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Innovations in scintillator materials for X-ray detection

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Since their introduction in the early 20th century, scintillators have become essential components of a wide range of applications, including high-energy physics, medical imaging, cryptography, and nuclear detection. As the demand for high-performance scintillating materials continues to rise in particle physics experiments and medical imaging technologies, the development of novel scintillator materials has become a critical area of research. In recent years, advancements in scintillators have flourished, presenting new opportunities for practical applications. This review presents a comprehensive overview of standard performance parameters for scintillators, aimed at enhancing our understanding and evaluating their advancements. Unlike previous reviews focusing on isolated material categories, this work provides a cross-comparative analysis of emerging scintillators, with particular emphasis on challenges for high-precision detection and low-dose imaging. We highlight the latest developments in scintillator materials, emphasizing research from the past three years and focusing on their intrinsic properties. Our analysis covers the perovskite scintillators, nanocluster scintillators, rare-earth ion-doped scintillators, organic scintillators, and scintillators with specialized structures. This classification offers a scientific perspective on the overall progress in the field of scintillators, and several forward-looking insights into the future development of scintillators are proposed, employing a problem-oriented approach. Future scintillator development requires synergistic material design integrating computational modeling and scalable fabrication techniques to enhance stability, radiation tolerance, and light yields. Prioritizing lead-free systems and defect-tolerant lattice engineering will address environmental and operational challenges, and advancements in hybrid architectures, and novel optical structures promise breakthroughs in low-dose imaging, industrial nondestructive testing and sustainable radiation detection technologies. Eventually, we discuss the challenges encountered in scintillator development, explore future prospects, and provide valuable insights for improving their performances and expanding their applications.

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1. Introduction

X-rays are a form of electromagnetic radiation with exceedingly short wavelengths, typically ranging from 0.01 to 10 nm. Their remarkable penetrating power and high energy enable them to traverse various materials, including soft tissues in the human body and denser substances. Due to their exceptional capacity for penetration and imaging, X-rays find extensive application across multiple domains, such as medical imaging, industrial non-destructive testing, encryption and security screenings.^{1–5} Since the groundbreaking discovery of X-rays by Wilhelm Röntgen in 1895,⁶ this technology has rapidly evolved and made significant strides, gradually becoming an indispensable part of modern science and technology. Röntgen's discovery

not only spurred in-depth investigations into the properties and behavior of X-rays but also ushered in a new era in imaging science, providing a novel tool for medical diagnosis.⁷ Through modalities such as X-ray films, computed tomography (CT), and digital imaging, physicians can observe the internal structures and lesions of patients with remarkable clarity, vastly improving the quality and efficacy of medical services. Consequently, the development of materials capable of detecting X-rays at low radiation doses is crucial.⁸ Furthermore, this technology adeptly identifies welding defects, internal cracks, and other structural issues, ensuring product quality and operational safety.

X-ray detection can be categorized into direct and indirect methods based on different detection principles (Fig. 1). Direct detection involves the direct absorption of X-rays, which are then converted into electronic or biochemical signals.⁹ In contrast, indirect detection involves the conversion of X-rays into visible light or ultraviolet light using scintillation materials, followed by detection with optical devices such as

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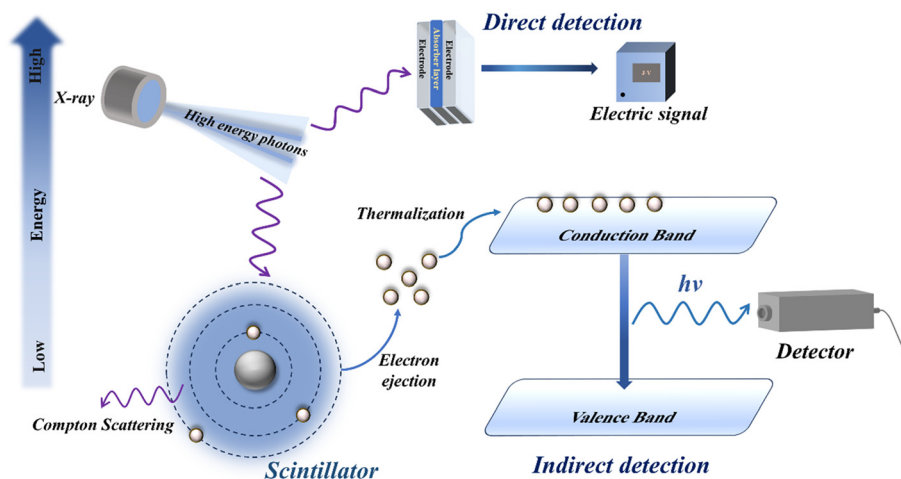


Fig. 1 Schematic diagram of the principles of direct and indirect X-ray detection.

photomultiplier tubes, charge-coupled devices, or complementary metal–oxide–semiconductor detectors.¹⁰ Indirect detection is generally more cost-effective than direct detection, and the availability of various scintillators allows for diverse applications, making them increasingly popular. Practically, take biological X-ray imaging as an example, X-rays pass through the biological specimen and create an image on the scintillator film. Then the scintillator film converts the high-energy X-ray image into a signal that can be detected by conventional imaging devices. Subsequently, this signal is processed by the corresponding software and transformed into an image that is recognizable by the human eye. Therefore, high-quality scintillator films are crucial in the application process.

Scintillators are indispensable components of indirect detection. When high-energy particles or electromagnetic radiation traverse the scintillators, they interact with the atoms within the material, exciting them to excited states. As these atoms return to their ground states, they release energy in the form of photons, emitting visible light or radiation of other wavelengths. After a long period of development, numerous traditional scintillators, characterized by their high light yield (LY) and exceptional energy resolution, have achieved commercial viability for their respective applications. For instance, due to their exceptionally high LYs, NaI:Tl and CsI:Tl have been utilized in medical imaging, X-ray detection, and medical radiography.^{11–13} Bi₄Ge₃O₁₂ (BGO) is widely accepted as a commercial-applicable inorganic scintillator due to its mechanically stable oxide with proportional performance.^{14,15} Moreover, single-crystalline Lu_{1.8}Y_{0.2}SiO₅:Ce is employed in positron emission tomography (PET), while GdWO₄ and Gd₂O₂S:Tb are used in CT.^{16–18} Additionally, Y₃Al₅O₁₂:Ce and BaF₂ are utilized in high energy physics experiments.^{19,20} Despite their widespread application, these traditional scintillators still suffer from some drawbacks. The preparation of traditional scintillator materials typically entails a complex process, high temperatures, and the use of capital-intensive equipment, resulting in considerable costs.^{16,21,22}

Furthermore, due to their intrinsic characteristics, these materials inevitably experience interference from a long after-glow, light scattering, and relatively low detection limit.²³ Additionally, their inflexible nature and biotoxicity limit their potential for advancement. Moreover, the challenges in converting the radiation emission wavelength into the visible light spectrum further hinder their applicability.²⁴

Recently, the rapid advancement of scintillator materials, driven by extensive and in-depth research, has captivated more researchers each year. Among these, metal halides have garnered significant attention due to their intriguing luminescence properties and high LYs. Since Chen *et al.* utilized fully inorganic perovskite nanocrystals as scintillators in 2018,²⁴ achieving strong X-ray absorption, intense radioluminescence (RL) at visible wavelengths, flexibility, and ultra-low detection limits, the development of metal halides has progressed by leaps and bounds. Accordingly, many studies have comprehensively summarized recent research advancements in scintillator materials. For instance, in 2023, Wang *et al.* summarized strategies for enhancing the LY of halide perovskite scintillators and presented a roadmap for improving X-ray imaging performance.²⁵ Zhou *et al.* summarized the latest advancements in low-dimensional metal halide scintillators and proposed future development directions.²⁶ Some studies have predominantly concentrated on specific categories of scintillator materials. Fan *et al.* focused on reviewing the progress of organic–inorganic hybrid Mn(II) halide scintillators²⁷ and Zhao *et al.* summarized advanced techniques for preparing rare-earth nanocrystalline scintillators and provided essential guidance for their application in biomedical research.²⁸ Besides, rare-earth-doped nano-scintillators have emerged as prominent scintillators in the field of biomedical applications. For example, Hong *et al.* presented a summary of the latest developments in advanced X-ray luminescence, focusing on its applications in imaging, biosensing, theragnostics, and neuromodulation through optogenetics.⁸ In 2024, Md. Helal Miah *et al.* summarized advancements in X-ray detection technologies across various materials, with a focus on meth-

ologies, structural designs, and device architectures. Notably, the review provides crucial insights into advances in perovskite materials for X-ray detection and imaging applications, emphasizing strategies to optimize efficiency, stability, and scalability for next-generation radiation sensing systems. To conclude, the development of scintillators is both extensive and profound. However, there is a noticeable absence of a comprehensive return to the development of advanced scintillator materials.

In this review, we initially summarize a set of standard performance parameters for scintillators to facilitate a deeper understanding and evaluation of their development. We update the latest developments in scintillator materials based on their inherent properties, with a particular focus on research from the past three years. Then we focus on scintillators containing lead and lead-free scintillators, Cu-based clusters, rare-earth-ion-doped scintillators, organic scintillators as well as certain specialized structures. Our classification provides a scientific overview of the overall development of scintillators, aiding researchers in gaining a comprehensive understanding of the latest advancements in the field. Ultimately, this review also addresses the challenges faced in the development of scintillators, analyzes the prospects for their future advancement, and offers insightful visions for enhancing the performance of scintillators and their applications. Unlike prior reviews that adopt siloed discussions of conventional material systems, this study pioneers a systematic cross-disciplinary assessment of next-generation scintillators, encompassing metal halide perovskites, copper-based nanoclusters, rare-earth nanocrystallites and triplet-harvesting organic matrices. By critically evaluating their interdependent photophysical limitations, such as ion migration in perovskites, environmental instability in Cu-based architectures, and triplet exciton quenching in organic systems, we have a clear and comprehensive understanding of the advantages and disadvantages of each material in the new generation of scintillators.

2. Principle

In this section, we focus on the intricate details of both material and device characteristics, with particular emphasis on the interdependence between them. The performance and features of the device are inevitably shaped by inherent properties of the materials employed. This comprehensive exploration aims to elucidate the complex relationship between material characteristics and the corresponding performance metrics of the device, thereby providing the reader with a deeper understanding of the subsequent content.

2.1 Light yield and overall conversion efficiency for X-rays

The LY represents the ability of a scintillator to convert high-energy rays into low-energy visible light, which can be expressed as follows:

$$LY = 1\,000\,000 \frac{SQ}{\beta E_g} \quad (1)$$

where S represents the transmission efficiency of electron-hole pairs to the optical emission centers, Q denotes the photoluminescence (PL) efficiency, β is a constant typically valued at 2.5, and E_g is the energy band gap. In scintillators, the standard time gate is usually set between 100 ns and 10 μ s, and the exact value is determined through practical experimentation. A high-energy LY is essential for scintillators. Moreover, the overall conversion efficiency of X-rays in the scintillator can be defined as the number of UV/visible photons (N_{ph}) generated during the scintillation conversion process per unit energy (E) of an incoming X-ray photon:

$$N_{ph} = \frac{E/\beta E_g}{SQ} \quad (2)$$

2.2 Attenuation coefficient

The attenuation coefficient (AC) indicates how effectively the detection material can absorb high-energy rays. It has a positive correlation with both its density ρ and effective atomic number Z_{eff} . The attenuation coefficient is directly proportional to these factors and can be expressed as:

$$AC = \frac{\rho Z_{eff}^4}{AE^2} \quad (3)$$

where A is the atomic mass and E is the radiation energy. This equation indicates that heavy elements and a high crystal density prevail in scintillators.

2.3 Afterglow

Afterglow refers to the time required for the intensity of RL in scintillators to diminish to a certain level after X-ray exposure. Due to the superimposition of previously exposed signals, prolonged afterglow can result in imaging artifacts. Therefore, a shorter afterglow duration is more desirable for common occasions. Yet along with the extension of scintillator applications, a long afterglow can be welcomed in some particular situations, like imaging scenes with memory properties, which are discussed in the following sections.

2.4 Spatial resolution

Spatial resolution pertains to the capacity to discern adjacent details within an object and their relative clarity. Specifically, higher spatial resolution is indicative of superior image quality. In scintillators, spatial resolution is primarily influenced by the geometric shape and intrinsic morphology of the scintillator itself, including its shape, size, and thickness. It has been widely accepted that the modulation transfer function (MTF), equivalent to 0.2, is derived from X-ray imaging to characterize spatial resolution. This also reflects the limitations of human visual discernment, typically measured in line pairs per millimeter (lp mm⁻¹). It is important to note that the thickness of the scintillator significantly impacts imaging resolution. Thus, for discussions concerning scintillators in studies, it is essential to address both spatial resolution and thickness concurrently.

2.5 Linear response of scintillators and matching with RL and the photodetector's spectrum

RL refers to visible or near-visible light emitted by a scintillator under the excitation of ionizing radiation. For the RL of scintillators, the radiation intensity must have a linear relationship with the dose of X-rays. Moreover, the wavelength of the RL needs to match the efficient operational wavelength region of the detector. It is also very important to select an appropriate detector for different scintillators.

2.6 Detection limit

The detection limit is defined as the equivalent dose rate that generates a signal three times that of the noise level. The captured signal should be recognized as valid only when the incident X-ray dose exceeds this limit. To lower this threshold and enhance sensitivity, it is crucial to minimize the dark current and improve the signal-to-noise ratio. In medical imaging, a smaller detection limit reduces the exposure to X-rays and thereby minimizes the potential harm to biological tissues.

2.7 Chemical stability and radiation resistance

Chemical stability is a crucial indicator of a scintillator's ability to maintain its performance under various environ-

mental conditions, including moisture resistance, temperature fluctuations, and oxidation resistance. For instance, CsI and NaI:Tl may undergo degradation in high-humidity environments, resulting in decreased luminescence efficiency. As for the radiation resistance, it is a key factor when assessing the ability of scintillator materials to preserve their performance upon exposure to high-energy radiation. Under exposure to high doses of X-rays, the lattice structure of scintillator materials may sustain damage, leading to a reduction in light output and a decline in energy resolution. Enhancing both chemical stability and radiation resistance can significantly improve the performance of X-ray scintillators. This enables them to maintain excellent detection capabilities and application efficacy in various harsh environments.

3. Technology innovations

This section provides a categorized overview of commonly used scintillator materials, highlighting their types, characteristics, recent representative works, and potential issues (Fig. 2). First, we introduce lead-based perovskite scintillators, which have been the subject of extensive research due to the strong X-ray absorption properties of Pb. This category

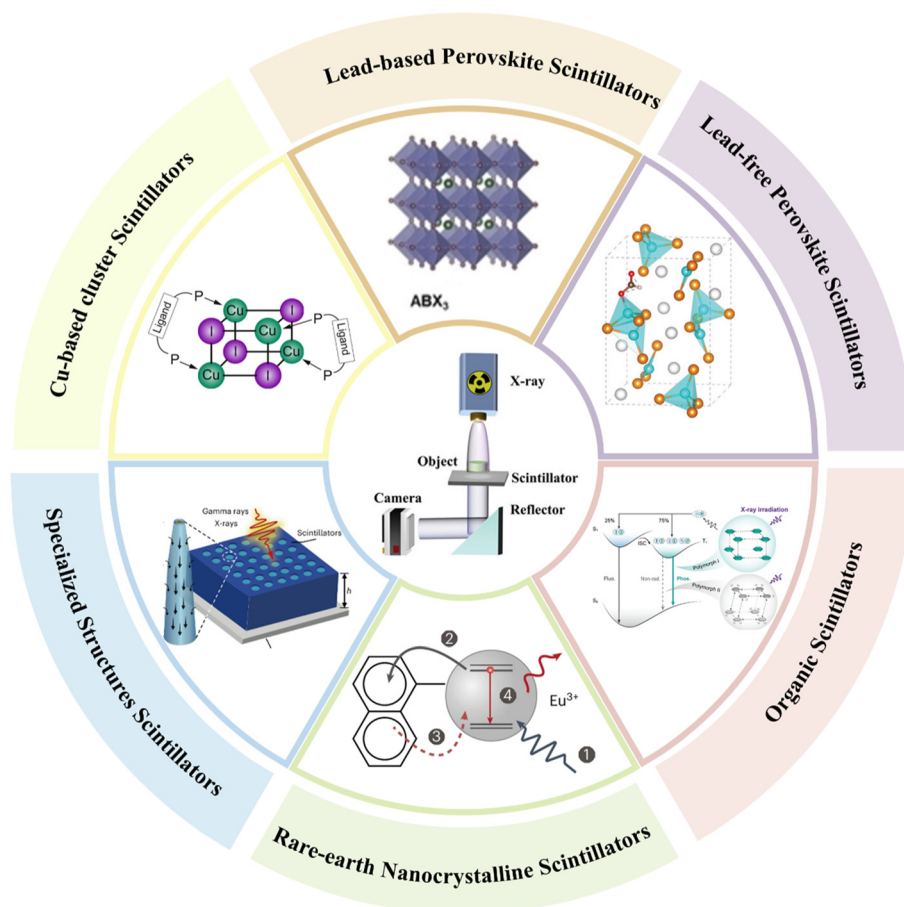


Fig. 2 Brief overview of this review.

remains active, with new studies continuously emerging. Next, we focus on lead-free scintillators, which have gained significant attention in recent years due to environmental and sustainable development concerns. Among these, copper-based scintillators are considered an excellent alternative to Pb and represent the main category within lead-free materials, as discussed in section 3.2.2. Additionally, there has been increasing interest in emerging copper-based nanoclusters, which are also being explored as scintillator materials, due to their exceptional optical properties. Rare earth elements, commonly used in optical materials, have attracted significant attention for their ability to regulate energy levels, with numerous applications in X-ray scintillators, which we address in detail in section 3.4. Furthermore, organic materials, due to their controllability and structural diversity, have also been extensively researched within the field of scintillators. Lastly, we present an overview of special structures in the scintillator domain, incorporating recent advancements and relevant studies.

3.1 Lead-based perovskite scintillators

Benefiting from adjustable bandgaps, high spatial resolution, fast response times, high LYs and considerable X-ray attenuation,^{24,29–32} lead-based perovskite scintillators have consistently been a major focus of research efforts and many valuable efforts have explored these charming materials (Table 1).^{24,33–52} Nevertheless, they still suffer from several drawbacks. The instability of a perovskite is always a non-negligible issue because of its hygroscopicity. Water and oxygen expand perovskite molecules, causing aggregation and affecting their optical properties, which will even be prominent under irradiation with X-rays,⁵³ and the typical method for improving a perovskite's stability is to passivate its surface or reduce inner defects. Favorable and accepted methods are to embed them in solid substances or encapsulate them with polymers like polyethylene glycol terephthalate (PET), polytetrafluoroethylene (PTFE) and poly(methyl methacrylate) (PMMA).⁵⁴

Based on similar methodologies, some groups have already completed related work,^{55,56} and in the past two years more noteworthy studies have been reported. Lv *et al.* combined CsPbBr₃ nanocrystals with PMMA to fabricate a perovskite film of over 100 cm² with a root-mean-square roughness of 1.06 nm, which retained 96% initial luminous intensity in water and an ambient environment for 480 h and 2880 h

(Fig. 3a).⁵⁷ Zheng *et al.* selected poly(maleic anhydride-*alt*-1-octadecene) (PMAO) with a low molecular weight ($M_n = 1911$) as the multi-dentate ligands and synthesized PMAO–CsPbBr₃ lead perovskite nanocrystals (PNCs) *via* hot injection. It shows preminent water stability, maintaining its fluorescence for 2600 min while traditional PNCs became completely dysfunctional after 100 min, as observed in Fig. 3b.⁵⁸ For the embedded-in-matrix strategy, Shen *et al.* applied cold-sintered technology to bulk scintillators to fabricate CsPbBr₃ nanocrystals embedded in the Cs₄PbBr₆ structure based on an “emitter-in-matrix” principle (Fig. 3c).⁵⁹ This group employed a rather low temperature of 90 °C during the sintering process and DMSO was introduced as the antisolvent to promote its densification (Fig. 3d). Compared to solid-state sintering at high temperature, cold sintering demonstrates highly compacted grains with enhanced densification and transparency, thus scattering can be well controlled. These bulk scintillators exhibit a high LY of 33 800 photons per MeV, a low detection limit of 79 nGy_{air} s⁻¹ and outstanding spatial resolution of 8.9 lp mm⁻¹.

Apart from those tactics mentioned for enhancing perovskite stability, some other interesting methods have appeared. Chen *et al.* found it detrimental to cross-linking of the organosilicon matrix that the silicon-based CsPbBr₃ encapsulated nanocrystals were linked with long-carbon-chain ligands, thus suffering from poor transparency and surface roughness.⁶⁰ Yet this can be improved by substituting long chain ligands for short ones (Fig. 3e). Then a dual-organosilicon ligand system was proposed, consisting of (3-aminopropyl)triethoxysilane (APTES) and (3-aminopropyl)-triethoxysilane with pentanedioic anhydride (APTES-PA), and the scintillator showed a satisfying spatial resolution above 20 lp mm⁻¹ and high stability in various harsh environments. Wu *et al.* applied the host-guest strategy, involving confining MAPbBr₃ nanocrystals in metal-organic frameworks (MOFs) (Fig. 3f).⁶¹ The *in situ* spatial confinement provides quantum confinement and surface passivation, resulting in significant stability against air, heat and irradiation. It exhibits a detection limit of 170 nGy_{air} s⁻¹ and an impressive spatial resolution of 14.7 lp mm⁻¹. Likewise, Li *et al.* presented a glass-ceramic CsPbBr₃ perovskite film with high transparency, in which CsPbBr₃ crystals grew *in situ*.⁶² The thickness was controlled by the crystal coordination-topology growth and *in situ* film formation strategy, resulting in little light scattering and a uniform distribution of the perovs-

Table 1 The parameters of lead-based scintillators

Material	Type	Light yield (photons per MeV)	Detection limit (nGy _{air} s ⁻¹)	Spatial resolution (lp mm ⁻¹)	Ref.
CsPbBr ₃	Nanocrystals	9001	—	9	57
CsPbBr ₃	Nanocrystals	—	—	—	58
CsPbBr ₃ /Cs ₄ PbBr ₆	Nanocrystals	33 800	79	8.9	59
CsPbBr ₃	Nanocrystals	—	—	20	60
MAPbBr ₃	MOFs	—	170	14.7	61
CsPbBr ₃	Nanocrystals	—	326	13.9	62
MAPbMgZnCdBr ₃	Nanocrystals	21 200	462	11.7	63
CsPb ₂ Br ₅ :K ⁺	Nanocrystals	25 160	162.3	21	64

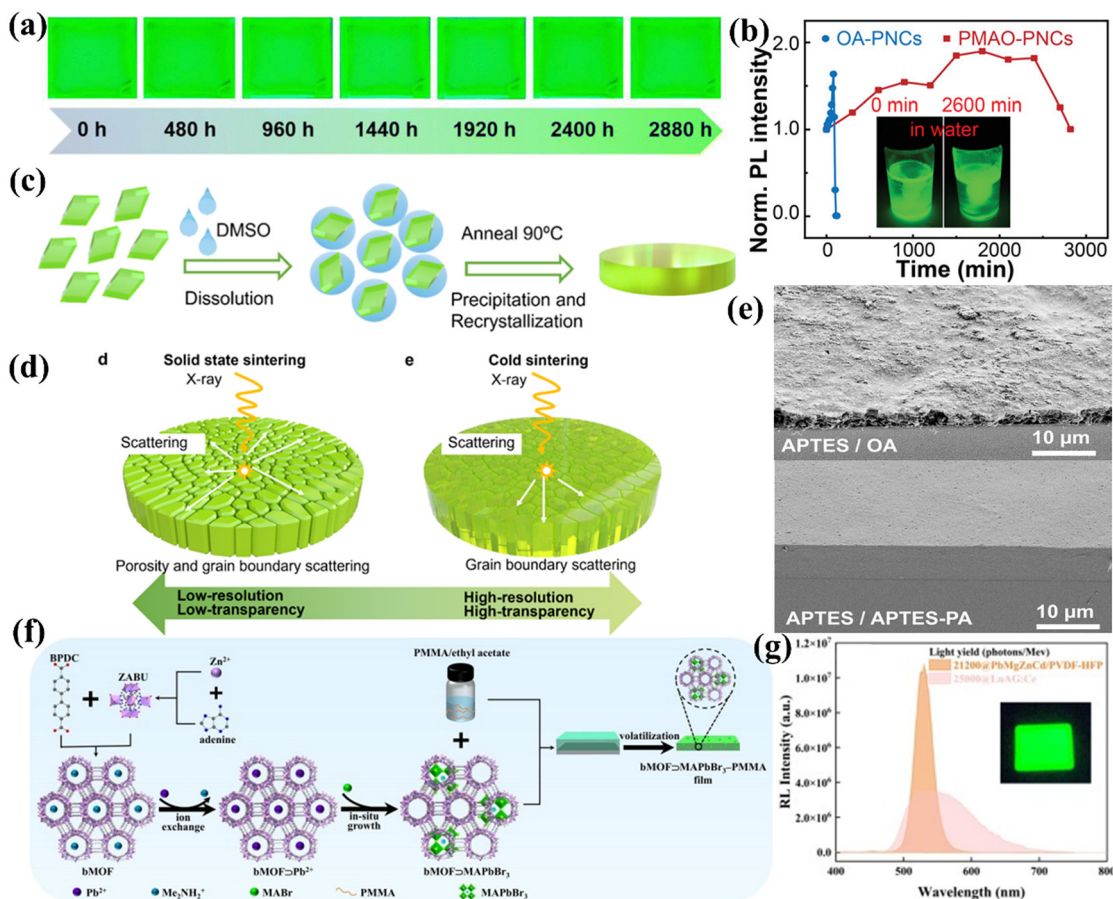


Fig. 3 (a) Photographs of the $\text{CsPbBr}_3\text{@PMMA@VmB1}$ film in air at various times to show its stability. (b) PL evolution of PNCs after being immersed in water for different times (inset: optical images of the PMAO-PNCs in water under vigorous stirring). (c) Design of the fabrication process to obtain bulk $\text{CsPbBr}_3/\text{Cs}_4\text{PbBr}_6$ ceramic composite pellets *via* a CSP. (d) Light scattering of residual pores and grain boundaries in conventional solid-state sintered ceramics, reducing energy resolution and transparency (left). Reduction of light scattering from grain boundaries in cold-sintered ceramics to enhance resolution and transparency (right). (e) SEM images of the PNC/silicone films with APTES/OA ligands (upper) and APTES/APTES-PA ligands (bottom). (f) Schematic illustration of the synthesis of bMOF-MAPbBr_3 and $\text{bMOF-MAPbBr}_3\text{-PMMA}$ films. (g) Comparison of LY between the $\text{MAPbMgZnCdBr}_3/\text{PVDF-HFP}$ scintillator and LuAG:Ce .

kite crystals. The as synthesized $250\ \mu\text{m}$ film enables a detection limit of $326\ \text{nGy}_{\text{air}}\ \text{s}^{-1}$ and a high spatial resolution of $13.9\ \text{lp}\ \text{mm}^{-1}$ with incredible stability under exposure to long-term X-ray irradiation, water soaking and high temperature.

Besides the stability improvements, progress was also made in the energy utilization efficiency by doping with new elements. Xiang *et al.* performed calculations on the surface formation energy and the vacancy defect energy and co-doped several bivalent metal ions to substitute Pb in MAPbBr_3 to improve the photoluminescence quantum yield (PLQY) and decay time of the nanocrystal.⁶³ Furthermore, the scintillator was fabricated using MAPbMgZnCdBr_3 as the energy conversion layer with PVDF-HFP as the carrier and demonstrated a LY of 21 200 photons per MeV (Fig. 3g) with a spatial resolution of $11.7\ \text{lp}\ \text{mm}^{-1}$. Moreover, Qiu *et al.* doped non-emissive CsPb_2Br_5 with alkali metals, which caused lattice distortion and enhanced electron-phonon coupling.⁶⁴ This led to the formation of potential transient energy wells, resulting in the creation of self-trapped excitons (STEs) generated by X-rays. The

STE experiences radiative recombination, with a PLQY of 55.92%, thus X-rays can be transformed into red light. Furthermore, the K^+ -doped CsPb_2Br_5 based X-ray scintillator displays high stability and weak self-absorption with a detection limit of $162.3\ \text{nGy}_{\text{air}}\ \text{s}^{-1}$ and $21\ \text{lp}\ \text{mm}^{-1}$.

Despite being a long-investigated scintillator material, it still suffers from non-negligible drawbacks. The predominant emission mechanism *via* band-edge excitons introduces substantial self-absorption effects owing to spectral overlap between its emission and absorption profiles. This fundamental limitation persists as a pervasive challenge inherent to virtually all perovskite-based systems. While lead-based perovskite scintillators exhibit unparalleled performance metrics, the absence of innovative stabilization strategies remains a critical bottleneck. No groundbreaking processing techniques addressing environmental robustness have emerged in the latest literature, which confines their practical deployment to laboratory-scale demonstrations or strictly controlled environments, as evidenced by rapid performance degradation. In addition,

the potential environmental leakage risk of heavy metals further hinders large-scale applications.

3.2 Lead-free perovskite scintillators

Although Pb based scintillators exhibit promising absorption of X-rays, the heavy atom's intrinsic toxicity, doing harm to human health and environment, is still a non-negligible hindrance to its extensive applications. Therefore, the pursuit of 'lead-free' scintillators has been put forward, expecting materials with no toxic element yet retaining exceptional and controllable properties. In recent years, research on lead-free scintillators has significantly outpaced that on traditional lead-based scintillators, especially in the past two years (Table 2).^{65–67} These materials can be categorized into the following types: double perovskite scintillators, manganese-based halide scintillators and others; these are all introduced and discussed in the following sections.

3.2.1 Copper-free scintillators. One of the important forms of copper-free scintillator materials is the double perovskite material. Their development stemmed from the recognition that, while perovskite materials have long been a focus of scintillator research due to the advantages in their synthesis, the isolated network of metal-halide octahedral units leads to poor carrier mobility and predominantly indirect bandgap

characteristics.^{68–70} In contrast, the double perovskite materials, typically represented as $A_2B_3X_6$ (where A is a monovalent metal ion like Ag^+ and Cs^+ , B is a trivalent metal ion like In^{3+} and Sb^{3+} , X refers to a halide anion), though suffering from lower carrier mobility and a larger indirect bandgap compared to the perovskites, possess a cubic structure with interlinked $[B^+X_6]^{5-}$ and $[B^+X_6]^{3-}$ sites and do not contain toxic Pb elements.^{69,71–73} This objectively enhances the material's stability and environmental friendliness, making it an attractive alternative to perovskite materials for researchers. Efforts have been made to circumvent these drawbacks. For example, Varnakavi *et al.* managed to inject 4.04% of Mn(II) into a double perovskite $Cs_2NaBiCl_6$ nanocrystal (Fig. 4a), which turned out to have a PLQY of 10.6% at 586 nm, circumventing self-absorption.⁷⁴ The $Cs_2NaBiCl_6:4.04\%Mn@poly(methyl\ methacrylate)$ scintillator film demonstrates a satisfying LY of 28 350 photons per MeV and a detection limit of 45.2 nGy_{air} s⁻¹, along with a rather exceptional spatial resolution of 14.76 lp mm⁻¹. Recent advances in Bi³⁺-doped perovskite scintillators⁷⁵ highlight their critical role in performance enhancement. In 2025, Chen *et al.* engineered 3D bismuth chloride perovskites *via* monovalent Na⁺ incorporation, achieving a sensitivity of 354.5 mCGy⁻¹ cm⁻² and an ultralow detection limit of 59.4 nGy s⁻¹ with stable cycling over 4500 on-off cycles.⁷⁶

Table 2 The parameters of lead-free scintillators

Material	Type	Light yield (photons per MeV)	Detection limit (nGy _{air} s ⁻¹)	Spatial resolution (lp mm ⁻¹)	Ref.
Cs ₃ Cu ₂ I ₅ :Mn	Single-crystal	95 772	—	—	100
Rb ₂ AgI ₃ :Cu ⁺	Single-crystal	53 983	231	14.1	101
Cs ₃ Cu ₂ I ₅ :HCOO ⁻	Single crystals	61 500	—	10	102
Cs ₃ Cu ₂ I ₅	0D	—	12 760 000	10	103
Cs ₃ Cu ₂ I ₅	Nanocrystals	—	105	14.3	104
Cs ₃ Cu ₂ I ₅	Nanocrystals	—	—	—	105
Cs ₃ Cu ₂ Cl ₅	Single crystal	95 000	2700	105	106
CsCu ₂ I ₃	Crystal	—	88.9	15.6	107
Cs ₅ Cu ₃ Cl ₆ I ₂	1D	64 800–67 200	—	27	108
Cs ₅ Cu ₃ Cl ₆ I ₂	1D	—	—	—	109
Cs ₅ Cu ₃ Cl ₆ I ₂ :Rb	Crystal	64 868	—	14	110
Rb ₂ AgBr ₃ :Cu ⁺	Single crystal	79 250	714.83	5.8	111
Rb ₂ AgI ₃ :Cu ⁺	1D single crystals	36 293	1022	10.2	112
Rb ₂ AgBr ₃ :Cu ⁺	Single-crystal	—	—	10.2	113
Cs ₂ NaBiCl ₆	Nanocrystals	28 350	45.2	14.76	74
Cs ₂ TeCl ₆	Microcrystals	38 523	258	15.9	66
Cs ₂ HfCl ₆	Microcrystals	21 700	55	11.2	67
Cs ₄ Cd _{1-x} Mn _x Bi ₂ Cl ₁₂	2D	34 450	183.6	16.7	77
(C ₂₅ H ₂₂ P) ₂ MnCl ₄	Single crystals	78 937	—	25.61	82
(C ₃₈ H ₃₄ P ₂)MnBr ₄	Crystal	35 945	5500	16.7	83
BPP ₂ MnBr ₄	0D	35 000	250	20	84
(MTP) ₂ MnBr ₄	Crystal	67 000	82.4	6.2	85
PrPP ₂ MnBr ₄	Single crystal	43 511	13.9	53.5	86
(Br-PrTPP) ₂ MnBr ₄	0D	68 000	45	12.78	87
(BTPP) ₂ MnX ₄	0D	53 000	89.9	14.1	88
A ₂ MnBr ₄	0D	80 100	30	14.06	89
DMA ₄ InCl ₇ :Sb ³⁺	0D	23 500	175	—	90
TTA ₂ In _{0.92} Sb _{0.08} Cl ₅	0D	60 976	90	—	91
TpyBiCl ₅	Single crystal	—	196.31	—	92
Ag ₆ S ₆ L ₆ (SC-Ag)	Metal cluster	17 420	—	16	93
Ag ₂ Cl ₂ (dppb) ₂	Single crystal	79 970	59.8	25	94
Cs ₂ Ag ₆ Na ₄ In _{0.85} Bi _{0.15} Cl ₆	Nanocrystals	—	—	—	95
Cs ₂ CdCl ₄ :10%Mn	Single crystal	88 138	31.04	16.1	96
Tb ³⁺ -Cs ₂ NaGdCl ₆	Single crystals	39 100	41.32	10.75	78
Te ⁴⁺ doping Cs ₂ SnCl ₆	Nanocrystals	—	132	20	79

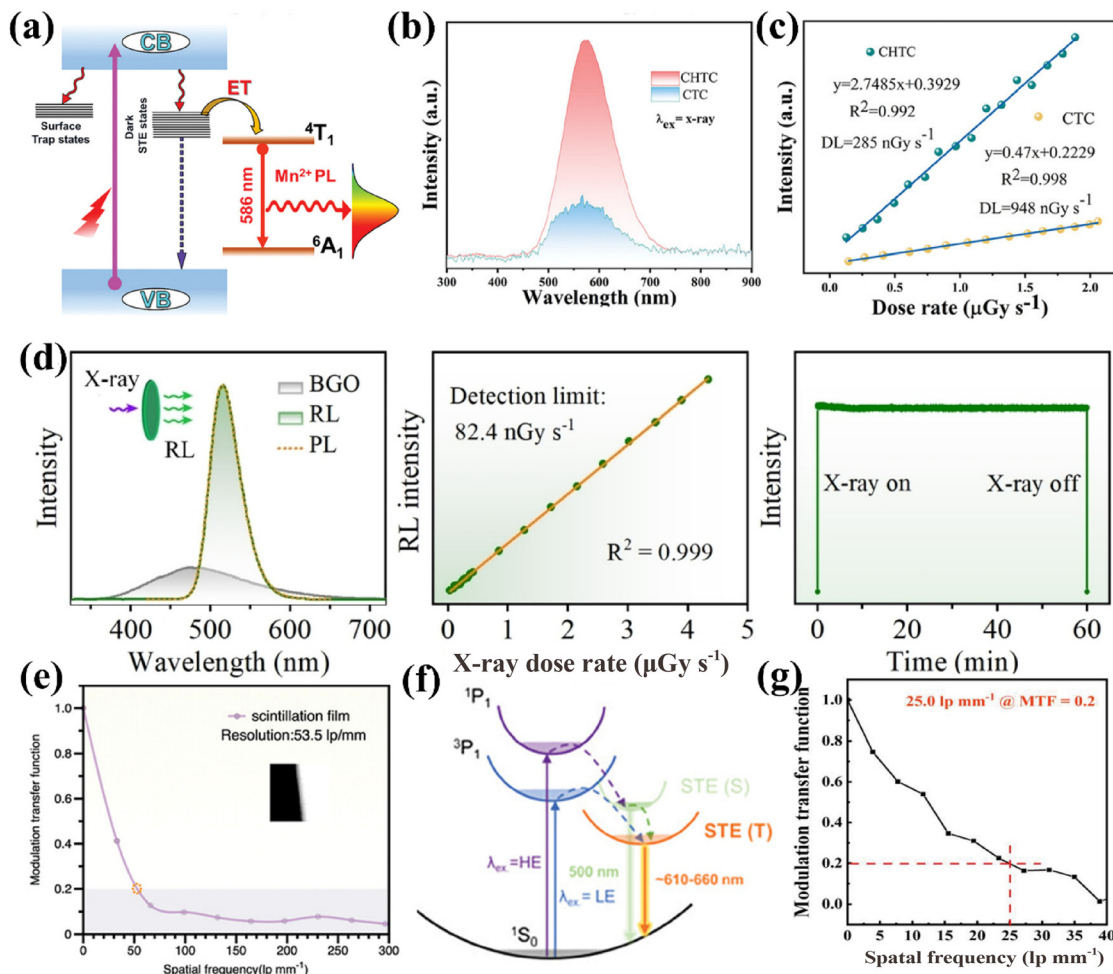


Fig. 4 (a) Schematic diagram of Mn^{2+} -doped $\text{Cs}_2\text{NaBiCl}_6$ showing STE dynamics and the Mn^{2+} PL mechanism. (b) RL of CHTC and CTC. (c) RL of CHTC and CTC as a function of dose rate. (d) RL spectra of $(\text{MTP})_2\text{MnBr}_4$ and BGO under X-ray irradiation (left). Integrated RL intensity of $(\text{MTP})_2\text{MnBr}_4$ as a function of X-ray dose rate (middle). Time-dependent RL intensity of $(\text{MTP})_2\text{MnBr}_4$ upon continuous X-ray irradiation at a dose rate of 42.29 mGy s^{-1} for 60 min (right). (e) The MTF of the $\text{PrPP}_2\text{MnBr}_4$ scintillation film beveled edge image. (f) Schematic diagram of the emission mechanism with STE (T) and STE (S). (g) The MTF of the $\text{Ag}_2\text{Cl}_2(\text{dppb})_2$ (Ag_1) scintillator edge images.

Complementing this, Subagyo *et al.* demonstrated the efficacy of Bi^{3+} in optimizing radiation hardness, LY, and defect dynamics, underscoring its potential for sustainable high-performance X-ray detectors. Together, these studies exemplify the versatility of Bi^{3+} in advancing perovskite scintillators for precision radiation sensing.⁶⁵ In parallel, Lai *et al.* discovered that the Te-based double perovskite, Cs_2TeCl_6 (CTC) microcrystal, showed a broadband yellow emission which was highly matched with CCD.⁶⁶ Furthermore, the hafnium alloyed CTC scintillator, with PDMS as the binder, demonstrates a LY of 38 523 photons per MeV (Fig. 4b), a detection limit of $258 \text{ nGy}_{\text{air}} \text{ s}^{-1}$ (Fig. 4c) and an outstanding spatial resolution of 15.9 lp mm^{-1} . Likewise, Zhang *et al.* came up with Cs_2HfCl_6 , and under ambient conditions for 60 days, there were no extra diffraction peaks observed in XRD spectra, indicating promising stability, which was possibly due to short Hf-Cl bonds, with a spatial resolution of 11.2 lp mm^{-1} .⁶⁷ Moreover, Li *et al.* reported a 2D layered double perovskite, namely $\text{Cs}_4\text{Cd}_{1-x}\text{Mn}_x\text{Bi}_2\text{Cl}_{12}$,

which demonstrated preminent scintillation properties, including exceptional emission response linearity and a high LY of $\sim 34\,450$ photons per MeV.⁷⁷ More admirably, the X-ray excited emission intensity remained at 92% and 94% respectively after being stored under an ambient atmosphere and exposed to a total dose of 11.4 Gy irradiation. After being mixed with PDMS, the scintillator screen exhibits an excellent spatial resolution of 16.7 lp mm^{-1} . Subsequently, Naresh *et al.* proposed the $\text{Cs}_2\text{NaGdCl}_6:5\%\text{Tb}^{3+}$ films with 0.4 mm thickness and achieved a low detection limit of $41.32 \text{ nGy}_{\text{air}} \text{ s}^{-1}$, an LY of 39 100 photons per MeV and excellent radiation stability.⁷⁸ After the introduction of Tb^{3+} , the PLQY increased from 8.4% to 72.6% due to efficient energy transfer from STE to Tb^{3+} . Likewise, Wang *et al.* introduced Te^{4+} into the double perovskite Cs_2SnCl_6 and fabricated flexible imaging screens with a resolution of 20 lp mm^{-1} and a low detection limit of $132 \text{ nGy}_{\text{air}} \text{ s}^{-1}$.⁷⁹

Manganese-based halides have only been utilized in the past five years. Due to their excellent LY and high energy radi-

ation absorption performance, they are designed to be potential candidates for efficient and environmentally friendly scintillator materials.^{80,81} Here we focus mainly on research published in the past year. For instance, Zhou *et al.* designed and synthesized the thickness-tunable inch-sized Mn-based halide single crystal, $(C_{25}H_{22}P)_2MnCl_4$, with a PLQY up to 99.10% and the scintillator film showed a LY of 78 937 photons per MeV and a spatial resolution of 25.61 lp mm⁻¹.⁸² Liu *et al.* synthesized the $(C_{38}H_{34}P_2)MnBr_4$ organogel scintillator and made it water-stable, stretchable and self-healing by carefully arranging and activating dynamic hydrogen bonds and coordination bonds.⁸³ The organogel scintillator is able to be stretched up to 13-fold while maintaining its property, and the imaging resolution reached 16.7 lp mm⁻¹. The scintillator film demonstrated seamless self-healing within 30 min, providing a robust solution for the self-repair of flexible films that may experience bending or cutting damage during usage. Wang *et al.* reported a method to control the crystallization process of manganese halides through organic cation modulation.⁸⁴ This approach aims to reduce uncontrolled crystallization during the fabrication of large-area flat-panel detectors, thereby enhancing their light-emitting properties. The large steric hindrance brought by the cation provides a high exciton binding energy and restrains non-radiative recombination. The as synthesized BPP_2MnBr_4 ($BPP = C_{25}H_{22}P^+$) exhibits a superior spatial resolution of more than 20 lp mm⁻¹, an exceptional LY of over 35 000 photons per MeV and an ultra-low detection limit of less than 250 nGy_{air} s⁻¹. Likewise, Zhang *et al.* reported the potential of applying $(MTP)_2MnBr_4$ (MTP refers to methyl-triphenylphosphonium) as an X-ray scintillator for its near-unity PLQY, high resistance to thermal quenching and decent stability.⁸⁵ They fabricated a thin film scintillator and determined its impressive LY of 67 000 photons per MeV and superior detection limit of 82.4 nGy s⁻¹ (Fig. 4d).

Moreover, Xiao *et al.* synthesized $PrPP_2MnBr_4$ ($PrPP_2$ refers to diisopropylammonium) by a dual-solvent evaporation method, and the 10 cm × 10 cm area scintillation screen based on the solidified melt salt exhibited a LY of 43 511 photons per MeV, an unprecedented high spatial resolution of 53.5 lp mm⁻¹ (Fig. 4e) and an outstanding detection limit of 13.9 nGy_{air} s⁻¹.⁸⁶ Also, Zhang *et al.*, recognizing the satisfying optical and scintillator properties of 0D organic-inorganic hybrid metal halides (OIMHs), synthesized $(Br-PrTPP)_2MnBr_4$ ($Br-PrTPP$ refers to (3-bromopropyl) triphenylphosphonium) *via* a facile saturated crystallization method.⁸⁷ Attributed to the tetrahedrally coordinated $[MnBr_4]^{2-}$ polyhedron, it demonstrates an excellent LY of up to 68 000 photons per MeV and lowest detection limit of 45 nGy_{air} s⁻¹. Li *et al.* prepared 0D $(BTPP)_2MnBr_4$ ($BTPP$ refers to benzyltriphenylphosphonium) as a promising scintillator with excellent air and radiation stability, which showed an appreciable LY of 53 000 photons per MeV, a low detection limit of 89.9 nGy_{air} s⁻¹ and an ultra-low afterglow comparable to the commercial BGO single crystal.⁸⁸ Furthermore, after fabricated as a flexible screen combined with PDMS, a rather high spatial resolution of 14.1 lp mm⁻¹ was achieved. In addition, Gong *et al.* presented a

novel group of 0D hybrid manganese halides of A_2MnBr_4 ($A = BzTPP, Br-BzTPP, \text{ and } F-BzTPP$).⁸⁹ Benefiting from near-unity PLQY, these hybrids show a state-of-art LY of 80 100 photons per MeV for hybrid manganese halides. The 30 nGy_{air} s⁻¹ ultra-low detection limit, 14.06 lp mm⁻¹ qualified spatial resolution and 0.3 ms short afterglow make it a promising scintillator for further applications.

Apart from all these copper-free scintillators mentioned, there are still several noteworthy studies we would like to present. For example, Wang *et al.* discovered the introduction of a small amount of Sb^{3+} into the indium halide DMA_4InCl_7 , resulting in $DMA_4InCl_7:10\%Sb^{3+}$, largely enhancing the PLQY by 9-fold, which originated from efficient energy transfer between Sb^{3+} ions and STE states.⁹⁰ The STE emission yields a high LY of 23 500 photons per MeV and a low detection limit of 175 nGy_{air} s⁻¹. Likewise, Huang *et al.* discovered the novel influence of the A-site on the band-edge structures and optoelectronic properties, and some Sb^{3+} -doped 0D halide perovskite derivatives were designed as experimental proof.⁹¹ Benefiting from the deformation of the lattice brought about by Sb^{3+} , STE states are harnessed properly (Fig. 4f), resulting in a qualified LY of up to 60 976 photons per MeV and a detection limit of 90 nGy_{air} s⁻¹. Moreover, Wang *et al.* designed and synthesized a molecular hybrid perovskite, $TpyBiCl_5$, which managed to circumvent thermal quenching *via* multi-excited state switching, originating from the inter-reaction among its counterparts in the perovskite.⁹² It also shows a rigid framework structure, resulting in 12 times higher PLQY than its organic ligand (Tpy) at room temperature. $TpyBiCl_5$ maintains an ideal detection limit of 196.31 nGy s⁻¹ even at 353 K through thermally activated delayed fluorescence (TADF), and a clear image at 413 K, effectively addressing potential annealing issues observed in other films. This characteristic offers a promising solution for the application of scintillators under extreme conditions, such as *in situ* imaging of thermal loss during mechanical operation. Furthermore, Wang *et al.* synthesized $Ag_6S_6L_6$ (SC-Ag) metal clusters through a simple solvothermal reaction.⁹³ The heavy atom guarantees an effective TADF mechanism, and satisfyingly, SC-Ag demonstrates a high PLQY of 91.6% and a LY of 17 420 photons per MeV, along with a preeminent spatial resolution of 16 lp mm⁻¹. Additionally, the rigid core structure endows SC-Ag with strong resistance to humidity and radiation. Additionally, Yuan *et al.* pioneered high-performance TADF Ag-based scintillators, namely $Ag_2Cl_2(dppb)_2$ ($dppb$ refers to 1,2-bis(diphenylphosphino)benzene), with an exceptionally high RL intensity and a low detection limit of 59.8 nGy s⁻¹.⁹⁴ Its superior performance, compared to the Cu series, mostly results from effective utilization of high excitons due to a small singlet-triplet energy gap. The flexible film fabricated has a high spatial resolution of 25 lp mm⁻¹ (Fig. 4g). Another work worth mentioning is that of O'Neill *et al.* who synthesized and characterized $Cs_2Ag_{0.6}Na_{0.4}In_{0.85}Bi_{0.15}Cl_6$ (CANIBIC), an inorganic mixed-cation double halide perovskite, and figured out its STE emission mechanism.⁹⁵ Moreover, the CANIBIC-PMMA film demonstrates higher X-ray luminescence intensity than that of

the pure pressed pellet of CANIBIC. Also, Wang *et al.* reported a Mn²⁺-activated all-inorganic 2D layered Ruddlesden–Popper perovskite, namely Cs₂CdCl₄:10%Mn.⁹⁶ It possesses a bright orange–red emission with 90.47% PLQY, strong X-ray absorption, an ultra-high LY up to 88 138 photons per MeV and 31.04 nGy_{air} s⁻¹. A 15 cm × 15 cm flexible screen was prepared by combining PDMS with Cs₂CdCl₄:10%, with a high spatial resolution of 16.1 lp mm⁻¹ and qualified imaging capability under an extreme low dose of 16 μGy_{air} s⁻¹.

While copper-free scintillators, as a subclass of their lead-free counterparts, enable elements with analogous elements, such as Mn, Bi, and Te, to better manifest their intrinsic properties for optimizing spectral characteristics and enhancing performance, critical challenges persist. These include the generation of toxic species from undesired byproducts during the synthesis of organic components in hybrid organic–inorganic systems, inherent parity-forbidden transitions in double perovskites necessitating additional strategies to improve the LY,⁹⁷ and the hygroscopic nature of lead-free variants imposing stringent encapsulation requirements that limit their practical deployment.⁹⁸

3.2.2 Copper-based scintillators. Cu substitution is the best accepted solution, and Cs₃Cu₂I₅ is a widely reported Cu-based scintillator with prominent feasibility, because the zero-dimensional (0D) structure of Cs₃Cu₂I₅ effectively localizes and segregates multiple STEs, thereby mitigating Auger recombination.⁹⁹ Several works have been reported on this appealing material;¹⁰⁰ here we focus only on work from recent years, for which researchers have focused their efforts on synthesis procedure progress or property enhancement. For instance, to avoid lattice distortion caused by heteromorphic homologs in vacuum-evaporated metal halides (MHs), Peng *et al.* brought out a single-source vacuum evaporation method.¹⁰¹ A ~10 μm thick film of Cu-MHs was synthesized by using this very strategy, to achieve an impressive LY of 53 983 photons per MeV and a spatial resolution of 14.1 lp mm⁻¹.

Doping is recognized as a potential method for enhancing efficiency and stability. For example, Wang *et al.* focused their efforts on multi-self-trapped excitons of HCOO⁻-doped Cs₃Cu₂I₅ single crystals with transient and steady-state spectroscopy, and figured out that it derived from the host lattice and external dopants separately, which could both be triggered independently.¹⁰² Then, after carefully tuning the excitation wavelength and selectively exciting different STEs, Cs₃Cu₂I₅:HCOO⁻ demonstrates a remarkable PLQY of 99.01% and little self-absorption, with LY being improved by 5.4 times, compared to that of pristine Cs₃Cu₂I₅ crystals, up to 61 500 photons per MeV (Fig. 5a). Also, like the lead-based perovskite scintillators, combination with poly-carriers is considered a way of enhancing stability. To illustrate this, Moon *et al.* reported novel technology that integrated Cs₃Cu₂I₅ perovskite nanoparticles into densified–delignified wood, namely Cs₃Cu₂I₅@D-Wood, for an eco-friendly and biodegradable X-ray scintillator film with a spatial resolution of 10 lp mm⁻¹.¹⁰³ Similarly, Hao *et al.* developed an innovative integration of Cs₃Cu₂I₅ with PVDF and PMMA, effectively prevent-

ing agglomeration and ensuring a more uniform size distribution, which, in turn, improved stability.¹⁰⁴ The as controlled flexible scintillator film displays a satisfying spatial resolution of 14.3 lp mm⁻¹ and a low detection limit of 105 nGy s⁻¹ was achieved.

To further exert the functionality of Cs₃Cu₂I₅ and its derivatives, some praiseworthy investigations have been conducted. For instance, considering that the absence of X-ray energy distinction shall result in insufficient image contrast, Ran *et al.* presented an innovative multi-energy X-ray linear-array detector based on Cs₃Cu₂I₅ (Fig. 5b).¹⁰⁵ Benefiting from the negligible self-absorption of the material and side-illuminated scintillation scenarios, the incident X-ray spectra could be reconstructed by analyzing the distribution of the scintillator intensity while illuminated from the side. As a result, the outcome error can be controlled to be below 5.63%. Xiang *et al.* presented centimeter-sized Cs₃Cu₂Cl₅ *via* a slow-cooling method, and the planar orientation, which was controlled in a space-confined chamber, required no further shaping before being fabricated as a scintillation screen.¹⁰⁶ The synthesized Cs₃Cu₂Cl₅ single crystal exhibits an impressive LY of up to 95 000 photons per MeV and a low detection limit of 2.7 μGy_{air} s⁻¹. Moreover, it could retain its original phase for 6 months in a vial after the surface passivation procedure. Intriguingly, Ran *et al.*, the same group mentioned above, also discovered a better match between CsCu₂I₃, once considered the oxidation product, and the spectral responsivity of regular flat-panel photodiode arrays than Cs₃Cu₂I₅, showing potential extension for further application in X-ray scintillators.¹⁰⁷

Another interesting and noteworthy Cu-based scintillator was presented in recent years, and has attracted attention in the frontier research field, namely Cs₅Cu₃Cl₆I₂. Its presenters, Wu *et al.*, discovered it possessed benign grain boundaries without dangling bonds, which were commonly considered to contribute to RL lifetime extension and nonradiative recombination loss increase.¹⁰⁸ After fabrication as a columnar morphology *via* close space sublimation, it exhibits negligible afterglow and a high LY (Fig. 5c). The imaging property was also determined; it demonstrated a spatial resolution of 27 lp mm⁻¹ with a frame rate up to 33 fps. Also, Zhang integrated a Cs₅Cu₃Cl₆I₂ event-based scintillator film with negligible ghosting artifacts (0.1%) and an impressive data compression ratio of 23.7%.¹⁰⁹ Shu *et al.* reported the Rb doped Cs₅Cu₃Cl₆I₂ crystal with considerable improvement in stability against heat and humidity and an outstanding LY.¹¹⁰ Combined with PDMS, the Cs₅Cu₃Cl₆I₂:Rb scintillator screen reaches a high spatial resolution of over 10 lp mm⁻¹.

Besides, copper(i) itself can also be used as a dopant to enhance the STE state. To illustrate, Hu *et al.* discovered that the introduction of 0.05% Cu⁺ into Rb₂AgBr₃:Cu⁺ could largely enhance the PLQY to 98.8% (Fig. 5d).¹¹¹ Consequently, a pre-eminent LY of 79 250 photons per MeV and rather low detection limit of 714.83 nGy s⁻¹ were achieved. Likewise, Yao *et al.* synthesized 1D Cu⁺-doped Rb₂AgI₃ with 76.48% PLQY, demonstrating a LY of 36 293 photons per MeV, a low detection limit of 1.022 μGy_{air} s⁻¹ and a decay time of 465 ns.¹¹² In very recent

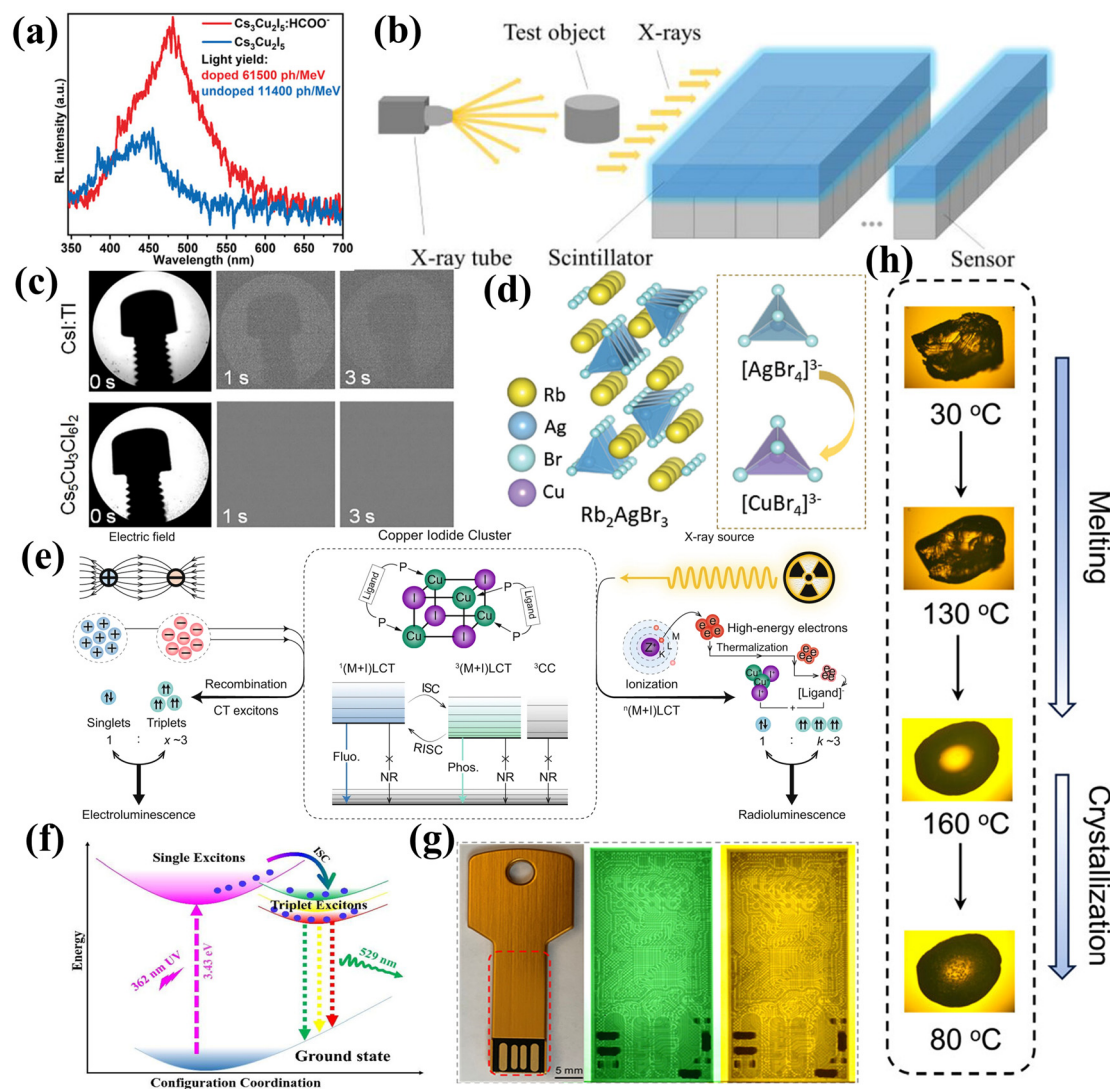


Fig. 5 (a) RL spectra of pristine and HCOO^- -doped $\text{Cs}_3\text{Cu}_2\text{I}_5$ single crystals. (b) Schematic of SIXS for multi-energy X-ray detection and imaging. (c) Dynamic imaging of a screw after 10 s of X-ray excitation using CsI:Tl and $\text{Cs}_5\text{Cu}_3\text{Cl}_6\text{I}_2$ films. The $\text{Cs}_5\text{Cu}_3\text{Cl}_6\text{I}_2$ film shows no residual image, while the image in the CsI:Tl film persists even after 3 s. (d) Crystal structure diagram of Rb_2AgBr_3 (left) and the schematic substitution model *via* Cu^+ doping (right). (e) Schematic of the mechanisms in Cu_4I_4 cubic clusters. (f) The PL mechanism in the configuration coordinate diagram of $[\text{BzTTPP}]_2\text{Cu}_2\text{I}_4$. (g) Bright-field and X-ray images of a USB drive. (h) POM images of M_3 at different temperatures.

work, Wu *et al.* further enhanced the performance of $\text{Rb}_2\text{AgBr}_3:\text{Cu}^+$, achieving single crystals with a PLQY of up to 99.2%.¹¹³ Through pulsed decay measurements and DFT calculations, they demonstrated that the incorporation of Cu increased the density of deep bound excitons, thereby significantly enhancing radiative recombination. By integrating the material with PDMS, they successfully fabricated a large-area flexible scintillator film, which exhibited a spatial resolution of 10.2 lp mm^{-1} , indicating its potential applications in radiation monitoring and biomedical imaging.

Copper-based systems, serving as a viable substitute for their lead-containing counterparts, have emerged as one of the most promising non-toxic scintillator candidates. Nevertheless, even though elemental doping strategies have

been employed to mitigate their degradation under ambient conditions, the degradation pathways, such as oxidation or segregation remain non-negligible.¹¹⁴ Furthermore, the spectral mismatch between their emission profile and the responsivity spectra of commercially available photodetectors imposes additional constraints on achieving optimal detection efficiency.¹¹⁵

3.3 Cu-based cluster scintillators

Another huge family of Cu-based scintillator materials are Cu_xI_x clusters, usually combined with organic cations or polymers, which no doubt are emerging as new X-ray scintillators (Table 3). Blessed with double STE emission, these clusters usually exhibit a considerable LY, a large Stokes shift and high-

Table 3 The parameters of Cu-based cluster scintillators

Material	Type	Light yield (photons per MeV)	Detection limit (nGy _{air} s ⁻¹)	Spatial resolution (lp mm ⁻¹)	Ref.
Cu ₄ I ₆ (bttmpe) ₂	Cluster	—	—	17	117
Cu ₆ I ₈ (bu-td) ₂	—	—	19.6	20	118
(IPP)CuI ₂	Cluster	75 793.83	62.29	10.2	119
(C ₁₂ H ₂₈ N) ₂ Cu ₂ I ₄	Single crystal	56 000	28.39	19.8	120
[DDPACDBFDP] ₂ Cu ₄ I ₄	Cluster	—	77	12	121
Cu ₄ I ₄ (C ₆ H ₁₄ N ₂) ₂	3D	48 538	30.68	—	122
Cu ₂ I ₂ (3,4-DMP) ₄	0D	38 031	106.71	10	123
(18-crown-6) ₂ Na ₂ (H ₂ O) ₃ Cu ₄ I ₆ (CNCl)	0D	109 000	59.4	16.3	124
(BzTPP) ₂ Cu ₂ I ₄	0D	27 706	352	4.928	125
(CISDM) ₄ [Cu ₄ I ₈] ₂ H ₂ O	2D	41 042	86.8	108	126
(4-bzpy) ₄ Cu ₄ I ₄	0D	60 948	155.2	5	127
TPA ₂ Cu ₂ I ₄	0D	40 124	126	5.5	128
Cu ₄ I ₆ (L1) ₂	0D	32 600	96.4	30	129
(MTP) ₂ CuI ₃ (M1)	0D	6400	72.6	20	130
Cu ₆ I ₈ (bu-td) ₂	—	—	32	17	131
1-rod	MOF	41 000	34.6	20	132
Cu ₄ I ₄ (R ₃ As) ₃ L	Cluster	15 000	18.1	—	133
Cu ₂ X ₂ (X = Cl, Br and I)	Nanoclusters	175 000	—	30	134
[CuI(PPh ₃) ₂ R] (R = PH, Cu-1 or PH-Br, Cu-2)	Cuprous complex	—	49.7	6.8	135

efficiency white light emission.^{116–118} As a highly regarded material, major efforts have been reported. For instance, Lai *et al.* prepared organic cuprous iodide clusters, (IPP)CuI₂ (IPP refers to iso-propyltriphenylphosphonium).¹¹⁹ These present the best scintillator property when IPP : CuI = 1 : 0.75, with a super low detection limit of 62.29 nGy s⁻¹ and an ultrahigh LY of 75 793.83 photons per MeV. They also applied it to CT and obtained a clear reconstructed 3D model, due to its superior spatial resolution of 10.2 lp mm⁻¹. Zhao *et al.* designed and fabricated a novel metal halide scintillator, (C₁₂H₂₈N)₂Cu₂I₄, with white emission.¹²⁰ Benefitting from the double self-trapped mechanism, it demonstrates a really high PLQY, which not only matches with the response of semiconductor-based sensors, but also decreases the doses to which the objects are exposed. The flexible transparent large-area scintillator film made from this shows a superior luminescence property, a high spatial resolution of 19.8 lp mm⁻¹ and an incredibly low detection limit of 28.39 nGy_{air} s⁻¹; these values are four times higher and 194 times lower than the typical value in medical usage. Moreover, recognizing the trivial factor that X-ray luminescence efficiency in metal clusters is governed by the competition between radiative states from organic ligands and non-radiative ones (Fig. 5e), Zhang *et al.* reported a class of Cu₄I₄ cubes, including [DDPACDBFDP]₂Cu₄I₄ (DDPACDBFDP refers to 10,10'-(4,6-bis(diphenylphosphanyl))). It presents an impressive emissive RL due to functionalizing the biphosphine ligands with acridine, which can effectively absorb ionization radiation to generate electron-hole pairs and subsequently transfer them to ligands during thermalization.¹²¹ It demonstrates a PLQY of 95%, benefiting from triplet-to-singlet conversion *via* a TADF matrix, the lowest detection limit of 77 nGy s⁻¹ and a high spatial resolution of up to 12 lp mm⁻¹. Similarly, Zou *et al.* employed a solvent-recyclable green process to synthesize stable three-dimensional

nanoclusters, Cu₄I₄(C₆H₁₄N₂)₂, achieving a PLQY close to 100%.¹²² The sturdy 3D framework ensures the morphological stability of the molecules at high temperatures and humidity. It retains 90% of its room-temperature luminescence intensity at 110 °C and 88.34% of its initial luminescence intensity after 100 days under ambient conditions. This demonstrates a promising approach for the green, low-toxicity synthesis of high-performance, lead-free nanoclusters. In work recently reported by the same group, Zou *et al.* synthesized hybrid copper iodide single crystals, Cu₂I₂(3,4-DMP)₄ (3,4-DMP = 3,4-dimethylpyridine), with a PLQY of 80%, utilizing ionic liquids as both the solvent and iodine source, and nitrogen-containing organic ligands for structural modulation.¹²³ Notably, the material overcomes the common issue of moisture-induced degradation, demonstrating exceptional stability over 90 days, thereby providing a novel solution for the inkjet printing of scintillators.

To extend the scintillator properties, Wang *et al.* synthesized (18-crown-6)₂Na₂(H₂O)₃Cu₄I₆ (CNCl) with little self-absorption and near-unity green light emission.¹²⁴ It possesses the highest values of LY ever reported, up to ~109 000 photons per MeV, and other impressive scintillator properties, like an ultralow detection limit of 59.4 nGy s⁻¹ and an appreciable spatial resolution of 16.3 lp mm⁻¹ for CNCl-polymer film screens, which are further improved after modification by a silicon wave guide structure to 24.8 lp mm⁻¹. Furthermore, Lin *et al.* designed (BzTPP)₂Cu₂I₄ (BzTPP refers to benzyltriphenylphosphonium) as a scintillator, exhibiting a substantial Stokes shift of 167 nm and 27 706 photons per MeV, the energy transfer mechanism of which is shown in Fig. 5f. This material stands out as the premier choice among copper-based scintillators.¹²⁵ Also, Yang *et al.* presented a 2D-copper-based cluster, (CISDM)₄[Cu₄I₈]₂H₂O (CISDM refers to *cis*-2,6-dimethylmorpholine), as a qualified scintillator material, with 87.2%

PLQY at 588 nm.¹²⁶ It also demonstrates a strong linear response to X-rays, with a LY of 41 042 photons per MeV, a low detection limit of 86.8 nGy s⁻¹, and an exceptional resolution of 108 lp mm⁻¹. These characteristics represent a state-of-the-art performance for lead-free metal halide hybrid materials in spatial detection. Also, Kong *et al.* combined a conjugated organic cation (4-benzylpyridine, 4-bzpy) with copper(I) iodide modules to grow organic copper halide single crystals, namely 0D cubane-like (4-bzpy)₄Cu₄I₄.¹²⁷ It demonstrates a high LY of 60 948 photons per MeV and an exceptional linear relationship with X-ray dose, originating from the ligand-to-core charge transfer state. Similarly, Lv *et al.* developed a copper halide scintillator, TPA₂Cu₂I₄, utilizing the principle of STE ultra-broadband emission.¹²⁸ The material achieves a remarkably high PLQY of 94.27%, a LY of 40 124 MeV⁻¹, and a spatial resolution of 5.5 lp mm⁻¹, thereby providing new insights into broadband-emission X-ray scintillators.

Also, a nano-rod strategy was employed to promote the performance. He *et al.* chose copper iodide ink, Cu₄I₆(L₁)₂ (L₁ refers to 1-propyl-1,4-diazabicyclo[2.2.2]octan-1-ium) 0D nano-rods, as the scintillator, which exhibited a PLQY of 95.3% for broadband green emission, and the imaging result of this scintillator film is shown in Fig. 5g.¹²⁹ The subsequent scintillator screen shows a low detection limit of 96.4 nGy s⁻¹, approximately 55 times lower than that required for standard medical diagnosis (5.5 μGy s⁻¹). Furthermore, the spatial resolution exceeding 30 lp mm⁻¹ was twice that of conventional scintillators. To enhance the stability, Bohan Li *et al.* fabricated three excellent glass phase 0D hybrid copper halides, (MTP)₂Cu₃(M₁) (MTP refers to methyltriphenylphosphonium), (MTP)₂Cu₄I₆-α (M₂)/(MTP)₂Cu₄I₆-β (M₃), which shared the same inorganic anions.¹³⁰ The prepared glass samples were heated in air to promote the crystal-glass phase transition, and the self-assembly enabled recrystallization, resulting in a hybrid bulk glass-ceramic; the procedure is illustrated by polarized optical microscopy (POM), as shown in Fig. 4h. It demonstrates an outstanding LY of 64 000 photons per MeV, a detection limit of 72.6 nGy s⁻¹ and high stability for real-time X-ray imaging with spatial resolution above 20 lp mm⁻¹.¹³⁰ Likewise, Gu *et al.* reported the synthesis of Cu₆I₈(bu-*ted*)₂ *via* a surfactant-assisted method, limiting the crystal size with the help of controlling surface tension.¹³¹ The material shows a near-unity PLQY, resulting in a light output 4.8 times higher than that of the commercial scintillator Lu₃Al₅O₁₂:Ce³⁺, as well as an impressively low detection limit of 32 nGy s⁻¹ and a spatial resolution of 17 lp mm⁻¹ while integrating SDBS into a flexible screen. Also, for the purpose of preparing a high quality, flexible and stable scintillator screen, Peng *et al.* selected a macrocyclic bridging ligand with an aggregation-induced emission feature as the MOF, which could endow the scintillator with high efficiency and preminent stability, combined with a copper iodide cluster.¹³² The as-fabricated scintillator film exhibits exceptional flexibility, enabling their integration with flexible OLEDs for the precise imaging of complex surfaces or internal structures of objects. This capability holds significant potential for applications in industrial inspection

and biological imaging. Moreover, the introduction of *in situ* polyvinyl pyrrolidone during the synthesis process leads to the regular rod-shaped microcrystal, enhancing the processibility even further. The scintillator realizes dynamic X-ray flexible imaging for the first time with a real time ultra-high spatial resolution of 20 lp mm⁻¹. Also, Demyanov *et al.* presented cubane Cu₄I₄-triarsenic clusters [Cu₄I₄(R₃As)₃L] (R refers to the organic functional group, L refers to no ligand or nitrile) with a record ultralow detection limit of 18.1 nGy s⁻¹ and an exquisite linear response over a wide range of X-ray doses (0–640 μGy_{air} s⁻¹).¹³³ Yuan *et al.* reported their discovery that Cu₂X₂ (X = Cl, Br and I) was a TADF nano-cluster scintillator suffering from no mechanochromism and blessed with a considerable X-ray absorption cross section, which contributed to a flexible scintillator screen with excellent radiation and humidity resistance, an ultra-high LY of 175 000 photons per MeV and a high spatial resolution of ~30 lp mm⁻¹.¹³⁴ Besides, for the purpose of being eco-friendly, Chen reported a mechanochemical method for synthesizing [CuI(PPh₃)₂R] (R = PH, Cu-1 or PH-Br, Cu-2), circumventing the harm brought by diverse organic solvents in traditional chemical experiments.¹³⁵ Cu-2 demonstrates 2.52 times higher RL intensity than that of the commercial BGO scintillator and 1.52 times larger than that of Cu-1 due to the introduction of PH-Br, which resulted in 37.75% higher PLQY.

Nanoclusters, emerging as a novel class of materials with dimensions intermediate between single atoms and quantum dots, exhibit near-unity PLQY and negligible Stokes shifts. However, key mechanistic aspects, such as the nature of halide ligand coordination dynamics, remain elusive,¹²⁷ while their prevalent mechanochemical grinding synthesis protocols face scalability challenges compared to solution-based methods commonly employed for other luminescent materials.¹³⁶

3.4 Rare-earth nanocrystalline scintillators

Rare-earth-doped scintillator materials demonstrate promising applications in nuclear medicine, radiographic imaging, encryption and particle detection. The high energy-transfer efficiency among rare-earth elements enhances the overall performance of the materials, yielding greater light output. Furthermore, doping with rare earth elements enhances the thermal and chemical stability of scintillator materials. Their unique properties also enable a wide range of luminous wavelengths, meeting the diverse needs of various applications.¹³⁷

In recent efforts, the doping of rare-earth elements into perovskite materials has yielded significant progress. For instance, in 2024, Wang *et al.* reported a series of Cs₂Ag_xNa_{1-x}In_yBi_{1-y}Cl₆ halide scintillators doped with rare-earth ions.¹³⁸ Among them, Cs₂Ag_{0.6}Na_{0.4}In_{0.95}Bi_{0.05}Cl₆:5%Tm has a high PLQY in NIR light of up to 76.3% and a LY of up to 33 500 photons per MeV. The flexible scintillators achieved a high spatial resolution of 11.2 lp mm⁻¹ and a low X-ray detection limit of 55.2 nGy_{air} s⁻¹. Moreover, after doping with Tm ions, they achieved clear imaging under bright external lighting by using an infrared camera with a 760 nm long-pass filter in front of the lens (Fig. 6a).

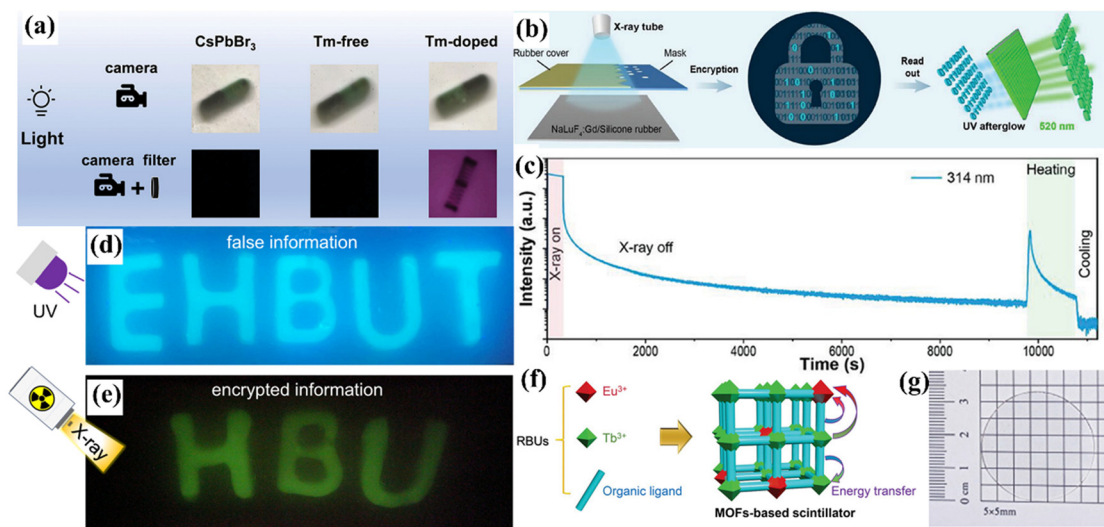


Fig. 6 (a) X-ray images of encapsulated springs under various conditions were captured using CsPbBr₃, Cs₂Ag_{0.6}Na_{0.4}In_{0.95}Bi_{0.05}Cl₆ (Tm-free), and Cs₂Ag_{0.6}Na_{0.4}In_{0.95}Bi_{0.05}Cl₆:5%Tm (Tm-doped) films. (b) Schematic diagram of writing X-ray imaging information and decryption by a layer of the perovskite nanocrystals. (c) Decay curve of the thermal-stimulated luminescence of NaLuF₄:Gd³⁺ (20 mol%) at 314 nm was recorded at room temperature after X-ray excitation was stopped for 5 min. The film was then heated to 80 °C for 160 min. (d) Photograph of the false information EHBUT in the prepared multilayer hydrogel under UV light (365 nm). (e) Photograph of the encrypted information HBU in the multilayer hydrogel after exposure to X-rays. (f) Schematic representation of radio-luminescent functional building units for constructing a thermo-responsive MOF scintillator. (g) Bright field photograph of glass scintillators.

Doping rare-earth ions into fluoride-containing nano-scintillators is also a recognized approach.^{8,139–144} In 2023, Yang *et al.* reported high-security X-ray imaging encryption technology by developing an ultra-long RL memory film, namely NaLuF₄:Gd³⁺ or Ce³⁺, which captured subtle pure UV emission characteristics.¹⁴⁵ The encrypted X-ray imaging information can be securely stored in memory films for over seven days and can be optically decoded using perovskite nanocrystals (Fig. 6b and c). Moreover, the as-fabricated flexible memory film exhibits a high spatial resolution of 20 lp mm⁻¹. The varying durations of fluorescence lifetime offer distinct advantages in different application contexts. To address the issue of strong hysteresis scintillation luminescence, Zhang *et al.* synthesized LiYF₄:15Tb/25Gd@LiYF₄ core/shell nano-scintillators to inhibit the generation of Frenkel defects.¹⁴¹ In the same year, Yang *et al.* reported NaLuF₄:Gd/Tb/Eu doped with 5%Tb with a high LY of 15 313 photons per MeV and a low detection limit of 84.1 nGy_{air} s⁻¹.¹⁴⁶ Subsequently, Zhang *et al.* exhibited Tb³⁺-doped Na₅Lu₉F₃₂ with good water dispersibility and highly sensitive luminescence to X-rays.¹⁴⁷ Remarkably, they employed multilayer hydrogels for information camouflage and multilayer encryption. The encrypted information can only be identified through X-ray exposure, while false information is revealed under ultraviolet light (Fig. 6d and e). In 2024, Lu *et al.* proposed a type of heterovalent Cu²⁺ ion co-doped LiLuF₄:Tb,Cu microcrystalline scintillation material with higher RL intensity, long afterglow and thermoluminescence intensity.¹⁴⁸ The detection limit of LiLuF₄:Tb,Cu can reach 2.7928 nGy s⁻¹ and a spatial resolution of 22 lp mm⁻¹@MTF = 0.2 is achieved.

In the field of scintillators materials, MOFs provide a tunable platform for the integration of X-ray absorption centers and luminescent chromophores in a dense framework manner and rare-earth-doped MOFs are widely studied recently, as it is in perovskite scintillators.¹⁴⁹ In 2024, Li *et al.* proposed thermo-responsive lanthanide MOF scintillators (Tb_{0.95}Eu_{0.05}-BPTC), presenting a low dose rate detection limit of 156.1 nGy_{air} s⁻¹ (ref. 150) (Fig. 6f). Tb_{0.95}Eu_{0.05}-BPTC has a high relative LY of 39 000 photons per MeV, imaging spatial resolution of 18 lp mm⁻¹, good irradiation stability and giant color transformation visualization, benefiting the applications. Similarly, Jin *et al.* reported [1-ethyl-3-methylimidazolium] EuBr₃MeOH as scintillators with a LY of 43 000 ± 100 photons per MeV and achieved spatial resolution above 10 lp mm⁻¹.¹⁵¹ Additionally, Yang *et al.* reported Eu-pba and Tb-pba, featuring excellent radiation, hydrolytic, and thermal stabilities. Using them as scintillators can result in a linear response to the X-ray dose rate with detection limits of 4.92 and 3.17 μGy s⁻¹, respectively.¹⁵² Recently, Xu *et al.* firstly proposed organo-lanthanide scintillators that offered diverse emission colors from ultraviolet to near-infrared and enabled precise control of luminescence lifetimes from nanoseconds to microseconds.¹⁵³ Their materials can efficiently capture the dark triplet excitons generated during the absorption of secondary X-rays, leading to the production of highly stable and efficient RL that surpasses those of commercialized scintillators. Among these scintillators, for instance, remarkably, Eu(NTA)₃DPEPO (where NTA refers to 4,4,4-trifluoro-1-(2-naphthyl)-1,3-butanedione and DPEPO refers to bis(2-(diphenylphosphino)phenyl) ether oxide) showed a more than 1000-fold enhancement in RL com-

pared with EuCl_3 salts with an equal molar amount of Eu^{3+} . In a recent study, Wang *et al.* reported the successful synthesis of a novel terbium-based MOF (Tb-MOF-1), the enhanced RL performance of which was attributed to the efficient sensitization effect of the HDOBPDC³⁻ ligand.¹⁵⁴ This material demonstrated a remarkably low X-ray detection limit of $1.71 \mu\text{Gy s}^{-1}$, significantly surpassing conventional scintillation thresholds. Furthermore, the researchers developed a flexible composite scintillator screen through polymer matrix integration, which exhibited exceptional imaging capabilities with a spatial resolution of 7.7 lp mm^{-1} , meeting clinical radiography requirements.

Rare-earth-doped glass scintillators have also attracted significant attention in recent years due to outstanding optical performances such as high transmittance (Fig. 6g) and their ability to absorb high energy radiation and convert the energy into visible light.^{155,156} In 2023, Sun *et al.* fabricated Tb^{3+} -doped and $\text{Ce}^{3+}/\text{Tb}^{3+}$ -co-doped oxyfluoride scintillator (C_4T_9) glass samples with a spatial resolution of 16 lp mm^{-1} .¹⁵⁷ Then in 2024, Li *et al.* synthesized Tb^{3+} -doped oxyfluoride aluminosilicate glass scintillators with a spatial resolution of 20 lp mm^{-1} and higher RL of up to 224% compared to the $\text{Bi}_4\text{Ge}_3\text{O}_{12}$ crystal.¹⁵⁸ Subsequently, Dai *et al.* reported a transparent glass-ceramic scintillator by embedding Tb^{3+} -doped NaLu_2F_7 nano-crystals in a glass matrix.¹⁵⁹ Importantly, most decreases in performance, due to prolonged exposure to high doses of X-rays, can be completely restored by heat treatment at $350 \text{ }^\circ\text{C}$ for 30 min.

Additionally, several other rare-earth-doped scintillators have been reported.^{160,161} For instance, in 2023, Ling *et al.* prepared $\text{YF}_3:30\%\text{Gd}^{3+}/10\%\text{Tb}^{3+}$ micro-particles with fine environmental stability and superb X-ray RL, which achieved an

imaging spatial resolution of 16.8 lp mm^{-1} .¹⁶² Subsequently, in 2024, Wang *et al.* reported zero-thermal-quenching $\text{Lu}_3\text{Al}_5\text{O}_{12}:\text{Ce}^{3+}$ transparent ceramics (0.5 mm) with excellent spatial resolution of up to 112 lp mm^{-1} .¹⁶³ In the same year, Yang *et al.* synthesized single crystals of rare earth ion-doped ternary chalcogenides, NaGaS_2/Eu , with a low detection limit of 250 nGy s^{-1} and a spatial resolution of 13.2 lp mm^{-1} .¹⁶⁴ Doping with rare-earth elements imparts some materials with unexpectedly excellent properties, making the continued exploration of the potential of rare-earth-doped scintillators very promising. Furthermore, Wang *et al.* recently advanced the frontier of high-temperature scintillator technology through the development of a Tb^{3+} -doped nanocrystalline glass composite system.¹⁶⁵ This material possesses a record LY of 54 900 photons per MeV and a sensitivity of $635.31 \text{ nGy}_{\text{air}} \text{ s}^{-1}$. Remarkably, the material maintains exceptional thermal stability with 28.1 lp mm^{-1} spatial resolution at $500 \text{ }^\circ\text{C}$ and robust moisture resistance, surpassing current high-temperature scintillators.

The utilization of rare earth elements in optical materials has been extensively investigated (Table 4), as their intricate electronic configurations offer a compelling avenue for tailoring material band structures. However, the environmental and biological implications of these elements demand critical scrutiny, as exemplified by nephrotoxicity associated with gadolinium exposure, pulmonary fibrosis risks linked to samarium compounds,¹⁶⁶ and inflammatory responses triggered by elevated cerium concentrations.¹⁶⁷ Furthermore, the strikingly similar physicochemical properties among rare earth elements complicates purification processes, which objectively inflate raw material costs and ultimately constrain their large-scale industrial implementation.

Table 4 The parameters of rare-earth nanocrystalline scintillators

Material	Type	Light yield (photons per MeV)	Detection limit ($\text{nGy}_{\text{air}} \text{ s}^{-1}$)	Spatial resolution (lp mm^{-1})	Ref.
$\text{Cs}_2\text{Ag}_{0.6}\text{Na}_{0.4}\text{In}_{0.95}\text{Bi}_{0.05}\text{Cl}_6:5\%\text{Tm}$	Single crystals	33 500	55.2	11.2	138
$\text{LiYF}_4:15\text{Tb}/25\text{Gd}@/\text{LiYF}_4$	Nanocrystals	—	—	20	141
$\text{NaLuF}_4:\text{Gd}/\text{Tb}/\text{Eu}$ with 5%Tb-doping	Nanocrystals	15 313	84.1	8.7	146
Tb^{3+} -doped $\text{Na}_3\text{Lu}_9\text{F}_{32}$	Nanocrystals	15 800	—	—	147
$\text{LiLuF}_4:\text{Tb},\text{Cu}$	Microcrystals	—	2.7928	22	148
$\text{Tb}_{0.95}\text{Eu}_{0.05}\text{-BPTC}$	MOFs	39 000	156.1	18	150
[1-Ethyl-3-methylimidazolium]EuBr ₃ MeOH	Organic-inorganic hybrids	43 000	405.4	>10	151
Eu-pba/Tb-pba	Organic-inorganic hybrids	—	4920/3170	8/12.6	152
Eu(NTA) ₃ DPEPO	Organic-inorganic hybrids	38 589	30.8	20	153
Ln-MOF [Tb(HDOBPDC)(DMF)(H ₂ O)] _n	MOF	—	1710	7.7	154
SiBNaBaGd-xTb	Glass	—	—	20	155
Ce^{3+} -doped $55\text{SiO}_2\text{-}20\text{Al}_2\text{O}_3\text{-}15\text{SrF}_2\text{-}10\text{NaF}$	Glass	—	—	18	156
$\text{Ce}^{3+}/\text{Tb}^{3+}$ -co-doped	Glass	—	343.9	16	157
$55\text{SiO}_2\text{-}14.5\text{SrF}_2\text{-}4.5\text{SrO}\text{-}15\text{LuF}_3$	—	—	—	—	—
$10\text{Al}_2\text{O}_3\text{-}0.5\text{Sb}_2\text{O}_3\text{-xTb}_4\text{O}_7\text{-yCeF}_3$	Glass	—	—	20	158
$55\text{SiO}_2\text{-}20\text{Al}_2\text{O}_3\text{-}15\text{BaF}_2\text{-}10\text{NaF-xTbF}_3$	—	—	—	—	—
Tb^{3+} -doped NaLu_2F_7	Nanocrystals	—	—	20	159
$\text{SrS}:\text{Ce}^{3+}/\text{CaS}:\text{Ce}^{3+}$	Nanocrystals	—	660	—	160
$2\text{BaO}\text{-}3\text{SiO}_2$ doped with Eu_2O_3	Glass	26 000	—	12	161
$\text{YF}_3:\text{Gd}^{3+}/\text{Tb}^{3+}$	Microparticles	—	—	16.8	162
$\text{Lu}_3\text{Al}_5\text{O}_{12}:\text{Ce}^{3+}$	Ceramics	40 200	—	112	163
NaGaS_2/Eu	Crystal	8188	250	13.2	164
$\text{KTb}_{3-x}\text{Gd}_x\text{F}_{10}$	Crystal	54 900	635.31	28.1	165

3.5 Organic scintillators

In addition to the scintillators mentioned above, organic scintillators have garnered significant attention^{168–170} due to their rapid response times, diverse chemical structures, low processing temperatures, and cost-effectiveness (Table 5). During the luminescence phase, the charge carriers recombine to produce singlet states (25%) and triplet states (75%), resulting in light emission. However, triplet excitons may be lost due to the dark state characteristics of fluorescent organic scintillators.¹⁷¹ TADF and room-temperature phosphorescence (RTP) materials have garnered significant attention due to their ability to exhibit bright triplet excitons under X-ray excitation.^{172,173} Additionally, hybridized local and charge-transfer (HLCT) excited states in organic donor–acceptor systems can produce efficient PL through high-efficiency triplet–singlet reverse intersystem crossing (RISC) transitions, achieving nearly 100% utilization of excitons.¹⁷⁴ To enhance efficiency and achieve rapid RISC, a common strategy involves increasing the spin-orbital coupling constant (SOC) in organic emitters by exploiting heavy-atom effects.¹⁷⁵ Researchers have proposed innovative methods and applications based on these principles and have identified numerous new, high-performance purely organic scintillators.

The RL performance of organic fluorescence scintillators could be enhanced by improving carrier transport capabilities.¹⁷⁶ Following this guidance, in 2023, Chen *et al.* introduced a highly efficient organic fluorescent scintillator, 4,4'-bis(9-carbazolyl) biphenyl (CBP), which demonstrated a high PLQY of 61.92% and excellent carrier transport ability (Fig. 7a), and biphenyl (BP) showed an average hole mobility of $0.055 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ while CBP achieved a hole mobility of $0.094 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$.¹⁷⁷

In the same year, Chen *et al.* proposed a molecular design strategy that incorporated supramolecular halogen bonding in fluorescence and room-temperature phosphorescence systems containing pyridine rings.¹⁷⁸ Upon exposure to X-ray radiation, the introduction of halogen heavy atoms ($X = \text{Cl}, \text{Br}, \text{I}$) into 9,10-bis(4-pyridyl)anthracene (BPA) and the formation of halogen bonding interactions in bromophenyl–methyl-

pyridinium iodide (PYI) crystals facilitate the generation of numerous hot electrons and deep holes (Fig. 7b). These charge carriers undergo a series of conversion and transport processes, including electron–electron scattering, Auger processes, and thermalization, resulting in a substantial number of low-energy carriers. Subsequently, these carriers recombined to form electron–hole pairs, leading to the production of 25% singlet and 75% triplet excitons, consistent with spin statistics. During the luminescence process, BPA-X crystals produced rapid fluorescent light emission, primarily utilizing 25% singlet excitons, while PYI crystals exhibited phosphorescent light emission, harnessing both singlet and triplet excitons. Similarly, Chen *et al.* utilized the strategy of incorporating heavy atoms in their latest publication.¹⁷⁹ They developed a new category of organogold(III) complexes, namely Tp-Au-1 and Tp-Au-2, by adopting a through-space interaction motif to achieve high X-ray attenuation efficiency and efficient harvesting of triplet excitons for emission. Gold is a promising transition metal due to its significant spin–orbit coupling constant, derived from its large atomic number ($Z = 79$) and greater abundance compared to widely used metals like iridium or platinum.¹⁸⁰ Consequently, neat films of Tp-Au-1 and Tp-Au-2 exhibit notable TADF properties with significant triplet–singlet charge transfer features, short decay lifetimes of 1.7–2.2 μs and a high PLQY of up to 77%. Remarkably, the emission mechanism of Tp-Au-1 and Tp-Au-2 can be tuned from TADF to phosphorescence by adjusting the polarity of the host matrix. Under X-ray irradiation, Tp-Au-2 demonstrated intense RL and a record-high scintillation LY of 77 600 photons per MeV for organic scintillators. The resulting scintillator screens provided high-quality X-ray imaging with a spatial resolution exceeding 16.0 lp mm^{-1} (Fig. 7c).

In 2024, Dong *et al.* reported two phenothiazine derivatives exhibiting polymorphism-dependent phosphorescence RL.¹⁸¹ These derivatives demonstrate high radio stability and a low detection limit of 278 nGy s^{-1} . The study not only introduced sulfur atoms to enhance X-ray absorption through the heavy atom effect but also confirmed that incorporating pyrimidine and pyridine groups containing heteroatoms into the phenothiazine skeleton promoted $n\text{-}\pi^*$ transitions, improving the

Table 5 The parameters of organic scintillators

Material	Light yield (photons per MeV)	Detection limit ($\text{nGy}_{\text{air}} \text{ s}^{-1}$)	Spatial resolution (lp mm^{-1})	Ref.
$\text{Py}_2\text{TTz-I}_2\text{F}_4$	32 583	70.49	26.8	168
$(\text{TPPcarz})_2\text{MnBr}_4$	44 600	32.42	—	169
Tetra(<i>p</i> -bromophenyl)ethene@poly(vinyltoluene)	14 443	—	13.9	170
4,4'-Bis(9-carbazolyl)biphenyl	—	25.5	14.3	177
BPA-X ($X = \text{Cl}, \text{Br}, \text{I}$)	—	25.95 ± 2.49	20	178
Tp-Au-2	77 600	—	16	179
10-(Pyridine-2-yl)-10 <i>H</i> -phenothiazine	—	278	—	181
Sulfone-based organic molecules C_6	16 558	—	15	182
DCB@C[3]A	—	77.1	20	183
BiND-OMe	—	70.68	11	186
OMNI-PTZ 2	—	97	20	187
1,1,2,2-Tetrakis(4-bromophenyl)ethylene	34 600	447	18.69	188

while achieving a high spatial resolution of 11.0 lp mm^{-1} . Additionally, Zhang *et al.* demonstrated that the oxidized 1,8-naphthalimide-phenothiazine dyad (OMNI-PTZ 2) with HLCT-excited states showed an enhanced overlap integral between the highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO) on MNI π -orbitals, as well as moderate donor-acceptor electron interactions.¹⁸⁷ Consequently, OMNI-PTZ 2(G) exhibits significantly increased RL and greatly reduced decay lifetime, achieving an X-ray dose sensitivity of 97 nGy s^{-1} and an exceptional spatial resolution of 20 lp mm^{-1} (Fig. 7g). Furthermore, Du *et al.* developed a new family of hot exciton scintillators, which were characterized by a large energy gap between the high-lying and lowest triplet states and a small energy difference between triplet and singlet states.¹⁸⁸ 1,1,2,2-Tetrakis(4-bromophenyl) ethylene demonstrates an ultrafast radiative lifetime of 1.79 ns and a LY of approximately 34 600 photons per MeV, showing an excellent combination of high LY and short decay time.

Organic materials, leveraging their structural diversity and tunability, have been extensively investigated for scintillation applications. However, reports on purely organic molecular scintillators remain relatively scarce compared to inorganic or hybrid organic-inorganic systems. This disparity stems from

intrinsic limitations: spin-forbidden transitions impose fundamental constraints on LY optimization through molecular design alone, while their inherently low Z_{eff} further limits their efficacy in high-energy radiation absorption.¹⁸⁹

3.6 Specialized structure scintillators

The previously discussed advancements in scintillators focus on innovative, high-performance compounds with the goal of synthesizing even more superior scintillators. Furthermore, unique structures have been incorporated into the design of scintillator materials to enhance their performance.^{190–192} In this regard, Lin *et al.* proposed the concept of ‘structural engineering’, which, through internal or external design at scales ranging from micro to macro, offered a promising strategy that not only enhanced the performance of scintillators but also broadened their functionality in specific applications such as radiation imaging and therapy, opening up new avenues for candidate materials.¹⁹³ Likewise, Roques-Carnes *et al.* proposed a first-principles theory of nanophotonic scintillators and demonstrated order-of-magnitude scintillation enhancements in two different platforms.¹⁹⁴ Furthermore, the arraying of scintillators through specialized structures to enhance photon yield or resolution is a common approach. In

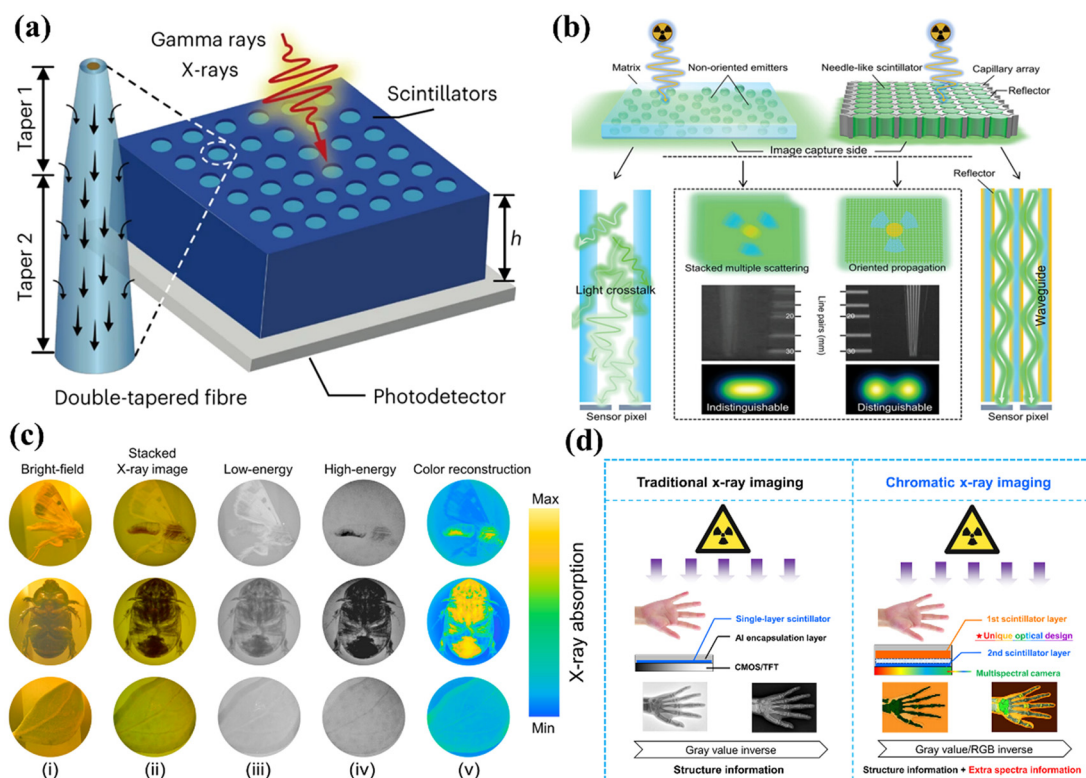


Fig. 8 (a) Schematic representation of scintillator-encapsulated fiber arrays for the detection and imaging of X-rays and gamma rays. Within the fiber array, radioluminescence can propagate over several centimeters. Without the fiber array, however, the radioluminescence becomes attenuated and undetectable due to absorption and scattering within the scintillator layer. (b) Mechanisms of light propagation and the performance of conventional unstructured scintillators versus structured capillary array scintillators in X-ray imaging. (c) Bright-field (i), stacked (ii), decomposed (iii and iv), and color-reconstructed (v) dual-energy X-ray images of a moth, a beetle, and a leaf, utilizing the top-filter-bottom sandwich structure scintillator. (d) Comparison of traditional X-ray imaging approaches and chromatic X-ray imaging strategies.

the design of such structures, three primary objectives are typically pursued: the growth of high-quality scintillators of appropriate thickness, increasing the refractive index contrast between the scintillator and its matrix, and optimizing the photon structure within the scintillator array.¹⁹⁵ In 2023, Dierks *et al.* reported that they synthesized CsPbBr₃ nanowires in anodized aluminum oxide and achieved sufficient stability for X-ray micro-tomography under ambient conditions, as well as high spatial resolution.¹⁹⁶ In the same year, Yi *et al.* proposed the design of double-tapered optical-fiber arrays, which could drastically increase the light-collection efficiency and imaging resolution (Fig. 8a).¹⁹⁷ They achieved a spatial resolution of 22 lp mm⁻¹ and pixelated gamma-ray imaging with a thickness of 4 mm and ~10 000 pixels under focused 6 MeV irradiation. Moreover, the dose rate (3.5 nGy s⁻¹ to 96 mGy s⁻¹) can be conveniently monitored. Then in 2024, considering that the absence of X-ray energy distinction resulted in insufficient image contrast, Ran *et al.* presented an innovative multi-energy X-ray linear-array detector based on Cs₃Cu₂I₅.¹⁰⁵ Benefiting from negligible self-absorption of the material and side-illuminated scintillation scenarios, the incident X-ray spectra could be reconstructed by analyzing the distribution of scintillator intensity while illuminated from the side, and the outcome error could be kept below 5.63%. Subsequently, Shao *et al.* fabricated a thick pixelated needle-like array scintillator capable of micrometer resolution (Fig. 8b).¹⁹⁸ They achieved ultrahigh spatial resolutions of 60.8 lp mm⁻¹ (the thickness of the scintillators is 0.5 mm) and 51.7 lp mm⁻¹ (the thickness of the scintillators is 1 mm) at a MTF of 0.2 due to isolated light-crosstalk channels and robust light output. More recently, Song *et al.* proposed an anti-scattering CsPbBr₃ scintillator array embedded within a polyurethane acrylate matrix, which could suppress light scattering and enhance the light collection efficiency by nearly two times compared to the monolithic film.¹⁹⁹ Due to the large refractive index contrast between the scintillator and matrix, the scintillator array exhibits a low dosage and high resolution.

In addition to the specialized structures mentioned above, dual-energy X-ray imaging (DEXI) is cutting-edge technology that provides more detailed information about specific materials compared to traditional single-energy X-ray imaging strategies. In 2023, Shao *et al.* demonstrated a top-filter-bottom sandwich structure scintillator for high-resolution DEXI within a single exposure, achieving an excellent resolution of approximately 18 lp mm⁻¹ on stacked images (Fig. 8c).²⁰⁰ Subsequently, Hu *et al.* successfully achieved color recognition of objects with different densities by using the scintillators of mutually inactive (BA)₂PbBr₄:Mn and Cs₃Cu₂I₅:TI, which contributed to the application of color X-ray imaging in real-world scenarios (Fig. 8d).²⁰¹

4. Conclusion and outlook

In conclusion, we first provide a comprehensive overview of the various parameters and performance standards of scintilla-

tors to facilitate the assessment of their efficacy in recent studies. In recent years, the field of scintillators has garnered significant attention and experienced rapid development.

Lead-based perovskite scintillators exhibit exceptional potential for X-ray detection due to their tunable optoelectronic properties, high spatial resolution, and rapid responses. However, their practical application is hindered by intrinsic challenges: environmental and health concerns from lead toxicity, structural instability under exposure to moisture and oxygen, and luminescence degradation under operational stress. Recent strategies, such as polymer encapsulation, metal-organic framework confinement, and defect-passivating ligand systems, have improved the stability by suppressing ion migration and environmental degradation. Concurrently, elemental doping and lattice engineering enhance energy transfer efficiency while mitigating self-absorption. On the other hand, lead-free perovskite scintillators, particularly those that are copper-based, double perovskites, and manganese-based halides, have emerged as promising alternatives to their toxic lead-based counterparts, offering a balance of environmental sustainability and high-performance X-ray detection. For instance, copper-based systems, such as Cs₃Cu₂I₅ and its derivatives, leverage zero-dimensional structures to localize STEs, suppressing non-radiative recombination while enabling tunable emission and flexible film integration. Innovations in doping, composite engineering, and morphology control further enhance their stability and LYs. Despite these advances, challenges persist, including insufficient carrier mobility in the double perovskites, long-term environmental degradation in certain hybrids, and scalability limitations in synthesis protocols. Future research should prioritize synergistic material designs—combining computational modeling with experimental validation—to engineer defect-tolerant lattices, optimize STE dynamics, and develop hydrophobic interfaces. Scalable fabrication techniques, such as inkjet printing and roll-to-roll processing, must be refined to produce large-area, flexible scintillator films for real-world applications. Additionally, exploring multifunctional systems that integrate radiation detection with stimuli-responsive properties could unlock novel applications in low-dose medical imaging, wearable diagnostics, and industrial nondestructive testing. By bridging material innovation with industrial compatibility, lead-free scintillators are poised to redefine radiation detection technologies, aligning high performance with ecological and operational sustainability.

Copper-based cluster scintillators, particularly those centered on Cu_xI_x architectures, represent a groundbreaking advancement in X-ray detection technologies. These materials harness the unique advantages of STE mechanisms, tunable organic-inorganic hybrid frameworks, and efficient radiative recombination, enabling the development of high-performance, low-toxicity scintillators. By optimizing ligand-core charge transfer states and structural engineering, they achieve near-unity PLQYs, ultralow detection limits, and exceptional spatial resolution, while demonstrating remarkable flexibility and environmental stability. Innovations in green synthesis,

such as ionic liquid solvents and mechanochemical approaches, further promote lead-free, energy-efficient fabrication processes, laying the foundation for scalable applications. Nevertheless, critical challenges persist. Moisture and oxygen sensitivity in certain materials compromise long-term stability and self-absorption effects and intricate synthesis protocols hinder large-scale production. Future research is expected to prioritize multifunctional solutions: enhancing stability through hydrophobic ligand design, glass-ceramic encapsulation, or MOF-based confinement; leveraging high-throughput computational models to refine STE energy levels and carrier transport dynamics; and advancing scalable manufacturing techniques like inkjet printing and roll-to-roll processing to bridge laboratory innovations with industrial deployment. Furthermore, it is necessary to increase the solubility of materials to reduce the scattering problem of scintillator films. The transformative potential of copper-based nanoclusters lies in their synergistic integration of performance, flexibility and sustainability. Their application in low-dose medical imaging, industrial nondestructive testing, and deep-space radiation monitoring holds immense promise.

Rare-earth-doped scintillator materials have garnered considerable attention as next-generation candidates for radiation detection and imaging applications. The introduction of rare-earth ions, acting as efficient luminescence centers, markedly enhances energy transfer efficiency and scintillation yield, owing to their distinctive 4f–5d electronic transitions. This quantum efficiency improvement translates to superior radiation sensitivity and spectral resolution, which are particularly crucial for low-dose radiography and nuclear medicine diagnostics. The incorporation of rare-earth dopants not only improves radiation hardness but also enhances thermal stability through optimized lattice interactions, ensuring operational reliability in extreme environments ranging from cryogenic detectors to aerospace systems. Furthermore, the engineered energy level configurations of rare-earth ions enable tunable emission spectra spanning near-UV to near-IR wavelengths, facilitating spectral matching with various photodetectors. It is recommended to strengthen mechanistic investigations into multivalent rare-earth activator configurations to elucidate quantitative structure–property relationships between complex energy level structures and optoelectronic characteristics, thereby formulating design principles for targeted applications including medical imaging and particle physics instrumentation. Moreover, exploring longer fluorescence lifetime modulation techniques expands innovative implementations in emerging optoelectronics such as temporal information encryption and anti-counterfeiting. A multidisciplinary approach integrating multiscale simulations with advanced characterization methodologies should be prioritized to overcome current limitations in understanding structure–performance correlations of rare-earth scintillators. It is crucial to emphasize that research on lanthanum-doped scintillator materials requires the systematic reporting of comprehensive performance metrics. The prevalent practice of overemphasizing isolated parameters while neglecting other

critical performance indicators constitutes a detrimental research tendency that could impede the healthy development of this field.

Organic scintillators have undergone transformative advancements through strategic manipulation of excited-state dynamics and molecular architectures, positioning them as viable alternatives to conventional inorganic systems. Recent breakthroughs in TADF and RTP materials have effectively addressed the intrinsic limitations of triplet exciton utilization in organic systems. Some innovations, such as introducing heavy-atom effects to amplify spin–orbit coupling, achieve record-breaking LYs while enabling tunable emission mechanisms between TADF and phosphorescence. Despite these achievements, critical challenges persist, including radiation-induced degradation, oxygen quenching of triplet states, and scalability limitations arising from batch-to-batch performance variations. Looking ahead, the field must prioritize the development of hybrid organic–inorganic architectures to reconcile the low-Z limitations of organic matrices with the high stopping power of embedded heavy-element nanostructures. Furthermore, engineering environmentally robust systems, such as oxygen-insensitive RTP scintillators *via* steric encapsulation or crosslinked polymer networks, will be essential for real-world deployment in medical imaging and aerospace environments. To bridge the gap between laboratory innovation and industrial adoption, concerted efforts must focus on establishing standardized benchmarking protocols and elucidating universal structure–property relationships governing triplet harvesting efficiency. Success in these endeavors could redefine performance paradigms across radiation detection technologies, ultimately enabling organic scintillators to transcend niche applications and compete at the forefront of high-energy physics, next-generation radiography, and portable radiation monitoring systems.

Beyond the development of high-performance scintillators, employing specialized structures to enhance the performance of scintillators has also emerged as a highly viable solution. The integration of scintillator materials with matrix materials featuring unique structures can significantly elevate the LY, stability, and detection resolution of the scintillators. This approach further enhances the potential of scintillators for practical applications. Moreover, DEXI offers a remarkably innovative and promising direction for the advancement of X-ray detection. This design provides the possibility of color recognition between objects of varying densities, thereby promoting the advancement of chromatic X-ray imaging in practical applications. However, studies on the impact of structural factors on optical performance have not been sufficiently developed. Future studies should prioritize the exploration of innovative optical architectures in X-ray scintillator systems, with particular emphasis on harnessing micro–nano-structural engineering to enhance photonic characteristics, thereby advancing scintillation performance and diversifying potential applications across radiation detection domains.

In summary, the shared purposes are to develop scintillators with flexibility, high LY, excellent stability, and high

resolution. We aspire for this review to enhance researchers' understanding of the overall progress in scintillator development, serving as a foundation for the creation of materials with superior performance, and more importantly, offering recommendations for future research directions. The primary purpose of this review is to revisit and update recent advancements in scintillators, while offering reliable recommendations for future developments in this field.

Data availability

No primary research results, software or code have been included and no new data were generated or analysed as part of this review.

Conflicts of interest

There are no conflicts to declare.

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