Highly Concentrated, Zwitterionic Ligand-Capped Mn$^{2+}$:CsPb(Br$_x$Cl$_{1-x}$)$_3$ Nanocrystals as Bright Scintillators for Fast Neutron Imaging

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ABSTRACT: Fast neutron imaging is a nondestructive technique for large-scale objects such as nuclear fuel rods. However, present detectors are based on conventional phosphors (typically microcrystalline ZnS:Cu) that have intrinsic drawbacks, including light scattering, γ-ray sensitivity, and afterglow. Fast neutron imaging with colloidal nanocrystals (NCs) was demonstrated to eliminate light scattering. While lead halide perovskite (LHP) FAPbBr$_3$ NCs emitting brightly showed poor spatial resolution due to reabsorption, the Mn$^{2+}$-doped CsPb(BrCl)$_3$ NCs with oleyl ligands had higher resolution because of large apparent Stokes shift but insufficient concentration for high light yield. In this work, we demonstrate a NC scintillator that features simultaneously high quantum yields, high concentrations, and a large apparent Stokes shift. In particular, we use long-chain zwitterionic ligand capping in the synthesis of Mn$^{2+}$-doped CsPb(BrCl)$_3$ NCs that allows for attaining very high concentrations (>100 mg/mL) of colloids. The emissive behavior of these ASC18-capped NCs was carefully controlled by compositional tuning that permitted us to select for high quantum yields (>50%) coinciding with Mn-dominated emission for minimal self-absorption. These tailored Mn$^{2+}$:CsPb(BrCl)$_3$ NCs demonstrated over 8 times brighter light yield than their oleyl-capped variants under fast neutron irradiation, which is competitive with that of near-unity FAPbBr$_3$ NCs, while essentially eliminating self-absorption. Because of their rare combination of concentrations above 100 mg/mL and high quantum yields, along with minimal self-absorption for good spatial resolution, Mn$^{2+}$:CsPb(BrCl)$_3$ NCs have the potential to displace ZnS:Cu as the leading scintillator for fast neutron imaging.

Radiographic imaging with fast neutrons (>1 MeV) is increasingly sought as a nondestructive probe for large-scale objects that even high-energy X/γ-rays or thermal neutrons (~25 meV) cannot penetrate effectively.1−9 This is because such high-energy neutrons offer low interaction cross sections for both high- and low-Z elements, whereas X/γ-rays are very sensitive to high-Z elements while thermal neutrons are especially interactive with low-Z elements such as Li or H.1 However, fast neutron imaging is limited by the detector performance, where trade-offs are routinely made in efficiency, spatial resolution, and the decay time required between measurements.10−12 Fast neutron detection for imaging relies on the elastic scattering of neutrons off nuclei, which then generates a recoil nucleus with small penetration depth; this recoil nucleus ionizes charge carriers and excites the scintillator.1,13−15 Traditionally, the community has utilized standard phosphors as the indirect detectors for recoil protons generated by scattering of fast neutrons.10,11,16

The leading material in the field is ZnS:Cu microparticles embedded in polypropylene (hereafter denoted ZnS:Cu(PP)) for high hydrogen density, which offers superior light yield10,11 but suffers from scattering at the plastic−phosphor interface that limits spatial resolution with increasing thickness as well as deleterious minutes-long afterglow under fast neutron irradiation, greatly impacting the time needed to conduct the

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applications spanning from LEDs for displays to fast neutron imaging scintillators. These intrinsic drawbacks of scattering and afterglow require a new approach for fast neutron imaging scintillators. One particularly appealing class of phosphors are colloidal semiconducting nanocrystals (NCs), which naturally comprise a two-component, scattering-free system that includes emissive and hydrogen-dense components (NCs and solvent, respectively) in close contact. Among colloidal nanocrystals, lead halide perovskite (LHP) NCs have recently emerged as the material of choice for fast neutron-for applications spanning from LEDs for displays to fast neutron imaging scintillators. These intrinsic drawbacks of scattering and afterglow require a new approach for fast neutron imaging scintillators. One particularly appealing class of phosphors are colloidal semiconducting nanocrystals (NCs), which naturally comprise a two-component, scattering-free system that includes emissive and hydrogen-dense components (NCs and solvent, respectively) in close contact. Among colloidal nanocrystals, lead halide perovskite (LHP) NCs have recently emerged as the material of choice for fast neutron imaging scintillators.1,2

While pure LHP NCs have bright emission, it is crucial to also achieve high concentrations and high Stokes shifts with suppressed optical reabsorption. For LHP NCs, these optical characteristics are typically achieved via the doping of CsPbBr3 nanocrystals with Mn2+ ions, in which optical reabsorption is minimized by decoupling the Mn2+ emission from the absorption of the perovskite matrix, resulting in an “apparent” large Stokes shift. However, the relatively poor chemical stability and the intrinsically low concentration limit which can be achieved by using oleyl chains as ligands hamper the efficiency of Mn2+:CsPbBr3 NCs as fast neutron scintillators.3

In this work, the development of a new synthesis yields Mn2+-doped LHP NCs that pair excellent optical quality with drastically higher concentrations. This approach improves the light yield in this material system under fast neutron irradiation over 8 times that of the oleyl-capped Mn2+-doped LHP NCs, permitting high light yields on par with the unity PL QY system FAPBr3, without the detrimental self-absorption. These ASC18-capped Mn2+:CsPbBr3 NCs represent a substantial advance in this field, combining the beneficial properties of colloidal NCs (lack of scattering, no long-term afterglow) with both high light yield and good spatial resolution for fast neutron imaging.

Building upon the remarkable tunability of perovskite NCs, recent work in our group has demonstrated that enhanced stability and a broad range of concentrations can be achieved in undoped CsPbX3 (X = Br, Cl) NCs through the appropriate choice of long-chain zwitterionic ligands. Here, we adapted these syntheses to generate zwitterionic-capped-CsPbBr3 NCs doped with Mn2+ utilizing 3-(N,N-dimethyloctadecylammonio)propanesulfonate (ACS18) as the ligand (see the Supporting Information for further details about the synthesis). Their optical properties (Figure 1a) are similar to those observed for their oleyl-capped counterparts, with a well-defined absorption edge (with clear excitonic features) and photoluminescence emission (PL) characterized by two contributions: one centered between 400 and 500 nm, associated with band-edge exciton recombination, and another centered between 580 and 650 nm corresponding to recombination from Mn2+ states. PL QY values of up to 60% can be achieved through appropriate compositional tuning. From a structural point of view, X-ray diffraction patterns (Figure 1b) are consistent with mixed Br−−Cl− perovskite composition, and they show the occasional presence of CsMnCl3 impurities when high concentrations of Mn2+ precursor are used in the synthesis. Transmission electron microscopy (TEM) images (Figure 1d) reveal that the NCs are characterized by cubic shape with size of roughly 10 nm. The

Figure 1. Optical and structural characterization of Mn2+:CsPb(BrCl)3 NCs. (a) Photograph of Mn2+:CsPb(BrCl)3 NCs solution under ultraviolet illumination. (b) Representative absorption (dashed line) and PL emission (solid line) spectrum of Mn2+:CsPb(BrCl)3 NCs. (c) Representative X-ray diffraction pattern of Mn2+:CsPb(BrCl)3 NCs (blue). XRD references correspond to CsMnCl3 (green, ICSD 2525), CsPbCl3 (purple, ICSD 243734), CsPbBr3 (red, ICSD 231017), and CsMnBr3 (orange, ICSD 2782). (d) Representative transmission electron micrograph of Mn2+:CsPb(BrCl)3 NCs. The average size is 9.5 ± 0.8 nm.
Figure 2. Tuning the optical properties of Mn$^{2+}$:CsPb(BrCl)$_3$ NCs. (a) Absorption (dashed lines) and PL emission (solid lines) spectra of 50% Mn$^{2+}$:CsPb(BrCl)$_3$ NCs with increasing Br$^-$ content (from bottom to top): 0, 25, 33, 50, 66, 75, and 100%. The nominal Mn$^{2+}$ content is constant at 50%. (b) Absolute QY of the total 50% Mn$^{2+}$:CsPb(BrCl)$_3$ PL emission (blue dots) and of the Mn$^{2+}$ emission (orange dots) as a function of Br$^-$ content. The green dashed line marks the reference sample. (c) Percentage of the emission from Mn$^{2+}$ levels on the overall emission of the 50% Mn$^{2+}$:CsPb(BrCl)$_3$ NCs as a function of Br$^-$ content. The green dashed line marks the reference sample. (d) XRD spectra of the 50% Mn$^{2+}$:CsPb(BrCl)$_3$ samples of panels a–c. Br$^-$ content increases from bottom to top. The red dashed lines mark the positions of the main reflections of CsPbBr$_3$ (ICSD 231017), while the purple dashed lines mark the positions of the main reflections of CsPbCl$_3$ (ICSD 243734). (e) Absorption (dashed lines) and PL emission (solid lines) spectra of Mn$^{2+}$:CsPbBrCl$_3$ NCs with increasing...
nominal Mn$^{2+}$ content (from bottom to top): 0, 20, 40, 50, 66, and 80%. The nominal Br$^{-}$ content is constant at 33%. (f) Absolute QY of the total Mn$^{2+}$-CsPbBrCl$_3$ PL emission (blue dots) and of the Mn$^{2+}$ emission (orange dots) as a function of Mn$^{2+}$ content. The green dashed line marks the reference sample. (g) Percentage of the emission from Mn$^{2+}$ levels on the overall emission of the Mn$^{2+}$-CsPbBrCl$_3$ NCs as a function of Mn$^{2+}$ content. The green dashed line marks the reference sample. (h) XRD spectra of the Mn$^{2+}$-CsPbBrCl$_3$ samples of panels e–g. Mn$^{2+}$ content increases from bottom to top.

The presence of zwitterionic capping ligands on the surface of the NCs greatly enhances the chemical stability and enables significantly higher concentrations of NCs in solution compared to their counterparts stabilized by oleyl. Typically, with ASC18 as stabilizing agent, concentrations of ~100 mg/mL were achieved (Table S1), in agreement with previous reports; this is an increase of 2 orders of magnitude compared to the oleyl-stabilized NCs from the previous study. We also attempted to utilize lecithin as the capping ligand to obtain even higher concentrations as reported for undoped NCs, but while we achieved concentrations of ~400 mg/mL (Figure S1), the PL QY of these NCs never exceeded 10%; this evidence implies that the ligand shell also plays a pivotal role in determining the final optical properties of the NCs.

We then sought to leverage the tunability of these doped perovskite NCs to achieve the high (apparent) Stokes shift and PL QYs required for efficient fast neutron scintillation. The optical properties of the NCs can be widely tuned by varying the ratios between the halide precursors during the synthesis. When increasing the Br$^{-}$ content at the expense of the Cl$^{-}$ content while keeping the Mn$^{2+}$ content constant (nominal 50%), we observe that the absorption band edge and the band-edge PL emission shift toward longer wavelengths, while the Mn$^{2+}$ PL shifts toward shorter wavelengths (Figure 2a–d), consistent with previous observations. In particular, the PL emission (Figure 2a) associated with band-edge exciton recombination shifts from 403 nm for pure Mn$^{2+}$-CsPbCl$_3$ to 515 nm for pure Mn$^{2+}$-CsPbBr$_3$, while the PL emission from Mn$^{2+}$ states shifts from 604 to 595 nm; this latter effect has been attributed to a weakening of the crystal field induced by the expansion of the crystal lattice when Cl$^{-}$ is substituted for Br$^{-}$. At the same time, we also observe the disappearance of the excitonic feature in the absorption spectra of the NCs due to a relaxation of the quantum confinement, consistent with previous observations in anion exchange reactions. Interestingly, the Br$^{-}$ content strongly affects the ratio between the two components of the PL emission: the emission originating from the band-edge exciton recombination becomes dominant for a nominal Br$^{-}$ content larger than 60%. This behavior is even more evident when considering the variation of the PL QY (Figure 2b) and of the amount of emission from the Mn$^{2+}$ states (relative to the overall emission; Figure 2c). The overall PL QY scales with the Br$^{-}$ content, increasing from a value of 9%, for pure Mn$^{2+}$-CsPbCl$_3$ to 62%, for nominal 50% Br$^{-}$ content; for larger values of Br$^{-}$ content, the overall PL QY remains constant. In contrast, the proportion (%) of the emission from Mn$^{2+}$ states slightly increases, while being close to unity for increasing Br$^{-}$ content, until it linearly decreases for Br$^{-}$ content greater than 33%. This behavior can be understood when considering the competition between the two recombination processes (from band-edge states and Mn$^{2+}$ states) and the exciton-to-activator (Mn$^{2+}$) forward and backward energy transfer. As the Br$^{-}$ content increases, the exciton radiative recombination rate decreases while the energy transfer rate increases, resulting in an increase in the emission from Mn$^{2+}$ states. At the same time, with increasing Br$^{-}$ content, the energy difference between the band-edge and the Mn$^{2+}$ states decreases, hence making the backward energy transfer more efficient. Furthermore, because the radiative recombination from band-edge states is much faster (~ns) than the recombination from the Mn$^{2+}$ states (~ms), this recombination process will be more favorable, effectively decreasing the proportion of excitons recombining from the Mn$^{2+}$ states (for Br$^{-}$ content greater than 33%). From a structural point of view, the XRD patterns (Figure 2d) are consistent with the evolution from pure Mn$^{2+}$-CsPbCl$_3$ perovskite NCs to pure Mn$^{2+}$-CsPbBr$_3$ via mixed halide composition.

Beyond the traditional halide tuning that has helped popularize perovskite NCs, the addition of the Mn$^{2+}$ dopant adds a further knob for controlling the resulting optical properties of Mn$^{2+}$-CsPbBrCl$_3$, NCs by varying the ratio between Pb$^{2+}$ and Mn$^{2+}$ precursors during the synthesis. Keeping the Br$^{-}$ content constant at 33% nominal (hereafter denoted Mn$^{2+}$-CsPbBrCl$_3$) to maximize both the PL QY and the proportion of Mn$^{2+}$ emission, we observe that the emission from Mn$^{2+}$ states scales with the nominal Mn$^{2+}$ content (Figure 2e–h). Furthermore, while the band edge emission remains constant at 427 nm, the Mn$^{2+}$ emission shifts from 596 nm (for 20% nominal Mn$^{2+}$ content) to 614 nm (for Mn$^{2+}$ contents larger than 50%); this effect has previously been attributed to an increase of the crystal field. Furthermore, we observe that for nominal Mn$^{2+}$ content greater than 50%, both the absorption band edge and the band-edge emission are shifted toward longer wavelengths (absorption, from 423 nm; PL, from 427 to 452 nm); we attribute this shift to the larger size of the NCs in this doping regime, as we will show later. The increasingly dominant contribution of emission from Mn$^{2+}$ states is clear when inspecting the dependence of the PL QY on the nominal Mn$^{2+}$ content (Figure 2fg). The PL QY steadily increases with increasing nominal Mn$^{2+}$ content, from a value of 4%, for pure CsPbBrCl$_3$, to a maximum of 54%, for 66% nominal Mn$^{2+}$ content; for higher Mn$^{2+}$ contents a decrease in the PL QY is observed. Analogously, when considering the contribution of the Mn$^{2+}$-emission to the overall emission (Figure 2g), we observe that a maximum is reached for 50% nominal Mn$^{2+}$ content, while for higher contents the values are lower. The peculiar optical behavior of Mn$^{2+}$-CsPbCl$_3$Br for nominal Mn$^{2+}$ content greater than 50% can be understood by complementing the optical data with XRD structural data (Figure 2h). The XRD patterns for the nominal Mn$^{2+}$ contents 0–50% are rather similar, highlighting the presence of mainly Mn$^{2+}$-CsPbCl$_3$Br with some minor impurities ascribable to the presence of CsMnCl$_3$ and CsMnBr$_3$, phases, which are the main byproducts of the synthesis. However, for samples with nominal Mn$^{2+}$ content greater than 50%, the sharpness of the XRD reflections implies the average domain size to be much larger compared to the other samples. Inspection of electron micrographs of these samples (Figure S2 in the Supporting Information) show the
The presence of large cubic (\(\sim 50\)–100 nm) \(\text{Mn}^{2+}:\text{CsPb(BrCl)}_3\) NCs. A difference in reactivity between the \(\text{Mn}^{2+}\) and the \(\text{Pb}^{2+}\) precursors, \(\text{Mn(oleate)}_2\) and \(\text{Pb(oleate)}_2\), respectively, is invoked as the origin of the much larger dimensions of the NCs synthesized in the presence of large quantities of \(\text{Mn}^{2+}\) precursor during the synthesis: the higher bond dissociation energy of the \(\text{Mn}−\text{O}\) bond (402 kJ/mol)\(^70\) compared to the \(\text{Pb}−\text{O}\) bond (378 kJ/mol)\(^70\) implies a lower reactivity of the \(\text{Mn(oleate)}_2\) precursor, resulting therefore in larger NCs. This difference in reactivity can also explain the discrepancy between the nominal \(\text{Mn}^{2+}\) content introduced in the reaction vessel and the actual \(\text{Mn}^{2+}\) content incorporated in the NCs. Energy-dispersive X-ray analysis of the NCs with different nominal \(\text{Mn}^{2+}\) content emphasizes how only a marginal percentage of \(\text{Mn}^{2+}\) is incorporated in the NCs: for nominal 50% \(\text{Mn}^{2+}\) content, only 2.0 atom % is detected in the NCs, while for nominal 20% \(\text{Mn}^{2+}\) content, only 0.1 atom % is detected. We also remark that the exact position of the dopants (in the bulk or on the surface) was not determined, but further analysis, e.g., with EPR,\(^{34}\) could discriminate between these two cases. Nevertheless, these relatively small, incorporated amounts have a clear and important impact on the optical properties of the NCs.

Now that these tunable NCs have delivered high concentrations coinciding with excellent PL QYs and a large apparent Stokes shift, we explored the potential for these \(\text{Mn}^{2+}:\text{CsPb(BrCl)}_3\) NCs as scintillators for fast neutron imaging. Given the importance of the Stokes shift in this application,\(^1\) NC samples with the fixed Br/Cl ratio of 1:2 (where \(\text{Mn}^{2+}\)-emission dominates the emission spectrum, Figure 2e–h) with varying nominal \(\text{Mn}^{2+}\) content were compared, and the basic characteristics of these samples are detailed in Table S1. Fast neutron imaging measurements were conducted at the thermal neutron imaging beamline \(\text{NEUTRA}^{71}\) of the \(\text{SINQ}\) spallation neutron source at the Paul Scherrer Institut. The thermal neutrons (\(\sim 25\) meV) were screened out of the beam by leaving the experimental shutter (composed of B\(_4\)C and Cd blocking layers) closed and adding a 2 mm Cd filter, leaving only the intermediate (\(>1\) eV) and fast neutrons (\(>1\) MeV) needed for the imaging experiments.
Samples were sealed in a light-tight box of the camera detector system that collected the incident light from the excited NC solutions, with a mirror ensuring that the camera is not in line with the beam to reduce background noise and camera irradiation.

A series of radiographs were measured for these Mn\textsuperscript{2+}:CsPbBr\textsubscript{3} NCs under 147.2 s fast neutron exposure, with FAPbBr\textsubscript{3} NCs and a commercial RC Tritec AG ZnS:Cu(PP) screen used for reference (Figure 3a). The background and asymmetry-corrected (Figure S3) light output are shown in Figure 3b as a percentage of that of the ZnS:Cu(PP) reference screen, and we observe that the scintillation intensity of these Mn\textsuperscript{2+}-containing NCs is comparable to the state-of-the-art FAPbBr\textsubscript{3} NCs\textsuperscript{1}, which have a quantum yield near unity. This validates the design principles described above, namely, that high Stokes shift and high concentration can overcome limitations of lower PL QQY. The light output of these ASC18-capped Mn\textsuperscript{2+}:CsPbBrCl\textsubscript{3} NCs is 8.65 times greater than that previously measured for oleyl-capped Mn\textsuperscript{2+}:CsPbBrCl\textsubscript{3} NCs\textsuperscript{2}, as determined using the relative light outputs of the FAPbBr\textsubscript{3} reference NCs to account for the different beam characteristics (see the Supporting Information for details).\textsuperscript{3} Notably, the undoped sample shows essentially no light output under the same conditions (likely due to the optical density of the highly concentrated colloid), further demonstrating the importance of high Stokes shift in achieving efficient scintillation in such dense samples. No afterglow was observed in 147.2 s images collected immediately after beam closure (Figure S4), though afterglow effects are reduced here because of the lower fast neutron flux of this beamline; therefore, this is not conclusive evidence against afterglow in these specific NCs. However, previous measurements for this system with oleyl ligands showed no afterglow, in stark contrast to ZnS:Cu(PP).\textsuperscript{4}

The brightest sample (nominally 66% Mn\textsuperscript{2+}) was selected for further characterization of the thickness- and concentration-dependence of the fast neutron scintillation performance (Figure 3c). The concentration-dependent light output shows a continuous increase with increasing concentration (Figure S5a), indicating that the charge collection efficiency increases as we obtain more NCs per unit volume. Compared to prior efforts,\textsuperscript{1,5} this increase is significantly closer to linearity, indicating that the self-absorption of these Stokes-shifted NCs is well below that of FAPbBr\textsubscript{3} NCs and closer to self-trapped exciton systems such as the ionic liquid THTDPCI-PbCl\textsubscript{3} that recently demonstrated spatial resolution superior to that of the ZnS:Cu reference screens.\textsuperscript{6}

Thickness-dependent measurements further confirm that the effect of self-absorption is substantially reduced in these doped NCs, as the light output is essentially linear and reaches 91.9% of the expected value (estimated using the stopping power of the solvent to 1 MeV neutrons) for a 10-fold increase in thickness (Figure 3d). Note that this expected value at 5 and 10 mm is overestimated as the beam includes both fast neutrons and intermediate neutrons, where the latter have smaller penetration depths and therefore would contribute to more light output at lower thicknesses; hence, the 91.9% value is a lower limit of the actual efficiency versus the expected values. This is a dramatic improvement over established NCs as FAPbBr\textsubscript{3} and CdSe/CdS NCs suffered substantial self-absorption losses as the thickness increased, reaching only 56.7% and 37.0% of the expected value, respectively.\textsuperscript{3} The scalability of the Mn\textsuperscript{2+}:CsPbBrCl\textsubscript{3} system is thus vastly superior to the prior NCs as both the concentration and thickness can be increased without reducing the signal, and the spatial resolution will similarly benefit as problematic reabsorption will not impact the sharpness of the measured radiographs.

In conclusion, a new synthesis based on zwitterionic ASC18 ligands yielded Mn\textsuperscript{2+}:CsPb(BrCl)\textsubscript{3} NCs which are colloidally stable to high concentrations of ∼100 mg/mL while maintaining excellent optical quality. This synthesis extends the current technology of zwitterionic capped LHP NCs to doped systems, opening the doors to robust doped NCs in a wider concentration range. Furthermore, this system is immensely tunable through the halide and metal precursors, enabling us to rationally select the Mn\textsuperscript{2+}:CsPbBrCl\textsubscript{3} system as having the ideal combination of high quantum yield (up to 53%) and dominant Mn\textsuperscript{2+} emission (97% of total emission). Fast neutron imaging experiments showed that these dense ASC18-capped NCs achieved light yields over 8 times greater than those of their oleyl-capped counterparts, matching that of near-unity FAPbBr\textsubscript{3} NCs, despite being characterized by a quantum yield which was only half that. Furthermore, thickness and concentration-dependent measurements under fast neutron irradiation showcased the lack of self-absorption in this doped system, with essentially linear concentration dependence and drastically enhanced scalability with greater than 91.9% of the expected light yield achieved for a 10-fold increase in thickness. These results validate the design principles laid out by us previously, demonstrating that the Mn\textsuperscript{2+}:CsPbBrCl\textsubscript{3} NCs offer an unparalleled combination of high concentrations (>100 mg/mL), quantum yields (>50%), and apparent Stokes shift (~1 eV) that permits efficient fast neutron scintillation without the problematic reabsorption that precludes FAPbBr\textsubscript{3} NCs from achieving sufficient spatial resolution to displace ZnS:Cu(PP). Future work will continue development of these NCs, including maximizing the concentration, utilizing solvents with higher hydrogen density than toluene, and preparing larger-scale detector screens.

### ASSOCIATED CONTENT

**Supporting Information**

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acsenergylett.1c01923.

- Nanocrystal synthesis, optical and structural characterization methods, and fast neutron imaging (PDF)

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REFERENCES

(1) McCall, K. M.; Sakhatksiy, K.; Lehmann, E.; Walfort, B.; Losko, A. S.; Montanarella, F.; Bodnarchuk, M. I.; Krieger, F.; Kestemur, Y.; Mannes, D.; Shynkarenko, Y.; Yakunin, S.; Kovalenko, M. V. Fast Neutron Imaging with Semiconductor Nanocrystal Scintillators. ACS Nano 2020, 14 (11), 14686–14697.

(2) Kardjilov, N.; Manke, I.; Hilger, A.; Strobl, M.; Banhart, J. Neutron Imaging in Materials Science. Mater. Today 2011, 14 (6), 248–256.

(3) Schilling, B.; Beaudet, A.; Fedrigo, A.; Grazzi, F.; Kullmer, O.; Laiß, M.; Makowska, M.; Werneburg, I.; Zanolli, C. Neutron Imaging in Cultural Heritage Research at the Heinz Maier-Leibnitz Center. J. Imaging 2018, 4 (1), 22.

(4) Kardjilov, N.; Manke, I.; Woracek, R.; Hilger, A.; Banhart, J. Advances in Neutron Imaging. Mater. Today 2018, 21 (6), 652–672.

(5) Büchler, T.; von Gostomski, C. L.; Baldauf, T. Fission Neutron Tomography of a 280-L Waste Package. Materials Research Proceedings 2018, 15, 299–304.

(6) Büchler, T.; Kalthoff, O.; von Gostomski, C. L. A Feasibility Study on Reactor Based Fission Neutron Radiography of 200-L Waste Packages. Phys. Procedia 2017, 88 (September 2016), 64–72.

(7) Börries, S.; Metz, O.; Pranzas, P. K.; Bellosta von Colbe, J. M.; Büchler, T.; Dornheim, M.; Klässen, T.; Schreyer, A. Optimization and Comprehensive Characterization of Metal Hydride Based Hydrogen Storage Systems Using In-Situ Neutron Radiography. J. Power Sources 2016, 328, 567–577.

(8) Mühlbauer, M. J.; Büchler, T.; Kellermeier, M.; Knapp, M.; Makowska, M.; Schulz, M.; Zinnik, S.; Ehrenberg, H. Neutron Imaging with Fission and Thermal Neutrons at NECTAR at MLZ. Phys. B 2018, 551, 359–363.

(9) Nelson, R.; Vogel, S.; Hunter, J.; Watkins, E.; Losko, A.; Tremsin, A.; Borges, N.; Cutler, T.; Dickinson, L.; Espy, M.; Gautier, D.; Madden, A.; Majewski, J.; Malone, M.; Mayo, D.; McClellan, K.; Montgomery, D.; Mosby, S.; Nelson, A.; Ramos, K.; Schirato, R.; Schroeder, K.; Sawano, S.; Swift, A.; Vo, L.; Williamion, T.; Winch, N. Neutron Imaging at LANSCE—From Cold to Ultrafast. J. Imaging 2018, 4 (2), 45.

(10) Makowska, M.; Walfort, B.; Zeller, A.; Grünzweig, C.; Büchler, T. Performance of the Commercial PP/ZnS:Cu and PP/ZnS:Ag Scintillation Screens for Fast Neutron Imaging. J. Imaging 2017, 3 (4), 60.

(11) Zboray, R.; Adams, R.; Morgan, M.; Kis, Z. Qualification and Development of Fast Neutron Imaging Scintillator Screens. Nucl. Instrum. Methods Phys. Res., Sect. A 2019, 930 (December 2018), 142–150.

(12) Zboray, R.; Adams, R.; Kis, Z. Scintillator Screen Development for Fast Neutron Radiography and Tomography and Its Application at the Beamline of the 10 MW BNC Research Reactor. Appl. Radiat. Isot. 2018, 140 (April), 215–223.

(13) Lehmann, E. H.; Vontobel, P.; Frei, G.; Brönnimann, C. Neutron Imaging—Detector Options and Practical Results. Nucl. Instrum. Methods Phys. Res., Sect. A 2004, 531 (1–2), 228–237.

(14) van Eijk, C. W. E. Inorganic Scintillators for Thermal Neutron Detection. Radiat. Meas. 2004, 38 (4–6), 337–342.

(15) Caruso, A. N. The Physics of Solid-State Neutron Detector Materials and Geometries. J. Phys.: Condens. Matter 2010, 22 (44), 443201.

(16) Bravar, U.; Brullaard, P. J.; Flekiger, E. O.; Macrì, J. R.; McConnell, M. L.; Moser, M. R.; Ryan, J. M.; Woolf, R. S. Design and Testing of a Position-Sensitive Plastic Scintillator Detector for Fast Neutron Imaging. IEEE Trans. Nucl. Sci. 2006, 53 (6), 3894–3903.
(47) Pan, G.; Bai, X.; Yang, D.; Chen, X.; Jing, P.; Qu, S.; Zhang, L.; Zhou, D.; Zhu, J.; Xu, W.; Dong, B.; Song, H. Doping Lanthanide into Perovskite Nanocrystals: Highly Improved and Expanded Optical Properties. *Nano Lett.*, 2017, 17 (12), 8005–8011.

(48) Shapiro, A.; Heindl, M. W.; Horani, F.; Dahan, M.-H.; Tang, J.; Amouyal, Y.; Lifshitz, E. Significance of Ni Doping in CsPbX₃ Nanocrystals via Postsynthesis Cation–Anion Coexchange. *J. Phys. Chem. C* 2019, 123 (40), 24979–24987.

(49) Akkerman, Q. A.; Meggiolaro, D.; Dang, Z.; De Angelis, F.; Manna, L. Fluorescent Alloy CsPbₓBr₁₋ₓ Nanocrystals with High Structural and Optical Stability. *ACS Energy Lett.* 2017, 2 (9), 2183–2186.

(50) Zou, S.; Liu, Y.; Li, J.; Liu, C.; Feng, R.; Jiang, F.; Li, Y.; Song, J.; Zeng, H.; Hong, M.; Chen, X. Stabilizing Cesium Lead Halide Perovskite Lattice through Mn(II) Substitution for Air-Stable Light-Emitting Diodes. *J. Am. Chem. Soc.* 2017, 139 (33), 11443–11450.

(51) Coropeanu, I.; Bawendi, M. G. Core/Shell Quantum Dot Based Luminescent Solar Concentrators with Reduced Reabsorption and Enhanced Efficiency. *Nan Lett.* 2014, 14 (7), 4097–4101.

(52) Meinardi, F.; Colombo, A.; Velizhanin, K. A.; Simonutti, R.; Lorenzon, M.; Beverina, L.; Viswanatha, R.; Klimov, V.; Brovelli, S. Large-Area Luminescent Solar Concentrators Based on ‘Stokes-Shift-Engineered’ Nanocrystals in a Mass-Polymerized PMMA Matrix. *Nat. Photonics* 2014, 8, 392–399.

(53) Meinardi, F.; McDaniel, H.; Carulli, F.; Colombo, A.; Velizhanin, K. A.; Makarov, N. S.; Simonutti, R.; Klimov, V.; Brovelli, S. Highly Efficient Large-Area Colourless Luminescent Solar Concentrators Using Heavy-Metal-Free Colloidal Quantum Dots. *Nat. Nanotechnol.* 2015, 10, 878–885.

(54) Meinardi, F.; Akkerman, Q. A.; Bruni, F.; Park, S.; Mauri, M.; Dang, Z.; Manna, L.; Brovelli, S. Doped Halide Perovskite Nanocrystals for Reabsorption-Free Luminescent Solar Concentrators. *ACS Energy Lett.* 2017, 2 (10), 2368–2377.

(55) Liu, W.; Lin, Q.; Li, H.; Wu, K.; Robel, I.; Pietyoga, J. M.; Klimov, V. I. Mn2+-Doped Lead Halide Perovskite Nanocrystals with Dual-Color Emission Controlled by Halide Content. *J. Am. Chem. Soc.* 2016, 138 (45), 14954–14961.

(56) Das Adhikari, S.; Guria, A. K.; Pradhan, N. Insights of Doping and the Photoluminescence Properties of Mn-Doped Perovskite Nanocrystals. *J. Phys. Chem. Lett.* 2019, 10 (9), 2250–2257.

(57) Guria, A. K.; Dutta, S. K.; Adhikari, S. Das; Pradhan, N. Doping Mn2+ in Lead Halide Perovskite Nanocrystals: Successes and Challenges. *ACS Energy Lett.* 2017, 2 (5), 1014–1021.

(58) Huang, G.; Wang, C.; Xu, S.; Zong, S.; Lu, J.; Wang, Z.; Lu, C.; Cui, Y. Postsynthetic Doping of MnCl₂ Molecules into Preformed CsPbBr₃ Perovskite Nanocrystals via a Halide Exchange-Driven Cation Exchange. *Adv. Mater.* 2017, 29 (29), 1700095.

(59) Mir, W. J.; Jagadeeswararao, M.; Das, S.; Nag, A. Colloidal Mn-Doped Cesium Lead Halide Perovskite Nanoplatelets. *ACS Energy Lett.* 2017, 2 (3), S37–S43.

(60) Ji, S.; Yuan, X.; Cao, S.; Ji, W.; Zhang, H.; Wang, Y.; Li, H.; Zhao, J.; Zou, B. Near-Unity Red Mn2+ Photoluminescence Quantum Yield of Doped CsPbCl3 Nanocrystals with Cd Incorporation. *J. Phys. Chem. Lett.* 2020, 11 (6), 2142–2149.

(61) Das Adhikari, S.; Behera, R. K.; Bera, S.; Pradhan, N. Presence of Metal Chloride for Minimizing the Halide Deficiency and Maximizing the Doping Efficiency in Mn(II)-Doped CsPbCl₃ Nanocrystals. *J. Phys. Chem. Lett.* 2019, 10 (7), 1530–1536.

(62) Paul, S.; Bladt, E.; Richter, A. F.; Döblinger, M.; Tong, Y.; Huang, H.; Dey, A.; Bals, S.; Debnath, T.; Polavarapu, L.; Feldmann, J. Manganese-Doping-Induced Quantum Confinement within Host Perovskite Nanocrystals through Ruddlesden–Popper Defects. *Angew. Chem., Int. Ed.* 2020, 59 (17), 6794–6799.

(63) Krieg, F.; Ochsenbein, S. T.; Yakunin, S.; ten Brinck, S.; Aellen, P.; Siess, A.; Clerc, B.; Guggisberg, D.; Nazarenko, O.; Shynkarenko, Y.; Kumar, S.; Shih, C.-J.; Infante, I.; Kovalenko, M. V. Colloidal CsPbX₃ (X = Cl, Br, I) Nanocrystals 2.0: Zwitterionic Capping Ligands for Improved Durability and Stability. *ACS Energy Lett.* 2018, 3 (3), 641–646.

(64) Xu, K.; Meijerink, A. Tuning Exciton–Mn2+ Energy Transfer in Mixed Halide Perovskite Nanocrystals. *Chem. Mater.* 2018, 30 (15), 5346–5352.

(65) De Siena, M. C.; Sommer, D. E.; Creutz, S. E.; Dunham, S. T.; Gamelin, D. R. Spinodal Decomposition During Anion Exchange in Colloidal Mn2+ -Doped CsPbX₃ Nanocrystals. *Chem. Mater.* 2019, 31 (18), 7711–7722.

(66) Liu, H.; Wa, Z.; Shao, J.; Yao, D.; Gao, H.; Liu, Y.; Yu, W.; Zhang, H.; Yang, B. CsPbₓBr₁₋ₓ Nanocrystals with High Mn Substitution Ratio. *ACS Nano* 2017, 11 (2), 2239–2247.

(67) Mahamuni, S.; Lad, A. D.; Patole, S. Photoluminescence Properties of Manganese-Doped Zinc Selenide Nanoparticles. *J. Phys. Chem. C* 2008, 112 (7), 2271–2277.

(68) Song, P.; Qiao, B.; Song, D.; Cao, J.; Shen, Z.; Xu, Z.; Zhao, S.; Wagh, A.; Al-Ghamdi, A. Modifying the Crystal Field of CsPbCl₃ 3-Mn2+ Nanocrystals by Co-Doping to Enhance Its Red Emission by a Hundredfold. *ACS Appl. Mater. Interfaces* 2020, 12 (27), 30711–30719.

(69) Cottrell, T. L. The Strengths of Chemical Bonds, 2nd ed.; Butterworth: London, 1958.

(70) Lehmann, E. H.; Vontobel, P.; Wielz, L. Properties of the Radiography Facility NEUTRA at SINQ and Its Potential for Use as European Reference Facility. *Nondestr. Test. Eval.* 2001, 16 (2–6), 191–202.

(71) Morad, V.; McCall, K. M.; Sakhatskyi, K.; Lehmann, E.; Walford, B.; Losko, A. S.; Tritt, P.; Strobl, M.; Yakunin, S.; Kovalenko, M. V. Luminescent Lead Halide Ionic Liquids for High-Spatial-Resolution Fast Neutron Imaging. *ACS Photonics* 2021, DOI: 10.1021/acsphotonics.1c01348.