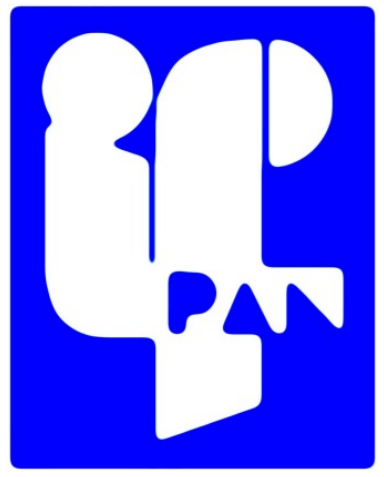


Luminescence and energy transfer processes in LuNbO₄:Bi,Eu

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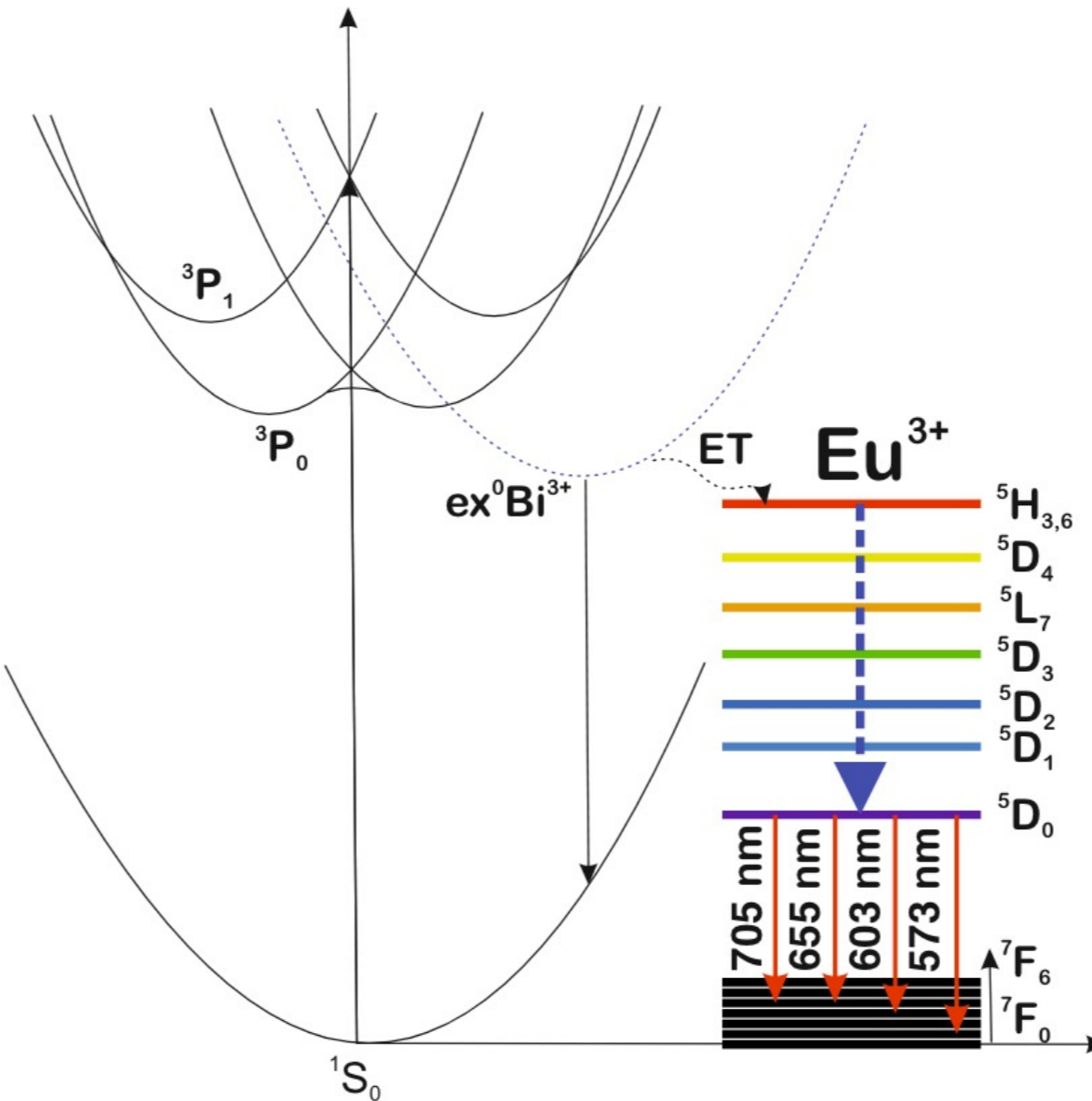
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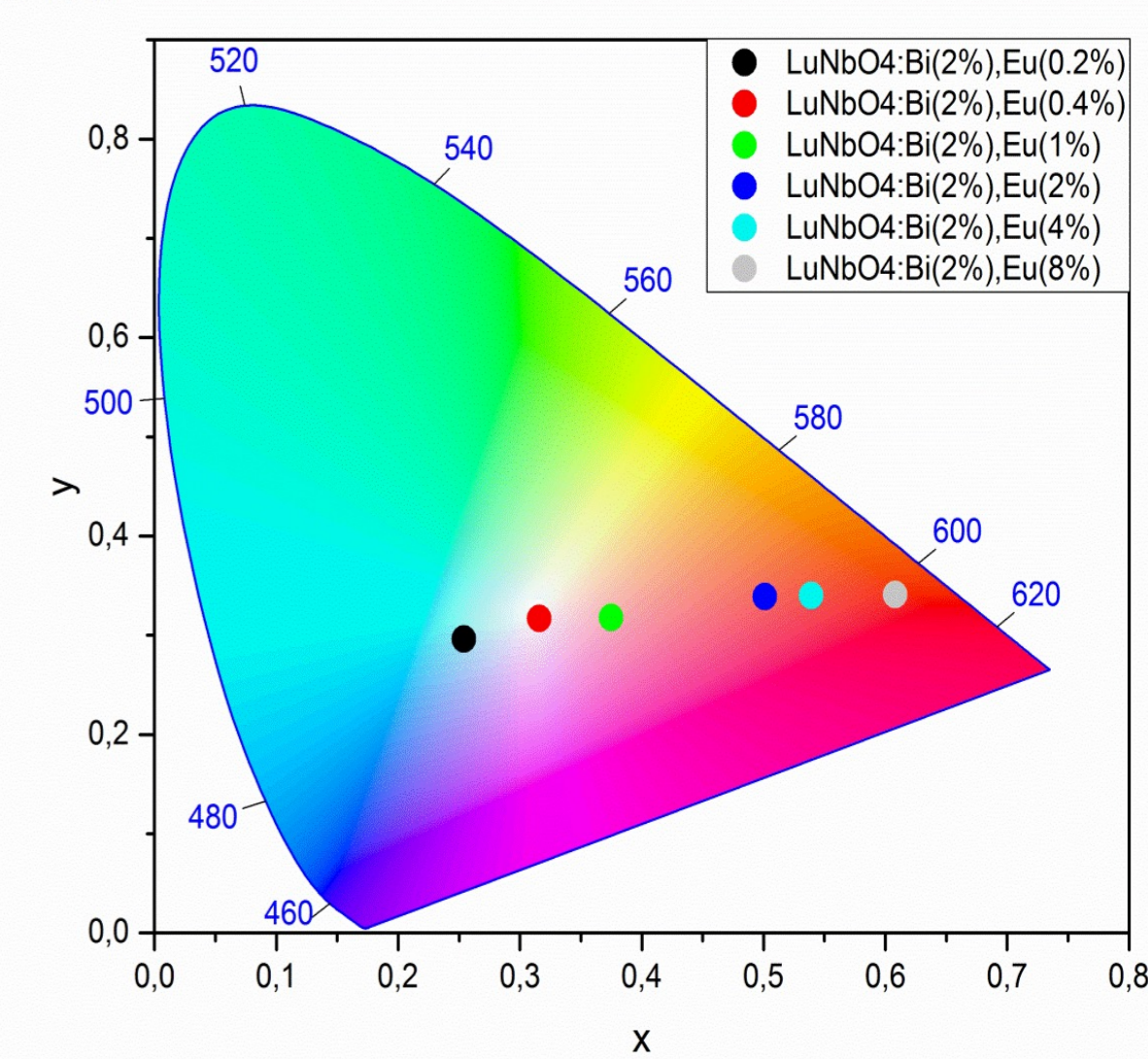
Motivation and aim of the work

The interest in Bi³⁺-doped compounds increased drastically in recent years due to their possible applications as scintillator and phosphor materials suitable for X-ray screens, white light-emitting diodes, solar cells, dosimeters, etc. The compounds co-doped with Bi³⁺ and different trivalent rare-earth (RE³⁺) ions were found to be potentially applicable as spectral converters for solar cells and solid-state light sources of new generation, so-called white light-emitting diodes (WLED), owing to strong absorption in the ultraviolet spectral region, intense broad visible Bi³⁺-related emission bands, and effective Bi³⁺ → RE³⁺ energy transfer, resulting in the appearance of the luminescence covering a wide spectral range from blue to red.

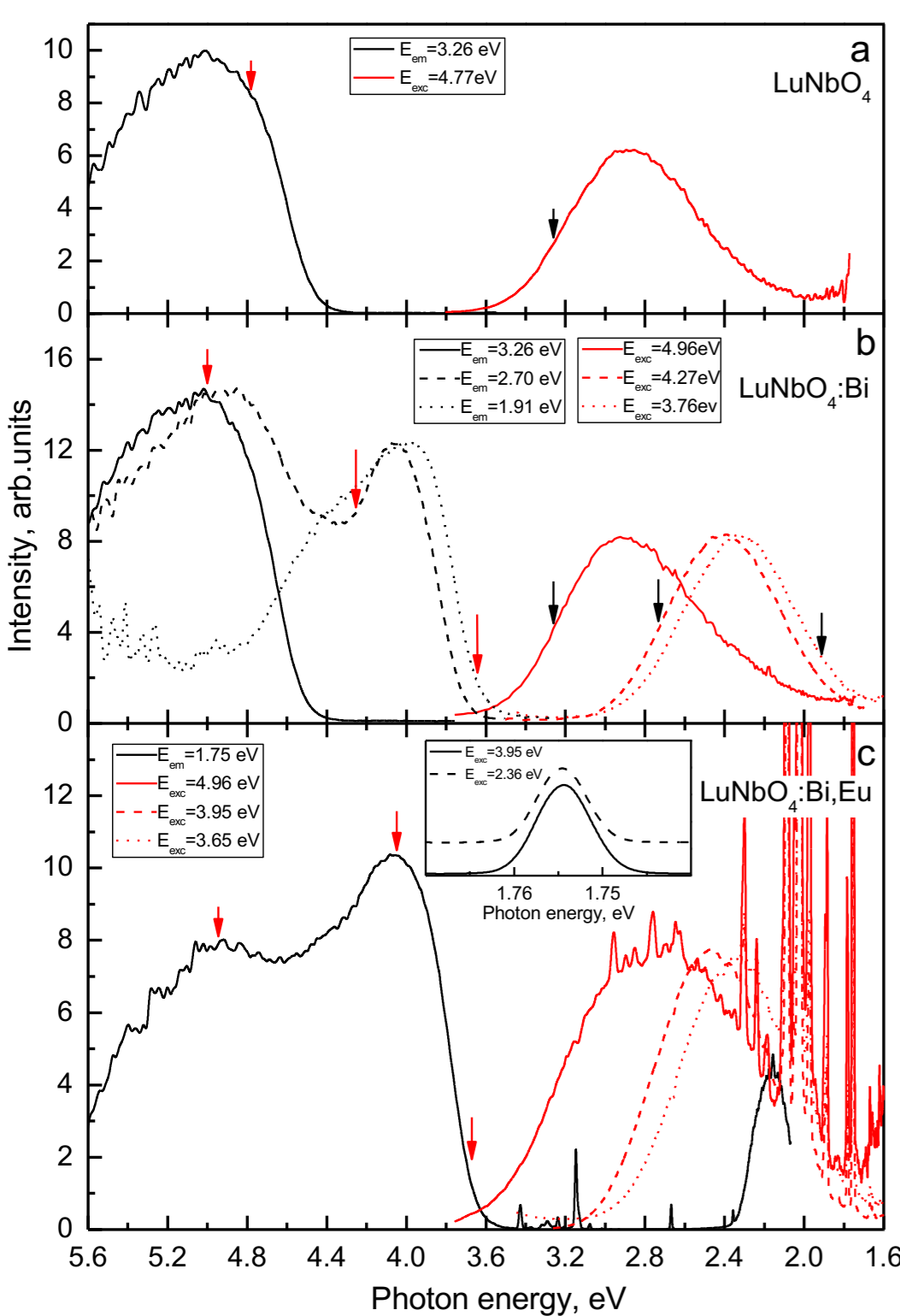
Recently, a detailed and systematic investigation of the Bi³⁺-related luminescence in YNbO₄:Bi, LuNbO₄:Bi, and GdNbO₄:Bi microcrystalline powders with different Bi³⁺ contents was carried out [1]. Two broad visible Bi³⁺-related emission bands were observed in the emission spectrum of the Bi³⁺-doped niobates. The analysis of temperature dependences of their decay time and the values of the corresponding relaxed excited states (RES) parameters allowed us to conclude their exciton-like origin. The absence of the ultraviolet emission, arising from the radiative decay of the triplet RES of a Bi³⁺ ion, and the exciton-like origin of the visible Bi³⁺-related emissions indicate that the triplet RES of both the Bi³⁺ ion and the {Bi³⁺-Bi³⁺} dimer are located inside the conduction band of the investigated niobates.



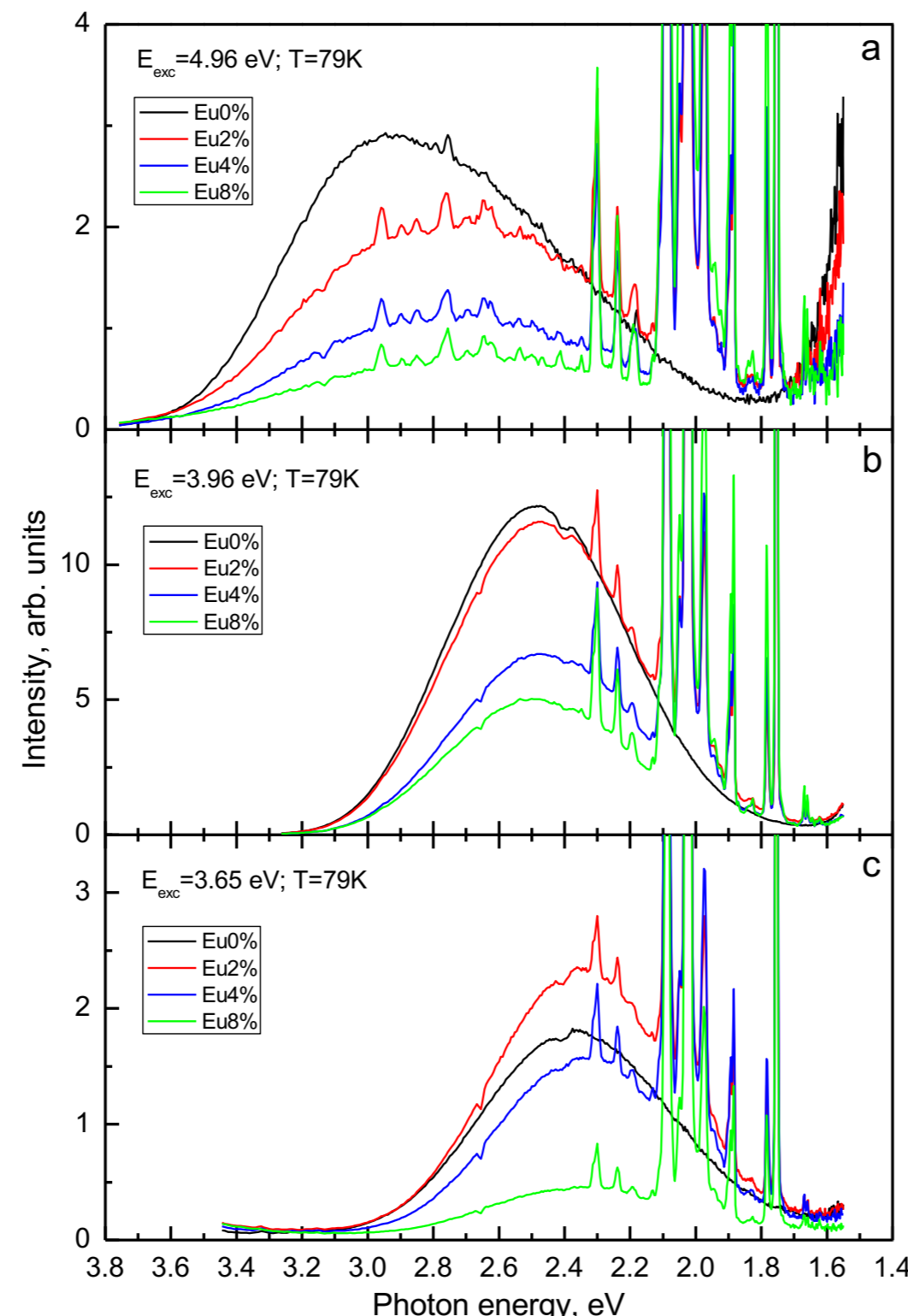
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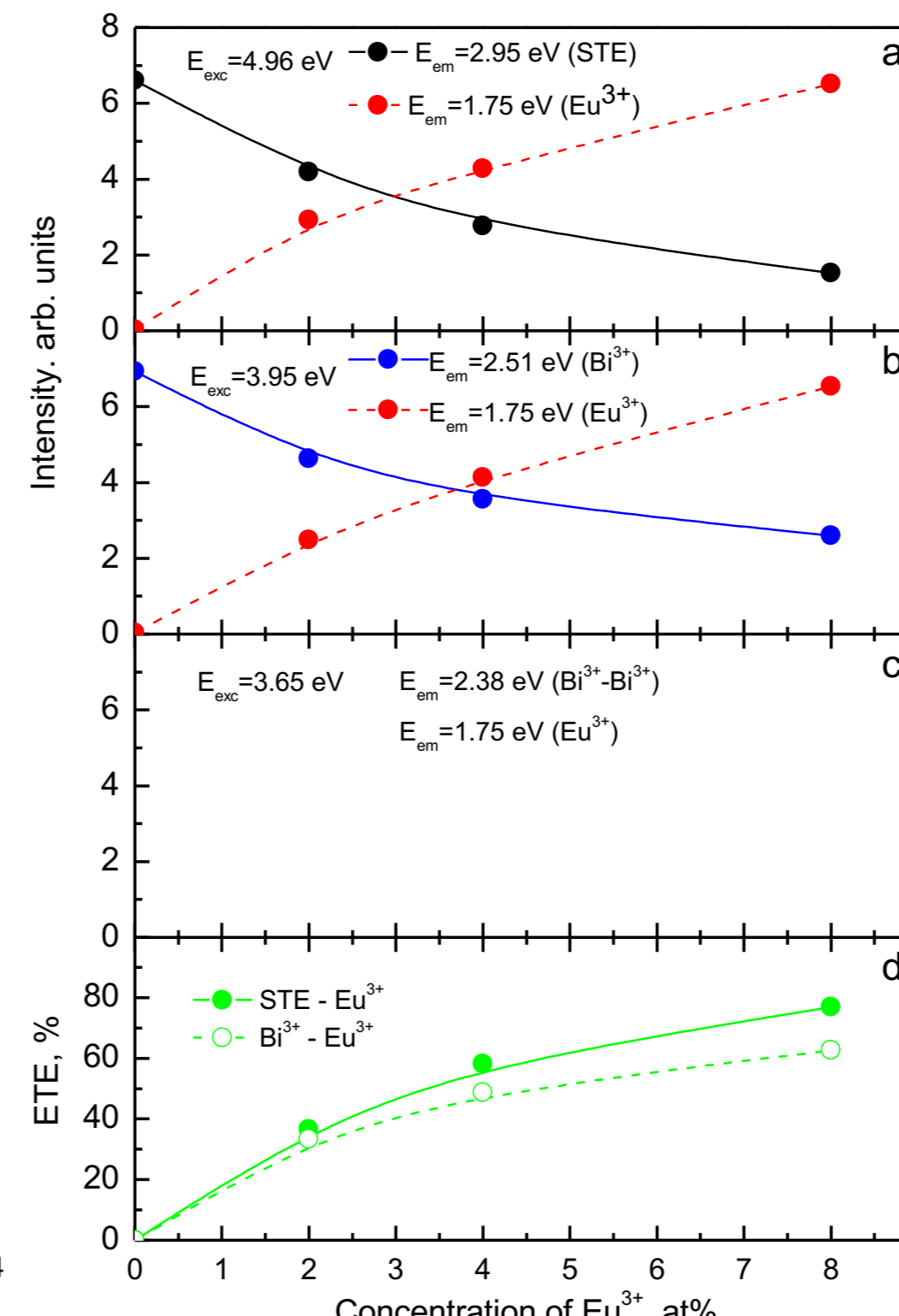
Experimental results



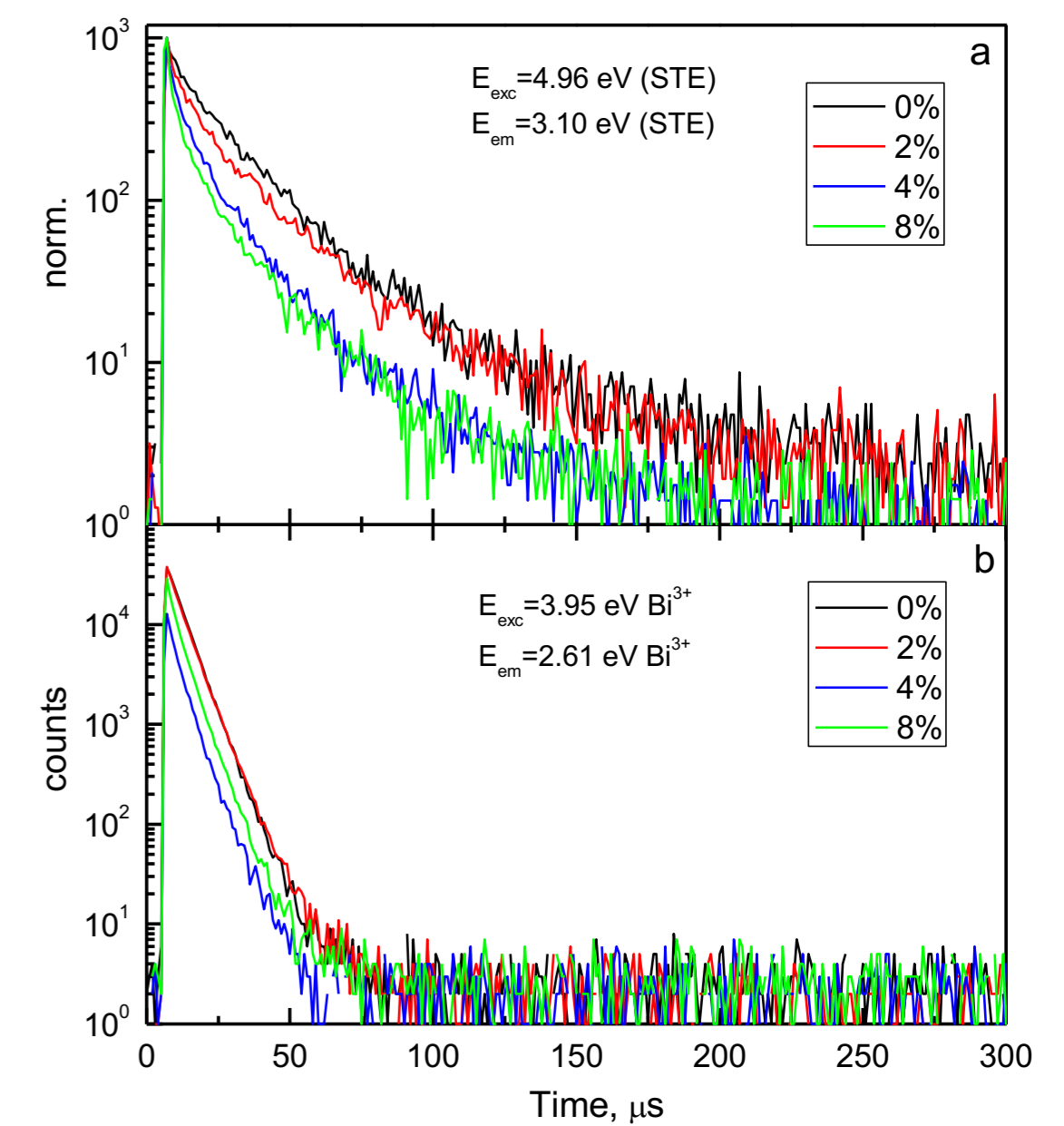
Emission (red lines) and excitation (black lines) spectra of (a) LuNbO₄, (b) LuNbO₄:Bi, and (c) LuNbO₄:Bi,Eu measured at 79 K. In the inset the 1.76 eV emission of Eu³⁺ measured under excitation in the absorption bands of Bi³⁺ ($E_{exc} = 3.95$ eV, solid line) and Eu³⁺ ($E_{exc} = 2.36$ eV, dashed line).



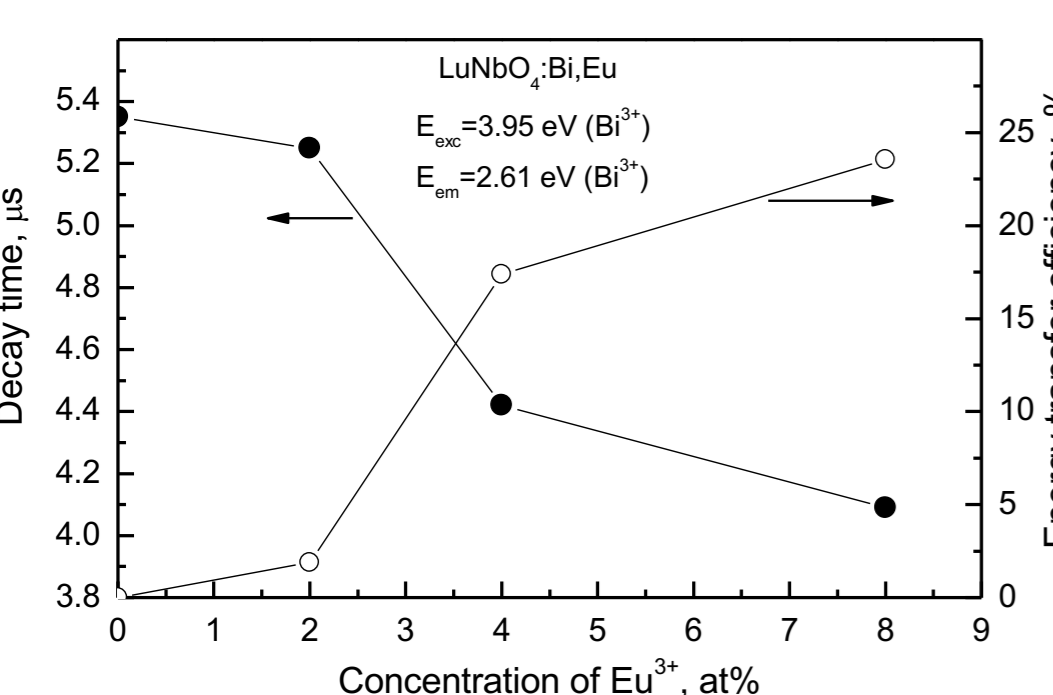
Emission spectra of LuNbO₄:Bi, Eu measured at 79 K for the samples with different nominal Eu³⁺ concentrations. (a) $E_{exc} = 4.96$ eV, (b) $E_{exc} = 3.96$ eV, (c) $E_{exc} = 3.65$ eV.



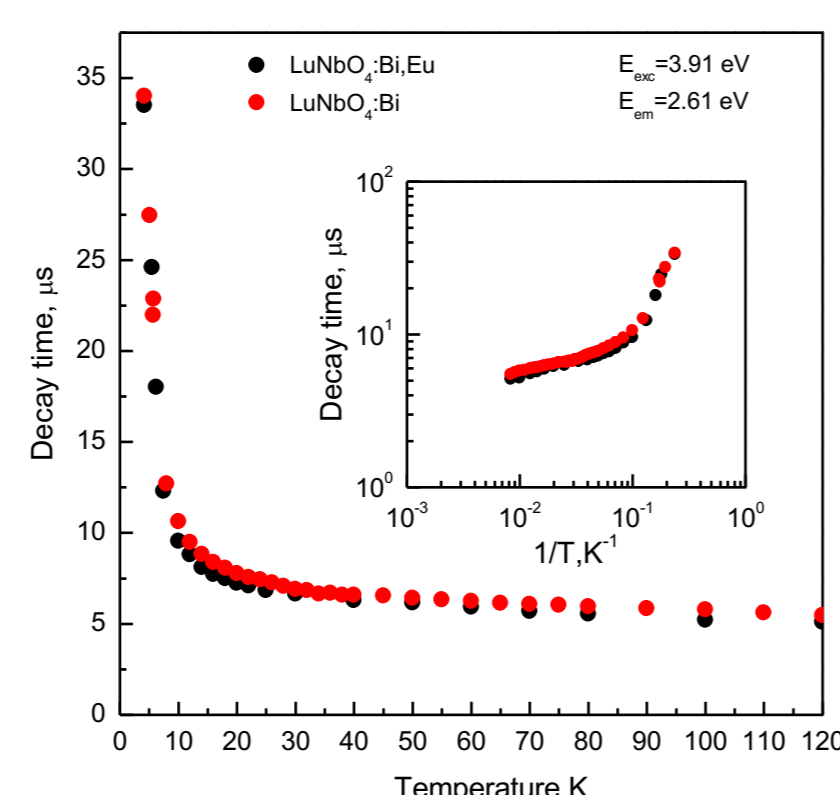
Dependences on the nominal Eu³⁺ concentration measured for (a,b,c) the maximum intensity of the intrinsic (black circles), Bi³⁺-related (blue circles) and Eu³⁺ (red circles) emissions of LuNbO₄:Bi,Eu. Dependences of the energy transfer efficiency (ETE) on the nominal Eu³⁺ concentration measured for the STE → Eu³⁺ (green solid line), ex⁰Bi³⁺ → Eu³⁺ (green dashed line) and ex⁰{Bi³⁺-Bi³⁺} → Eu³⁺ energy transfer in LuNbO₄:Bi,Eu. T = 79 K.



Decay curves of (a) the intrinsic and (b) the Bi³⁺-related emission measured at 79 K for the single-phase LuNbO₄:Bi, Eu samples with different nominal Eu³⁺ concentrations.



Dependences on the nominal Eu³⁺ concentration measured for the decay time of the Bi³⁺-related luminescence (filled circles) and for the corresponding energy transfer efficiencies (empty circles) in LuNbO₄:Bi, Eu. T = 79 K.



Temperature dependences of the decay times of the Bi³⁺-related emission in LuNbO₄:Bi (red filled circles) and LuNbO₄:Bi, 4%Eu (black filled circles).

Conclusion

In LuNbO₄:Bi,Eu, sample obtained results indicate the presence of energy transfer from the intrinsic and Bi³⁺-related centers to Eu³⁺ ions by the non-radiative (resonance) energy transfer process.

The values of a color rendering index (CRI) of 76%, quantum efficiency of ~18%, and correlated color temperature (CCT) of 5206 K are achieved for the LuNbO₄:2% Bi³⁺,0.4 % Eu³⁺ powder with the best CIE chromaticity coordinates (x = 0.316, y = 0.317).

References

1) M. Baran, K. N. Belikov, A. Kissabekova, A. Krasnikov, A. Lushchik, E. Mihokova, V. Tsiurma, L. Vasylechko, S. Zazubovich, Ya. Zhydachevskyy, J. Alloys and Compounds 859, 157800.

The values of color rendering indexes (CRI), quantum yields (QY), Bi³⁺ → Eu³⁺ energy transfer efficiencies (ETE), correlated color temperatures (CCT) and CIE chromaticity coordinates (x,y) obtained for the investigated LuNbO₄:Bi, Eu samples with different nominal Eu³⁺ concentrations.

Eu ³⁺ (%)	QY, %	ETE, %	CRI, %	CIE, x,y	CTT, K
0.2	18.8 ± 0.7	-	90	0.254, 0.296	9511
0.4	17.9 ± 0.6	-	76	0.316, 0.317	5206
1	25.1 ± 0.7	-	58	0.375, 0.318	2874
2	26.2 ± 0.6	32	53	0.501, 0.339	1529
4	29.1 ± 2.0	47	45	0.539, 0.340	1237
5	41.9 ± 2.1	60	35	0.608, 0.341	1091