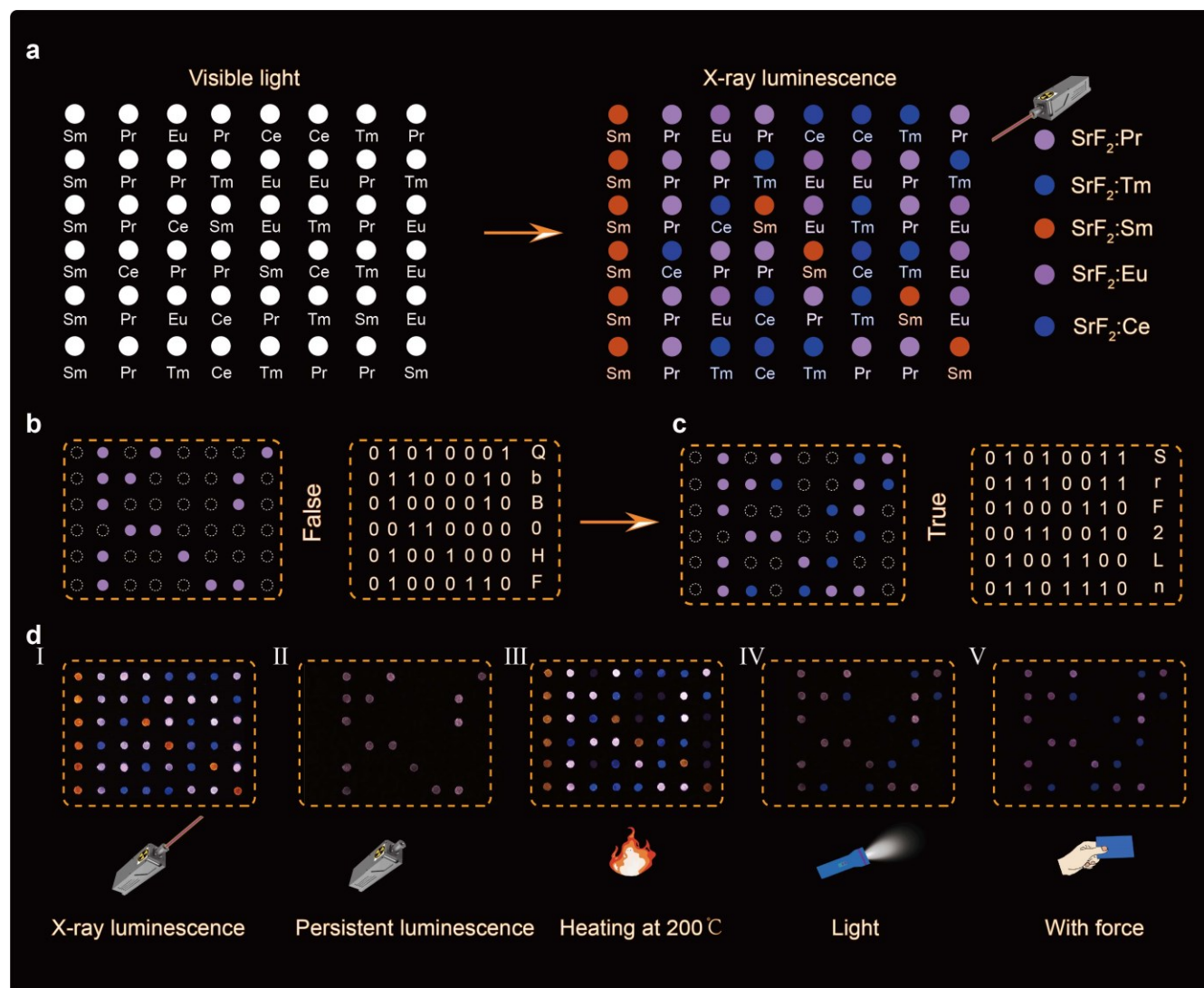
Supplementary Fig. 46 | Photograph of $\text{ZnS}:\text{Cu}^{2+}$ @polydimethylsiloxane in dark environment upon stretching, taken from a distance of 6 m.



Supplementary Fig. 47 | Demonstration of outdoor emergency rescue by using UVC photon. A person is trapped in an outdoor hazardous situation where visibility is limited. In this case, it is difficult to identify the trapped person. At this moment, the stretchable elastic film made of $\text{SrF}_2:\text{Pr}^{3+}$ @polydimethylsiloxane might work. The trapped person pulls the $\text{SrF}_2:\text{Pr}^{3+}$ @polydimethylsiloxane film and the UVC mechanoluminescence can be located using a solar blind camera by rescue personnel. Such strategy, together with other rescue technologies, is expected to effectively enhance the efficiency of rescue operations.



Supplementary Fig. 48 | $\text{SrF}_2\text{:Ln}^{2+,3+}\text{@polydimethylsiloxane}$ array. **a** Photograph of $\text{SrF}_2\text{:Ln}^{2+,3+}\text{@polydimethylsiloxane}$ arranged in an 8*6 array. **b** Diameter demonstration, and **c** Thickness demonstration of a single $\text{SrF}_2\text{:Ln}^{2+,3+}\text{@polydimethylsiloxane}$ cylinder.



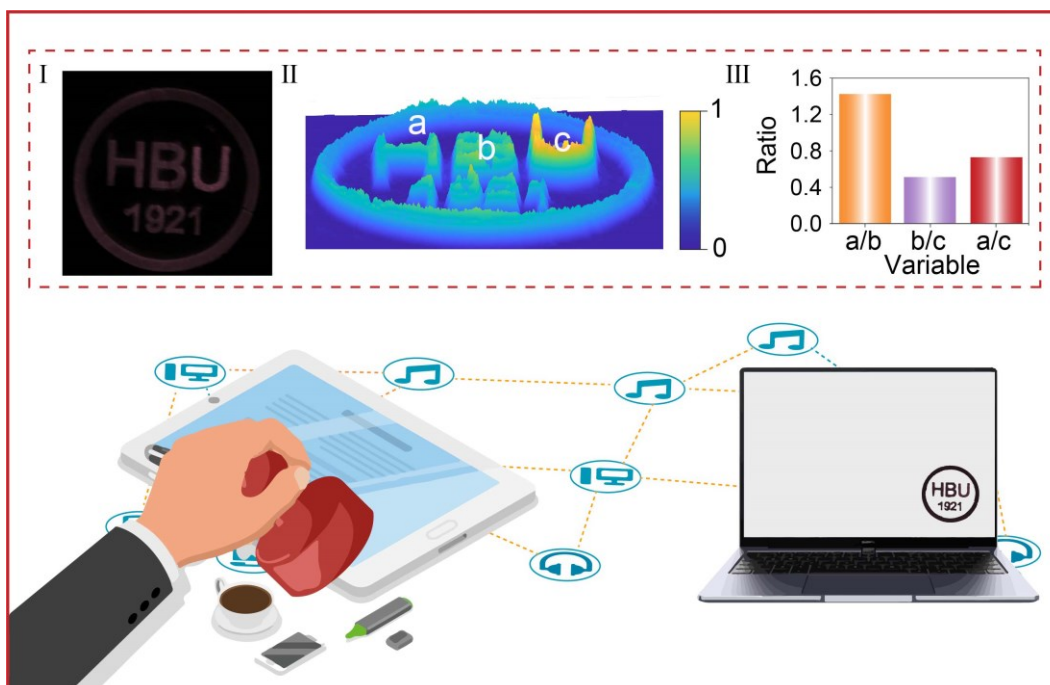
Supplementary Fig. 49 | Information storage and encryption based on $\text{SrF}_2\text{:Ln}^{2+,3+}$. **a** Schematic of the mechanoluminescent array using $\text{SrF}_2\text{:Ln}^{2+,3+}$ @polydimethylsiloxane. **b** Luminescence array with false information. **c** Luminescence array with correct information. **d** Multimodal luminescence array: (I) X-ray on; (II) X-ray off; (III) Heat treatment; (IV) Optical stimulation; (V) Force stimulation.

Supplementary Note 2 | Applications based on visible light.

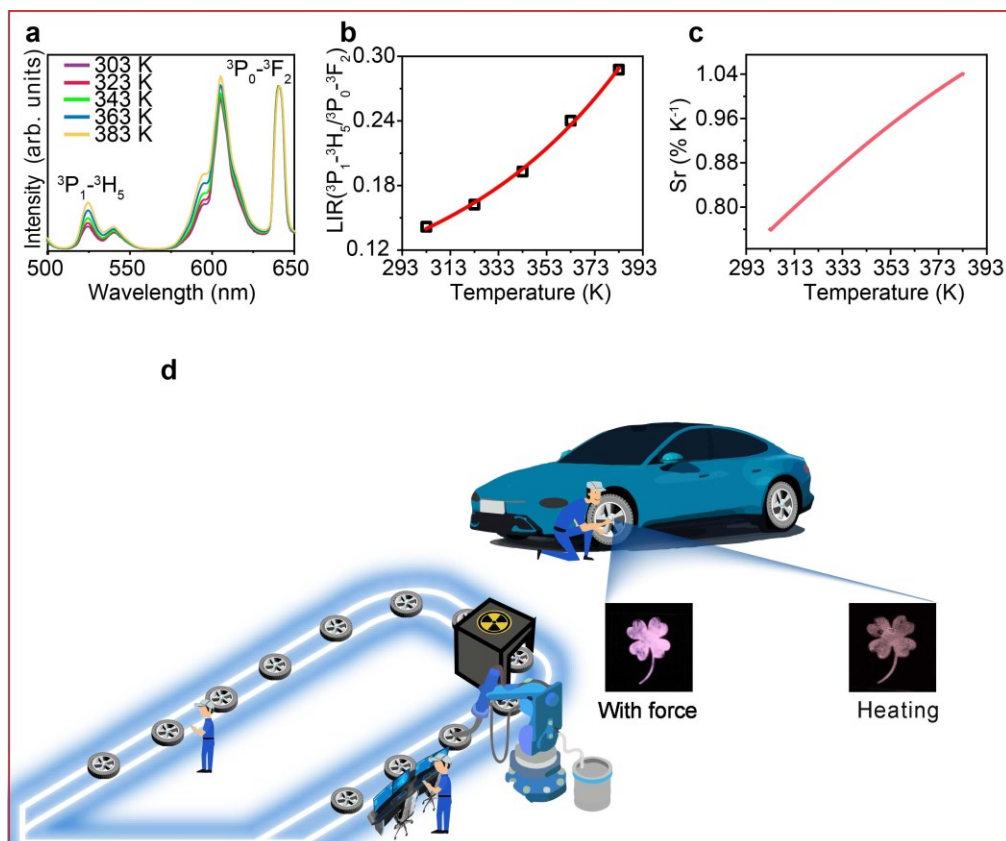
We introduced a novel binary encoding method by using $\text{SrF}_2:\text{Ln}^{2+,3+}$. By incorporating $\text{SrF}_2:\text{Ln}^{2+,3+}$ ($\text{Ln} = \text{Pr}^{3+}, \text{Tm}^{3+}, \text{Sm}^{3+}, \text{Eu}^{2+,3+}$ and Ce^{3+}) into polydimethylsiloxane (PDMS), we designed an 8*6 cylindrical array (Supplementary Figs. 48 and 49a). Under daylight, these cylinders all appeared the same. After exposure to X-ray, the cylindrical array became colorful due to the characteristic emissions of dopant luminescent centers. The correct information was hidden by different combinations of luminescence (Supplementary Figs. 49b and 49c). When the array was subjected to X-ray irradiation or heated after exposure to X-ray irradiation, all samples emitted light without conveying any useful message (Supplementary Figs. 49d-I and 49d-III). After switching off the X-ray source for 5 min, the persistent luminescence (PersL) of $\text{SrF}_2:\text{Pr}^{3+}$ was far stronger than that of others, resulting in the false information ‘QbB0HF’ (Supplementary Fig. 49d-II). Under the stimulation of photon or force, the luminescence of $\text{SrF}_2:\text{Pr}^{3+}$ and $\text{SrF}_2:\text{Tm}^{3+}$ provided the correct information ‘SrF2Ln’ (Supplementary Figs. 49d-IV and 49d-V). These results reveal that our binary coding based on $\text{SrF}_2:\text{Ln}^{2+,3+}$ has great potential in advanced information encryption and anti-counterfeiting technology.

Electronic signature (*e*-signature) has been widely used nowadays. However, it is easy to imitate the profile of an *e*-signature. Therefore, adding more information on *e*-signature is essential. $\text{SrF}_2:\text{Ln}^{2+,3+}$ can play a useful role in improving the security of *e*-signature. We designed a unique ‘HBU’ stamp and pressed it onto the $\text{SrF}_2:\text{Pr}^{3+}$ @PDMS film (Supplementary Fig. 50), leaving there a ‘HBU’ pattern with different luminescence distribution. We fetched the mechanoluminescence (ML) information from the points of a, b and c and then calculated the intensity ratio of a/b, a/c and b/c. By defining a unit as a threshold, the obtained intensity ratios were found to be [100]. In this way, we added information to *e*-signature.

We also expanded the application potential of $\text{SrF}_2:\text{Pr}^{3+}$ for non-destructive examination. In the beginning, $\text{SrF}_2:\text{Pr}^{3+}$ was coated on the surface of automotive tires. During non-destructive inspection by X-ray, the coated $\text{SrF}_2:\text{Pr}^{3+}$ was pre-charged. Thereafter, by means of PersL or ML from the $\text{SrF}_2:\text{Pr}^{3+}$, the worker and consumer can confirm whether or not the tire had passed the inspection before leaving the factory and after buying the tire, respectively (Supplementary Fig. 51). To ensure high quality, tires usually undergo friction test in factory. Since friction generates heat, the PersL intensity ratio of $^3\text{P}_1 \rightarrow ^3\text{H}_5$ to $^3\text{P}_0 \rightarrow ^3\text{F}_2$ transitions of $\text{SrF}_2:\text{Pr}^{3+}$ can also be used to monitor the temperature during friction testing.



Supplementary Fig. 50 | Electronic seal. (I) Visualization of 'HBU' stamp through UVC mechanoluminescence (ML). (II) Distribution of ML intensity of 'HBU' stamp extracted from (I). (III) ML intensity ratios between different points at a, b and c.



Supplementary Fig. 51 | Nondestructive testing application. **a** Emission spectra of $\text{SrF}_2:0.5\%\text{Pr}^{3+}$ upon different thermal stimulations after 5 min of X-ray irradiation. **b** Luminescence intensity ratio (LIR) between the $^3\text{P}_1 \rightarrow ^3\text{H}_5$ and $^3\text{P}_0 \rightarrow ^3\text{F}_2$ transition of Pr^{3+} , and **c** Relative sensitivity (S_r) of designed ratiometric thermometry as a function of temperature. **d** Schematic diagram of non-destructive testing by using $\text{SrF}_2:0.5\%\text{Pr}^{3+}$.

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