

Supporting Information

Strong sensitivity enhancement in lifetime-based luminescence thermometry by co-doping of SrTiO₃:Mn⁴⁺ nanocrystals with trivalent lanthanide ions

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The average lifetime of the excited states were calculated with the equation Eq. S1:

$$\tau_{avr} = \frac{A_1\tau_1^2 + A_2\tau_2^2}{A_1\tau_1 + A_2\tau_2} \quad (\text{Eq. S1})$$

where: τ_1 , τ_2 – the average time, which is in accordance with the relation $\tau = t \cdot \ln(2)$ and A_1, A_2 – amplitude, which are the parameters of the double exponential function:

$$y = y_0 + A_1 \cdot \exp\left(-\frac{x}{t_1}\right) + A_2 \cdot \exp\left(-\frac{x}{t_2}\right) \quad (\text{Eq. S2})$$

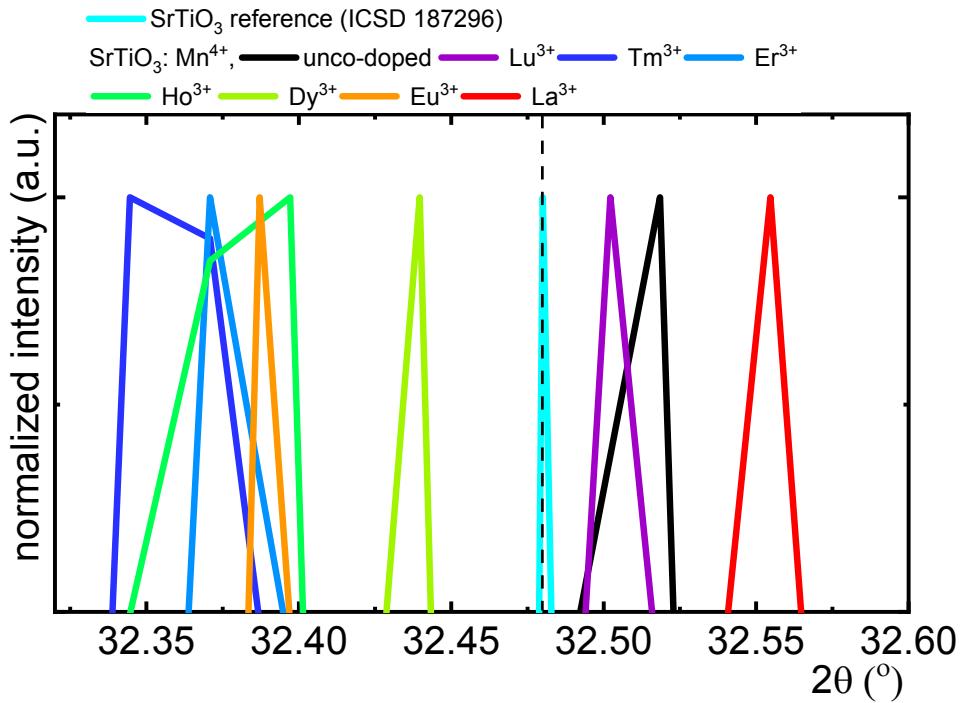


Figure S1. The zoom of the maximum of X-ray diffraction patterns of $\text{SrTiO}_3:\text{Mn}^{4+}$ with different Ln^{3+} dopants.

Table S1. Shannon effective ionic radii of SrTiO_3 lattice and dopant ions.

Site	Coordination number (CN)	Shannon effective ionic radii (EIR) (pm)							
		Ti^{4+}		Mn^{4+}				Ti^{3+}	
octahedral	VI (6-fold)	60.5		53				67	
		Sr^{2+}	Lu^{3+}	Tm^{3+}	Er^{3+}	Ho^{3+}	Dy^{3+}	Eu^{3+}	La^{3+}
preferred	VIII/X (8/10-fold)	97.7	99.4	100.4	101.5	102.7	106.6	127	
cuboctahedral	XII (12-fold)	144	120.4 °	122.4 °	123.4 °	123.4 °	125.5 °	129.5 °	136

e – extrapolated linearly (in the all cases $R^2 > 0.9987$)

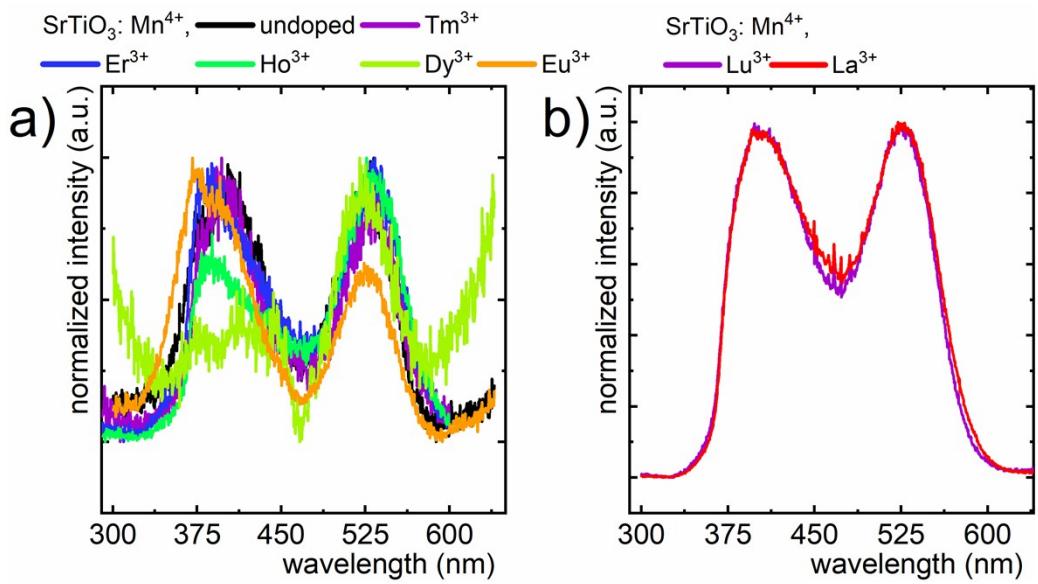


Figure S2. Excitation spectra of $\text{SrTiO}_3:\text{Mn}^{4+}, \text{Ln}^{3+}$ with different optically active-a) and passive -b) Ln^{3+} ions measured at 123 K for emission of Mn^{4+} ($\lambda_{\text{em}} = 725$ nm).

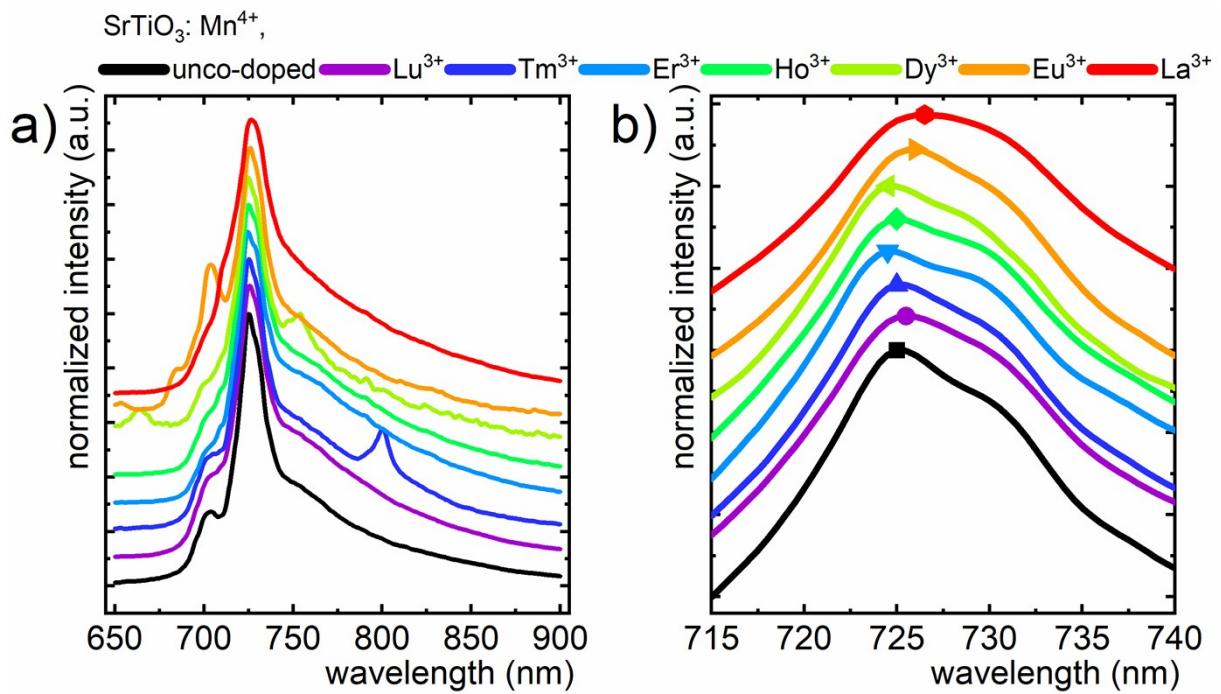


Figure S3. Emission spectra of $\text{SrTiO}_3:\text{Mn}^{4+}, \text{Ln}^{3+}$ with different Ln^{3+} ions measured with $\lambda_{\text{exc}} = 400$ nm at 123 K – a) with the zoom of the same spectra in the 715-740 nm spectral range – b).

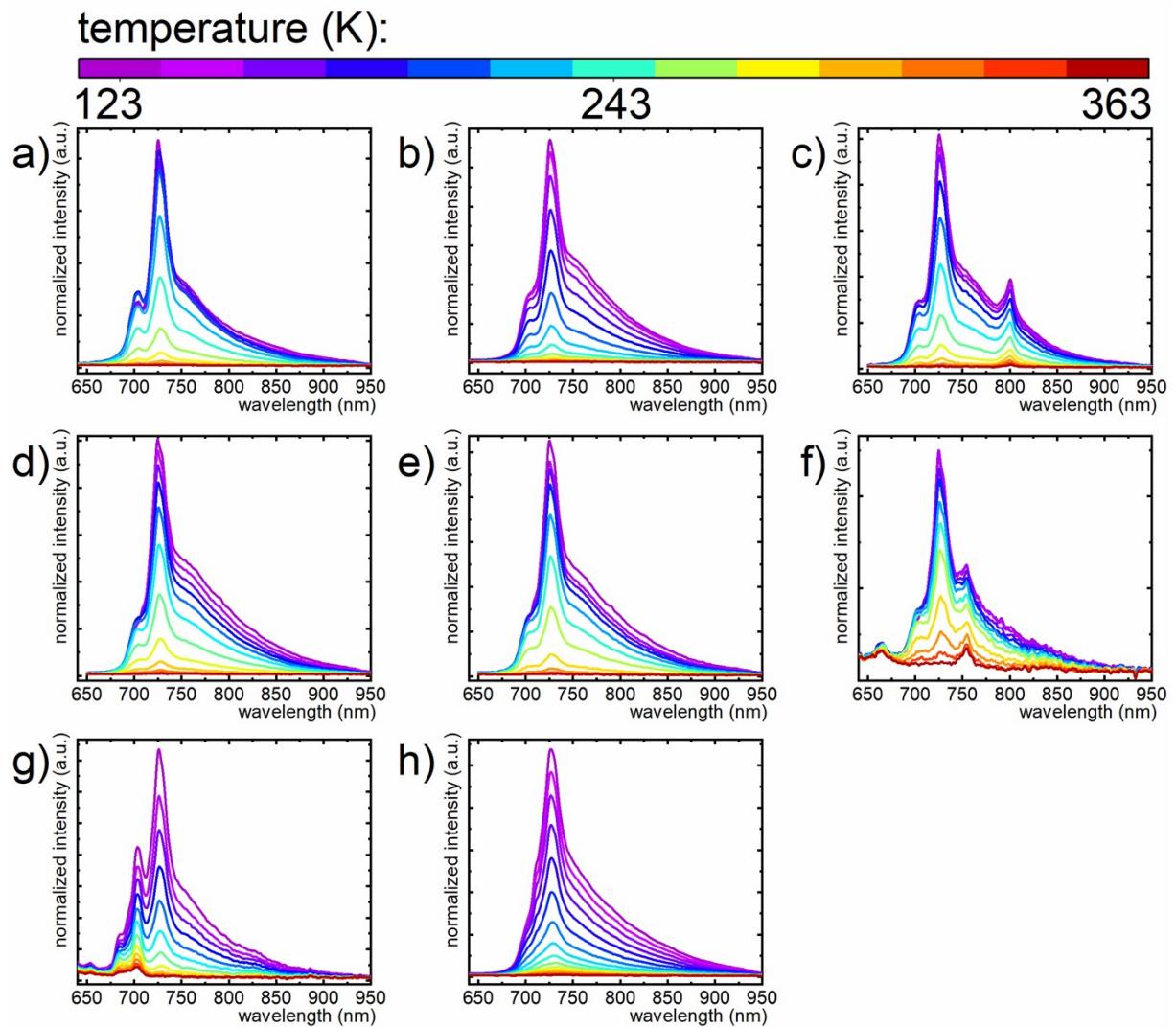


Figure S4. Thermal evolution of emission spectra excited by $\lambda_{\text{exc}} = 400$ nm for the SrTiO₃:0.1% Mn⁴⁺ – a) and SrTiO₃:0.1% Mn⁴⁺, Ln³⁺, where Ln³⁺ = Lu³⁺ – b), Tm³⁺ – c), Er³⁺ – d), Ho³⁺ – e), Dy³⁺ – f), Eu³⁺ – g), La³⁺ – h).

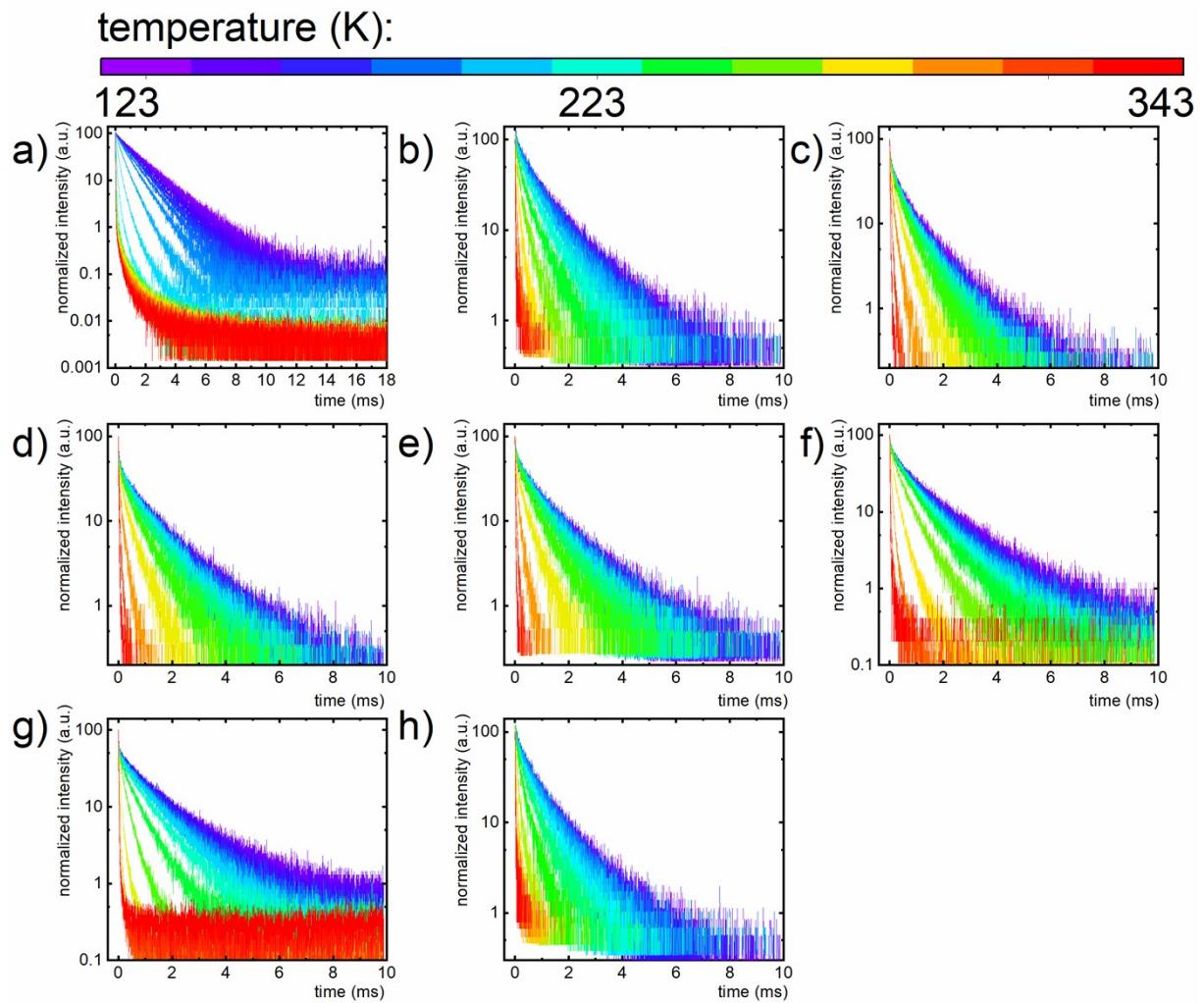


Figure S5. Thermal evolution of luminescent decays of ^2E excited state of Mn^{4+} for the $\text{SrTiO}_3:0.1\% \text{Mn}^{4+}$ – a) and $\text{SrTiO}_3:0.1\% \text{Mn}^{4+}$, Ln^{3+} , where $\text{Ln}^{3+} = \text{Lu}^{3+}$ – b), Tm^{3+} – c), Er^{3+} – d), Ho^{3+} – e), Dy^{3+} – f), Eu^{3+} – g), La^{3+} – h).

Table S2. The comparison of energy distances of Ln^{3+} excited states from ^2E of Mn^{4+} ions and thermometric parameters.

co-dopant ion	Eu^{3+}	Dy^{3+}	Er^{3+}	Ho^{3+}	Lu^{3+}	La^{3+}	Tm^{3+}
energy distance of excited state above ^2E (cm^{-1})	3436	7228	1591	1658	-	-	689
energy distance of excited state below ^2E (cm^{-1})	8144	560	11 07	299	-	-	940
mean energy distance of excited state from ^2E (cm^{-1})	5790	3894	1349	978.5	--	--	814.5
τ_{avr} at 123 K (ms)	0.90	0.86	0.78	0.71	0.71	0.61	0.54
local minimum value of S_A	2.5331	2.3892	1.9774	1.6879	2.066	~0.8894	~1.373

	$(\mu\text{s K}^{-1})$							
temperature of $S_A \text{ min (K)}$	196.9	199.8	182.4	164.9	173.15	~131.4	~190.05	
maximum value of $S_A (\mu\text{s K}^{-1})$	10.959	10.003	10.155	9.4355	7.5298	6.3693	7.1677	
temperature of $S_A \text{ max (K)}$	263.5	265.2	254.7	251.9	249.15	248.9	261.9	

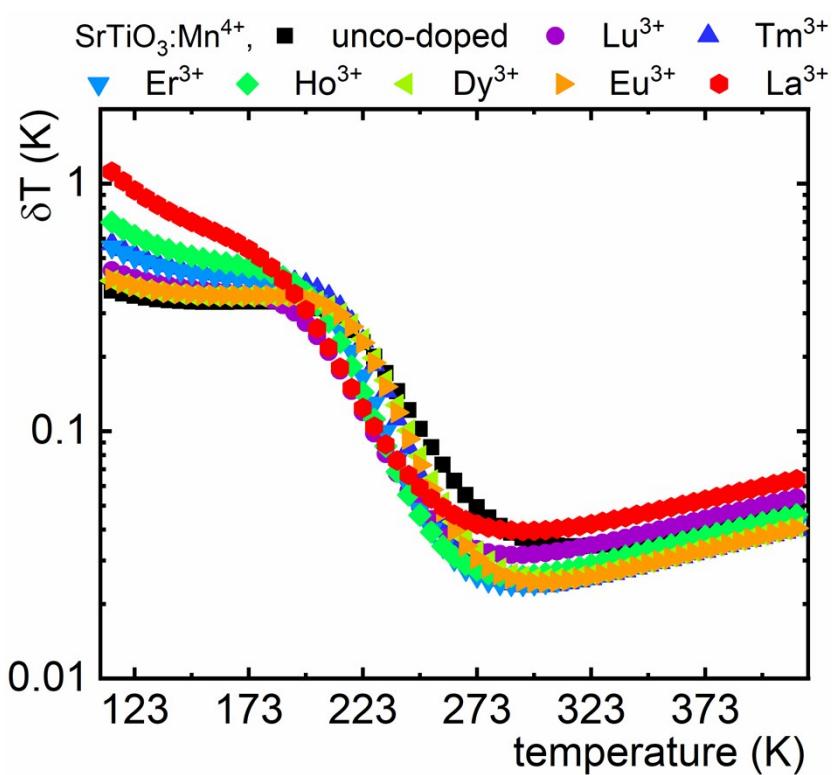


Figure S6. Thermal dependence of temperature estimation uncertainty for different Ln^{3+} in $\text{SrTiO}_3:\text{Mn}^{4+}, \text{Ln}^{3+}$ samples.

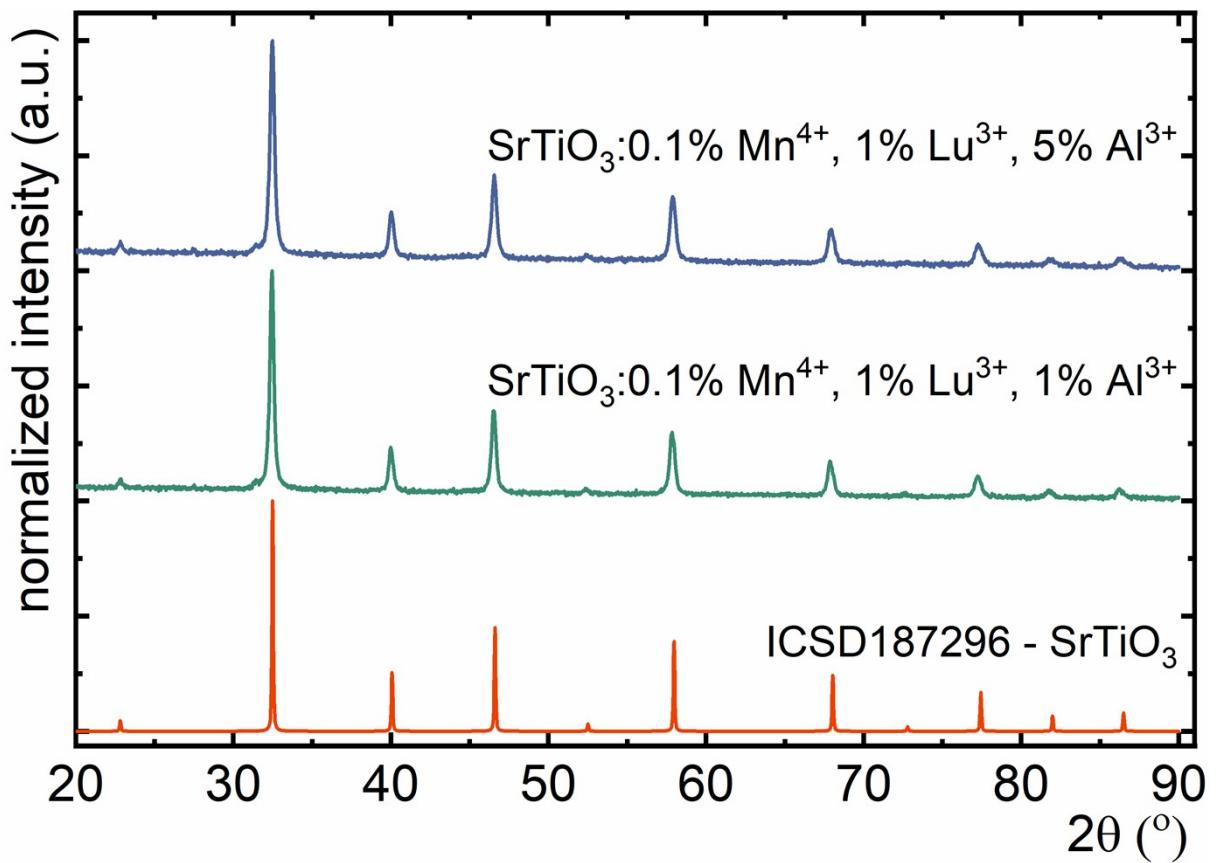


Figure S7. X-ray diffraction patterns of $\text{SrTiO}_3:\text{Mn}^{4+}$, Lu^{3+} co-doped with 1% and 5% of Al^{3+} ions.

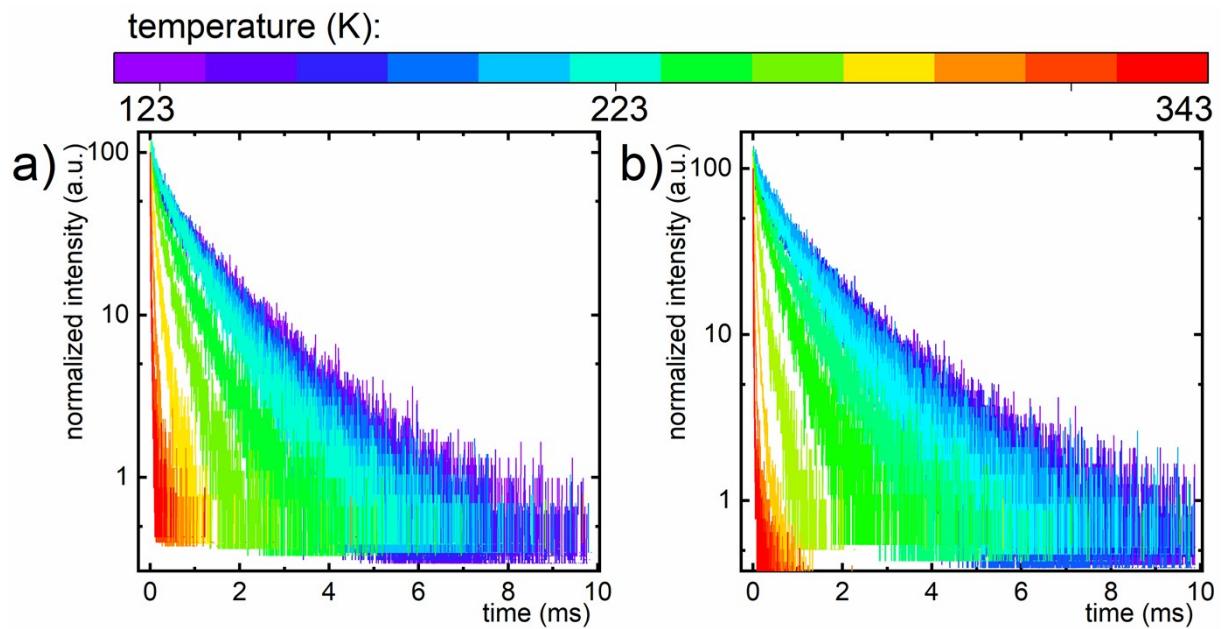


Figure S8. Thermal evolution of luminescent decays of ^2E excited state of Mn^{4+} for the $\text{SrTiO}_3:0.1\% \text{Mn}^{4+}, 1\% \text{Lu}^{3+}, 1\% \text{Al}^{3+}$ - a and $\text{SrTiO}_3:0.1\% \text{Mn}^{4+}, 1\% \text{Lu}^{3+}, 5\% \text{Al}^{3+}$ - b.