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Solution-processed one-dimensional CsCu₂I₃ nanowires for polarization-sensitive and flexible ultraviolet photodetectors†

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Recently, much progress has been made in polarization-sensitive photodetectors based on the newly-emerging metal-halide perovskites. However, compared to extensive work on visible/infrared-light photodetection, polarization-sensitive perovskite photodetectors targeted at the ultraviolet (UV) region have not been reported. In addition, most of the previously reported photodetectors concentrated on lead-halide perovskites, but the lead-toxicity and poor stability severely restricted their practical applications. In this study, for the first time, one-dimensional CsCu₂I₃ nanowires (NWs) were used as the light absorber to fabricate a polarization-sensitive UV photodetector because of their electric and optical anisotropy of the asymmetric structure and external morphology anisotropy, and a high photocurrent anisotropy ratio of ~3.16 was achieved. Owing to the high crystallinity of the CsCu₂I₃ NWs, the device exhibits remarkable photodetection ability with a photoresponsivity of ~32.3 A W⁻¹, a specific detectivity of 1.89 × 10¹² Jones, and a fast response speed of 6.94/214 μs. Moreover, such a device fabricated on a flexible substrate shows almost no photodetection degradation after extreme bending for 1000 cycles, demonstrating good flexibility and bending endurance. The results obtained highlight the great potential of such copper halides as a stable and environmentally-friendly candidate for polarization-sensitive UV photodetectors, rendering them potentially useful for assembly of optoelectronic systems in the future.

Introduction

Recently, polarization-sensitive photodetectors have increasingly attracted research interest because of their significant

New concepts

Polarization-sensitive ultraviolet (UV) photodetectors are highly desired due to their great significance in many applications. As newly-emerging functional semiconductors, metal-halide perovskites have attracted huge attention in photodetection due to their excellent optoelectronic features. However, the previously reported perovskite polarized photodetectors mainly focused on the visible/infrared-light region, and devices targeted at the UV region have not been reported. Here, for the first time, ternary copper halide CsCu₂I₃ nanowires were used as the light absorber to achieve polarization-sensitive UV photodetection. By combining the intrinsic anisotropy of the asymmetric structure and external morphology anisotropy of CsCu₂I₃ nanowires, a record photocurrent anisotropy ratio of ~3.16 was achieved, and also the shortcomings of lead-toxicity and environmental instability for conventional lead-halide perovskites were overcome effectively. Moreover, such photodetectors fabricated on a flexible substrate demonstrate good flexibility and stability, showing almost no photoresponse degradation after extreme bending for 1000 cycles. The results suggest that the lead-free CsCu₂I₃ nanowires are potentially an attractive candidate for high-performance polarization-sensitive UV photodetectors compatible for practical applications.

applications like high-contrast polarizers, optical switches and interconnects, *etc.*,^{1,2} which require photoactive materials with outstanding optoelectronic characteristics, and anisotropic crystal structures or morphology. On the one hand, materials with unique anisotropic properties originating from asymmetric crystal structures have always been applied in polarization-sensitive photodetectors, such as black phosphorus,^{3–5} germanene,⁶ and

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antimony selenide.^{7,8} On the other hand, materials with one-dimensional (1D) morphology can be also used as photoactive materials for polarization-sensitive photodetection, resulting in linear polarization sensitivity.^{2,9,10} For example, by means of the order-of-magnitude polarization anisotropy of InP nanowires (NWs) resulting from the great dielectric contrast between the NWs and the surrounding environment, Wang *et al.* demonstrated a polarization-sensitive nanoscale photodetector based on isolated InP NWs.² A similar polarization-sensitive photoreponse was also observed in an aligned CdSe quantum-wire solid by Singh *et al.*¹¹

As newly-emerging functional semiconductors, the intrinsically large light absorption coefficient and long charge diffusion length provide metal-halide perovskites with broad prospects for high-performance photodetectors,^{12–20} and much progress in this field has been witnessed in the past three years.^{21–24} Very recently, polarization-sensitive photodetectors based on such perovskite materials have attracted much attention.^{9,25–32} For example, by using the anisotropic geometry of CH₃NH₃PbI₃ NWs, polarization-sensitive light detection was achieved by Tang's and Jiang's groups.^{9,32} Zhang *et al.* reported a polarization-dependent photodetector based on CsPbX₃ single-crystal NW arrays,²⁵ and the minimum photocurrent appears in the direction perpendicular to the axial direction, while the maximum photocurrent appears when the polarization angle is parallel to the NWs. For the above three studies, the polarization-sensitive photodetection was realized through the external anisotropy of the 1D geometry of perovskite NWs, which is due to the fact that tetragonal CH₃NH₃PbI₃ and cubic CsPbX₃ have symmetric crystal structures lacking intrinsic anisotropy. In reality, polarization-sensitive photodetection realized by the intrinsic anisotropy of the asymmetry crystal structure was also proved to be an effective strategy. For example, Ding *et al.* demonstrated a CH₃NH₃PbCl₃ perovskite single crystal with an anisotropic crystal structure for polarized light detection.²⁶ By using anisotropic orthorhombic CsPbI₃ NWs, Zhou *et al.* fabricated a linearly polarized visible photodetector with a photocurrent anisotropy ratio of 2.68.²⁷ More recently, Ghoshal *et al.* demonstrated the potential of 2D layered (BA)₂PbI₄ perovskite NWs with anisotropic crystal structure for polarized light detection.²⁸ But, so far, the previously reported devices have mainly focused on visible/infrared-light detection, and polarization-sensitive perovskite photodetectors targeted at the ultraviolet (UV) region are rarely reported. Moreover, another important factor is that most of the previous studies use conventional lead-halide perovskites, and their environmental instability and toxicity limit their further commercialization.^{33–35} Thus, it is imperative to search for environmentally stable and lead-free alternatives as the photoactive material targeted at the UV region to solve the above problems.

Herein, we demonstrated a polarization-sensitive UV photodetector based on solution-processed 1D CsCu₂I₃ NWs. By combining the intrinsic anisotropy of the asymmetric structure and external morphology anisotropy of the CsCu₂I₃ NWs, a record photocurrent anisotropy ratio of ~3.16 was achieved

among perovskite-based polarized photodetectors. Owing to the high crystallinity of the CsCu₂I₃ NWs, the device performance is remarkable in terms of a photoresponsivity of ~32.3 A W⁻¹, a specific detectivity of 1.89 × 10¹² Jones, and a fast response speed of 6.94/214 μs. Moreover, such devices fabricated on a flexible polyethylene terephthalate (PET) substrate demonstrate good flexibility and stability, showing almost no photoresponse degradation after 1000 bending cycles. Given the record polarization sensitivity and high stability, the lead-free copper halides might hold promise for high-performance polarization-sensitive UV photodetectors compatible for practical applications.

Results and discussion

Fig. 1a illustrates the preparation procedures of 1D CsCu₂I₃ NWs by using antisolvent engineering, as reported for conventional lead-halide perovskites, and diethyl ether was employed as the antisolvent. Note that the diffusion of diethyl ether in acetonitrile will consume some acetonitrile solvent in the precursor solution. Since CsI and CuI cannot dissolve in diethyl ether, the consumption of acetonitrile will increase the concentration of CsI and CuI in the solution. This will result in the crystallization and formation of CsCu₂I₃ crystals, and detailed parameters are shown in the Experimental section. The right panes in Fig. 1a present the corresponding dried powders under sunlight and UV light irradiation, which are composed of isolated CsCu₂I₃ NWs with a large scale. Since the crystallization rate for antisolvent engineering is very fast, so the dropping speed of diethyl ether is very important for the crystallinity and morphology characteristics of the products. In the present case, the width and length of the CsCu₂I₃ NWs can be adjusted within a large range from the nanometer (35 nm) to the micrometer (6.0 μm) scale by changing the dropping speed of diethyl ether, as displayed in Fig. S1 (ESI†), and the length of the wires could be up to 0.1 mm. Fig. 1b shows a typical morphology of the as-synthesized CsCu₂I₃ NWs, which were measured by scanning electron microscopy (SEM) to have a length of 7.4 μm and width of 350 nm. Fig. 1b shows the elemental mapping examined by energy-dispersive X-ray spectroscopy (EDS), in which the elements are homogeneously distributed throughout an individual wire. Fig. S2 (ESI†) shows the corresponding EDS spectrum, which gives a quantified atomic ratio (%) of 1:1.95:2.83 for Cs:Cu:I, close to the stoichiometry of CsCu₂I₃. Atomic force microscopy (AFM) measurements were further performed to investigate the morphology characteristics of the as-synthesized CsCu₂I₃ NWs. As shown in Fig. 1c, the 1D CsCu₂I₃ NW is characterized by a homogeneous height of ~100 nm and a smooth top surface. Fig. 1d and g present typical transmission electron microscopy (TEM) images of CsCu₂I₃ wires with a straight boundary and uniform widths of ~35 nm and 4.0 μm. The high-resolution TEM image (Fig. 1e) taken from the main body marked in the wire reveals clear lattice fringes, and the interplanar distance is 0.341 nm, corresponding to the (221) lattice plane of the orthorhombic CsCu₂I₃ crystal. The above results together with



Fig. 1 (a) Schematic of the processing procedures of the CsCu_2I_3 NWs. (b) SEM image of an individual NW, and the corresponding EDS mapping showing the uniform distribution of Cs, Cu, and I elements. (c) AFM image of an individual CsCu_2I_3 wire. TEM images of the CsCu_2I_3 NWs with a width of (d) 35 nm, and (g) 2 μm . (e) High-resolution TEM image of the CsCu_2I_3 NW as marked in (d), and the corresponding (f) SAED pattern.

the selected area electron diffraction (SAED) pattern shown in Fig. 1f confirm the single-crystalline property of the CsCu_2I_3 NWs.

The orthorhombic structural characteristics of the produced CsCu_2I_3 NWs were also tested by X-ray diffraction (XRD). As seen in Fig. 2a, four dominant diffraction peaks at 10.88° , 21.71° , 32.74° , and 44.09° were observed, which can be assigned to the (110), (220), (330), and (242) planes of orthorhombic CsCu_2I_3 (space group $Cmcm$, $a = 10.505 \text{ \AA}$, $b = 13.147 \text{ \AA}$, $c = 6.072 \text{ \AA}$, $\alpha = \beta = \gamma = 90^\circ$). No other impurity phases were found, further indicating the good crystallinity of the produced CsCu_2I_3 NWs. Fig. 2b shows the crystal structure diagram of CsCu_2I_3 , which is composed of a 1D chain structure formed by edge-sharing $[\text{CuI}_4]^-$ dimerized units. Each $[\text{CuI}_4]^-$ tetrahedron is linked by three common edges, forming parallel rod bundles. The Cs^+ ions are intercalated between the chains, which are coordinated by eight I^- ions in the form of double-capped

trigonal prisms. The crystal structures of CsCu_2I_3 from the top and left view indicate that CsCu_2I_3 has an anisotropic crystal structure and could easily grow into a 1D linear morphology, which provides the possibility of designing a photodetector by using its anisotropic character. Further, the optical properties of the CsCu_2I_3 NWs were studied by UV-visible absorption and photoluminescence (PL) spectra. As can be seen from Fig. 2c, a clear absorption edge at around 332.4 nm was observed, and the corresponding Tauc plot of the absorption spectrum (Fig. S3, ESI †) reveals that CsCu_2I_3 possesses a direct bandgap of 3.73 eV, which is consistent with other articles.^{36–38} The blue-dotted line displays the PL spectrum of a single CsCu_2I_3 NW at an excitation wavelength of 330 nm, and an emission peak centered at 542 nm was observed, and the linewidth is 112 nm. It should be mentioned that there is a large Stokes shift of $\sim 209.6 \text{ nm}$, which implies that the PL mechanism can be

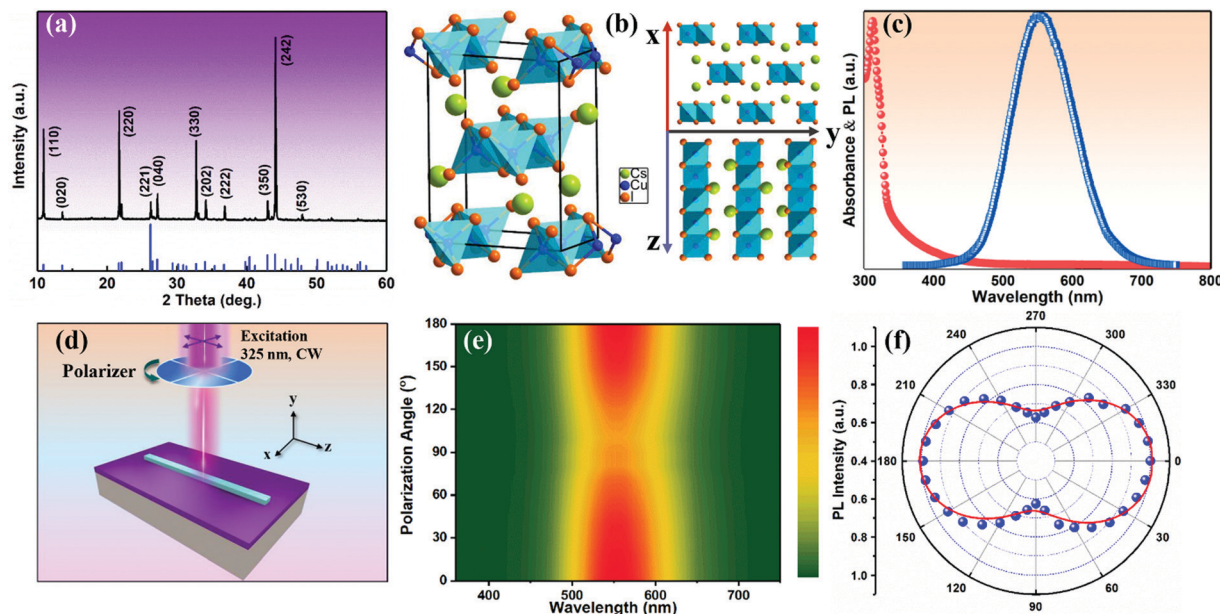


Fig. 2 (a) Experimental XRD spectrum (top) of the as-grown CsCu_2I_3 NWs and the standard XRD pattern (bottom) for the orthorhombic CsCu_2I_3 . (b) Crystal structure of CsCu_2I_3 from side, top, and left views. (c) Steady-state PL and absorption spectra of the CsCu_2I_3 NWs. (d) Schematic diagram of polarization PL measurements. (e) PL spectra as a function of the incident light polarization. The color represents the emission intensity. (f) Polarization-dependent PL intensity of 1D CsCu_2I_3 NWs. The data (blue dots) are fitted with a cosine (red) function.

explained by excited-state structural reorganization or exciton self-trapping rather than a direct band-to-band transition, and this behavior was frequently observed in low-dimensional metal-halide perovskites.^{39–41} To better study the optical recombination mechanisms of CsCu_2I_3 , temperature-dependent PL measurements were conducted. As shown in Fig. S4 (ESI[†]), an obvious thermal quenching behavior of the PL intensity with increasing the temperature can be observed. Fig. S4b (ESI[†]) shows the relationship between the integrated PL intensity and the reciprocal temperature, and the thermal quenching behavior can be fitted by the following formula:

$$I(T) = \frac{I_0}{1 + A \exp\left(-\frac{E_b}{k_B T}\right)} \quad (1)$$

where I_0 is the emission intensity at 0 K, A is a proportionality constant, and k_B is the Boltzmann constant. The fitting curve produces a value of 201.7 ± 6.5 meV for the exciton binding energy (E_b). Further, we measured the PL excitation spectra of CsCu_2I_3 NWs at different emission wavelengths from 470 to 670 nm. As shown in Fig. S5 (ESI[†]), the PL excitation spectra show the same properties and shapes, indicating that the PL at 542 nm is caused by the relaxation of the same excited state instead of the defect recombination centers. Considering the anisotropic properties of the CsCu_2I_3 NWs, polarization-dependent PL measurements were therefore performed (one NW with a width of 100 nm was taken as an example) by using a continuous-wave laser (325 nm) as the excitation source, and the schematic diagram of such a measurement configuration is drawn in Fig. 2d. Note that 0° was defined as the polarization angle parallel to the NWs (the z -axis labeled in Fig. 2b).

Fig. 2e shows a color plot showing the variation in the PL intensity *versus* the polarization angle. One can see that the emission at around 542 nm shows the maximum intensity at an angle of 0° , and the PL intensity decreases gradually with changing the polarization angle and reaches the minimum at 90° polarization (perpendicular to the wire). The red curve in Fig. 2f shows the cosine fitting of the PL intensity with a period of 2π , and the PL intensity shows a polarization ratio of $\sim 1.7:1$. It should be noted that the polarization ratio herein results from the anisotropic crystal structure and anisotropic morphology simultaneously. An additional observation is that the polarization ratio is inversely proportional to the width of the NWs. As shown in Fig. S6 (ESI[†]), a relatively small polarization ratio of ~ 1.37 was achieved for the CsCu_2I_3 NW with a width of $6.0 \mu\text{m}$, which may be due to the effect of electrostatic dielectric confinement.²⁸

The as-synthesized 1D CsCu_2I_3 NWs with high crystallinity and strong emission anisotropy suggest their potential as a photoactive material for polarization-sensitive photodetectors. In our case, a metal–semiconductor–metal type photodetector was constructed by transferring a single CsCu_2I_3 NW to a silicon oxide substrate with Au as the contact electrode, as schematically illustrated in Fig. 3a, and the real device photograph is shown Fig. 3b. Note that the length of the NW used for device fabrication is about $90 \mu\text{m}$, and the width is about 600 nm . The energy band alignment of Au/ CsCu_2I_3 was drawn according to the obtained UV-visible absorption spectra together with the UV photoelectron spectroscopy (UPS) data (Fig. S7, ESI[†]), and a Schottky barrier formed at the hetero-interface. The formation of the Schottky barrier can suppress the leakage current, reduce the dark current, and improve the response speed of the photodetector.

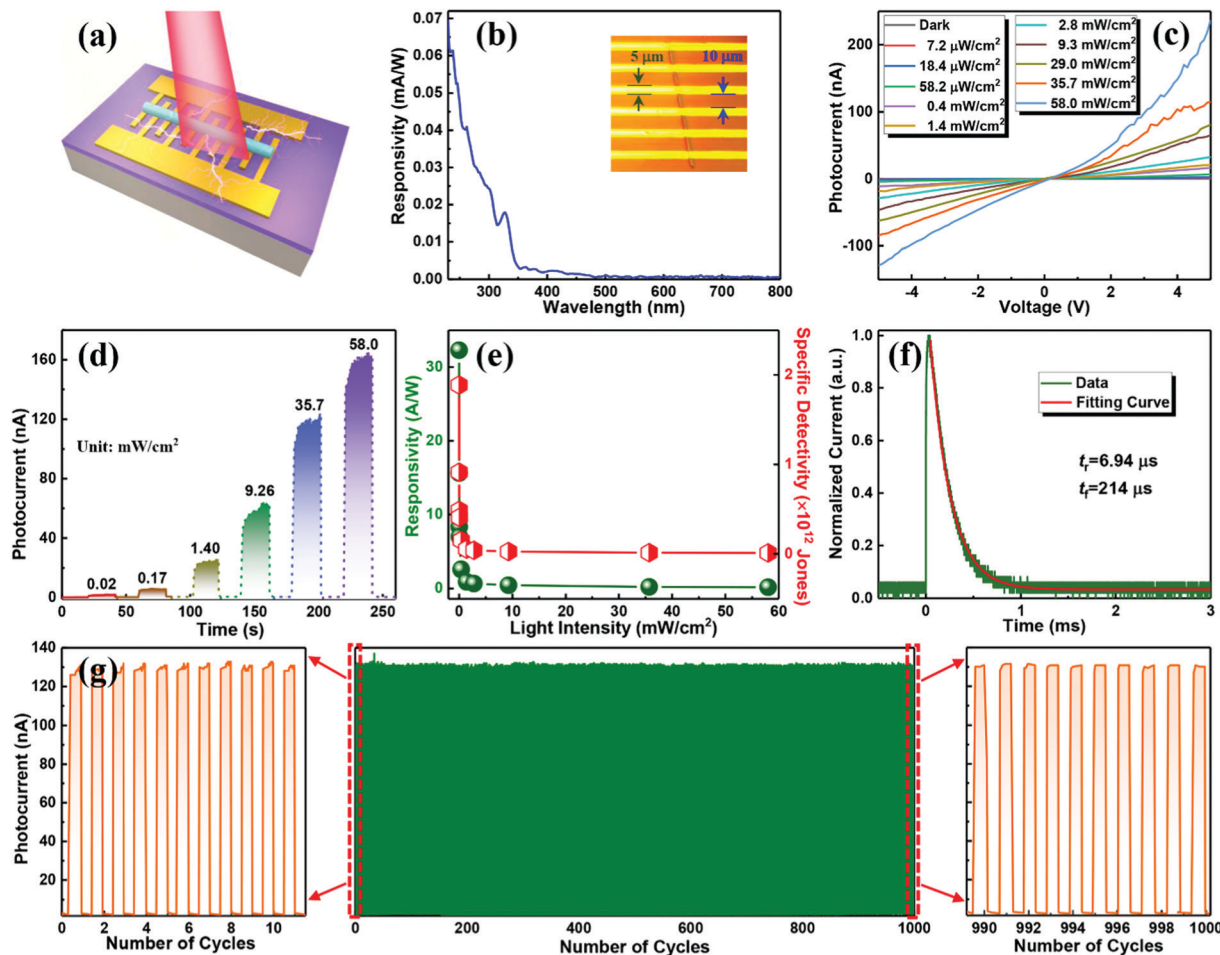


Fig. 3 (a) Schematic illustration of the photodetector based on an individual CsCu_2I_3 NW. (b) Spectral response of the photodetector. The inset shows an optical microscope image of the photodetector. (c) I - V and (d) I - t curves of the devices with different light irradiation intensities. (e) Responsivity and specific detectivity of the device *versus* light irradiation power. (f) Temporal response under 266 nm pulsed laser excitation with a pulse width of 1 ns. (g) One thousand continuous response cycles of the device with a period of 1 s.

The spectral response in the range of 230–800 nm was firstly tested to achieve the response sensitivity of the photodetector. As seen in Fig. 3b, the device exhibits a photoresponse ranging from 230 to 350 nm, demonstrating the same trend as the absorption spectrum of CsCu_2I_3 and also suggesting a bandgap excitation process of CsCu_2I_3 .^{42,43} Fig. 3c exhibits the current *versus* voltage (I - V) characteristics of the photodetector tested in the dark and under light excitation (325 nm) with different irradiation intensities. Owing to the intrinsic low carrier concentration of copper halides, the device displays a rather low dark current of 6.3×10^{-11} A under a bias of 5 V, which is beneficial for high detectivity of such device. Following the light illumination, the current increases gradually because the number of photogenerated carriers increases with the absorbed photon flux. In the present case, the I - V curves possessed an asymmetrical behavior, which may be caused by the undesired interface states at the $\text{CsCu}_2\text{I}_3/\text{Au}$ interface and also the formation of a Schottky barrier. In addition, the curves between photocurrent (I_p) and the light intensity can be fitted by the power law of $I_p = \alpha \cdot P^\beta$, where α represents a proportionality constant,

P is the light intensity, and β is an empirical value that reflects the recombination of photoexcited carriers. As plotted in Fig. S8 (ESI[†]), fitting the curve in a wide light intensity range gives rise to a value of $\beta = 0.71$. The nonunity exponent of the law is due to the complex process of electron-hole generation, trapping, and recombination activity within the CsCu_2I_3 NW photodetector.⁴⁴ The current-time (I - t) curves of the photodetector *versus* varied excitation intensity were also measured, where the pulse bandwidth is 20 s per cycle and the bias voltage is 5 V. As shown in Fig. 3d, the current reversibly switches between high and low conductance with excellent reproducibility and stability, generating a series of tunable on/off ratios. Under a light power of 58.0 mW cm^{-2} , the on/off ratio reaches 2.6×10^3 , which is comparable with the photodetectors based on conventional lead-halide perovskites.^{45–48} Besides, the photoresponsivity (R) and specific detectivity (D^*) are key parameters to quantitatively characterize the photoresponse ability to an optical signal, which can be expressed using the following equations:

$$R = (I_p - I_d)/P_{\text{opt}} \cdot A \quad (2)$$

$$D^* = \frac{A^{1/2}R}{(2eI_d)^{1/2}} \quad (3)$$

in which I_p represents the photocurrent, A represents the active area ($A = 54 \mu\text{m}^2$), and P_{opt} is the incident light intensity. Based on the equations, the relationships of R and D^* with the light intensity at a bias of 5 V are summarized in Fig. 3e, and the maximum values of R and D^* for such a photodetector were calculated to be 32.3 A W^{-1} and 1.89×10^{12} Jones ($1 \text{ Jones} = 1 \text{ cm Hz}^{1/2} \text{ W}^{-1}$), respectively. Note that both R and D^* decrease with increasing light intensity; this may be due to the enhanced carrier recombination possibility at higher light intensity.^{49–51} Besides, we investigated the response time of the CsCu_2I_3 NW-based photodetector by illuminating it with a 266 nm pulsed laser (pulse width 1 ns, 1000 Hz). One can observe that the device could monitor and respond to each pulsed signal (Fig. S9, ESI[†]), producing a rise time (t_r) of 6.94 μs defined as the time for the current to increase from 10% to 90% of the peak value, as seen in Fig. 3f. In addition, by fitting a time-dependent photocurrent curve at the falling edge using an exponential decay curve, the fall time (t_f) of the device is

estimated to be $\sim 214 \mu\text{s}$. The fast response speed for such a device is believed to be related to the high crystallinity of the CsCu_2I_3 NWs, which ensures few grain boundaries, a low defect concentration, and a small carrier recombination rate, and therefore facilitates the fast transport of charged photo-generated carriers.^{47,52,53} Moreover, the long-term working stability of the CsCu_2I_3 NW-based device was investigated. By using a mechanical chopper with a frequency of 1 Hz to switch the incident light (325 nm , 36 mW cm^{-2}), the photocurrent signals were obtained by an oscilloscope in real time with a bias voltage of 5 V. As can be seen in Fig. 3g, the photodetector can maintain its photoresponse ability after 1000 consecutive response cycles, revealing good stability and repeatability.

By virtue of the flexible and deformable structural framework of the 1D NWs, the electrical stability of such a photodetector fabricated on a flexible PET substrate in the bending state was further studied for practical application. As displayed in the insets of Fig. 4a, the device fixed in a vernier caliper was bent at different angles (0° , 60° , 90° , 120° , and 160°) to test its flexibility, and the light intensity was kept at 0.2 mW cm^{-2} for each cycle (100 s). As shown in Fig. 4a, the current remains almost unchanged under

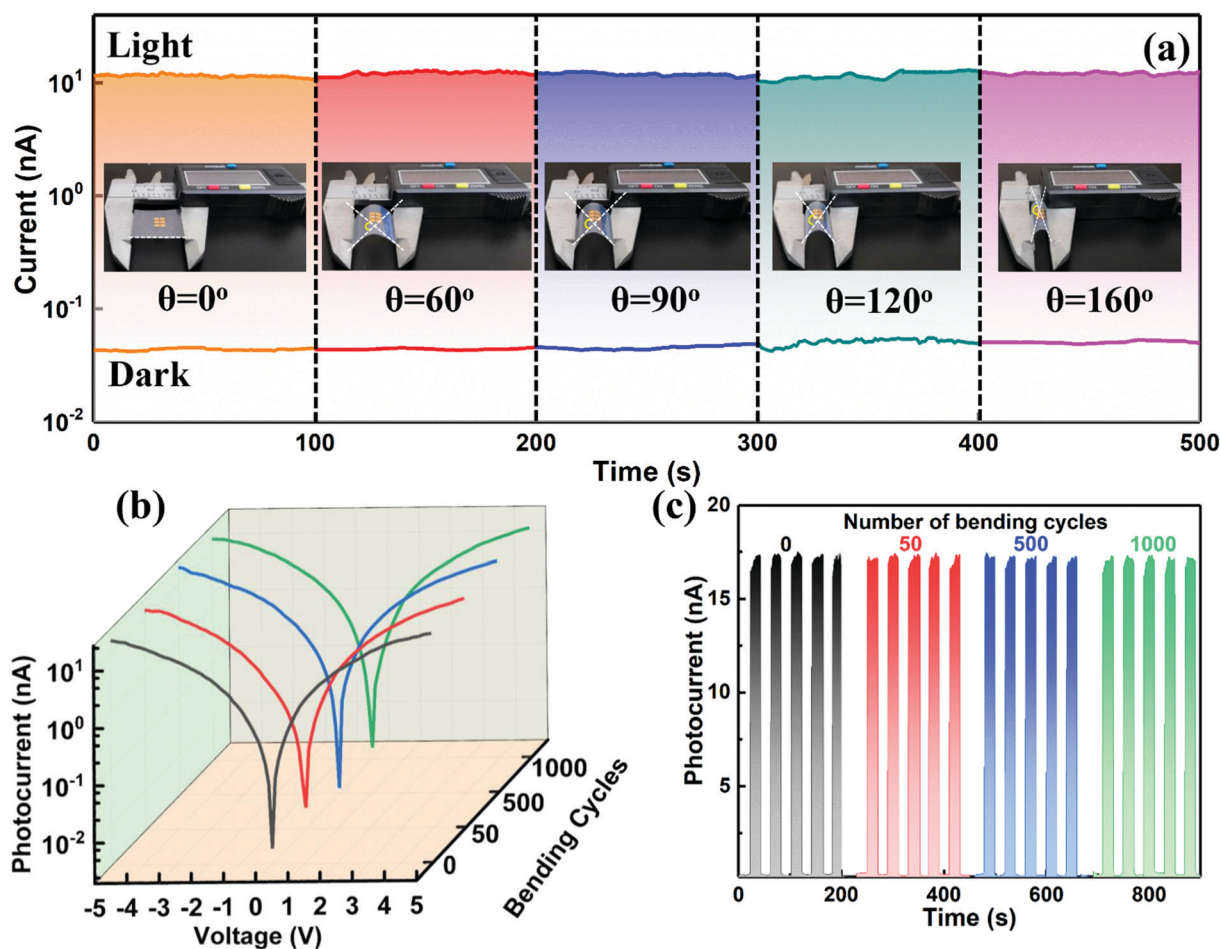


Fig. 4 (a) Output currents of the device with different bending angles under dark and light illumination with a light intensity of 0.2 mW cm^{-2} at a bias voltage of 5 V. The insets are the corresponding optical images of the device under five bending states. θ is the bending angle. (b) I - V and (c) I - t curves of the device under different bending cycles.

various bending angles, revealing that this flexible photodetector was not influenced by the external bending stress. The excellent electrical stability of the device is also evidenced by the logarithmic I - V curves and I - t period curves under various bending angles, as illustrated in Fig. S10 (ESI[†]). To further investigate the electrical stability and folding endurance of the flexible photodetector, the logarithmic I - V and I - t curves after different bending cycles with a bending angle of 90° were acquired and put together for comparison. As shown in Fig. 4b and c, the photocurrent, dark current, and photo-switching characteristic of the device were kept very well even after an extreme 1000 bending cycles. The highly flexible photodetector thus opens up opportunities for fundamental research on 1D lead-free perovskites and may greatly expand their application fields especially in portable devices.

As we all know, in order to realize high-precision detection in complex environments, more information on the light is needed, such as polarimetric information.^{54–56} The strong anisotropy of 1D CsCu_2I_3 NWs promotes their applications as a suitable photoactive material for polarization-sensitive photodetectors. In view of this, the incident light was set to pass through a polarizer firstly and then illuminate the NW, as illustrated in Fig. 5a, and the measurements were performed by rotating the polarizer every 10° to change the polarization direction while keeping the light power (58 mW cm^{-2}) constant. Fig. 5b depicts the remarkable anisotropic photo-response of the photodetector *via* a 2D colormap within a bias from 0 to 1.0 V. Obviously, upon linearly polarized light

excitation, the photocurrent of the device is a strong function of the angle between the polarization of the incident light and the orientation of the NW, as well as the bias voltage. Similar to the polarization-dependent PL spectra, the photocurrent reaches the maximum when the polarization of the light is parallel to the wire (0°) and reaches the minimum when it is perpendicular to it (90°). Then the photocurrent increases with the polarization angle and recovers totally at 180° polarization. The evolution of current at a bias voltage of 1.0 V was further extracted as a function of the polarization angle in polar coordinates, as plotted in Fig. 5c, and a high photocurrent anisotropy ratio of about 3.16 was achieved (defined as the photocurrent tested at a polarization of 0° divided by the photocurrent tested at 90°), which is the maximum value among the perovskite-based polarized photodetectors as far as we know, and also superior to previous studies on 2D materials, such as GeSe (2.16), GeS_2 (2.1), GeAs_2 (2.0), and so on,^{54,57–59} as summarized in Table 1. Fig. 5d depicts the corresponding I - t curves of the photodetector as a function of the polarization angle, and an obvious polarization dependence was observed, consistent with the above discussions. Besides, another photodetector with a wider CsCu_2I_3 wire of $6.0 \mu\text{m}$ was used for identical measurements. From the polarization angle-dependent curve plotted in Fig. 5e, a relatively smaller photocurrent anisotropy ratio of ~ 1.24 was obtained. This trend with smaller width showing higher polarization sensitivity is well consistent with the above polarization-dependent PL results.



Fig. 5 (a) Schematic illustration of the polarization-sensitive photodetection based on 1D CsCu_2I_3 NWs. (b) Anisotropic response in photocurrent under 325 nm light excitation described *via* a 2D colormap (photocurrent is denoted by the color bar, with voltage as the x -axis and polarization angle as the y -axis). Polarization dependence of the photocurrent for NWs with different widths of (c) 600 nm, and (e) $6.0 \mu\text{m}$. (d) Photocurrent response of a 1D CsCu_2I_3 NW device under incident light with different polarization angles.

Table 1 Summary of the performances of the polarization-sensitive photodetectors

| Materials | Response range (nm) | Lead-free (yes/no) | Responsivity ($A W^{-1}$) | D^* ($\times 10^{12}$ Jones) | Polarization ratio | Ref. |
|----------------------------------|---------------------|--------------------|-----------------------------|---------------------------------|--------------------|-----------|
| $CH_3NH_3PbI_3$ nanowires | 400–800 | No | 4.95 | 20 | 1.3 | 9 |
| $CH_3NH_3PbCl_3$ single crystals | 350–450 | No | 0.0011 | — | 2.25 | 26 |
| $CH_3NH_3PbI_3$ nanowires | 400–800 | No | 1000 | — | 2.6 | 32 |
| $CsPbBr_3$ nanowires | 400–530 | No | 0.0115 | — | 0.78 | 25 |
| $CsPbI_3$ nanowires | 400–800 | No | 0.745 | 0.035 | 2.68 | 27 |
| $(BA)_2PbI_4$ nanowires | 480–600 | No | — | — | 3.62 | 28 |
| $CsPbCl_3$ microcrystals | 300–520 | No | 0.11 | 0.024 | 3 | 29 |
| Black phosphorus flake | 400–3750 | Yes | 3.5×10^{-4} | — | 3.5 | 4 |
| $GeAs_2$ flake | 400–700 | Yes | — | — | 2 | 57 |
| $GeSe$ flake | 400–950 | Yes | — | — | 2.16 | 58 |
| GeS_2 flake | 250–350 | Yes | — | — | 2.1 | 59 |
| ZnO | 400–600 | Yes | — | — | 1.13 | 60 |
| $CsCu_2I_3$ nanowires | 250–350 | Yes | 32.2 | 1.89 | 3.16 | This work |

Conclusions

In conclusion, high-quality lead-free 1D $CsCu_2I_3$ NWs were successfully prepared by an antisolvent engineering approach. The polarization-dependent PL spectra confirm the optical anisotropy of the $CsCu_2I_3$ NWs, which results from their intrinsic anisotropy of the asymmetric structure and external morphology anisotropy. Furthermore, we demonstrated a polarization-sensitive photoconductive detector with an individual $CsCu_2I_3$ NW as the photoactive material. A record photocurrent anisotropy ratio of ~ 3.16 was achieved, which is the maximum value among perovskite-based polarized photodetectors. Moreover, the photodetector exhibits a good detection performance, a large photoresponsivity of $\sim 32.2 A W^{-1}$, a high specific detectivity of 1.89×10^{12} Jones, and a fast response speed of $6.94/214 \mu s$. In addition, the photodetector fabricated on a flexible substrate demonstrates good flexibility and stability, showing almost no photoresponse degradation after 1000 bending cycles. The results obtained highlight the great potential of such lead-free $CsCu_2I_3$ NWs as an attractive candidate to be applied in polarization-sensitive UV photodetectors.

Experimental section

Preparation of $CsCu_2I_3$ NWs

0.7 mmol CsI and 1.4 mmol CuI were added to 4 mL anhydrous acetonitrile. The precursor solution was prepared by stirring at $60^\circ C$ for 5 h. Then, the precursor solution was naturally cooled down to room temperature. After that, the saturated solution was filtered using a syringe filter of PTFE ($0.45 \mu m$). Then, the solution was titrated with anti-solvent (diethyl ether) to precipitate $CsCu_2I_3$ wires. By changing the dropping speed of diethyl ether, high-quality 1D $CsCu_2I_3$ NWs with different sizes can be obtained.

Fabrication of devices

For polarization-sensitive photodetector fabrication, the as-prepared $CsCu_2I_3$ NWs were transferred onto a silicon oxide substrate with Au patterned fingers as the contact electrode, and the Au contact electrode fingers were $5 \mu m$ wide and $500 \mu m$ long, and the channel

spacing was $10 \mu m$. For the flexible photodetector fabrication, a commercially available PET substrate was employed, and other fabrication procedures are the same as above.

Characterization of materials and devices

The morphology of the produced $CsCu_2I_3$ wires was studied by field-emission SEM (JEOL-7500F, 15 keV). The element composition was analyzed using an EDS attached to the SEM. XRD (Panalytical X'Pert Pro) was conducted to examine the crystallinity of the products. AFM (Dimension Icon, Bruker) was used to study the morphology and height profiles of the $CsCu_2I_3$ NWs. The PL spectra were studied using a spectrofluorometer (Horiba; Fluorolog-3). A closed-cycle helium cryostat (Jannis; CCS-110) was used to carry out the PL measurements at different temperatures. The absorption spectra were measured using a Shimadzu UV-3150 spectrophotometer. UPS was conducted to study the energy band properties, and the UPS radiation was generated by a He-gas discharge lamp (He $I = 21.22 eV$). The optoelectrical and electrical measurements of the photodetectors were conducted by using a testing system that includes an oscilloscope (Tektronix, DPO2012B), an optical chopper (SRS, SR540), a digital SourceMeter (Keithley 2636B), a monochromator, and light sources in ambient air. A Xe lamp (Gloria-X150A, Zolix) equipped with a monochromator (Omni- $\lambda 300$, Zolix) was used to generate the monochromatic light to conduct the spectral response measurements.

Conflicts of interest

There are no conflicts to declare.

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