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Ultrasensitive 1D field-effect phototransistor: CH₃NH₃PbI₃ nanowire sensitized individual carbon nanotube

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Abstract: Field-effect phototransistors were fabricated based on individual carbon nanotubes (CNTs) sensitized by CH₃NH₃PbI₃ nanowires (MAPbI₃NW). These devices represent light responsivities of $R=7.7 \times 10^5$ A/W at low-lighting conditions in the nWmm⁻² range, unprecedented among CNT-based photo detectors. At high incident power (~ 1 mWmm⁻²), light soaking results in a negative photocurrent, turning the device insulating. We interpret the phenomenon as a result of efficient free photoexcited charge generation and charge transfer of photoexcited holes from the perovskite to the carbon nanotube. The charge transfer improves conductance by increasing the number of carriers, but leaves electrons behind. At high illumination intensity their random electrostatic potential quench the mobility in the nanotube.

Keywords: individual carbon nanotube based photodetector, infrared spectroscopy, CH₃NH₃PbI₃ nanowire, optical switch, perovskite carbon nanotube interface

In various optoelectronic applications, like light-emitting diodes, photodetectors and photovoltaic cells, semiconducting carbon nanotubes (CNTs) have been successfully used due to their direct band gap and outstanding electronic and mechanical properties.¹ Photodetection of individual CNTs excited by infrared²⁻⁴ (IR) or visible light⁵ has been achieved by separating the excitons with large enough electric fields generated locally by asymmetric Schottky contacts², p-n junctions³ or local charge defects⁵. However, their performance has been limited to quantum efficiencies of about 10%⁴. This is mainly limited by the high binding energy and long lifetime of excitons in CNTs.^{1,6}

CH₃NH₃PbI₃ (MAPbI₃) is efficiently used as photosensitizer in many optoelectronic hybrid devices in conjunction with carbon nanomaterials, due to its exceptional, but still not completely explained, physical properties favorable for light harvesting (*i.e.* direct bandgap, large absorption coefficient, long charge diffusion lengths), chemical flexibility and low-cost solution-based processability⁷⁻¹⁰. Several types of heterostructures have been made by combining MAPbI₃ and different carbon materials. Fullerenes have been reported to enhance the stability and to reduce drifts and hysteretic effects of MAPbI₃ solar cells^{11,12}. Incorporation of graphene and carbon nanotube films resulted in semitransparent flexible solar cells^{13,14}. The graphene lead-halide interface as a hybrid phototransistor was used as a high-sensitivity phototransistor owing to the successful photo-gating of graphene¹⁵⁻¹⁷.

Here we studied the light induced transfer characteristics of micro-fabricated field-effect transistors, built from individual metallic and semiconducting CNTs and CH₃NH₃PbI₃ nanowires (hereafter MAPbI₃NW). The sensitization of individual CNT-FETs with a network of MAPbI₃ nanowires resulted in responsivities as high as 7.7×10^5 A/W and external quantum efficiencies of 1.5×10^6 owing to the successful doping and gating of CNT-FETs. According to our knowledge, our microfabricated hybrid devices attained best-in-class responsivity in low-intensity visible-light detection. The extremely high sensitivity of the present MAPbI₃NW/CNT field-effect phototransistors (FEpT) is related to the photodoping of the carbon structure by photoexcited carriers of MAPbI₃NW, a mechanism reported on graphene/MAPbI₃ hybrid photodetectors as well¹⁵⁻¹⁷. Importantly, however, because of the unipolar nature of CNT FETs, the present MAPbI₃NW/CNT FEpT photodetectors can be switched off unlike the graphene/MAPbI₃ counterparts¹⁵⁻¹⁷.

The fabricated field-effect transistors are appropriate tools to obtain valuable information about the light induced charge transfer phenomena at the interface by means of fairly simple electrical transport measurements, as FETs use an electric field to control the conductivity of a channel of one type of charge carrier in a semiconductor material. Detailed analysis of the device characteristics unraveled the charge transfer process between the intimate contact of MAPbI₃ and metallic or semiconducting CNTs. Despite the remarkable progress in prototype building, however, there is a lack of knowledge about the fundamental chemical and photo-physical characteristics of the interfaces formed between the carbon nanomaterials and the organometal halide perovskites. Infrared (IR) and Raman spectroscopy of semitransparent highly purified single-walled nanotube buckypapers and MAPbI₃ nanowire composites confirmed the observed photo-induced charge transfer process.

Results and discussion

The fabrication of our MAPbI₃NW/CNT photo-FET started with the fabrication of an individual CNT based FET. The fabrication of CNT-FETs begins with the metal catalyst deposition to lithographically predefined positions. The cobalt-containing resist was spin-coated on a highly p-doped Si substrate with 200 nm thick SiO₂ thermally grown on top. Patterning the resist by electron-beam lithography created dots of metal ion doped resist as small as 100 nm (Figure 1a). The catalytic nanoparticles were formed by burning the organic resist at 800°C in oxygen (Figure 1b). Carbon nanotubes were grown by CVD at 800°C using ethanol as a carbon source (Figure 1c, S7). Next, CNT-FETs were fabricated by patterning and evaporating the source and drain metal contacts (Ti/Pd 1 nm/70 nm, Figure 1d).

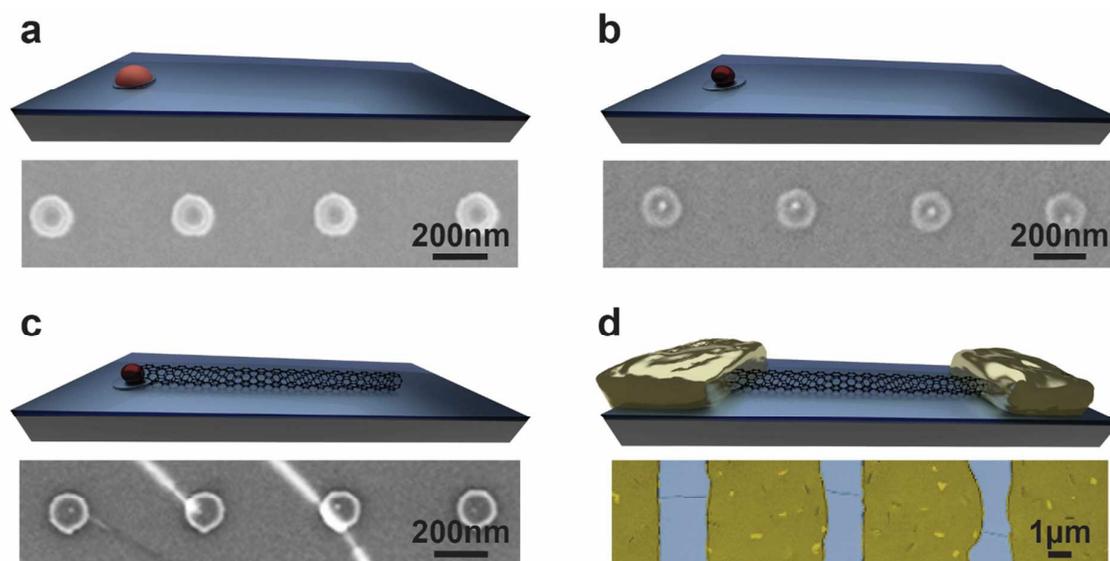


Figure 1. Schematic representation and corresponding false-color high-resolution SEM micrographs of the process used for synthesizing the CNTs. (a) e-beam lithography patterning of the metal-doped negative-tone resist. (b) Catalyst nanoparticle formation by thermal oxidation. (c) CNT synthesis by ethanol-assisted CVD. (d) Metal contacts deposition by e-beam evaporation.

The CNT-FET (Figure 2a) was sensitized with a network of photoactive MAPbI₃ nanowires, deposited by the recently developed slip-coating method¹⁸ (Figure 2b). The hybrid device was subsequently covered with a 500 nm-thick polymethyl methacrylate (PMMA) layer to protect the organometal network from the detrimental effect of humidity.

In the pristine CNT-FET the high work function of the Pd contacts¹⁹ and the p-type doping induced by the exposure to air (O_2)^{20,21} led to dark transfer characteristics showing unipolar p-type behavior with a threshold voltage $V_{th} \approx 2$ V and 8.2×10^4 cm^2/Vs mobility (Figure 2c). The $MAPbI_3$ nanowire deposition caused both a shift of $V_{th} \approx -1$ V and a decrease of the CNT charge mobility by about 40% to 4.9×10^4 cm^2/Vs (Figure 2c).

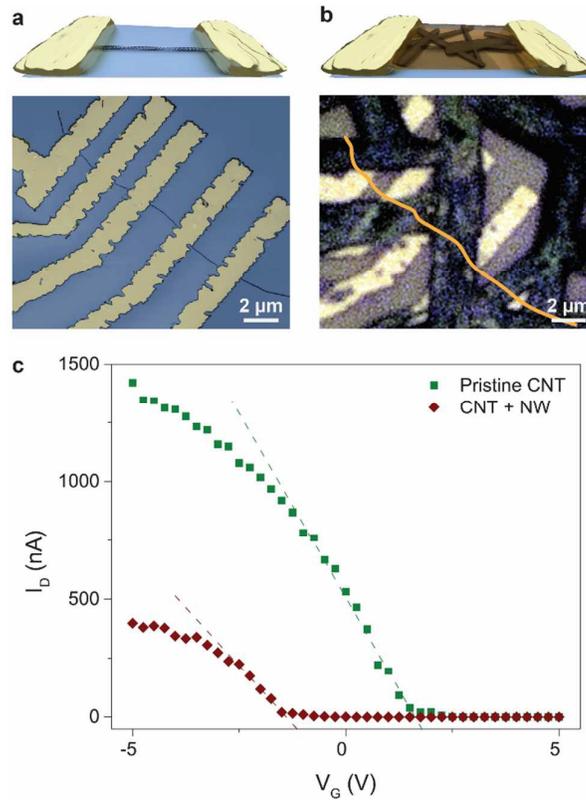


Figure 2: (a) Schematic representation and false color SEM micrograph of a series of a representative CNT-FET. (b) Schematic representation and optical micrograph of a representative $MAPbI_3$ NW/CNT-FET. (c) Transfer characteristic in dark of a representative device before (green curve) and after (red curve) sensitization ($V_D=0.2$ V). Dashed lines show the shift of the on-off threshold voltage V_{th} , and the reduction of the CNT mobility upon $MAPbI_3$ deposition.

The central finding of our paper is the remarkable photosensitivity of the hybrid $MAPbI_3$ NW/CNT-FETs with responsivity $R=7.7 \times 10^5$ A/W at low light conditions. The photoresponsivity of the hybrid device by the illumination was tested with a red laser ($\lambda=633$ nm) in the 62.5 $nWmm^{-2}$ to 2.5 $mWmm^{-2}$ intensity range.

Under illumination, electron-hole pairs are generated in MAPbI₃ nanowires. The holes are injected into the nanotube due to the chemical potential mismatch¹⁴, contributing to an increase in the output current in both the ON- and OFF-state of the MAPbI₃NW/CNT-FEpTs.

Above the threshold voltage, in the OFF-state of the MAPbI₃NW/CNT-FEpTs, the photocurrent, hence the total current, I_D , of the hybrid device, increases by increasing incident irradiation power (Figure 3b and S1). The I_D current in the OFF-state does not show gate voltage (V_G) dependence, thus it corresponds to the intrinsic photocurrent generation of the MAPbI₃ nanowire network shortcutting the source-drain contacts, as it was reported in our previous work¹⁸.

The evolution of the ON-state I_D current as a function of illumination intensity and V_G shows a markedly different behavior compared to the OFF-state (Figure 3 and Figure S1,S2). I_D -ON shows strong V_G dependence testifying that its origin is predominantly a CNT conduction channel. More interestingly, however, it shows a non-monotonous dependence on illumination intensity (Figure 3 and Figure S1, S2). At low light conditions below 100 nWmm⁻² intensity I_D increases monotonically by about a factor 2 relative to the dark current. By further increasing the light power, however, I_D rapidly falls and reaches I_D values observed for OFF-state (Figure S1, S2). Illuminating the device with light intensities higher than 95 μWmm⁻² resulted in a complete switch-off of the nanotube channel conductance over the whole range of positive and negative gate biases applied (Figure 3, S1). At the same time, V_{th} was independent of the light intensity.

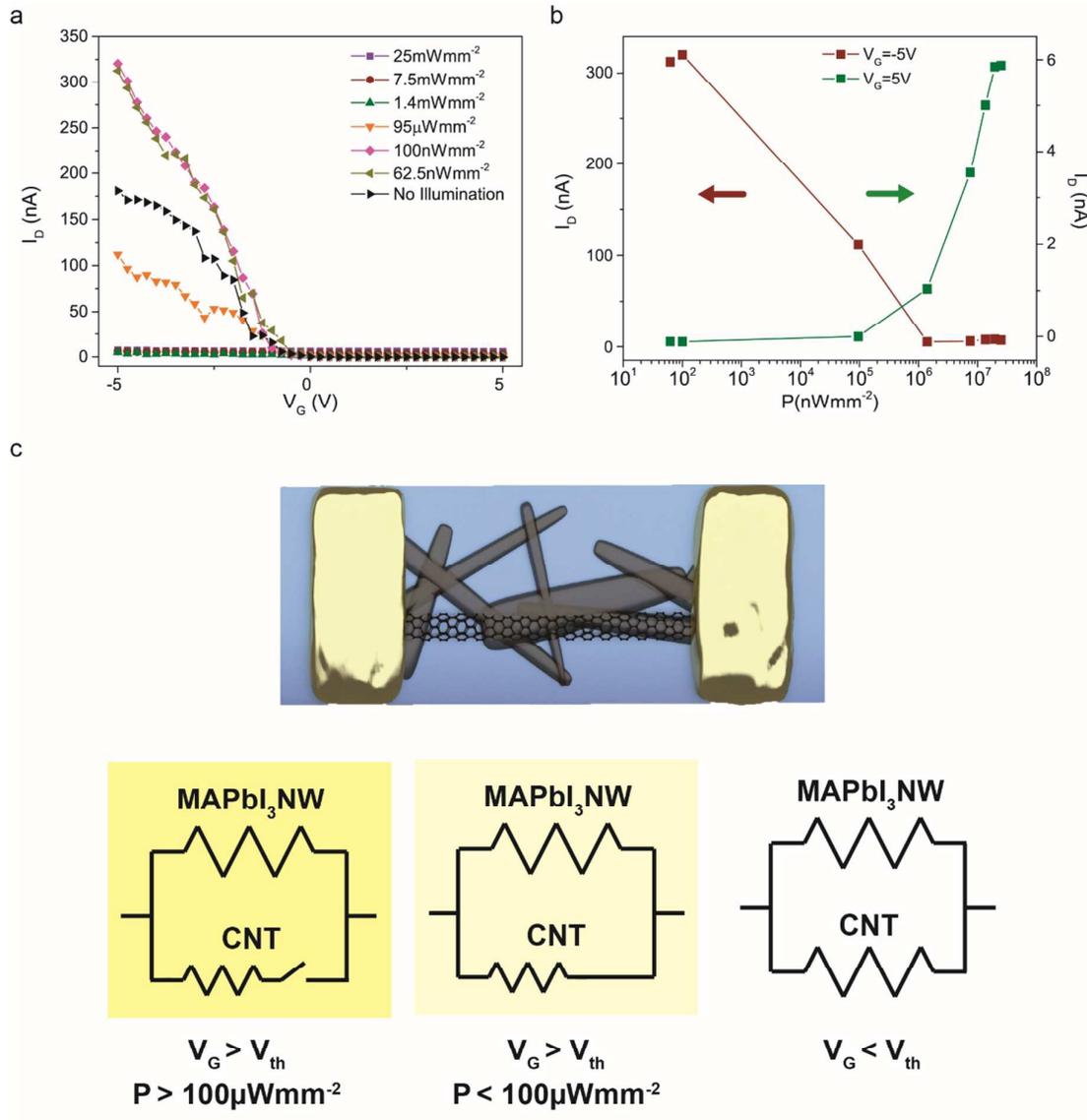


Figure 3. (a) Transfer characteristic of the hybrid phototransistor upon different light irradiation intensities. (b) I_D at $V_G=5$ and -5 V as a function of light power. (c) Schematic representation of the proposed two-parallel-resistor model used to describe our system.

The responsivity (R), the magnitude of the electrical signal output in response to a given light power, is one of the most important performance parameters of a photodetector. For the calculation of R the active area of the photodetector is needed. In order to conservatively estimate R of our device, we considered an active area equal to the distance between the contacts ($3 \mu\text{m}$) multiplied by the carrier diffusion length of photogenerated charge carriers in MAPbI₃ reported in the literature (*i.e.* $\sim 1 \mu\text{m}$)^{22,23}. In the ON-state ($V_G=-5$ V) and at extremely low light intensities ($6.25 \text{ nWmm}^{-2} \sim 375 \text{ fW}$) responsivity as high as $7.7 \times 10^5 \text{ A/W}$ with an

external quantum efficiency of 1.5×10^8 % was measured (Figure 4a). It is worth noting that the responsivities of the substituent CNT-FET and MAPbI₃NWs are 1×10^{-4} A/W (Figure S9) and 5×10^{-3} A/W¹⁸ respectively. The synergy of these two nanomaterials brings about eight orders of magnitude increase in responsivity. According to our best knowledge this outperforms by about 7 orders of magnitude the best carbon-nanotube based photodetectors reported so far⁴. Moreover, the device gain is highly linear as a function of both V_D (Figure 4a inset and S8) and V_G (Figure 4b) further facilitating applications.

The responsivity in accordance with the device characteristics presented in Figure 3a rapidly drops by increasing the light power and reaches zero when the MAPbI₃NW/CNT-FETs reaches the light-induced OFF state.

Apart from responsivity, the other important benchmark of photodetector performance is the response time. For the hybrid device the response time to illumination is less than 1 s, (limited by the time resolution of our measurement setup) under all operating conditions tested (Figure 4). On the other hand, the fall-time lasts between ~ 15 s ($P < 95 \mu\text{Wmm}^{-2}$) and ~ 35 s ($P > 95 \mu\text{Wmm}^{-2}$).

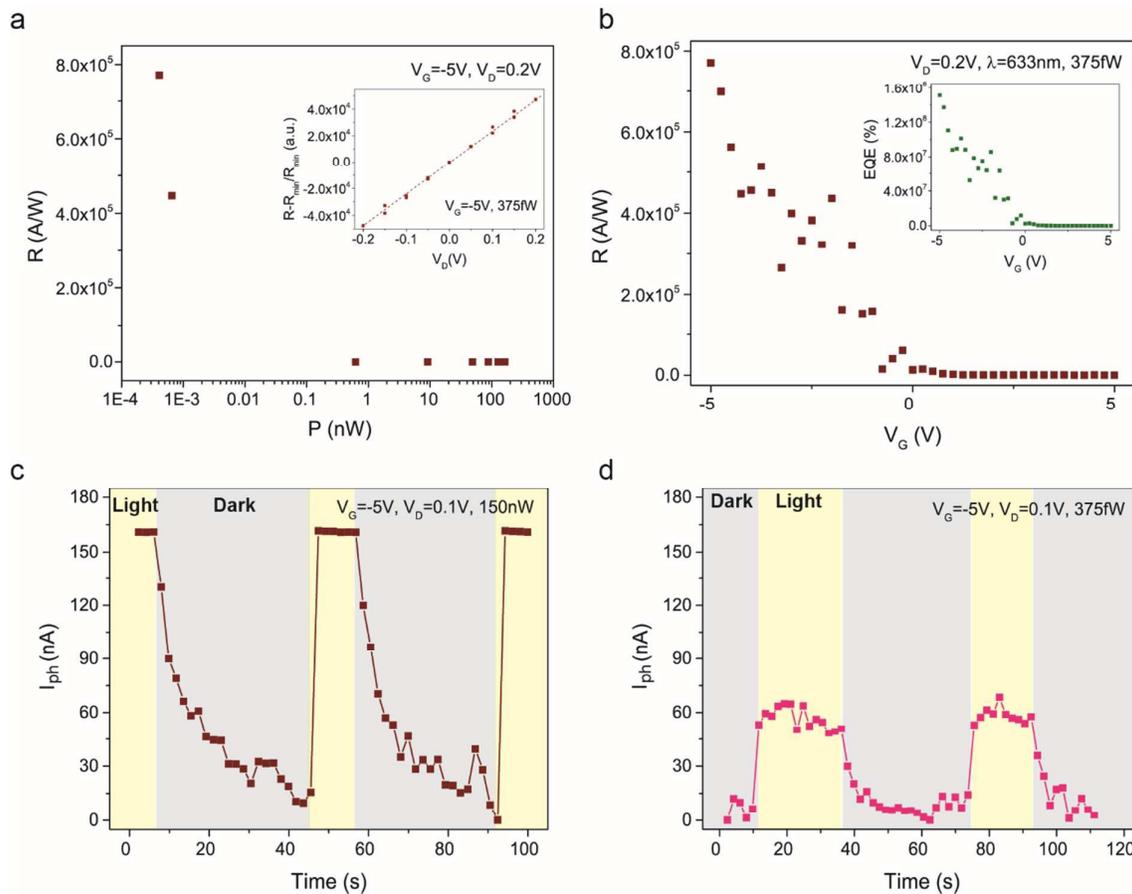


Figure 4. (a) Responsivity of the hybrid device as a function of the light intensity and source-to-drain voltage (inset). (b) Responsivity and external quantum efficiency under different gate biases. (c-d) Response time under high, 150 nW, (left) and low, 375 fW, (right) light intensity.

The photodiode characteristics of the individual CNT based MAPbI₃NW/CNT-FET give valuable insight to the interface behavior of CNT and MAPbI₃NW. Complications due to the intricate internal behavior of CNT films do not mask intrinsic interface properties, as we use single CNT devices. Moreover, metallic and semiconducting CNTs can be tested separately. In the case of MAPbI₃NW/CNT, hybrid devices fabricated from metallic carbon nanotubes the devices showed metallic behavior. Schottky barrier formation was not observed (Figure S2). For semiconducting CNT/MAPbI₃ devices the observed shift of V_{th} and the drop of CNT mobility (Figure 2) upon the exposure of CNT-FET to the concentrated MAPbI₃-DMF (Dimethylformamide) solution indicate changes of the CNT chemical potential and increased effective disorder along the tubes, respectively.

In order to reveal the potential corrosive effects of the MAPbI₃-DMF solution on carbon nanotubes, a free-standing semi-transparent film enriched in semiconducting single walled CNTs was prepared and used as a 3D scaffold for the growth of MAPbI₃ nanowires (Figure S3-S5). We studied the interaction of the MAPbI₃ nanowires with the CNTs with and without illumination (633 nm LED source) by infrared, near-infrared and Raman spectroscopy. Depositing the nanowires affected only the IR, but not the Raman spectra of the CNT films. (See supporting information Figure S3-S5). We conclude by spectroscopic methods, that under illumination no other significant reaction can be detected between the nanowires and the nanotubes but charge transfer, resulting in mobile carriers. It should be noted, however, that the current running through a functional device can induce ion migration or additional electrochemical redox reactions at the carbon nanotube-MAPbI₃NWs interface which can increase the number of defects, hence reduce the mobility, thus further optical measurements under operating conditions needs to be done to clarify the origin of the mobility drop.

The photodiode characteristics of our MAPbI₃NW/CNT-FET can be described by a parallel resistor model (Figure 3b). When the hybrid CNT-FET is electrostatically switched off, the resistance of both 1D nanostructures (CNT and MAPbI₃NW) are in the GΩ range, the current is low. Due to the closed conduction channel of the CNT, the photocurrent is essentially equal to the photo-generated charges in the MAPbI₃ nanowires. If the CNT-FET is in the ON regime due to the electrostatic gating, its high conductivity dominates the performance of the hybrid device. Upon illumination the photo generated positive charges enter the CNT and acts as chemical doping, in agreement with the IR and Raman spectroscopy. The photo-doping, however, does not shift the chemical potential of the CNTs indicating a nearby Van Hove singularity. The photo-induced negative charges, which are not injected in the CNT due to the work function mismatch¹⁴, constitute scattering centers inhomogeneously distributed along the carbon nanotube. The resulting random potential reduces the charge carrier mobility, thus the overall current. These two effects compete and at high light intensities the detrimental effects of the random potential overcompensate the doping and switch off the CNT conduction channel. Thus the MAPbI₃NW/CNT-FET acts as a light switch at high powers.

Conclusion

In conclusion, we demonstrated gate voltage-dependent visible light photo-response of microfabricated individual MAPbI₃NW/CNT photo-FETs for the first time. In the mWmm⁻² power range light soaking resulted in quenching the conductance of the ON-state p-channel of the individual CNT-FET, effectively making the device an optical switch. Exposure of these

hybrid devices to sub nWmm^{-2} light intensities, however, manifested a strong positive photocurrent. The best devices showed as high as 7.7×10^5 A/W responsivity and external quantum efficiencies of 1.5×10^8 %, indicating that the device can be used as a low-intensity visible-light detector. We attributed this unconventional photocurrent transfer characteristic of the unique charge distribution over the 1D semiconductor nanotubes. Analysis of the gate dependent transfer characteristics in the dark and under illumination allowed the underlying photon induced charge transfer mechanisms between MAPbI_3 and metallic and semiconductor CNTs to be probed. The results have important implications in the fundamental understanding of the photo-physical picture of MAPbI_3 and CNT interfaces and in the development and fabrication of organometallic halide perovskite based optoelectronic devices such as solar cells, LEDs, photodetectors, single photon-detectors and optical switches.

Experimental Section

Resist preparation. A high-resolution Cobalt-containing negative-tone resist was prepared by dissolving 0.2 wt% of 4-Methyl-1-acetoxycalix[6]arene (Synchem OHG) in monochlorobenzene and 0.2 wt% of Co(III) acetylacetonate, (Sigma-Aldrich GmbH, 99%). After stirring for 1 hour at 700 rpm the solution was filtered through a 0.2-mm Teflon membrane to remove potential solid residues.

Nanoparticle localization. The resist was patterned by e-beam lithography with a Vistec EBPG5000 operating at 100kV and 1nA. The nucleation centers were localized by a reactive ion etch step of 10 seconds with an Adixen AMS200 and a gas mixture of Ar and C_4F_8 .

Carbon Nanotube synthesis. The deposited nanoparticles are catalytically activated by a 10 min reduction at 800 °C under controlled atmosphere (Ar/H_2 8:1 vol%). Next, ethanol vapor was introduced in the quartz tube using Argon and Hydrogen (1:2 vol%) as carrier gas. After 5 minutes the carbon source was evacuated and the samples were cooled down to room temperature.

Carbon Nanotube film synthesis. Films of single walled carbon nanotubes were prepared from P2 and semiconductor enriched nanotubes as described by Borondics et al.²⁴

MAPbI_3 nanowires synthesis. The network of MAPbI_3 nanowires was subsequently deposited by the slip-coating technique reported by Horváth et al.¹⁸

Photoelectrical characterization. The photoelectric response measurements of the fabricated hybrid devices were performed using a standard DC technique. The light sources used were a red laser beam ($\lambda=633$ nm) with a spot size of about 4 mm. All the measurements were performed at room temperature and in ambient environment.

High-resolution scanning electron microscopy (SEM) was performed with a MERLIN Zeiss electron microscope.

Conflict of Interests: The authors declare no competing financial interest.

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Supporting Information Available: Infrared and Raman spectroscopy with additional electronic transfer characterizations are included. This material is available free of charge *via* the internet at <http://pubs.acs.org>

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