

Cite this: *J. Mater. Chem. C*, 2023, 11, 970

Crystallinity-modulated hollow CeO_{2-x} nanorods as free radical scavengers for long-term photostability in organic photovoltaics†

Seongwon Yoon,[†] Taehyun Kwon,[‡] Sohyun Kim,^b So Hyun Park,^{ad} Youngjoon Lim,^b Jihun Kim,^e Jun Kim,^b Kwangjin An,^e Sungmin Park,^f Jin Young Kim^{†*} and Hae Jung Son^{†*}

Here we investigated the effects of CeO_{2-x} nanostructures as free radical scavengers on the long-term photostability of an organic photovoltaic (OPV) structure. From powder X-ray diffraction, Raman spectroscopy, X-ray photoelectron spectroscopy, and N₂ adsorption experiments, it was determined that the single-crystalline hollow CeO_{2-x} nanorods were very effective as hydroxyl radical scavengers. This was attributed to their having more Ce³⁺ states and a wider surface area than other types of CeO₂ nanostructures. Time-dependent UV-visible absorption spectra analyses also revealed that the improved scavenging of hydroxyl radicals in the OPV device was related to the better interfacial compatibility between the organic active and ZnO layers, resulting in improved OPV photostability.

Received 30th September 2022,
Accepted 18th December 2022

DOI: 10.1039/d2tc04152a

rsc.li/materials-c

1. Introduction

Organic photovoltaics (OPVs) are considered one of the key technologies for future sources of clean energy.¹ OPV devices can be easily prepared on a flexible substrate using a low-cost roll-to-roll solution process.^{2,3} As a result, OPVs have received a lot of research interest following the intensive development of active organic materials such as non-fullerene acceptors. OPVs have recently achieved remarkable efficiency enhancements of close to 20%.⁴⁻⁸

However, the long-term stability of OPV devices remains low, and is the main barrier to its future commercialization.^{9,10} OPVs are vulnerable to several degradation mechanisms. π -conjugated small molecules and polymers in particular suffer from degradation because they are typically susceptible to

environmental factors, especially moisture, oxygen and UV-light.¹¹ Among these conditions, light-induced degradation, also known as photo-degradation, is considered as the main obstacle hindering the long-term stability of such devices.^{12,13} Photo-oxidation of the donor or acceptor materials by photocatalytic reaction under illumination occurs when metal oxides such as ZnO or TiO_x are used as interlayer materials. SnO_x has also been employed to prevent photocatalysis with its wider optical bandgap than ZnO.¹⁴ However, due to a high-temperature annealing process and comparatively low performance,¹⁵ SnO_x film was not widely used in OPV devices. After excitation under UV light irradiation, electrons or holes in the ZnO film can be trapped by surface defect states, which are related to oxygen vacancies. They subsequently undergo redox reactions with H₂O adsorbed on the ZnO film, leading to the formation of hydroxyl radicals (\bullet OH).¹⁶ To reduce the \bullet OH generation through defect states of ZnO, a few research was reported such as molecular passivation and doping of ZnO,¹⁷⁻¹⁹ but the \bullet OH generation issue still remains.

Once generated, hydroxyl radicals are destructive to nearly all organic compounds. For example, the vinyl enone group in high performing non-fullerene acceptors such as Y6 and ITIC can easily undergo a radical addition reaction, which is a critical factor in the low stability of non-fullerene acceptor-based OPV devices.¹⁷ Therefore, to protect the active materials and develop high-performing OPV devices with high stability under light soaking, it is highly important to quench the \bullet OH radicals.

Herein, we investigated CeO_{2-x}-based radical scavengers with different morphologies and crystallinity to enhance the

^a Advanced Photovoltaics Research Center, Korea Institute of Science and Technology (KIST), Seoul 02792, Republic of Korea. E-mail: hjson@kist.re.kr

^b Hydrogen Fuel Cell Research Center, KIST, Seoul 02792, Republic of Korea. E-mail: jinykim@kist.re.kr

^c Department of Chemistry and Research Institute of Basic Sciences, Incheon National University, Incheon 22012, Republic of Korea

^d Graduate School of Energy and Environment (KU-KIST Green School), Korea University, Seoul 02841, Republic of Korea

^e School of Energy and Chemical Engineering, Ulsan National Institute of Science and Technology (UNIST), Ulsan 44919, Republic of Korea

^f Realistic Media Research Center, ICT Convergence Research Division, Gumi Electronics & Information Technology Research Institute (GERI), Gumi, Gyeongbuk 39253, Republic of Korea

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d2tc04152a>

* These authors contributed equally.

long-term photostability of OPV devices. Importantly, the addition of CeO_{2-x}-based free radical scavengers between the active layer and ZnO layer in the OPVs rapidly reacted with harmful •OH radicals before they attacked the active organic material in the OPV device. More importantly, among various radical scavengers, single-crystalline hollow nanorod structures were found to be the most effective at improving the overall photostability of the high-performing OPV devices. Their positive effect was mainly attributed to their more efficient radical quenching rate and interfacial compatibility between the constituent layers in the OPVs, compared to other types.

2. Experimental section

2.1. Reagents

Cerium(III) nitrate hexahydrate (Ce(NO₃)₃·6H₂O, 99%), methyl violet 2B, iron(II) sulphate heptahydrate (Fe(SO₄)₂·7H₂O, ≥99.0%), and hydrogen peroxide solution (H₂O₂, 30% (w/w) in H₂O), zinc acetate dihydrate, ethanolamine, 2-methoxyethanol, chloroform, 1-chloronaphthalene, ethylene glycol, and molybdenum(VI) trioxide (MoO₃) were purchased from Merck. PM6 and N3 were purchased from 1-Material. Sodium hydroxide (NaOH, 98%) was purchased from Samchun Chemicals. All reagents were used as received without further purification.

2.2. Synthesis

The synthetic procedures of the CeO_{2-x} hollow nanorods were adopted and slightly modified from the previous literature.^{20,21} Before preparing CeO_{2-x} single-crystalline hollow nanorods (S-hNR), non-porous CeO₂ nanorods (NR) were prepared by using a previously reported method.²² Briefly, NaOH (19.2 g) was dissolved in deionized (DI) water (75 mL) and Ce(NO₃)₃·6H₂O (1.736 g) was added. The resulting solution was placed in a Teflon-lined reactor and kept in an oven at 100 °C for 24 h. The precipitate was collected by centrifugation with DI water and ethanol alternatively for 3 times and dried at 80 °C overnight. Then, the prepared CeO₂ NR (0.516 g) was dispersed in 160 mL of DI water, followed by adding Ce(NO₃)₃·6H₂O (2.605 g) into the CeO₂ NR suspension under stirring. Then, the resulting suspension was placed in a Teflon-lined reactor and kept at 100 °C for 3 h. Afterwards, the precipitate was collected by centrifugation and washed by DI water for 3 times and dried at 60 °C for 12 h.

For polycrystalline hollow nanorods (P-hNR), the synthetic procedure was adopted and slightly modified from the previous literature.²¹ Before preparing CeO_{2-x} P-hNR, non-porous Ce(OH)₃/CeO₂ nanorod precursor was prepared by using a previously reported methods with slight modification.²³ Ce(NO₃)₃·6H₂O (4.25 g) and NaOH (48 g) were dissolved in 25 and 75 mL of DI water, respectively. The two solutions were thoroughly mixed in a Pyrex bottle and the resulting solution was aged with continuous stirring for 30 min. Subsequently, the Pyrex bottle was transferred into an oven at 100 °C for 24 h. After cooling to room temperature, the precipitate was collected by centrifugation, and washed with DI and ethanol until pH 7–8 and dried at 80 °C overnight. Then, the non-porous nanorod

precursor (20 mg) were dispersed in 10 mL of DI water by sonication for 10 min. The mixture was stirred for 30 min and then transferred to a 50 mL Teflon-lined autoclave reactor and heated at 160 °C for 12 h. After cooling to room temperature, the pale-yellow solid products were collected by centrifugation, washed with DI water and ethanol once, respectively. The resulting powder was dried at 80 °C.

2.3. Material characterization

Transmission electron microscopy (TEM). TEM, high-resolution TEM (HRTEM), and scanning TEM (STEM) images were obtained using FEI Tecnai G2 F30ST microscopy operated at 200 kV. The fast Fourier transform (FFT) patterns were obtained and analysed from the HRTEM images using Gatan Digital Micrograph software.

Powder X-ray diffraction (PXRD). PXRD patterns were collected with Rigaku Ultima III diffractometer system using graphite-monochromatized Cu-Kα radiation at 40 kV and 30 mA.

X-Ray photoelectron spectroscopy (XPS). XPS spectra were acquired using an X-ray photoelectron spectrometer (ESCALAB 250XI, Thermo Fisher Scientific) with a monochromatic Al-Kα X-ray source (1486.6 eV).

Raman spectroscopy. Raman spectra were obtained using *inVia* Raman microscopes (Renishaw) operated with a 532 nm laser at 10 mW focused at 0.41 μm² (100× objective), and collection was carried out an acquisition time of 10 s with 15 times of accumulation.

Gas sorption measurement. The N₂ adsorption experiment was performed at −196 °C using a BELSORP-max (Microtrac MRB) system to estimate the Brunauer–Emmett–Teller (BET) surface area. Before the analysis, all samples were pretreated under vacuum at 100 °C for 12 h.

Radical scavenging efficiency test. The test procedure was adopted and slightly modified from the previous literature.²⁴ The hydroxyl radical scavenging activity of the CeO_{2-x} samples were evaluated by tracking the changes in UV/vis absorption spectrum of methyl violet 2B (MV) dye in the presence of •OH radicals generated by Fenton's reaction. The reaction solution for photometric determination contained 2.85 mL of 14.3 μM MV aqueous solution, 60 μL of 3 wt% H₂O₂, 90 μL of 2 ppm FeSO₄, and 0.3 mL of CeO_{2-x} dispersion (1 mg_{Ce} mL⁻¹). The overall concentration of the radical scavengers in the solution was 0.091 mg_{Ce} mL⁻¹. The absorption spectra were measured by Agilent Cary 100 (Agilent Technologies) UV-vis spectrometer while incubating the reaction solution for 90 min.

Scanning electron microscopy (SEM). The morphologies of the CeO_{2-x} coated surface of the ZnO films and corresponding energy dispersive spectroscopy (EDS) elemental mapping analyses were obtained with field emission SEM (FE-SEM, FEI Inspect F, FEI Company) operated at an activation voltage of 15 kV.

2.4. Fabrication of OPV cells

The indium tin oxide (ITO)-patterned glass substrates were sequentially sonicated in distilled water, acetone, and 2-propanol

for 10 min each. Then the remaining solvents on the cleaned substrates were eliminated by using nitrogen flow and hot oven (80 °C). Prior to spin-coating the ZnO solution, the dried and cleaned substrates were treated with UV-ozone to make the substrates more hydrophilic. The sol-gel-based ZnO solution can be formed by blending 1 g of zinc acetate dihydrate, 280 mg of ethanolamine in 10 mL of 2-methoxyethanol. The ZnO solution was vigorously stirred at 60 °C for >2 h before use. The ZnO solution was spin-coated onto the UV-ozone-treated substrates at 4000 rpm for 30 s and the ZnO-coated substrates were annealed at 200 °C for 30 min. The CeO_{2-x} NP, CeO_{2-x} P-hNR, and CeO_{2-x} S-hNR layers were formed by spin-coating using corresponding aqueous solutions with a concentration of 1 mg mL⁻¹ and no surfactants were added to the solutions. The aqueous cerium oxide solutions were spin-coated onto the ZnO films at 4000 rpm for 60 s and the substrates were annealed at 120 °C for 10 min to eliminate the residual solvents. The fabricated ITO/ZnO or ITO/ZnO/cerium oxide substrates were moved in nitrogen-filled glove box for forming the photoactive layer. The PM6 and N3 were mixed with a weight ratio of PM6:N3 = 1:1.2 w/w in chloroform-based solvent with the concentration of PM6 of 7 mg mL⁻¹. The chloroform-based solvent can be formed by mixing chloroform and 1-chloronaphthalene with a ratio of chloroform:1-chloronaphthalene = 100:0.5 v/v. The formed solution for active films was spin-coated on the ITO/ZnO substrates to form a nominal thickness of ~110 nm. Then 3 nm of MoO₃ and 100 nm of silver were sequentially deposited on the active film-coated substrates by using thermal evaporator with a pressure under ~3.0 × 10⁻⁶ torr. The active area defined by shadow mask was 0.0422 cm².

2.5. OPV characterization

The current density–voltage (*J*–*V*) characteristics were measured using laptop-controlled Keithley 2400 sourcemeter with a solar simulator (YSS-200A, Yamashita Denso) with AM 1.5G spectra at 100 mW cm⁻². We calibrated the light intensity of the solar simulator using a certified silicon solar cell (91150-KG5, Newport). The external quantum efficiency spectra were measured using a quantum efficiency measurement system (QuantX-300, Oriel). The thickness information of films was obtained from a surface profiler (Alpha-Step IQ, KLA-Tencor). The UV-visible absorption spectra were collected from an UV/VIS spectrometer (Lambda 35, PerkinElmer). The atomic force microscopy (AFM) images were obtained from an XE-100 (Park Systems).

2.6. Photostability measurements

The fabricated OPV cells were encapsulated using cover glass and UV-curable epoxy resin to avoid other degradation factors such as oxygen and moisture. The encapsulated OPV cells were exposed to 1 sun condition using a solar cell reliability test system (Polaronix K3600, McScience) in air and the *J*–*V* characteristics of each cell were collected at selected times. For measuring the evolution of UV-visible absorption spectra of PM6:N3 blend films, the thin PM6:N3 blend films were exposed to UV in a glove box.

3. Results and discussions

3.1. Materials synthesis and characterization

Various CeO_{2-x} nanostructures with different morphology and crystallinity were modulated using two-step hydrothermal synthetic methods, by adopting and slightly modifying previously reported studies.^{20,21} Hollow structures were prepared by structural transformation from solid CeO_{2-x} nanorod (NR) structures under hydrothermal conditions (Fig. S1, ESI†). During the secondary hydrothermal reaction, a morphological evolution from solid to hollow nanorods occurs due to the partial dissolution and redeposition of Ce-based species.^{20,21,23} The final crystallinity of the hollow CeO_{2-x} nanorods could be changed by the reaction temperature and time.

Representative STEM images of single-crystalline hollow CeO_{2-x} nanorods (CeO_{2-x} S-hNR) and polycrystalline hollow CeO_{2-x} nanorods (CeO_{2-x} P-hNR) are shown in Fig. 1a and c. The low magnification TEM images of the two CeO_{2-x} hollow nanorods showed that the lengths of CeO_{2-x} S-hNR and CeO_{2-x} P-hNR were 133.4 ± 31.8 and 63.6 ± 15.2 nm, respectively (Fig. S2, ESI†); the CeO_{2-x} P-hNR was shorter than the CeO_{2-x} S-hNR. However, the diameter of the two nanorods were similar, around ~9 nm.

The crystallinity of the hollow nanotube structures was further investigated using high-resolution TEM (HRTEM) images and corresponding fast Fourier transform (FFT) patterns, as shown in Fig. 1b and d. The HRTEM analysis shown in Fig. 1b showed lattice distances of 0.312 and 0.271 nm, which correspond to the lattice distances of the (002) and (111) facets of the cubic CeO₂, respectively. Also, the corresponding FFT pattern indicated that the observed zone axis of the HRTEM image is [110], verifying that most of the exposed facets of CeO_{2-x} S-hNR are (110) and the overall nanostructure possess single-crystalline nature. On the other hand, according to the HRTEM image (Fig. 1d) and the corresponding FFT patterns (inset) of CeO_{2-x} P-hNR, grain boundaries (yellow dotted lines) were observed, while the measured lattice distances (0.312 and 0.271 nm) were the same as those of CeO_{2-x} S-hNR.

The powder X-ray diffraction (PXRD) patterns of CeO_{2-x} S-hNR and CeO_{2-x} P-hNR demonstrated that the phase of those two nanorods were the cubic CeO₂ phase (Fig. 1e). The average crystallite domain sizes of the samples were calculated using the Scherrer equation, with PXRD peaks near 28.5°. The calculated average crystallite sizes of CeO_{2-x} S-hNR, CeO_{2-x} P-hNR, and commercial CeO₂ nanoparticles (NPs) were 13.4 ± 0.3, 11.1 ± 0.3, and 39.0 ± 0.9 nm, respectively. The calculated crystallite sizes of commercial CeO₂ NPs by Scherrer equation were similar to the sizes of those observed in the TEM image, as shown in Fig. S3 (ESI†).

Also, the PXRD patterns of S-hNR and P-hNR were slightly negative-shifted compared to that of the NP, which indicates the crystal lattices in the hollow structure were slightly expanded to bulk phase. The expanded crystal lattice was attributed to the decrease in crystallite domain size, which increased the surface concentration of oxygen vacancies and Ce³⁺ states in the nanocrystal.^{26,27}

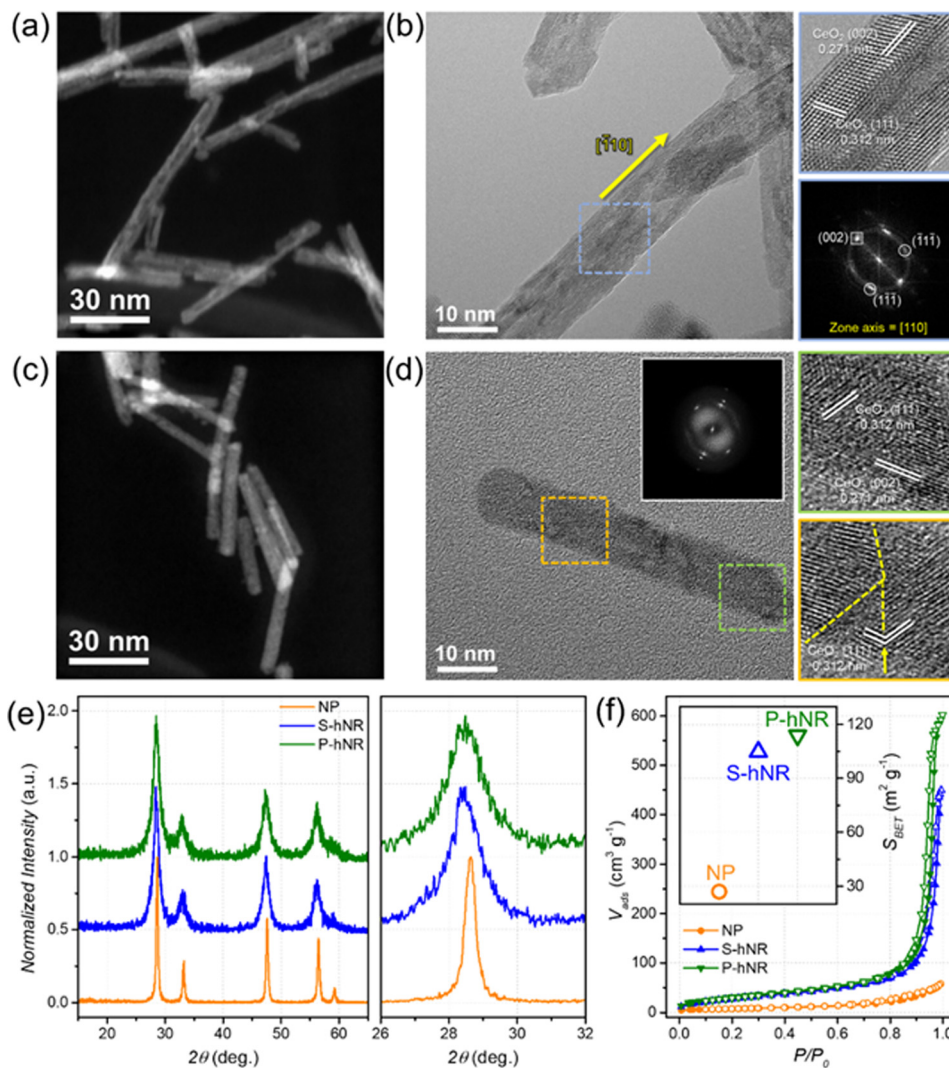


Fig. 1 (a) STEM image of CeO_{2-x} S-hNR. (b) HRTEM image of CeO_{2-x} S-hNR (left) and enlarged HRTEM image and corresponding FFT patterns of the marked area (right). (c) STEM image of CeO_{2-x} P-hNR. (d) HRTEM image of CeO_{2-x} P-hNR (left) and enlarged HRTEM images of the marked area (right). (e) PXRD patterns of CeO_{2-x} S-hNR, P-hNR, and commercial CeO_2 nanoparticles (NP) and enlarged PXRD patterns from 26° to 32° (right). (f) N_2 -isotherm curves of CeO_{2-x} S-hNR, P-hNR, and NP and the calculated BET surface area of the samples (inset).

N_2 isotherms (Fig. 1f) of CeO_{2-x} S-hNR, CeO_{2-x} P-hNR, and commercial CeO_2 NP samples were measured at -196°C to calculate their BET surface areas (S_{BET}). The surface areas of the CeO_{2-x} S-hNR ($104.80\text{ m}^2\text{ g}^{-1}$) and CeO_{2-x} P-hNR ($113.87\text{ m}^2\text{ g}^{-1}$) were much higher than that of CeO_2 NP ($26.99\text{ m}^2\text{ g}^{-1}$). Also, the S_{BET} of those hNRs was higher than the non-porous CeO_{2-x} nanorods ($94.26\text{ m}^2\text{ g}^{-1}$, Fig. S4, ESI[†]), indicating that the increased surface areas originate from the inner cavity in the nanorod structure. In particular, because the CeO_{2-x} P-hNR has a larger number of defect structures and pores along multiple grain boundaries, it showed higher S_{BET} than the CeO_{2-x} S-hNR.

The radical scavenging efficiencies of the CeO_{2-x} samples were evaluated by dye tests, using the chemical degradation of methyl violet 2B (MV) in Fenton's solution ($\bullet\text{OH}$ radical source) over 90 min (Fig. S5, ESI[†]).²⁴ The concentration of radical scavenger in the reaction solution was fixed to $0.091\text{ mg}_{\text{Ce}}\text{ mL}^{-1}$ (see experimental section for details). The radical scavenging

efficiency was evaluated by the change in the absorption peak intensity of MV dye at 575 nm . The normalized peak intensity change over time is summarized in Fig. 2a. The addition of CeO_2 or CeO_{2-x} decreased the degradation rate of the MV dye regardless of their structure, confirming that they can efficiently scavenge $\bullet\text{OH}$ radicals. Without radical scavengers, the absorption intensity of MV dye in the solution remained at 34.6% of the initial state after 90 min. However, the final absorption intensity remained at 69.7%, 70.3%, and 58.1% of the initial state following the addition of CeO_{2-x} S-hNR, CeO_{2-x} P-hNR, and commercial CeO_2 NP, respectively. This indicates that the hollow nanorod structures have higher radical scavenging efficiency than commercial NPs regardless of their initial crystallinity.

The chemical states of the CeO_{2-x} structures were next investigated by Raman spectroscopy (Fig. 2b) to determine the origin of the improved radical scavenging activity of the CeO_{2-x} S-hNR and CeO_{2-x} P-hNR. It has been reported that perfect CeO_2 crystal

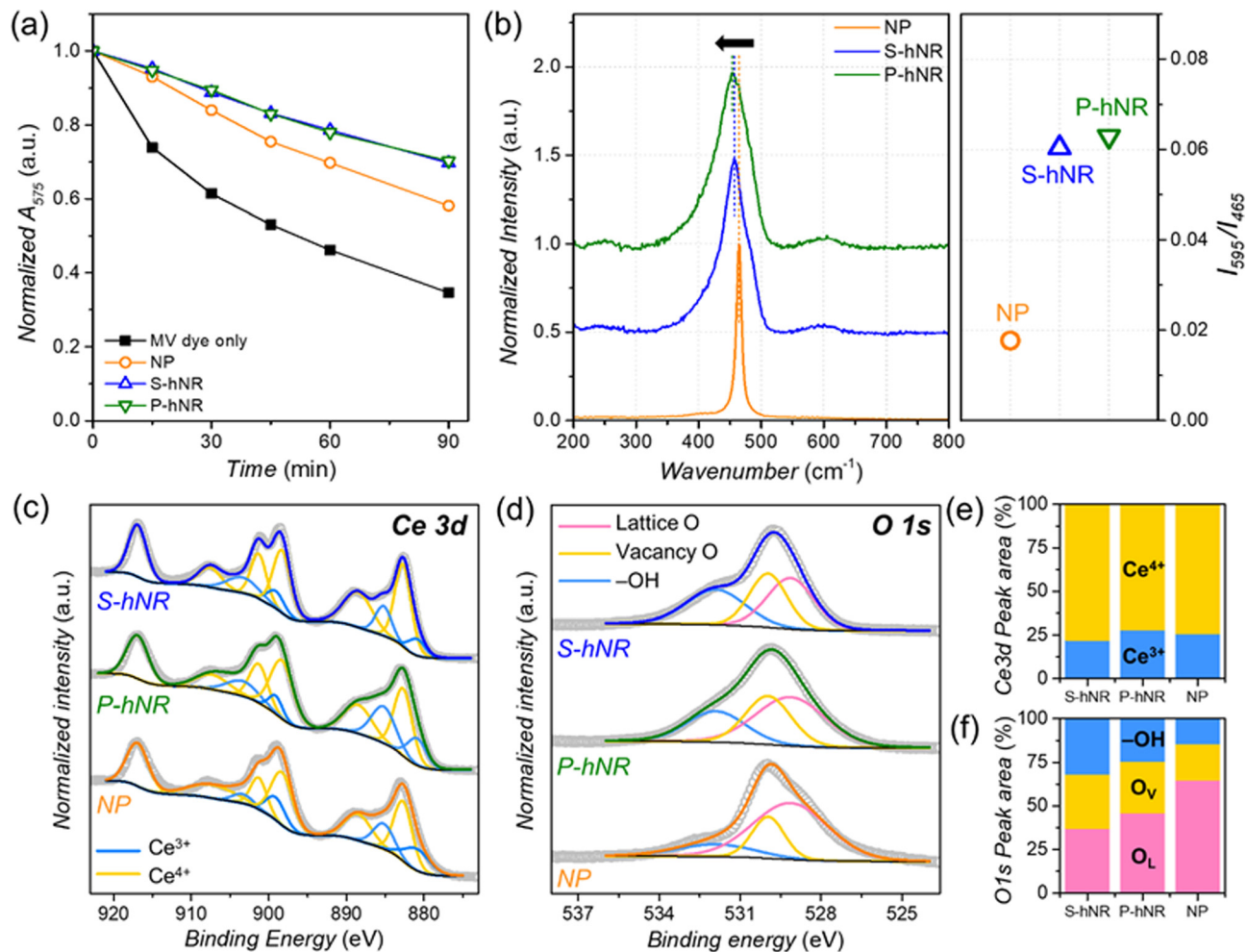


Fig. 2 (a) Hydroxyl radical scavenging efficiency measurement of the CeO_{2-x} -based radical scavengers. Change in absorption intensity at 575 nm over time in the presence of MV dye and radical scavengers after injecting Fenton's reagent. (b) Raman spectra of CeO_{2-x} S-hNR, CeO_{2-x} P-hNR, and commercial CeO_2 NP, normalized to F_{2g} band near 465 cm^{-1} for clarity. The right graph indicates the peak intensity ratio between 595 cm^{-1} and 465 cm^{-1} . (c) Ce 3d XPS spectra (d) O 1s XPS spectra of CeO_{2-x} S-hNR, CeO_{2-x} P-hNR, and commercial CeO_2 NP. (e and f) Relative peak area from the deconvoluted (e) Ce 3d and (f) O 1s XPS spectra in (c) and (d), respectively.

without any oxygen vacancies shows only a single Raman active mode with F_{2g} symmetry at 465 cm^{-1} .²⁸ For the three CeO_{2-x} samples, a single major peak near 465 cm^{-1} was observed. However, a red shift of the major peak was observed in both the CeO_{2-x} S-hNR and CeO_{2-x} P-hNR samples, compared to the commercial CeO_2 NP. This red shift largely originates from partially reduced states (Ce^{3+}) and the lattice expansion of the CeO_2 lattice,²⁹ which corroborates with the PXRD results in Fig. 1e. Moreover, in the Raman spectra of CeO_{2-x} S-hNR and CeO_{2-x} P-hNR showed asymmetric broadening of the 465 cm^{-1} peak and minor peak around 595 cm^{-1} . These features are related to the breaking of F_{2g} symmetry by the formation of oxygen vacancies in the CeO_{2-x} crystal.^{28,30} Because the Raman peak at 595 cm^{-1} is a defect-induced mode,³¹ the relative intensities (I_{595}/I_{465}) of the 465 and 595 cm^{-1} bands were compared. The I_{595}/I_{465} value of CeO_{2-x} P-hNR (0.063) was slightly larger than that of CeO_{2-x} S-hNR (0.060) and much larger than that of CeO_2 NP (0.018). Therefore, the multiple grain boundaries in CeO_{2-x}

P-hNR enabled a higher concentration of surface oxygen vacancies in their structure. However, because the surface area and small crystallite size were comparable, the surface oxygen vacancy concentration of CeO_{2-x} S-hNR is comparable to that of CeO_{2-x} P-hNR.

XPS spectra were also measured to further investigate the chemical states of the CeO_{2-x} hNRs (Fig. 2c–f). The surface concentration of Ce^{3+} and Ce^{4+} states were evaluated by deconvoluting the Ce 3d XPS spectra (Fig. 2c and Table S1, ESI[†]) of the samples. The calculated ratios of the Ce^{3+} state were in the following order (Fig. 2e): CeO_{2-x} S-hNR (21.9%) < CeO_2 NP (25.5%) < CeO_{2-x} P-hNR (28.0%). Because the CeO_{2-x} P-hNRs had multiple grain boundaries and defect sites in their structure, they possessed the highest concentration of Ce^{3+} states, which are known to be active species toward $\bullet\text{OH}$ radical scavenging. The states of the surface oxygen species in the samples were also evaluated by O 1s XPS spectra (Fig. 2d). The deconvolution result (Fig. 2f) showed the ratio of oxygen species

near the vacant sites (vacancy O, O_v) in CeO_{2-x} S-hNR (31.3%) and CeO_{2-x} P-hNR (29.9%) was higher than that of CeO_2 NP (20.6%), corroborating the Raman spectra (Fig. 2b).

It has been well known that the Ce^{3+} state is an active species for $\bullet OH$ radical scavenging.^{32–37} However, while the CeO_{2-x} S-hNRs possessed the lowest surface concentration of Ce^{3+} species, they showed radical scavenging efficiency that was comparable to the CeO_{2-x} P-hNRs. Therefore, the superior $\bullet OH$ radical scavenging performances of CeO_{2-x} S-hNR and CeO_{2-x} P-hNR are mainly due to their high surface area.

However, the change in surface states with the increased surface concentration of oxygen vacancies and Ce^{3+} states could also have an effect on their radical scavenging activity. Because the CeO_{2-x} S-hNR samples still possess a higher surface concentration of oxygen vacancies than the other samples, these can also improve the $\bullet OH$ radical scavenging activity.

3.2. OPV device preparation and characterization

Zinc oxide (ZnO) is one of the most widely used materials for the electron transport layer in the inverted OPV cell architecture.^{38–41} ZnO layers are typically developed using a sol-gel-based solution⁴² and ZnO nanoparticle solution⁴³ due to the ease of film formation with low annealing temperature and highly reproducible film quality. However, it is also widely known that hydroxyl radicals ($\bullet OH$) can form when ZnO film with H_2O on its surface is exposed to UV irradiation,¹⁶ and the generated radicals can decompose the active film, especially the non-fullerene acceptor (NFA) molecules, at the ZnO/active layer interface.¹⁷ Therefore, if the hydroxyl radicals generated by UV exposure can be efficiently eliminated, one could see a considerable enhancement in the operational stability of the resulting ZnO-based inverted OPVs.

We introduced CeO_{2-x} S-hNR, CeO_{2-x} P-hNR, and commercial CeO_2 NP to the ZnO film in the inverted OPV structure of indium tin oxide (ITO)/ZnO/active layer/ MoO_3/Ag . Prior to application to the OPV architecture, we first investigated the coating behaviour of the CeO_{2-x} films on ZnO using a scanning electron microscope (SEM). Fig. S6 (ESI†) show the SEM images of NP, S-hNR, and P-hNR films on ZnO films, respectively, and Fig. S7 (ESI†) are the elemental mapping images of the Ce line in the above SEM images. As observed, there are aggregates consisting of cerium oxides, regardless of the crystallinity of cerium oxide. However, the CeO_{2-x} S-hNR-based film had the most homogeneous coverage of the ZnO film with the smallest aggregates. Considering that such aggregations can act as pin-holes in active film, and this could lead to a short circuit in an OPV device, we expect that CeO_{2-x} S-hNR-based OPVs show better and more reproducible photovoltaic properties.

Then we introduced various cerium oxide layers between the ZnO layer and active layer in the OPV structure. As seen in Fig. S8 (ESI†), the cerium oxide layers fully cover the spiky surface of ZnO with reduced root-mean-square roughness (R_{rms}) values, irrespective of the crystallinity of cerium oxide. The measured thickness values of cerium oxide layers were ~ 10 nm, regardless of their crystallinity (Fig. S9, ESI†). To confirm the effect of CeO_{2-x} nanoparticle aggregations on

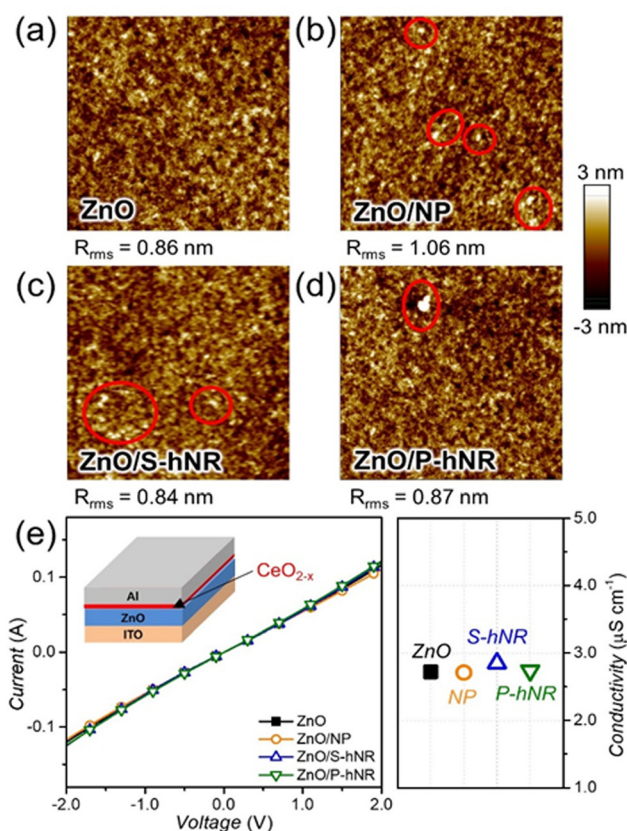


Fig. 3 (a–d) AFM topography images ($5 \mu m \times 5 \mu m$) of thin (~ 40 nm) PM6:N3 blend films on (a) ZnO, (b) ZnO/ CeO_2 NP, (c) ZnO/ CeO_{2-x} S-hNR, and (d) ZnO/ CeO_{2-x} P-hNR layers. The red circles in the images indicate the aggregations in the films. (e) The $I-V$ curves of ITO/ZnO/Al or ITO/ZnO/ CeO_{2-x} /Al sandwiched devices for measuring the conductivity of ZnO or ZnO/ CeO_{2-x} layers. Inset: Schematic illustration of the device configuration for conductivity measurements. The right graph indicates the calculated conductivity by the $I-V$ curve.

the morphology of active films, we observed the morphology of a thin (~ 40 nm) active film on ZnO or ZnO/ CeO_{2-x} films using an atomic force microscope (AFM). The topography images can be found in Fig. 3a–d. Although small aggregates are shown in all the active films on the CeO_{2-x} layers, their root-mean-square roughness values were nearly the same, ranging from 0.84 to 1.05 nm. Therefore, we speculated that the aggregates in the active films would not significantly affect the device performance.

Before fabricating the OPV cells, we tested the effect of CeO_{2-x} layers on the conductivity of the ZnO layers. The current–voltage ($I-V$) curves of the ITO/ZnO/Al or ITO/ZnO/ CeO_{2-x} /Al sandwiched devices were measured, and the $I-V$ curves and the calculated conductivity values can be found in Fig. 3e. The calculated conductivity values were similar, with a range of 2.7 to 2.8 $\mu S cm^{-1}$, regardless of the presence of CeO_{2-x} layers. Therefore, we can expect that the introduction of CeO_{2-x} layers would not lead to a decrease in the electrical properties of the OPV structure. However, it was also found that the introduction of too thick cerium oxide layer can lead to a slight deterioration in electrical conductivity, because the

electrical property of cerium oxide is not as good as that of ZnO (Fig. S10, ESI†).

Then, we fabricated inverted OPV cells with a structure of ITO/ZnO/CeO_{2-x}/active layer/MoO₃/Ag (Fig. 4a). The *J*-*V* characteristics and external quantum efficiency (EQE) spectra of the resultant OPVs can be found in Fig. 4b and c, respectively, and the photovoltaic parameters are summarized in Table S2 (ESI†). As we expected above, the fabricated OPVs showed device performance similar to the PCEs, ranging from 14.6 to 15.0% with similar shapes of EQE spectra (Fig. 4c), irrespective of the types of CeO_{2-x} nanostructures.

Based on the fabricated CeO_{2-x}-integrated OPV cells, we tested the photostability of the OPV cells. The OPVs were encapsulated to prevent any additional degradation by oxygen and water in air. The encapsulated OPVs were exposed to 1 sun condition up to 36 hours and the device performances were recorded at selected light soaking times (Fig. S11, ESI†). The dependency of PCEs on light soaking time is summarized in Fig. 4d and dependencies of other photovoltaic parameters such as *J*_{SC}, *V*_{OC} and FF can be found in Fig. S12 (ESI†).

Interestingly, only the OPV with CeO_{2-x} S-hNR layer showed better photostability than that of ZnO-based reference OPV, and the device performances of CeO₂ NP- and CeO_{2-x} P-hNR-based OPVs were even worse than that of the reference OPV. The ZnO only, CeO₂ NP-, CeO_{2-x} P-hNR- and CeO_{2-x} S-hNR-based OPV devices decreased their average PCEs to -35.3%, -57.3%, -48.6%, and -24.4% of their initial efficiencies after 36 h of light exposure, respectively. All the photovoltaic parameters of the CeO₂ NP- or CeO_{2-x} P-hNR- based OPV cells showed faster decreases than those of ZnO- or CeO_{2-x} S-hNR-based OPVs (Fig. S12, ESI†). To confirm the effect of UV light on photostability of the OPVs, the photostability test was repeated using ZnO- and ZnO/CeO_{2-x} S-hNR-based OPV cells under 1 sun

condition. By attaching polyimide films to backside of the OPV cells, the incident UV light in solar spectrum can be effectively excluded (Fig. S13a, ESI†). As shown in Fig. S13b (ESI†), this photostability was clearly improved regardless of the presence of CeO_{2-x} layers. This result indicates that the photostability issues are mainly related to the hydroxyl radical generated by photocatalysis under UV exposure. As mentioned above, since the generated hydroxyl radicals can attack active materials, the cerium oxide layers in the OPV structure acted as radical scavengers against hydroxyl radicals, thereby suppressing the decomposition of the active layer. However, as being revealed from the analyses using SEM (Fig. S6, ESI†) and AFM images (Fig. 3a-d and Fig. S8, ESI†), CeO_{2-x} S-hNR formed uniform deposition on the ZnO film, in contrast with CeO₂ NP and CeO_{2-x} P-hNR films which were rough and showed large aggregates. It is widely known that the morphology of the active film deposited on a rough surface are more labile than that on a smooth surface by external stimulus, which can accelerate the degradation of OPV devices, especially for the initial step of degradation (burn-in degradation).¹⁰

3.3. Stability enhancement mechanism of CeO_{2-x} in the interlayer

To elucidate the reason why the CeO_{2-x} S-hNR-based OPVs showed better photostability, we collected the absorption spectra of PM6:N3 blend films on ZnO or ZnO/CeO_{2-x} layers after various exposure times to UV. The PM6:N3 blend films were fabricated with an ultrathin thickness of ~10 nm to observe the aspects occurring at the ZnO/active layer or the CeO_{2-x}/active layer interface more directly. Fig. S14 (ESI†) shows the evolution of UV-visible absorption spectra under various UV exposure times up to 6 min. As UV exposure time increases, one can clearly see that the absorption peak near 780 nm gradually decreases. Considering the absorption spectra of PM6 and N3 (Fig. S15, ESI†), the absorption peak near 780 nm is related to the absorption of N3. The trend in absorption of blend films on various layers is summarized in Fig. 5a.

For the blend film on the ZnO layer, the absorption peak near 780 nm was decreased by -53.8% after 6 min of UV exposure. However, with the blend films on CeO_{2-x} layers, relatively small decreases of -50%, -45.9%, and -42.9% were exhibited by the blend films on the CeO₂ NP, CeO_{2-x} P-hNR and CeO_{2-x} S-hNR layers, respectively.

For the absorption peak near 620 nm, which is related to the absorption of PM6, the absorption decreased by less than 10%, irrespective of the presence of cerium oxide layers. Therefore, we can conclude that the decrease in light absorption efficiency due to the degradation of N3 molecules is one of the main reasons for deterioration in PCE. N3 can be decomposed by the attack of •OH radicals generated from ZnO layer to vinyl linkers. Since the conjugation in the decomposed N3 molecules are broken, they cannot act as electron acceptor anymore, resulting in poor electron transfer from N3 to ZnO. However, CeO_{2-x} can suppress those •OH radicals by radical scavenging mechanism,³⁷ thereby enhancing the photostability of the resultant OPV cells.

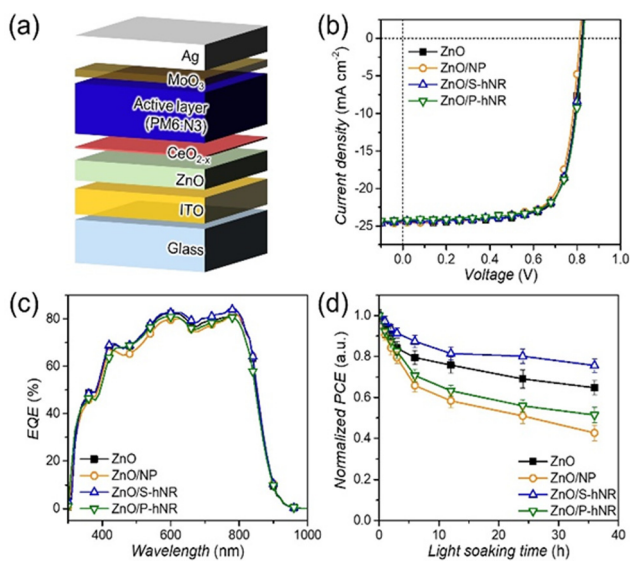


Fig. 4 (a) Schematic illustration of the OPV device. (b) *J*-*V* characteristics and (c) external quantum efficiency (EQE) spectra of the OPVs with various CeO_{2-x}-based interlayers and (d) their dependency of normalized average PCE on light soaking time under 1 sun condition.



Fig. 5 (a) Change in the intensity of 780 nm absorption peaks of the UV-visible absorption spectra (Fig. S14, ESI[†]) of thin PM6:N3 blend films on ZnO, ZnO/CeO₂ NP, ZnO/CeO_{2-x} S-hNR, and ZnO/CeO_{2-x} P-hNR layers under the UV exposure. (b) Raman spectra of ZnO, ZnO/CeO₂ NP, ZnO/CeO_{2-x} S-hNR, and ZnO/CeO_{2-x} P-hNR films, normalized to F_{2g} band near 465 cm⁻¹. (c) Wavelength change at the optical density of 0.30 of the UV-visible absorption spectra (Fig. S16, ESI[†]) of CeO_{2-x}-based aqueous dispersion (695 mM) after injection of H₂O₂ (5 μmol), related to the catalytic degradation of H₂O₂ by surface redox reaction of CeO_{2-x} nanostructures.

Although the CeO_{2-x} S-hNR and the CeO_{2-x} P-hNR showed similar radical scavenging activity in solution phase (Fig. 2a), the OPV device with the CeO_{2-x} S-hNR interlayer showed higher stability against •OH radicals. To elucidate the difference in radical scavenging activity between solution phase and device, we further investigated the states of the CeO_{2-x} based radical scavengers when they were located on ZnO film surfaces.

The Raman spectra (Fig. 5b) of ZnO/CeO_{2-x} films on ITO glass revealed that the ZnO/CeO_{2-x} S-hNR film had a much broader band shape and much larger red shift in the F_{2g} band (465 cm⁻¹) than the ZnO/CeO_{2-x} P-hNR film. Also, the intensity of the 595 cm⁻¹ bands in the ZnO/CeO_{2-x} S-hNR film was much higher than those of the ZnO/CeO_{2-x} P-hNR and ZnO/CeO₂ NP films. This indicates that the CeO_{2-x} S-hNR possessed more surface Ce³⁺ states with defect structures,⁴⁴ which are active toward •OH radical quenching, than the CeO_{2-x} P-hNR when they formed films on the ZnO surface.

Moreover, as observed in the SEM (Fig. S6, ESI[†]) and AFM (Fig. 3c) analyses, the CeO_{2-x} S-hNR-based film had the best coverage on ZnO film. Therefore, we suppose that on the ZnO film, the CeO_{2-x} S-hNR can effectively quench •OH radicals generated from photocatalytic reactions on the ZnO surfaces. This occurs because they can effectively form interfaces on the ZnO surface. For the ZnO/CeO_{2-x} P-hNR (Fig. 3d), on the other hand, several aggregations were observed due to the severe aggregation of CeO_{2-x} P-hNR radical scavengers. As a result, the interfacial CeO_{2-x} P-hNR on the ZnO film could not effectively form an interface with the ZnO, and failed to rapidly quench •OH radicals generated from the ZnO layer.

It has been reported that CeO_{2-x} nanostructures can also possess pro-oxidant properties generating reactive oxygen species under a certain environment rather showing radical-scavenging activity.^{35,45} Therefore, accelerated degradation of OPV devices with CeO₂ NP and CeO_{2-x} P-hNR interlayer might originate from the unwanted side redox reaction on the surface of the CeO_{2-x}. To elucidate this, we observed the catalytic H₂O₂ degradation activity of the three CeO_{2-x} nanostructures, which can be measured by the change in UV-vis spectra of an aqueous

dispersion of CeO_{2-x} after injection of H₂O₂ (Fig. S16, ESI[†]).⁴⁶ In particular, the change in wavelength at the optical density of 0.30, related to the generation of Ce³⁺ species from the surface Ce⁴⁺ states, is the indicator of the H₂O₂ decomposition activity. As shown in Fig. 5c, the wavelength change was found to be in the order of CeO₂ NP > CeO_{2-x} P-hNR > CeO_{2-x} S-hNR. This trend corroborates with OPV results (Fig. 4d), where the CeO₂ NP-based device showed the most rapid degradation rate of the average PCE.

While the exact reaction mechanism is still in debate, the reaction of CeO_{2-x} with H₂O₂ can undergo both anti-oxidative (superoxide dismutase- or catalase-like activity) and pro-oxidative (oxidase- or peroxidase-like activity).^{45,47} In our case, the trends observed in the change in UV-vis spectra are well with the OPV results – the CeO_{2-x} S-hNR, which showed the smallest change in optical density after injecting H₂O₂, only exhibited higher OPV device stability. However, the CeO₂ NP and CeO_{2-x} P-hNR-based OPV showed inferior stability to the ZnO-only OPV device without CeO_{2-x}-based interlayer. Those two nanostructures showed higher activity toward the generation of the Ce³⁺ state *via* catalytic H₂O₂ decomposition, which was observed by the UV-vis spectra. In this case, we suppose that the H₂O₂ decomposition reaction underwent oxidase- or peroxidase-like mechanisms, thereby generating reactive oxygen species. Therefore, it seems that the surface regeneration of the Ce⁴⁺ to Ce³⁺ state is also a key factor for the performance of radical-scavenging CeO_{2-x}-based interlayers for OPV applications.

4. Conclusions

In conclusion, we demonstrated the ability of CeO_{2-x} nanostructures to mitigate the photodegradation process in OPV devices by measuring the scavenging effect of hydroxyl radicals with a focus on the influence of morphology and crystallinity. Material characterization and analyses of device test results revealed that as a radical scavenger the S-hNR nanostructured

CeO_{2-x} was very effective in OPV devices and was responsible for the higher surface concentration of Ce³⁺ and the interfacial compatibility between constituent layers, which all contributed to radical scavenging.

Author contributions

Seongwon Yoon: conceptualization, methodology, investigation, visualization, writing-original draft. Taehyun Kwon: conceptualization, methodology, investigation, visualization, writing-original draft. Sohyun Kim: investigation, visualization. So Hyun Park: investigation. Youngjoon Lim: investigation, validation. Jihun Kim: investigation. Jun Kim: validation. Kwangjin An: validation. Sungmin Park: investigation, writing-original draft. Jin Young Kim: conceptualization, methodology, supervision, funding acquisition, writing-review & editing. Hae Jung Son: conceptualization, methodology, supervision, funding acquisition, project administration, writing-review & editing.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

This work was supported by the program of Future Hydrogen Original Technology Development (NRF-2021M3I3A1082879) through the National Research Foundation of Korea (NRF) funded by the Korean government (Ministry of Science and ICT), the Korea Institute Energy Technology Evaluation and Planning (KETEP) grant funded by the Korea government (MOTIE) (no. 20213030030260), and KIST Institutional Program (2E31871). This work was also supported by the National R & D Program through the NRF of Korea funded by the Ministry of Science and ICT (NRF-2020M3H4A1A02084910), and the Nano Future Material Technology Development Program through the NRF of Korea funded by the Ministry of Science and ICT (NRF-2020M3H4A3081817). T. K. acknowledges the NRF of Korea grant funded by the Korea government (MSIT) (No. 2022R1C1C2004703).

References

- G. Zhang, F. R. Lin, F. Qi, T. Heumuller, A. Distler, H. J. Egelhaaf, N. Li, P. C. Y. Chow, C. J. Brabec, A. K. Jen and H. L. Yip, *Chem. Rev.*, 2022, **122**, 14180–14274.
- P. Xue, P. Cheng, R. P. S. Han and X. Zhan, *Mater. Horiz.*, 2022, **9**, 194–219.
- S. Yoon, E. Y. Shin, N. K. Cho, S. Park, H. Y. Woo and H. J. Son, *J. Mater. Chem. A*, 2021, **9**, 24729–24758.
- Y. Cui, Y. Xu, H. Yao, P. Bi, L. Hong, J. Zhang, Y. Zu, T. Zhang, J. Qin, J. Ren, Z. Chen, C. He, X. Hao, Z. Wei and J. Hou, *Adv. Mater.*, 2021, **33**, e2102420.
- P. Bi, S. Zhang, Z. Chen, Y. Xu, Y. Cui, T. Zhang, J. Ren, J. Qin, L. Hong, X. Hao and J. Hou, *Joule*, 2021, **5**, 2408–2419.
- Y. Wei, Z. Chen, G. Lu, N. Yu, C. Li, J. Gao, X. Gu, X. Hao, G. Lu, Z. Tang, J. Zhang, Z. Wei, X. Zhang and H. Huang, *Adv. Mater.*, 2022, **34**, e2204718.
- W. Gao, F. Qi, Z. Peng, F. R. Lin, K. Jiang, C. Zhong, W. Kaminsky, Z. Guan, C. S. Lee, T. J. Marks, H. Ade and A. K. Jen, *Adv. Mater.*, 2022, **34**, e2202089.
- Z. Zheng, J. Wang, P. Bi, J. Ren, Y. Wang, Y. Yang, X. Liu, S. Zhang and J. Hou, *Joule*, 2022, **6**, 171–184.
- S. Park, T. Kim, S. Yoon, C. W. Koh, H. Y. Woo and H. J. Son, *Adv. Mater.*, 2020, **32**, e2002217.
- L. Duan and A. Uddin, *Adv. Sci.*, 2020, **7**, 1903259.
- P. Cheng and X. Zhan, *Chem. Soc. Rev.*, 2016, **45**, 2544–2582.
- J. Guo, Y. Wu, R. Sun, W. Wang, J. Guo, Q. Wu, X. F. Tang, C. K. Sun, Z. H. Luo, K. Chang, Z. H. Zhang, J. Yuan, T. F. Li, W. H. Tang, E. J. Zhou, Z. Xiao, L. M. Ding, Y. P. Zou, X. W. Zhan, C. L. Yang, Z. Li, C. J. Brabec, Y. F. Li and J. Min, *J. Mater. Chem. A*, 2019, **7**, 25088–25101.
- Y. S. Zhao, Z. N. Wu, X. Liu, Z. P. Zhong, R. H. Zhu and J. S. Yu, *J. Mater. Chem. C*, 2021, **9**, 13972–13980.
- Y. Y. Jiang, L. L. Sun, F. Y. Jiang, C. Xie, L. Hu, X. Y. Dong, F. Qin, T. F. Liu, L. Hu, X. S. Jiang and Y. H. Zhou, *Mater. Horiz.*, 2019, **6**, 1438–1443.
- A. Uddin and H. M. Yi, *Sol. RRL*, 2022, **6**, 2100983.
- Z. M. Wang, K. Wang, H. Wang, X. Chen, W. X. Dai and X. Z. Fu, *Catal. Sci. Technol.*, 2018, **8**, 3260–3277.
- S. Park and H. J. Son, *J. Mater. Chem. A*, 2019, **7**, 25830–25837.
- Y. Wang, Z. Zheng, J. Wang, X. Liu, J. Ren, C. An, S. Zhang and J. Hou, *Adv. Mater.*, 2022, e2208305.
- A. Soultati, A. Fakharuddin, E. Polydorou, C. Drivas, A. Kaltzoglou, M. I. Haider, F. Kournoutas, M. Fakis, L. C. Palilis, S. Kennou, D. Davazoglou, P. Falaras, P. Argitis, S. Gardelis, A. Kordatos, A. Chroneos, L. Schmidt-Mende and M. Vasilopoulou, *ACS Appl. Energy Mater.*, 2019, **2**, 1663–1675.
- F. Cao, M. Zhang, K. Yang, Z. Tian, J. Li and Y. Qu, *Nano Res.*, 2020, **14**, 715–719.
- S. Zhang, Z. Q. Huang, Y. Ma, W. Gao, J. Li, F. Cao, L. Li, C. R. Chang and Y. Qu, *Nat. Commun.*, 2017, **8**, 15266.
- H. X. Mai, L. D. Sun, Y. W. Zhang, R. Si, W. Feng, H. P. Zhang, H. C. Liu and C. H. Yan, *J. Phys. Chem. B*, 2005, **109**, 24380–24385.
- J. Li, Z. Y. Zhang, Z. M. Tian, X. M. Zhou, Z. P. Zheng, Y. Y. Ma and Y. Q. Qu, *J. Mater. Chem. A*, 2014, **2**, 16459–16466.
- Y. Xue, Q. F. Luan, D. Yang, X. Yao and K. B. Zhou, *J. Phys. Chem. C*, 2011, **115**, 4433–4438.
- A. L. Patterson, *Phys. Rev.*, 1939, **56**, 978–982.
- S. Deshpande, S. Patil, S. V. N. T. Kuchibhatla and S. Seal, *Appl. Phys. Lett.*, 2005, **87**, 133113.
- X. Hao, A. Yoko, C. Chen, K. Inoue, M. Saito, G. Seong, S. Takami, T. Adschiri and Y. Ikumura, *Small*, 2018, **14**, e1802915.
- R. Schmitt, A. Nennung, O. Kraynis, R. Korobko, A. I. Frenkel, I. Lubomirsky, S. M. Haile and J. L. M. Rupp, *Chem. Soc. Rev.*, 2020, **49**, 554–592.
- Y. Lee, G. He, A. J. Akey, R. Si, M. Flytzani-Stephanopoulos and I. P. Herman, *J. Am. Chem. Soc.*, 2011, **133**, 12952–12955.

- 30 C. Schilling, A. Hofmann, C. Hess and M. V. Ganduglia-Pirovano, *J. Phys. Chem. C*, 2017, **121**, 20834–20849.
- 31 Z. Wu, M. Li, J. Howe, H. M. Meyer, 3rd and S. H. Overbury, *Langmuir*, 2010, **26**, 16595–16606.
- 32 A. Filippi, F. Liu, J. Wilson, S. Lelieveld, K. Korschelt, T. Wang, Y. Wang, T. Reich, U. Poschl, W. Tremel and H. Tong, *RSC Adv.*, 2019, **9**, 11077–11081.
- 33 T. Kwon, Y. Lim, J. W. Cho, R. Lawler, B. J. Min, W. A. Goddard, S. S. Jang and J. Y. Kim, *Mater. Today*, 2022, **58**, 135–163.
- 34 T. Kwon, S. H. Park, B. J. Min, S. Park, S. Ramadhani, Y. Lim, S. S. Jang, H. Jeong, H. J. Son and J. Y. Kim, *Adv. Energy Sustainability Res.*, 2022, **3**, 2200011.
- 35 M. S. Lord, J. F. Berret, S. Singh, A. Vinu and A. S. Karakoti, *Small*, 2021, **17**, e2102342.
- 36 K. J. Mitchell, J. L. Goodsell, B. Russell-Webster, U. T. Twahir, A. Angerhofer, K. A. Abboud and G. Christou, *Inorg. Chem.*, 2021, **60**, 1641–1653.
- 37 V. Prabhakaran, C. G. Arges and V. Ramani, *Proc. Natl. Acad. Sci. U. S. A.*, 2012, **109**, 1029–1034.
- 38 J. Yuan, Y. Zhang, L. Zhou, G. Zhang, H.-L. Yip, T.-K. Lau, X. Lu, C. Zhu, H. Peng, P. A. Johnson, M. Leclerc, Y. Cao, J. Ulanski, Y. Li and Y. Zou, *Joule*, 2019, **3**, 1140–1151.
- 39 C. Li, J. Zhou, J. Song, J. Xu, H. Zhang, X. Zhang, J. Guo, L. Zhu, D. Wei, G. Han, J. Min, Y. Zhang, Z. Xie, Y. Yi, H. Yan, F. Gao, F. Liu and Y. Sun, *Nat. Energy*, 2021, **6**, 605–613.
- 40 Y. Cui, H. Yao, J. Zhang, T. Zhang, Y. Wang, L. Hong, K. Xian, B. Xu, S. Zhang, J. Peng, Z. Wei, F. Gao and J. Hou, *Nat. Commun.*, 2019, **10**, 2515.
- 41 S. Yoon, S. Park, S. H. Park, S. Nah, S. J. Lee, J. W. Lee, H. Ahn, H. Y. G. Yu, E. Y. Shin, B. J. Kim, B. K. Min, J. H. Noh and H. J. Son, *Joule*, 2022, **6**, 2406–2422.
- 42 Y. Sun, J. H. Seo, C. J. Takacs, J. Seifter and A. J. Heeger, *Adv. Mater.*, 2011, **23**, 1679–1683.
- 43 S. K. Hau, H. L. Yip, N. S. Baek, J. Y. Zou, K. O'Malley and A. K. Y. Jen, *Appl. Phys. Lett.*, 2008, **92**, 253301.
- 44 S. Loidant, *Catal. Today*, 2021, **373**, 98–111.
- 45 V. Baldim, N. Yadav, N. Bia, A. Graillot, C. Loubat, S. Singh, A. S. Karakoti and J.-F. Berret, *ACS Appl. Mater. Interfaces*, 2020, **12**, 42056–42066.
- 46 S. S. Lee, W. Song, M. Cho, H. L. Puppala, P. Nguyen, H. Zhu, L. Segatori and V. L. Colvin, *ACS Nano*, 2013, **7**, 9693–9703.
- 47 T. J. Fisher, Y. Zhou, T.-S. Wu, M. Wang, Y.-L. Soo and C. L. Cheung, *Nanoscale*, 2019, **11**, 4552–4561.