
Chapter 7: Conclusions and Future Perspective

7.1 Conclusion of the Present Investigation

In this thesis work, different compositions of inorganic halide perovskite (CsPbX_3 ; $\text{X}=\text{Cl, Br}$) have been synthesized using various synthesis methods, such as hot-injection, and ligand-assisted reprecipitation methods. A detailed discussion of the synthesis methods is given in Chapter 2. Lanthanide ions (Ln^{3+}) have their unique optical properties, including ladder-like energy levels, sharp emission, and long luminescence lifetimes, which make them suitable for various applications. The Ln^{3+} -ions have been used as a dopant to increase the optoelectronic properties of CsPbX_3 . Ln-doping can passivate defects within the perovskite structure, thereby impacting charge carrier dynamics and overall material stability, as discussed in Chapter 3. Ln^{3+} -ions doping into CsPbX_3 enhance their photoluminescence (PL) properties. Energy transfer occurs from the perovskite matrix to the lanthanide ions and vice-versa, resulting in characteristic Ln^{3+} -emissions along with CsPbX_3 emission. Interfacing CsPbX_3 with different host matrix such as lanthanide metal organic framework (Ln-MOF), increases the stability of CsPbX_3 . The Ln-MOFs have beautiful morphology, as bi-flower like morphology of Eu-MOF and bi-directional needle like morphology of Tb/Eu-MOF are shown in Chapter 4 and 5, respectively. These Ln-MOFs have a porous structure, that helps in encapsulating the CsPbX_3 . The $\text{CsPbBr}_3@$ Eu-MOF is explored for the anti-counterfeiting application, while $\text{CsPbCl}_{1.5}\text{Br}_{1.5}@$ Tb/Eu-MOF shows anti-counterfeiting and white light emission applications. Also, Yb^{3+} , Er^{3+} : GdScO_3 oxide perovskite NCs is synthesised via the gel-combustion method for the optical thermometry application. The complete work is based on the lanthanides doped CsPbX_3 and GdScO_3 materials, and inorganic halide perovskites

interfaced lanthanide metal organic frame works. The materials have been synthesized using different synthesis techniques for various optical studies and future applications, the overall conclusion of the thesis work can be summarized as follows:

The first work investigates the structural and optical properties of pristine and Europium (Eu)-doped CsPbBr₃ halide perovskite nanocrystals (NCs) synthesized via the hot-injection method. Eu-doping enhances the crystallinity of CsPbBr₃ NCs and significantly reduces the average particle size from 19 nm to approximately 7 nm. The bandgap (E_g) of the undoped CsPbBr₃ NCs is determined to be 2.31 eV, and it increases slightly to 2.35 eV upon Eu²⁺-ion doping. The undoped CsPbBr₃ NCs exhibit a sharp emission peak at 515 nm in the green spectral region with a full width at half maximum (FWHM) of about 24 nm. In contrast, Eu-doped CsPbBr₃ NCs show two distinct emission peaks at 498 nm (FWHM ~20 nm) and 442 nm (FWHM ~13 nm). The additional emission peak at 442 nm is attributed to the $4f^6(^7F_1)5d^1 \rightarrow 4f^7$ transition of Eu²⁺-ion. The shift in green emission for the doped NCs is linked to the reduced particle size from 19 nm to 7 nm. Detailed decay dynamics reveal that the average decay time for CsPbBr₃ monitored at 515 nm is 7.27 ns, while for Eu-doped CsPbBr₃, the decay times are 5.80 ns and 8.89 ns for the 498 nm and 442 nm emission peaks, respectively. The Commission Internationale de l'Eclairage (CIE 1931) chromaticity coordinates of Eu-doped CsPbBr₃ emission spectra were analyzed under different excitation wavelengths, showing minimal variation. The luminous efficacy of radiation (LER) for undoped CsPbBr₃ NCs is calculated to be 447 lm/W, while for Eu-doped CsPbBr₃ NCs, it ranges between 138-179 lm/W. Notably, the color purity of the emission spectra exceeds 85% under various excitations. These optical characteristics suggest that CsPbBr₃ and Eu-doped CsPbBr₃ NCs

have potential applications in blue LEDs, color displays, and anti-counterfeiting technologies.

As discussed, the stability issue of the CsPbX₃ is a major concern. Doping metal-ion, surface capping, growing stable cell on the CsPbX₃ core, etc., may increase the stability of CsPbX₃. Metal organic frameworks (MOFs) are porous materials, which provide room to the nanomaterials to intercalated into it. The Eu-MOF is synthesized by the hydrothermal method and further a hybrid of this is formed with CsPbBr₃ i.e. CsPbBr₃@ Eu-MOF. It has a tetragonal structure with space group *P4₃22*. Synthesized Eu-MOF has bi-flower morphology whose mid portion and conical part length both are below 10 μm. The conical part of the bi-flower consists of many squared rods of square size ~200 nm. In UV-visible-NIR absorption spectrum, characteristic absorption edge of CsPbBr₃ is present. The PL emission spectrum as well as elemental analysis using SEM-EDX, confirms successful formation of the CsPbBr₃@Eu-MOF. Excitation wavelength-dependent PL emission of CsPbBr₃@Eu-MOF shows variation in the emission peak intensity. The (x, y) coordinate covers the whole green to red region in the CIE 1931 diagram. The maximum LER value achieved is 359 lm/W, and the maximum color purity is 94.4% which is significant for optical applications. The CIE 1931 (x, y) coordinates for the different excitation wavelength show the suitability of the materials for the optical anti-counterfeiting application. The pattern “LMDD” is encrypted on four different substrates (white paper, butter paper, green plastic, and aluminum foil) and the color change of the patterns under different excitation wavelengths has been thoroughly studied. The overall visualization and the chemical stability of the encrypted patterns on green plastic [“LMDD”-G] is the best among all the four substrates studied. This clearly suggest the suitability of developed material for anti-counterfeiting application.

Further, in a luminescent guest center encapsulated Ln-MOFs, luminescence emission can occur from the guest molecules, Ln³⁺-ions, and organic linker. To understand the energy-transfer among these luminescent centers, Tb/Eu-MOF and CsPbCl_{1.5}Br_{1.5} (CPCB) encapsulated CPCB@ Tb/Eu-MOFs hybrid have been synthesized using hydrothermal method. The Tb/Eu-MOF form bi-directional needle-like morphology. The single tetragonal phase of Tb/Eu-MOFs and CPCB@ Tb/Eu-MOFs are verified by the LeBail fitting of the XRD data. Successful incorporation of the inorganic halide perovskite CPCB into Tb/Eu-MOF is evidenced by the characteristic absorption band observed in the UV-visible spectrum and the peak in the PL emission. The Tb³⁺/Eu³⁺-ion and O²⁻ form the CTB. This band acts as an antenna for sensitizing the Tb³⁺ and Eu³⁺-ions. The energy transfer scheme among Tb³⁺ and Eu³⁺-ions, as well as CPCB, is also studied in detail. The CTB effectively excites the Tb³⁺ and Eu³⁺-ions, but not the CPCB. At the same time, the energy transfers from Tb³⁺ to Eu³⁺-ions also takes place but the reverse is not observed. In the PL emission spectrum of CPCB@ Tb/Eu-MOFs for the RGB color mixing, the red part is observed from Eu³⁺-ion, the green part is observed from Tb³⁺-ion, and the CPCB contributes the blue part. CIE 1931 (x, y) color coordinate of the c-CPCB@ Tb/Eu-MOF emission is calculated, that is very close to the white light coordinate (0.333, 0.333) suitable candidate for white light applications. To the best of our knowledge, this is the first report on CPCB luminescent guest-centered Tb/Eu-MOF for WLED application. Excitation wavelength-dependent PL emission of CPCB@ Tb/Eu-MOFs is also recorded, and the CIE (x, y) color coordinate is calculated. It gives different color sensations to the human eyes. Integrating this idea, pattern "W" was developed on various surfaces (borosilicate glass, aluminum foil,

transparent plastic) for optical anti-counterfeiting application. The stability of the "W-BG" pattern is checked in ambient conditions and encounters with water and propanol solution. Results show that CPCB@ Tb/Eu-MOFs is very suitable for white light emission and for optical anti-counterfeiting applications.

The Ln³⁺-ions pairs such as Er³⁺-Yb³⁺, Ho³⁺-Yb³⁺, Tm³⁺-Yb³⁺ exhibit non-linear photon upconversion (UC) phenomena in certain host materials. The UC emission is somehow difficult to achieve in CsPbX₃ because of week crystal field effect. The oxide perovskite GdScO₃, known for its wide bandgap (~5 eV) and thermal stability, has been investigated as a host for Yb³⁺ and Er³⁺ ions. Nanocrystalline powders of GdScO₃ and Yb³⁺, Er³⁺: GdScO₃ were synthesized using the gel-combustion method. Optimization of the doping concentrations of Er³⁺ and Yb³⁺ ions in this host was performed to maximize luminescence intensity. Remarkably, this material exhibits strong near-infrared (NIR) to visible UC emission under 980 nm continuous-wave (CW) laser excitation, as well as down-shifting emission under UV-blue excitation. To the best of our knowledge, this study was the first to report on the UC properties of Yb³⁺, Er³⁺ co-doped GdScO₃. Additionally, Yb³⁺, Er³⁺: GdScO₃ demonstrates excellent temperature sensitivity, making it suitable for sensing applications. The maximum absolute sensitivity achieved is $10.49 \times 10^{-3} \text{ K}^{-1}$ at 423 K, and the maximum relative sensitivity is $11.79 \times 10^{-3} \text{ K}^{-1}$ at 299 K, with a thermal resolution of 0.4 K, which surpasses that of many other oxide perovskite materials. This material holds significant potential for use as a non-contact, luminescence-based optical temperature sensor, particularly in environments such as coal mines and high-temperature metal industries. It may also be applicable in the optoelectronic industry for monitoring heat-related damage in electronic circuits.

Overall, it can be concluded that the stability and optical properties of the CsPbX₃ can be enhanced by interfacing with the Ln³⁺-ions and metal organic frameworks for the different optoelectronic applications such as color and white LEDs, optical encryption and decryption, optical thermometry, etc. At the same time Ln³⁺-ions doped high bandgap material GdScO₃ is suitable for the non-invasive optical thermometry.

7.2 Outlook for Future Work

For the future perspective the following works are proposed for a deep understanding of the various interaction within the inorganic halide perovskites molecules for the various optoelectronic applications.

- ❖ Surface engineering to enhance the stability as well as properties of CsPbX₃.
- ❖ Doping Ln³⁺-ions pairs such as Er³⁺-Yb³⁺, Ho³⁺-Yb³⁺, Tm³⁺-Yb³⁺, etc., into CsPbX₃ for the UC emission.
- ❖ Optoelectronic behaviors study in CsPbX₃ encapsulated transition metals-MOFs.
- ❖ Exploring these materials for the evergreen applications such as solar cell, energy storage, sensing, etc.