High-entropy-alloy nanoparticles with 21 ultra-mixed elements for efficient photothermal conversion

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ABSTRACT

Multi-metallic nanoparticles have been proven to be an efficient photothermal conversion material, for which the optical absorption can be broadened through the interband transitions (IBTs), but it remains a challenge due to the strong immiscibility among the repelling combinations. Here, assisted by an extremely high evaporation temperature, ultra-fast cooling and vapor-pressure strategy, the arc-discharged plasma method was employed to synthesize ultra-mixed multi-metallic nanoparticles composed of 21 elements (FeCoNiCrYTiVCuAlNbMoTaWZnCdPbAgInMnSn), in which the strongly repelling combinations were uniformly distributed. Due to the reinforced lattice distortion effect and excellent IBTs, the nanoparticles can realize an average absorption of >92% in the entire solar spectrum (250 to 2500 nm). In particular, the 21-element nanoparticles achieve a considerably high solar steam efficiency of nearly 99% under one solar irradiation, with a water evaporation rate of 2.42 kg m⁻² h⁻¹, demonstrating a highly efficient photothermal conversion performance. The present approach creates a new strategy for uniformly mixing multi-metallic elements for exploring their unknown properties and various applications.

Keywords: high-entropy alloy, nanoparticle, ultra-mixing, arc-discharged method, photothermal conversion

INTRODUCTION

Photothermal conversion materials have been attracting considerable attention with regard to their water evaporation applications, as a result of the intensified consumption of clean water and severe pollution of water resources recently [1-5]. To achieve a satisfying photothermal conversion efficiency, the essential condition is to absorb solar energy with wavelengths of between 250 to 2500 nm as much as possible [6-8]. Consequently, a series of mechanisms, such as the plasmonic heating of metallic nanoparticles (e.g. Au [6], Ag [9] and Al [7]), non-radiative relaxation of semiconductors (e.g. copper selenide nanocrystals [10]) and thermal vibration of molecules [11], have been widely investigated. Despite the achievement of photothermal conversion performance, the aforementioned materials still lack the capacity to cover the entire solar spectrum, thus limiting the optimization of optical absorption and solar-driven water evaporation behavior [1,2,6,7,11].

In our recent research, it has been proposed that multi-metallic nanoparticles, so-called high-entropy-alloy nanoparticles (HEA-NPs) [12-15], are one of the most promising candidates for tailoring the optical absorption property [16,17]. Such a performance is induced by the reinforced interband transitions (IBTs), contributed by the fully filled energy regions around the Fermi level [12,18]. Therefore, the solar energy absorption property can be optimized through the addition of composited elements [17], resulting in a broadened working bandwidth and thus improving the photothermal conversion performance. However, when the composited elements exceed 3d transition metals, the related research for understanding the relationship between the HEA-NPs and photothermal conversion performance is still lacking, and the crucial issue is how to cross the immiscibility of composited elements to synthesize the HEAs [19]. Meanwhile, it has been found that phase separation has occurred when elements exceed eight in our previous experiments.
Thus, it is necessary to improve our synthesized method and meanwhile explore the limits of the method. Surprisingly, it has been reported that the HEA-NPs composed of 15 elements with strongly repelling combinations were synthesized through a carbon-thermal shock method, and the immiscibility can be overcome by the high-entropy effect and non-equilibrium synthesized process [20,21], promising a pathway for breaking the restrictions of elemental combination with strong repellency.

Herein, assisted by non-equilibrium synthesized conditions including extremely high evaporation temperature and ultra-fast cooling rate, we synthesized a series of HEA-NPs including 9, 13, 17 and 21 composited elements (denoted 9-, 13-, 17- and 21-HEA-NPs) through an arc-discharged method with vapor-pressure (V.P.) strategy. It has been found that the V.P. design could counterbalance the evaporation rates of composited elements, and plays a critical role in realizing the HEAs from the immiscible combinations. Meanwhile, the photothermal conversion performance can be improved along with the addition of composited elements. As a result, 21-HEA-NPs exhibit an excellent solar-steam generation performance with nearly 99% solar steam efficiency and 2.42 kg m⁻² h⁻¹ water evaporation rate under 1 sun irradiation (1.0 kW m⁻²). The present work opens a vast space for synthesizing HEA-NPs with immiscible elements, which broadens the family of HEA-NPs and provides a great platform for exploring the application of HEA-NPs.

RESULTS AND DISCUSSION

Fabrication and microstructure characterizations of HEA-NPs

HEA-NPs, composed of 9, 13, 17 and 21 metallic elements, were synthesized by arc-discharging the compressed mixture of metallic powders in a mixed atmosphere of argon and hydrogen (Fig. 1a) [22,23]; the ratio of different powders was controlled via the V.P. strategy to realize the uniform mixing of compositions (Supporting Text in Supplementary Data) [21]. The macroscopic contents for HEA-NPs determined by inductively coupled optical emission spectroscopy (ICP-OES) (Table S1, Supplementary Data) were used to estimate the mixing entropy (∆S_mix) of HEA-NPs. The ∆S_mix of 9-, 13-, 17- and 21-HEA-NPs was calculated to be 17.83, 19.79, 21.82 and 24.28 J mol⁻¹ K⁻¹, respectively [17,21,24]. It was found that the mixing entropy increases synchronously with the increasing of alloying combinations. The entropy of 21-HEA-NPs was 2.69 times higher than that of 3-HEA-NPs (9.12 J mol⁻¹ K⁻¹), while the increased entropy could be stabilized due to the high-entropy effect [20]. The elemental distribution of 21-HEA-NPs is illustrated in Fig. 1c, using the scanning transmission electron microscope (STEM) assisted with energy-dispersive X-ray spectroscopy (EDS). Meanwhile, the high-resolution TEM image and atomic EDS maps of 21-HEA-NPs have been exhibited in Fig. 2, in which a uniform elemental distribution of most of the composited elements can be observed, while the Sn and In exhibit a slightly constituent fluctuation. As for the phase structures in Fig. 1b, the crystalline structures were changed from face-centered-cubic (FCC) to body-centered-cubic (BCC) when the composited elements exceeded nine. In addition, it can be noted that the X-ray diffraction (XRD) peaks of 13-, 17- and 21-HEA-NPs was broadened and shifted to a lower 2θ value, which is ascribed to the lattice distortion of the high-entropy effect and the addition of larger atomic radius elements including Ta, W, Pb and Bi [25,26].

Here, the synthesized process has been divided into three parts: extremely high evaporation temperature, ultra-fast cooling rate and V.P. strategy, as illustrated in Fig. 3a. According to our previous study, the high evaporation temperature created from the arc-discharged plasma can result in a non-equilibrium synthesized atmosphere, fabricating the HEA-NPs with the composited elements of Fe, Co, Ni, Ti, V, Cr and/or Cu [17]. However, it has been found that the ratio of Ti and V in the 7-HEA-NPs is relatively lower than the other composited elements. Such a phenomenon is induced by the vapor pressure of different composited elements, as exhibited in Fig. S1 (Supplementary Data), in which a low vapor pressure of Ti/V can be observed compared to other elements, indicating a slower evaporation process during the synthesized process. Therefore, to confirm the effect of the V.P. strategy, the nanoparticles that are composed of ZnCdPbBi, AgInMnSn and NbMoTaW were synthesized (Fig. 3b and Fig. S2, Supplementary Data). The phase separation can be observed in the former two samples through the XRD patterns, and the uniform mixing behavior appears in the NbMoTaW nanoparticles (Fig. S2, Supplementary Data).

Consequently, based on the curves of vapor pressure [21,27] (Fig. S1, Supplementary Data), different metals used in this work could be divided into three groups: (i) metals with low vapor pressure (V.P.₁: Mo, Nb, Ta, W), (ii) metals with medium vapor pressure (V.P.₁Ι: Fe, Co, Ni, Y, Ti, V, Al, Cu, Cr, Sn), in which the Ti and V are sub-V.P.₁ metals and Sn, Al and Cu are sub-V.P.₁Ι metals; (iii) metals
with high vapor pressure (V.P., Ag, Mn, In, Pb, Bi, Zn, Cd). Following this guideline, the 9-HEA-NPs of FeCoNiCrYTiVCuAl were synthesized from the mixture with a 10-time content of Ti and V and a 0.5-time content of Cu and Al (marked as FeCoNiCrYT10V10Cu0.5Al0.5). The corresponding synthesized rule to fabricate the HEA-NPs through the V.P. strategy has been summarized in the Vapor pressure (V.P.) strategy for the preparation of HEA-NPs section in the Supplementary Data, and it can be recognized that the optimization of chemical composition is highly correlated to the difference in vapor pressure of the composited elements. The EDS maps reveal the uniform elemental distribution of the composited elements, while the crystalline structure exhibits a combination of FCC and BCC, indicating the lattice distortion (Figs 1b and 3c). Here, using the 9-HEA-NPs as the substrate, we explore the effectiveness of the V.P. strategy, and a series of 13- and 17-element nanoparticles were synthesized. Uniform mixing and phase separation can be realized in different nanoparticles (Figs S10–S14, Supplementary Data), and it can be concluded that the V.P. elements are key to optimizing the evaporation rate for alloying the immiscible combinations together (Supporting Text). In addition, the 13- and 17-HEA-NPs exhibit an FCC crystalline structure with uniform distribution of composited elements, as shown in Fig. 3d and e and Figs S3 and S4 (Supplementary Data).

Furthermore, several fundamental analyses were carried out to illustrate the characters of HEA-NPs.
Figure 2. High-resolution TEM image, fast Fourier transform (FFT) image and the corresponding atomic TEM-EDS maps of 21-HEA-NPs, scale bar: 1 nm.

The TEM analysis shows that all samples exhibit a sphere-like shape, and the diameter of 9-, 13-, 17- and 21-HEA-NPs was calculated to be 75, 72, 82 and 62 nm, respectively (Figs S5 and S6, Supplementary Data). It can be noted that the size of nanoparticles reduced noticeably when ultra-fast-cooling treatment was activated. The surface state of 21-HEA-NPs was characterized by X-ray photoelectron spectroscopy (XPS), and the result shows that the metals with an oxidation potential have already been oxidized under the storage process in ambient air (Figs S17–S19, Supplementary Data) [28–30].

Optical absorption property and photothermal conversion performance of HEA-NPs

The optical absorption performances of HEA-NPs have been investigated through a scattering spectrum in the wavelength region of 250 to 2500 nm (UV-Vis-NIR, ultraviolet-visible-near infrared), as shown in Fig. 4c, wherein 100 mg nanoparticles were placed in the circle area with a diameter of 24 mm (0.737 g/cm³). It can be recognized that, of all the samples, the 21-HEA-NPs exhibit the highest absorption property (>97%) at the NIR region, while absorption at the UV-Vis regions is reduced compared to the 9-HEA-NPs but still remains ~95%. As for the HEAs compositing with 3d transition metals, it has been reported that their optical absorptions at visible light can be reinforced through the addition of V and Cr, while at UV light can be enhanced through the composition of Ti and Cu [17]. Such a performance is correlated with the different 3d band localized energy and the d–d IBTs. Therefore, the absorption of entire solar spectrum can be realized in the combination of all 3d transition metals, but the absorption at visible light can be reduced through the addition of main group elements [31,32], which is ascribed to decreasing IBT behavior [17], and in agreement with the optical absorption property of the 21-HEA-NPs. Meanwhile, the optical absorption properties of all the HEA-NPs have been compared with carbon-based materials including graphene, graphite, carbon nanotubes and reduced graphene oxide (Table S3, ref. (S.M.) 23 to 29, Supplementary Data). It can be seen that the HEA-NPs demonstrated the best optical performance among all the materials.

Afterward, the photothermal conversion performances of all the samples were investigated in the wavelength regions of 250 to 2500 nm; 100 mg HEA-NPs were irradiated under simulated sunlight of 1.0 kW m⁻² (1 sun) and their surface temperatures were recorded through an infrared (IR) camera. As exhibited in Figs 3d and 4a, the surface temperature of the
21-HEA-NPs can increase to \( \sim 107^\circ\)C within 90 s and rapidly reduce to room temperature when the simulated light is removed. Such a phenomenon can also be recognized in other HEA-NPs under 1 sun irradiation, and it is found that the maximum surface temperatures of all the samples can exceed 100°C after \( \sim 150\) s irradiation (Fig. 4b), evidence for an excellent photothermal conversion performance.

**Water evaporation performance of HEA-NPs**

Owing to the great photothermal conversion performances, the water evaporation behavior of HEA-NPs was investigated by measuring the mass change of pure water under sunlight irradiation (Figs S20 and S21, Supplementary Data). To deeply understand the facilitation performance of HEA-NPs, the evaporation rates of pure water and original nylon membrane were calculated to be 0.67 and 0.69 kg m\(^{-2}\) h\(^{-1}\), respectively. Here, the evaporation rates of the 9-, 13-, 17- and 21-HEA-NPs under 1 solar irradiation (Fig. 5a) were analyzed to be 2.20, 2.26, 2.33 and 2.42 kg m\(^{-2}\) h\(^{-1}\), respectively, indicating that the evaporation rate of HEA-NPs can be improved through the ultra-mixing strategy of the composited elements. Moreover, the evaporation rate of 21-HEA-NPs has also been measured under the different simulated sunlight energy, in which the average water evaporation rate achieves 2.42, 4.84, 7.60, 11.10 and 15.12 kg m\(^{-2}\) h\(^{-1}\) under the light intensities of 1, 2, 3, 4 and 5 kW m\(^{-2}\), respectively (Fig. 5b and c, Fig. S22 and Table S4, Supplementary Data), indicating an excellent solar steam generation performance.

In addition, the evaporation rates of all the samples have been evaluated in saline water. The saline concentration was fixed at the average salinity of the world’s oceans (3.5 wt%). As shown in
Fig. 5d, the evaporation rates of the 9-, 13-, 17- and 21-HEA-NPs under 1 solar irradiation in the saline water were calculated to be 2.17, 2.27, 2.29 and 2.43 kg m\(^{-2}\) h\(^{-1}\), respectively. Such a performance strongly suggests that the photothermal conversion performance of HEA-NPs cannot be influenced by inorganic salt. In addition, the evaporation rates of 21-HEA-NPs under different energy intensities (1 to 5 kW m\(^{-2}\)) were calculated to be 2.43, 4.71, 7.70, 11.75 and 14.47 kg m\(^{-2}\) h\(^{-1}\), respectively (Fig. 5e and f, Fig. S22 and Table S4, Supplementary Data). It is noted that the HEA-NPs in saline water (3.5 wt%) have a similar photothermal conversion performance to those in pure water, indicating great potential for their application in seawater desalination.

The solar steam efficiency of solar light (\(\eta\)), another important value for evaluating photothermal conversion performance, has been calculated based on Equation S3 (Supplementary Data)\(^7,33\). The average steam generation efficiency of the 9-, 13-, 17- and 21-HEA-NPs in pure water is 93.55%, 94.64%, 96.15% and 97.97%, respectively, and in saline water (3.5 wt%) is 93.17%, 93.25%, 96.20% and 97.83%, respectively (Table S5, Supplementary Data). Such a result indicates that efficiency can be improved along with the increase of composited elements. In particular, the steam generation efficiency...
of the 21-HEA-NPs could even exceed 99% during measurement, exhibiting an excellent photothermal conversion performance. To illustrate the excellent evaporation performance, the evaporation rates of HEA-NPs was compared to other previously reported materials (Table S6, ref. (S.M.) 30 to 53 in Supplementary Data), and the samples, including 7-, 9-, 13-, 17- and 21-HEA-NPs, possessed satisfactory water evaporation performance in the related reports (Fig. 6d).

The desalination performance of 21-HEA-NPs was evaluated in four artificial seawater samples with a salinity of 0.8 wt% (Baltic Sea), 3.5 wt% (world ocean), 4 wt% (Red Sea) and 10 wt% (Dead Sea) (Fig. S23, Supplementary Data) [7,34]. The average water evaporation rates of 21-HEA-NPs in 0.8, 3.5, 4 and 10 wt% seawater were 2.38, 2.46, 2.48 and 2.37 kg m⁻² h⁻¹, respectively (Fig. S23, Table S7, Supplementary Data), and the corresponding solar steam efficiencies were calculated to be 97.13%, 98.04%, 98.06% and 96.77% (Table S7, Supplementary Data). The concentration ions, including Na⁺, Mg²⁺, Ca²⁺ and K⁺, in collected water, have been reduced by about three orders of magnitude compared to that of simulated saline water before the desalination (Fig. 6a and b and Fig. S24, Supplementary Data). The salinity of collected water perfectly fits the standards of the World Health Organization (WHO) and the US Environmental Protection Agency (EPA) [35,36]. Therefore, the HEA-NPs are good candidates for photothermal conversion materials used for solar steam generating and seawater desalinating applications. Meanwhile, the long-term stability of the evaporation performance of 21-HEA-NPs has been evaluated in 10 wt% saline water under 1 sun irradiation. It was recognized that the evaporation rate and solar steam efficiency were stabilized at ~2.36 kg m⁻² h⁻¹ and 95.52% after 100 h continuous measurement, respectively (Fig. 6e and Fig. S25, Supplementary Data), indicating that the HEA-NPs could serve as excellent desalination materials.

The photothermal conversion performance of the HEA-NPs was enhanced after the ultra-mixing of composites' elements. Such an enhancement can be attributed to two main factors, including the d–d IBTs and the lattice distortion effect (high-entropy induced). Commonly, the energy of d bands in transition metals is gradually moved below the Fermi level when electron is increasing [17,37]. For example, the 3d bands of Fe, Co and Ni are located around the Fermi level, Ti, V and Cr are above the Fermi level, and Mn and Cu are below the Fermi level. Therefore, through composed with the 3d elements from different energy regions in the 21-HEA-NPs, a stronger d–d IBT can be realized, in which the energy of d bands around...
the Fermi level can be fully filled [38,39], resulting in an improved solar energy absorption property [17]. In our previous report, we showed that the energy regions were fully filled in 7-HEA-NPs (FeCoNiTiVCrCu), which demonstrated an excellent photothermal conversion (water evaporation rate) of 2.26 kg m\(^{-2}\) h\(^{-1}\) under 1 sun irradiation [20]. It is known that the addition of main group elements can reduce the performance of IBTs [31,32], however, the performance of 21-HEA-NPs was increased to 2.43 kg m\(^{-2}\) h\(^{-1}\). Thus, the IBT is not the only factor dominating photothermal conversion performance; it can be optimized via the lattice distortion effect. In general, there are four types of high-entropy effect—the high-entropy, the lattice distortion, the sluggish diffusion and the ‘cocktail’ effect [13,14,40–42]—which result in excellent performances characterized by, for example, good ductility, corrosion resistance and high strength of the HEAs [25,43,44]. Here, the lattice distortion effect of HEAs could result in the random occupation of metallic atoms in a crystalline structure, reducing the electrical and thermal conductivity [42,45,46]. The reduction of thermal conductivity can result in the accumulation of heat under solar irradiation, increasing the surface temperature of nanoparticles and thus enhancing the water evaporation rate. As demonstrated in XRD results (Fig. 1b), it can be concluded that lattice distortion has occurred with the addition of larger-atomic-radius elements including Ta, W, Pb and Bi. Consequently, the thermal conductivity of all the samples was carried out (Table S8, Supplementary Data), and it can be noted that the values were reduced with the increase of composite elements. In particular, the thermal conductivity of 21-HEA-NPs was reduced by >15% compared to that of 7-HEA-NPs, showing that the enhancement of photothermal conversion performance is associated with the coupling between IBTs and lattice distortion effect.

**Outdoor application of 21-HEA-NPs**

Finally, to visually demonstrate the water evaporation performance of 21-HEA-NPs in the natural environment, the experimental equipment was placed on the terrace of the gymnasium at Shenyang (41.76° N, 123.42° E) for outdoor tests on a typical sunny day (28 August 2021) (Fig. S26, Supplementary Data). The temperature on that day was 17–28°C with an III-level southwesterly wind. The incident sunlight flux, evaporation rate, temperature and mass changing were recorded and are exhibited in Fig. 6c and Fig. S27 (Supplementary Data). The surface temperature of the HEA-NPs rose to the maximum value of 48.8°C at noon while...
the evaporation rate reached 4.13 kg m$^{-2}$ h$^{-1}$ under $\sim0.80$ sun irradiation. On a cloudy day, in comparison, the evaporation behavior was reduced, but still remained at a 2.24 kg m$^{-2}$ h$^{-1}$ rate under 0.57 sun irradiation (Fig. S27, Supplementary Data). In addition, the 21-HEA-NPs enable an evaporation rate of 0.72 and 0.50 kg m$^{-2}$ h$^{-1}$ at night on a sunny and cloudy day, respectively. It can be concluded that 21-HEA-NPs exhibit excellent photothermal conversion performance in the natural environment, and can be used in other photic applications.

**CONCLUSION**

In summary, a series of HEA-NPs with 9, 13, 17 and 21 composited elements have been synthesized through an arc-discharged plasma method with the assistance of the V.P. strategy. In particular, a record ultra-mixed 21-element HEA-NP was successfully synthesized from a strongly repelling system, which exhibited a BCC structure and uniformly elemental distribution. Meanwhile, due to the $d-d$ IBTs and the lattice distortion effect, such HEA-NPs demonstrated excellent photothermal conversion performance; the surface temperature reached 107$^\circ$C within 90 s under simulated sunlight irradiation (1 sun). The evaporation rate and photothermal conversion efficiency are 2.42 kg m$^{-2}$ h$^{-1}$ and 99%, respectively, under 1 solar irradiation, demonstrating the highest photothermal conversion performance among all the HEA-NPs. Our work provides new insights into HEA-NP discovery and optimization, introducing a new approach to manipulating the compositions of high-entropy alloy and thus achieving differentiated performance design. Meanwhile, the relationship between photothermal conversion performance and high-entropy effect has been clarified, and could be expanded to other solar-based applications.

**MATERIALS AND METHODS**

**Arc-discharged plasma method for synthesizing HEA-NPs**

The arc-discharged method was achieved by an ultrfast-cooling rate arc-discharged furnace designed by our research group [22,23]. The original metallic powders (purity in Table S9) were first pressed into the cylinder shape and then placed into the vacuum chamber. The reaction gas H$_2$ and Ar were introduced into the chamber after the vacuum degree was reduced to 5 $\times$ 10$^{-3}$ Pa. The arc-discharge process would be sustained for $\sim$5–10 min to make sure that the metallic powders had been reproduced into nanoparticles. The cooling collection plate was set at 300 and 150 K by employing water and liquid nitrogen as the cooling media. The as-made nanoparticles were collected after passivating for $\sim$6 h and then stored in an ambient environment.

**Structural characterizations**

The microstructure and morphology of the nanoparticles were characterized by field-emission transmission electron microscope (FE-TEM, JEOL JEM-2100F, JEOL ARM-200 and Thermo Scientific Talos F200S) with an accelerating voltage of 200 kV. As for the preparation of the TEM sample, the nanoparticles were firstly dispersed in the ethanol and then dropped on the copper mesh. The particle size was measured by ImageJ software. XRD was performed using a Smart Lab9KW (Rigaku) with a scan step of 0.04$^\circ$ in the 2$\theta$ region of 20$^\circ$ to 90$^\circ$. XPS was performed on a Thermo ESCALAB 250. The metal contents of HEA-NPs were analyzed by ICP-OES (PerkinElmer Optima 5300DV). The solutions were prepared by digesting the samples in a mixture of hydrochloric acid (HCl), hydrofluoric acid (HF) and perchloric acid (HClO$_4$). The thermal conductivity was tested through a thermal constantan analyzer (TPS2500S, Hot Disk) at 40$^\circ$C.

**Photothermal conversion performance tests**

Solar absorption performance was characterized by ultraviolet-visible spectroscopy (UV-3600, Shimadzu) with an attached integrating sphere (ISR-3100) in the UV-vis-NIR region. The absorption spectra were obtained by measuring the diffuse reflectance recorded ($A\% = 1-R\%$). IR thermal images were collected by the HIKMICROH36, and the solar conversion performance was measured via an IR thermometer (SA-D2580A, Wuxi Shiao Technology Co., Ltd.). The water evaporation rate was measured based on the mass loss using an electronic mass balance (FA3204C, 0.1-mg accuracy) with a communication model RS322 connected with the computer. A Xe light with AM 1.5 filter (1.0 to 5.0 kW m$^{-2}$, CEL-PF300L-3A, Beijing Education Au-light Co., Ltd.) was employed as the simulated sunlight. The light intensity was monitored by a full-spectrum optical power meter (CEL-FZ-A, Beijing Education Au-light Co., Ltd.). The surface temperature of HEA-NPs was recorded by the SA-D2580A and a recorder (SA-JLY01A4, Wuxi Shiao Technology Co., Ltd.) for 60 min with an interval of 1 s. The simulated seawater, with salinity equaling the average of the world’s oceans, was prepared by dissolving 1.314 g NaCl, 0.037 g KCl, 0.05 g CaCl$_2$, 0.305 g MgCl$_2$·6H$_2$O and 0.059 g MgSO$_4$ in 50 mL
deionized water. Other simulated seaways (Baltic Sea, Red Sea, Dead Sea) were prepared by scaling up and down the salinity of the simulated world’s oceans.

SUPPLEMENTARY DATA
Supplementary data are available at NSR online.

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AUTHOR CONTRIBUTIONS
Y.X.L. and X.F.Z. conceived the idea and designed the present work. Y.J.L., Z.Z.J., Z.Y.Z. and Y.X.L. carried out the synthesis. Y.X.L., L.Z.J., Y.J.L. and X.L.L. carried out the TEM characteristics. Y.J.L., Z.Y.Z. and Y.X.L. performed the photothermal conversion measurements. All authors discussed the results and contributed to the final manuscript.

Conflict of interest statement. None declared.

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