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Highly stable, substrate-free, and flexible broadband halide perovskite paper photodetectors†

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In this work, we aim to fabricate a highly stable and flexible perovskite paper photodetector based on a Zn-doped MA_{0.6}FA_{0.4}PbI₃ perovskite and CNC. The paper photodetector has been successfully synthesized by the vacuum filtration method and deposited with interdigitated electrodes. The paper photodetector exhibits a significant photoresponse with a responsivity of 0.23 A W⁻¹ under 650 nm light irradiation when operated at 5 V. The stability of the paper photodetector has also been tested and it shows high photoresponse after 30 days under ambient conditions. Therefore, this paper photodetector holds promise for developing efficient, stable, and flexible optoelectronic devices in the future.

Introduction

Halide perovskites have attracted extensive attention in recent years due to their outstanding properties in the development of various optoelectronic devices including photodetectors, solar cells, light-emitting diodes (LEDs), and field effect transistors (FETs).^{1–4} Despite their great performance in terms of a wide absorption range, narrow emission width, high wavelength tunability, ultrahigh quantum yield, and hysteresis-free behavior, the stability problem has remained a major challenge for their practical application.^{5–7} Halide perovskite materials suffer from various instability problems including structural, interfacial, and environmental instabilities. Recently, many studies have been carried out to improve the interface stability of perovskite light emitting/conversion or photodetection devices through various passivation methods. For perovskite nanocrystal-based light emitting devices, conventionally used capping ligands such as oleic acid and oleylamine exhibit a dynamic nature which makes them highly sensitive to moisture. Therefore, strategies such as phthalimide passivation, benzoic acid treatment, and ascorbic acid post-treatment have

been applied to increase the stability as well as the quantum yield of perovskite nanocrystals.^{8–14} For photodetection devices, small organic molecules such as *sec*-pentanol have been used as polar additives to improve the crystallization of the perovskite films and reduce the grain boundaries.¹⁵ A fluorinated polymer has also been used on the perovskite film as a light protective layer.¹⁶ Ye *et al.* used the amino groups of 1*H*,1*H*-perfluorooctylamine (PFA) to bind with the Pb²⁺ of perovskite microwires. The hydrophobic fluorocarbon alkyl chains in PFA can prevent the perovskite microwires from moisture. The PFA-modified device retained 80% of its initial performance after 30 days under ambient conditions.¹⁷ Ma *et al.* mixed poly[48-bis(5-(2-ethylhexyl)thiophen-2-yl)benzo[1,2-*b*;4,5-*b'*]dithiophene-2,6-diyl-*alt*-(4-(2-ethylhexyl)-3-fluorothiopheno[3,4-*b*]thiophene-2-carboxylate-2,6-diyl)] (PCE10) and polystyrene (PS) with the precursor solution. These polymers with a large dipole moment can suppress the ion migration of the device, resulting in stability enhancement.¹⁸

Paper is a flexible material that is light weight, biodegradable, and recyclable. Recently, various paper electronics including transistors, diodes, photodetectors, memory devices, photovoltaics, and even batteries have been developed.^{19–24} Therefore, with the development of paper electronics, the properties and processing of cellulose materials play important roles in the mechanical properties and electrical properties, as well as the biodegradability and environmental compatibility of the devices. Among various cellulose based materials, cellulose nanocrystals (CNCs) have been reported to exhibit a large surface area, a high aspect ratio, high mechanical strength, and adjustable surface chemistry.^{25,26} In addition, CNCs can be obtained through the acid hydrolysis of cellulose to form hybrids with other nanomaterials including nanocrystals, nanowires, and nanoplatelets. Previously, we have shown that CNC surface-passivated halide perovskites possess better stability and optical characteristics. The organosulfate groups on CNCs with high electronegativity can attach to the exposed Pb ions of halide perovskite nanocrystals and reduce the trapping states caused by halide vacancies.²⁷

In the development of flexible devices, most of the flexible optoelectronic devices are fabricated simply using flexible sub-

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strates such as polyimide (PI), polyethylene terephthalate (PET), polymethyl methacrylate (PMMA), or papers with electrodes, transport layers, and active layers on top of them.^{1,28} Therefore, additional packaging techniques should be applied to further protect the device from degradation. By adopting substrate-free fabrication techniques, electronic circuits and devices can be fully embedded within the matrix for further reducing the manufacturing waste, cost, and weight of the device. These properties are in high demand in applications such as e-skin and wearable electronics.²⁹ Recently, CNCs with controllable conductivity have been achieved by performing functionalization techniques to form functional groups.^{25,26} Moreover, hydrophilic and hydrophobic properties can also be adjusted through various functionalization techniques. Apart from the encapsulation of electronic devices, it is expected that CNCs can play an important role in improving the interface stability.

In our previous study, we have shown that the performance of CsPbI₃ perovskite quantum dots can be efficiently enhanced by doping Zn due to the improved structural stability of iodide-based perovskites.³⁰ Moreover, excess ZnI₂ can form an iodide-rich condition during paper fabrication to facilitate the formation of perovskite crystals. Herein, we combine the advantage of CNCs and Zn dopants to synthesize highly stable Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC papers. The technique can be applied to fabricate substrate-free and flexible photodetectors by directly depositing a metal electrode on the as-synthesized perovskite paper. The paper photodetector exhibits a significant photoresponse with a responsivity of 0.23 A W⁻¹ under 650 nm laser irradiation when operated at 5 V. The photoresponse retains 65% of its original value after 30 days under ambient conditions. In addition, the responsivity of the paper photodetector retains ~80% of its original value after 3000 bending cycles. Due to its high stability and flexibility, the perovskite paper photodetector shows great potential for long-term operation of photodetector devices.

Results and discussion

Fig. 1a shows the structure of the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper. The presence of organosulfate groups with high electronegativity on CNCs (–OSO₃[–]) can provide a strong binding strength to the exposed Pb on the surface of perovskite crystals and stabilize the structure of perovskites to achieve surface passivation. In addition, the long carbon chains and hydrogen bonds between CNCs can also prevent perovskites from oxygen and moisture-induced degradation. The photographic images of the bare CNC paper and the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper are shown in Fig. 1b. The bare CNC paper is highly transparent and the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper is extremely dark, showing great potential in light absorption. Fig. 1c shows the schematic of the fabrication process of the paper photodetector. The details of the experimental procedure are further explained in the Experimental section. The absorbance spectra and of the bare

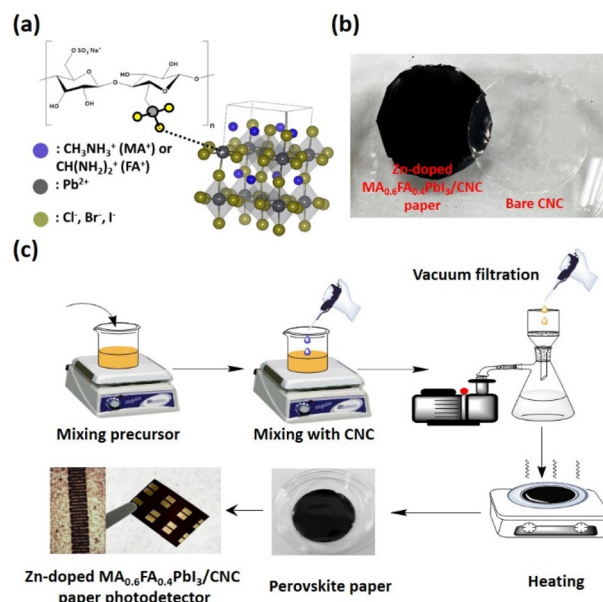


Fig. 1 (a) Schematic of a Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC structure. (b) Photographic images of bare CNC and Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC papers. (c) Experimental procedure and photograph of the paper photodetector.

CNC paper, the Zn-doped MAPbI₃/CNC paper, and the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper are shown in Fig. 2a. It can be observed that the absorbance of the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper exhibits a wider absorption wavelength range than those without FA⁺, indicating its broadband absorption behavior. The fluorescence spectra of Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC and Zn-doped MAPbI₃/CNC papers are also shown in Fig. 2a (dashed line). The emission peaks of Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC and Zn-doped MAPbI₃/CNC papers are around 809 nm and 762 nm, respectively. The optical band gap was determined using the Tauc plot (Fig. 2b) and the calculated value is around 1.49 eV. The X-ray diffraction (XRD) patterns of the bare CNC paper, Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper, and Zn-doped MA_{0.6}FA_{0.4}PbI₃ paper are shown in Fig. 2c. The diffraction peaks of the bare CNC paper at 15° and 22° correspond to the (110) and (200) planes of CNCs, respectively.^{27,30} The diffraction peaks of the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper are 15°, 21°, 25°, 28.4°, 31.8°, 35°, 40.5°, and 43°, which correspond to the (110), (112), (211), (220), (310), (312), (224), and (330) planes of the Zn-doped MA_{0.6}FA_{0.4}PbI₃ perovskite. However, the (110) and (211) peaks of the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper overlap with the (110) and (200) peaks of CNC due to the broader peak width of CNC.

The scanning electron microscopic (SEM) images of the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper and Zn-doped MA_{0.6}FA_{0.4}PbI₃ films are shown in Fig. 3a and b, respectively. The size of perovskite crystals in the Zn-doped MA_{0.6}FA_{0.4}PbI₃ films is around 2–8 μm. On the other hand, the size of perovskite crystals in the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper is

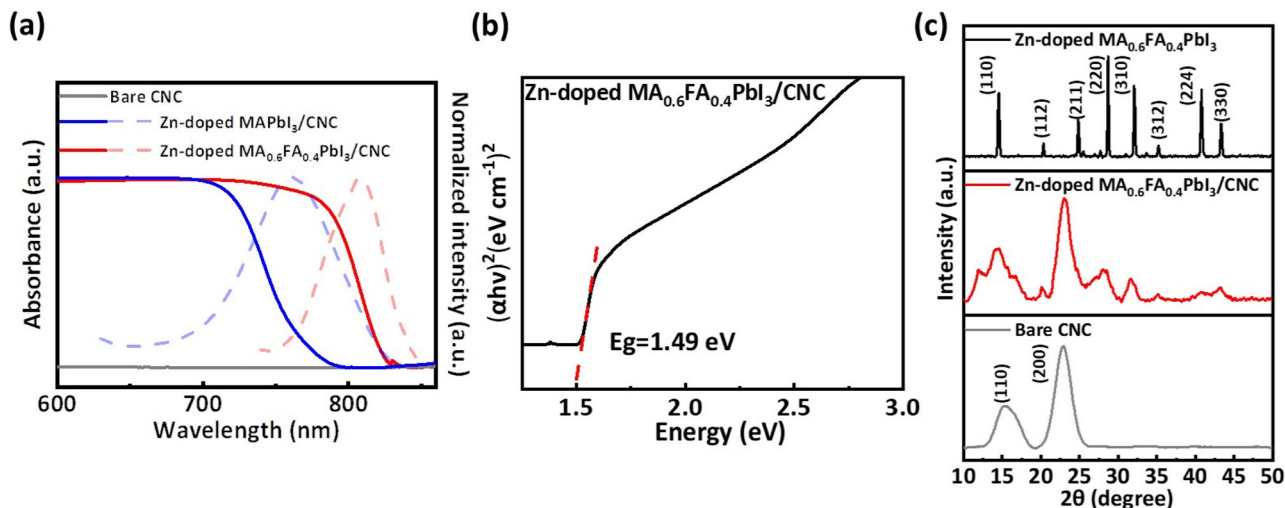


Fig. 2 (a) Absorbance spectra of bare CNC, Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC, and Zn-doped MAPbI₃/CNC papers and fluorescence spectra of Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC and Zn-doped MAPbI₃/CNC papers (dashed lines). (b) Tauc plot of the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper. (c) XRD patterns of bare CNC, Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC papers, and Zn-doped MA_{0.6}FA_{0.4}PbI₃ films.

<1 μm due to the growth limited by CNC. The transmission electron microscopy (TEM), high resolution transmission electron microscopy (HRTEM), and fast Fourier transform (FFT) images of the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper and Zn-doped MA_{0.6}FA_{0.4}PbI₃ are shown in Fig. S2.† The *d*-spacing of (220) in the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper has been found to be around 0.32 nm, which is consistent with that obtained in Zn-doped MA_{0.6}FA_{0.4}PbI₃ without CNC (0.318 nm).

For device fabrication, 5 nm/95 nm of Ti/Au was deposited on the paper through electron-beam evaporation with interdigitated electrode patterns defined by a shadow mask. The photographs of the paper before and after electron-beam evaporation are shown in Fig. S1a.† It can be observed that the appearance of the paper does not obviously change after the evaporation process. Moreover, the XRD patterns of the paper before and after electron beam evaporation were collected and are shown in Fig. S1b.† The diffraction peaks of the paper show similar patterns before and after the evaporation, indicating that the structure of the perovskite crystals does not change after the evaporation process.

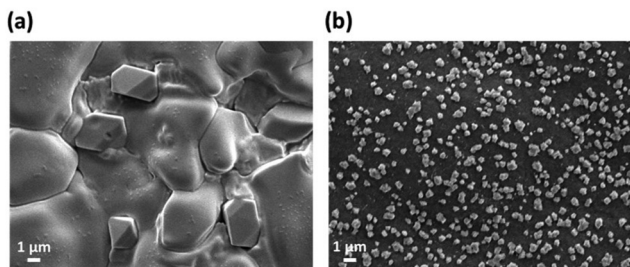


Fig. 3 (a) SEM image of the Zn-doped MA_{0.6}FA_{0.4}PbI₃ film. (b) SEM image of the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper.

Fig. 4a, b, and c show the characteristic curves of the paper photodetector under 450 nm, 520 nm, and 650 nm laser light irradiation, respectively. It can be observed that the current

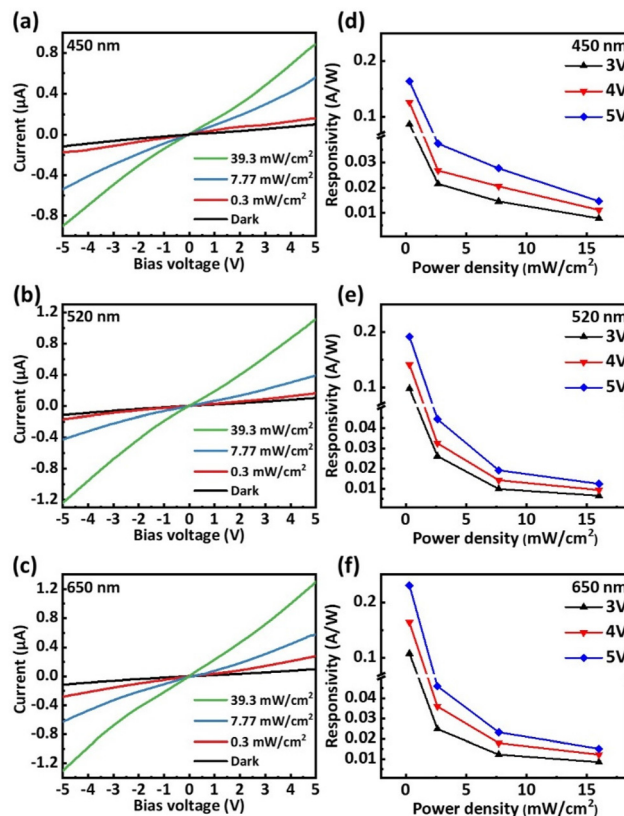


Fig. 4 *I*-*V* curves of the paper photodetector under dark and light illumination conditions at (a) 450 nm, (b) 520 nm, and (c) 650 nm laser irradiation. Power density dependent responsivity of the paper photodetector under (d) 450 nm, (e) 520 nm, and (f) 650 nm laser irradiation.

increases with increasing power density of the excitation laser. Fig. 4d, e, and f show the photoresponsivity as a function of power density and bias voltages (operated from 3 V to 5 V) under 450 nm, 520 nm, and 650 nm laser light irradiation, respectively. It can be observed that responsivity decreases rapidly with increasing power density due to the decreasing empty states in the conduction band. The highest photoresponse of 0.23 A W^{-1} (with 650 nm laser irradiation) can be observed under a power density of 0.3 mW cm^{-2} at an applied bias voltage of 5 V. As shown in Fig. 5a, the current–time ($I-t$) curves of the paper photodetector were obtained from 1 V to 5 V at an interval of 1 V under 650 nm (35 mW cm^{-2}) laser irradiation. Based on the results, the device exhibits good photocurrent stability, fast response, and consistent response in numerous ON–OFF cycles. Additionally, the paper photodetector exhibits a fast temporal photoresponse. The rise and fall time are $600 \mu\text{s}$ and $709 \mu\text{s}$, respectively (Fig. 5b).

In order to demonstrate the flexibility of the paper photodetector, bending test has been carried out and the schematic is shown in Fig. 6a. The paper photodetector is bended from flat to curved (angle of bend = 180°) and then back to flat as a bending cycle. The result of the bending test was recorded and is shown in Fig. 6b. The relative responsivity is defined as the responsivity divided by the initial responsivity. The responsivity of the paper photodetector remains at 94%, 87%, and $\sim 70\%$ of the original value after 500, 1000, and 3000 bending cycles, respectively. The curve shows a linear degradation after each bending cycle. Fig. 6c shows the stability test results of the bare Zn-doped $\text{MA}_{0.6}\text{FA}_{0.4}\text{PbI}_3$ film on a SiO_2/Si substrate and the paper photodetector. The responsivity of the paper photodetector retained 65% of its original value after 30 days under ambient conditions (temperature of 25°C and relative humidity of 70%), whereas the bare Zn-doped $\text{MA}_{0.6}\text{FA}_{0.4}\text{PbI}_3$ film on the substrate quickly degrades to 0% in only 3 days. Moreover, doping Zn into iodide-based perovskites can further enhance the stability. Fig. S3a and S3b† show the photographs of the Zn-doped $\text{MA}_{0.6}\text{FA}_{0.4}\text{PbI}_3$ film and the $\text{MA}_{0.6}\text{FA}_{0.4}\text{PbI}_3$ film under ambient conditions immediately after drop-coating (0 h) and after 20 h. The color of the

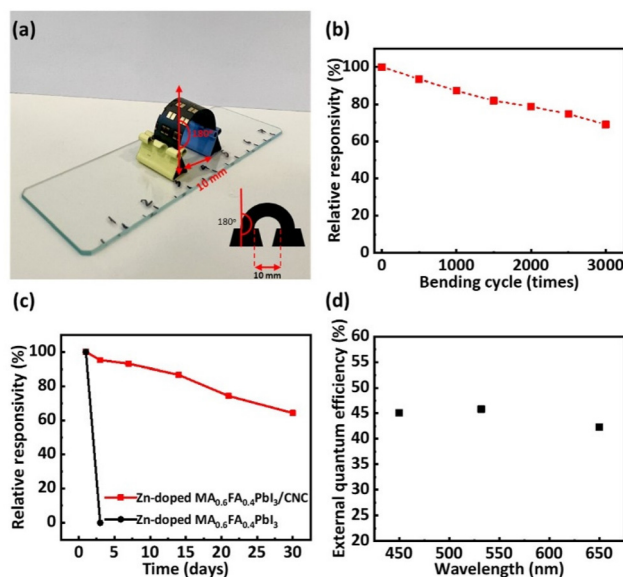


Fig. 6 (a) Schematic of the bending test of the paper photodetector. (b) Stability of the paper photodetector after various bending cycles. (c) Stabilities of the paper and bare Zn-doped $\text{MA}_{0.6}\text{FA}_{0.4}\text{PbI}_3$ photo-detectors under ambient conditions. (d) EQE of the paper photo-detectors under 450 nm, 520 nm, and 650 nm laser irradiation.

undoped $\text{MA}_{0.6}\text{FA}_{0.4}\text{PbI}_3$ film has changed after 20 h due to degradation. Therefore, the enhanced stability of the paper photodetector can be attributed to the combination of the long carbon chain of CNC, the strong binding strength between the organosulfate ($-\text{OSO}_3^-$) groups of CNC, and Zn dopants within the perovskite structure.

Fig. 6d shows the calculated external quantum efficiency (EQE) under 450 nm, 520 nm, and 650 nm laser irradiation using the following equation:

$$\text{EQE} = \frac{R \times hc}{\lambda \times e} \quad (1)$$

where R is the responsivity, h is the Planck constant, c is the light velocity, λ is the wavelength of incident light, and e is the elementary charge. The EQE values are 45.1%, 45.8%, and 42.2% under 450 nm, 520 nm, and 650 nm laser irradiation, respectively. Comparing the above-mentioned properties to those of other flexible photodetectors previously reported (Table 1), the paper photodetector in this work is substrate-free, it can save the cost of the substrate and also simplify the fabrication process. In addition, the paper photodetector shows a good value in responsivity comparing to the other works. In bending stability, the paper photodetector shows a perfect performance comparing to other works. Based on the result of comparison, the performance of the Zn-doped $\text{MA}_{0.6}\text{FA}_{0.4}\text{PbI}_3/\text{CNC}$ paper photodetector shows highly competitive and good potential for further applications.

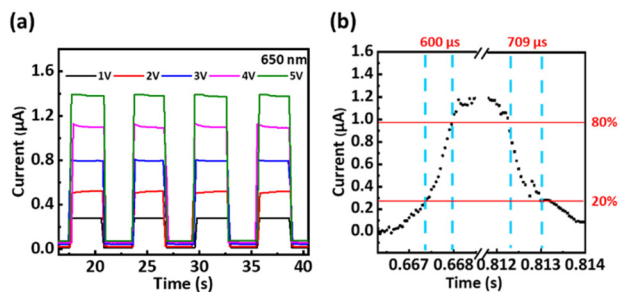


Fig. 5 (a) Time response at various applied voltages under 650 nm laser irradiation. (b) $I-t$ curve of the paper photodetector in a cycle.

Table 1 Photodetection properties and bending stability of various flexible photodetectors. T_r and T_f represent the rise time and the fall time of the devices, respectively

Substrate type	Devices	Bias voltage	Response range	R ($A W^{-1}$)	Bending cycles	Bending condition	Bending stability	Response time	Ref.
PI	Au/MAPbI ₃ /Au	0	300 nm	0.0022	—	$R = 3.7$ mm	—	$T_r = 27.2, T_f = 26.2$ (ms)	31
PET	Au/MAPbI ₃ /Au	10	650 nm	0.1	10 000	$\theta = 40^\circ$	90%	~0.3 ms	32
PEN	Au/P(VDF-TrFE)/MAPbI ₃ nanowires/Au	0	650 nm	0.012	100	$\theta = 45^\circ$	~75%	$T_r = 88, T_f = 154$ (μ s)	33
PET	ITO/MAPbI ₃ /Al	26	459 nm	0.004	—	$R = 5$ mm	66%	$T_r = 50, T_f = 120$ (μ s)	34
PET	Au/PbPc/MAPbI _{3-x} Cl _x /Au	50	655 nm	0.152	1000	$R = 6.5$ mm	~99%	$T_r = 632, T_f = 680$ (μ s)	35
						$\theta = 80^\circ$			
PEN	Au/MAPbI ₃ /RhB/Au	5	550 nm	0.0426	1000	$R = 9$ mm	92.7%	$T_r = 60, T_f = 40$ (ms)	36
PEN	Au/P(VDF-TrFE)/MAPbI ₃ /Au	0	650 nm	0.02	200	$\theta = 120^\circ$	~90%	$T_r = 92, T_f = 193$ (μ s)	37
Substrate-free	Au + Ti/MAPbI ₃ + CNC/Ti + Au	5	650 nm	0.23	1000	$R = 2$ mm	87%	$T_r = 600, T_f = 709$ (μ s)	This work
						$\theta = 180^\circ$			

Conclusions

In summary, we have demonstrated the synthesis, characterization, and fabrication of a Zn-doped MA_{0.6}FA_{0.4}PbI₃ perovskite paper photodetector. It is shown in the bending test that the paper photodetector retains 70% of its initial responsivity after 3000 bending cycles. In addition, the responsivity of the device retains 65% of its initial value after 30 days under ambient conditions. Moreover, the paper device exhibits a considerable responsivity of up to 0.23 A W⁻¹. As a result, the substrate-free, flexible, stable, broadband, and high responsivity paper photodetector holds the potential for the future development of optoelectronic and photosensing devices.

Experimental section

Chemicals

Methylammonium iodide (MAI, FMPV, 99%), formamidinium iodide (FAI, FMPV, 99%), zinc iodide (ZnI₂, SHOWA, 98%), lead iodide (PbI₂, ACROS, 99%), cellulose nanocrystals (CNC, Celluforce), dimethylformamide (DMF, Echo, 99%), and de-ionized (DI) water were used.

Preparation of the dispersed CNC solution

5 wt% of a CNC/DI water mixture was prepared by dissolving CNC powders in DI water. Then, an appropriate amount of the as-prepared CNC/DI water mixture was dispersed in DMF to prepare 1 wt% of dispersed CNC solution. The solution was kept at room temperature for subsequent procedures.

Preparation of the Zn-doped MA_{0.6}FA_{0.4}PbI₃ precursor solution

To prepare the Zn-doped MA_{0.6}FA_{0.4}PbI₃ precursor solution, 2.4 mmol of MAI, 1.6 mmol of FAI, 4 mmol of PbI₂, and 0.4 mmol of ZnI₂ were dissolved in 5 mL of DMF. The precursor solution was kept stirred until all the powders were completely dissolved. The precursor solution was then kept at room temperature for subsequent procedures.

Preparation of the Zn-doped MAPbI₃ precursor solution

To prepare the Zn-doped MAPbI₃ solution, 4 mmol of MAI, 4 mmol of PbI₂, and 0.4 mmol of ZnI₂ were dissolved in 5 mL of DMF. The precursor solution was kept stirred until all the powders were completely dissolved. The precursor solution was then kept at room temperature for subsequent procedures.

Preparation of the MA_{0.6}FA_{0.4}PbI₃ precursor solution

To prepare the MA_{0.6}FA_{0.4}PbI₃ precursor solution, 2.4 mmol of MAI, 1.6 mmol of FAI, and 4 mmol of PbI₂ were dissolved in 5 mL of DMF. The precursor solution was kept stirred until all the powders were completely dissolved. The precursor solution was then kept at room temperature for subsequent procedures.

Fabrication of the Zn-doped MA_{0.6}FA_{0.4}PbI₃ photodetector

1 mL of the Zn-doped MA_{0.6}FA_{0.4}PbI₃ precursor was directly dropped on the SiO₂/Si substrate and dried on a hot plate at 80 °C overnight. 5 nm/95 nm of Ni/Au was deposited on the paper through electron beam evaporation with interdigitated electrode patterns defined by a shadow mask.

Fabrication of the Zn-doped MA_{0.6}FA_{0.4}PbI₃/CNC paper photodetector

5 mL of the as-prepared precursor and 5 mL of the as-prepared dispersed CNC solution were mixed and stirred to obtain a homogeneous mixture. Afterwards, the mixture was poured into a Buchner funnel with a 0.02 μ m pore size ceramic filter to perform vacuum filtration. This procedure was kept on until all of the remaining solvent was drained for 30 min. Then the paper was dried at 60 °C in an oven for 24 h until the paper can be easily separated from the ceramic filter. For device fabrication, 5 nm/95 nm of Ti/Au was deposited on the paper through electron beam evaporation with interdigitated electrode patterns defined by a shadow mask. The active area of the device is ~0.27 mm² defined by the open area between the interdigitated fingers.

Characterization

The crystal structures of the papers were measured using a D2 phaser X-ray diffractometer (Bruker). The photoluminescence

spectra of the papers were recorded using a spectrofluorometer (JASCO FP8500). The transmittance and absorbance spectra of the papers were recorded using a UV-vis-NIR spectrophotometer (JASCO V670). The characteristic curves of paper photodetector were confirmed using a sourcemeter (Keithley 2612B) with a probestation and power tunable laser light sources (450 nm, 520 nm, and 650 nm). The light powers were measured using a power meter (LP10, SANWA, Japan). The transmission electron microscopy and high-resolution transmission electron microscopy images were collected using an FEI (Thermo Fisher Scientific), Talos F200XG2. The scanning electron microscope images were collected using a JEOL 7900F.

Author contributions

L. G. H. D. conceived this study and designed the experiments. L. G. H. D., C. H. C., and M. L. T. wrote the manuscript. L. G. H. D., S. M. S., T. Q. H. N., J. S., H. S. W., and Y. L. L. carried out the experiments. M. L. T. guided the project.

Conflicts of interest

There are no conflicts to declare.

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