5 nm NiCoP nanoparticles coupled with g-C₃N₄ as high-performance photocatalyst for hydrogen evolution

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ABSTRACT Graphitic carbon nitride (g-C₃N₄) coupled with NiCoP nanoparticles with sizes around 5 nm have been fabricated *via* a controllable alcoholothermal process. NiCoP is an excellent electron conductor and cocatalyst in photocatalytic reactions. The coupling between tiny NiCoP nanoparticles and g-C₃N₄ through *in-situ* fabrication strategy could be a promising way to eliminate the light screening effect, hinder the recombination of photo-induced charge carriers, and improve the charge transfer. The NiCoP/g-C₃N₄ nanohybrids exhibit an excellent photocatalytic activity in the hydrogen generation, with a significantly improved performance compared with original g-C₃N₄, CoP/g-C₃N₄ and Ni₂P/g-C₃N₄, respectively. This study paves a new way to design transition metal phosphides-based photocatalysts for hydrogen production.

Keywords: transition metal phosphides, photocatalytic hydrogen generation, carbon nitride, nanohybrids

INTRODUCTION

The solar power as a main source among renewable energies, has attracted a great deal of attention over the past years. Since the first report of water splitting on semiconductor-based photocatalysts by Fujishima and Honda [1] in the 1970s, photocatalysis with semiconductors has become a quite promising technique in solar energy harvesting [2]. Carbon nitride with a graphitic structure (g-C₃N₄) has great potential as non-metal and environment-friendly visible light-driven catalyst due to the tunable electronic structure [3,4]. However, due to the stacked structure, pristine g-C₃N₄ also has many drawbacks, such as low specific surface area, quick recombination of photo-induced electrons and holes, weak electrical conductivity, poor absorption of visible spec-

trum and utilization efficiency [5]. In order to boost the activity of g-C₃N₄, cocatalysts are introduced to avoid the rapid recombination of photo-generated charge carriers [6–12]. So far, precious metals like platinum (Pt), have been widely utilized as effective cocatalysts for photocatalytic H₂ evolution [13,14]. However, the high-cost and scarcity are the major obstacles for their practical use at a large scale. Therefore, exploring non-noble-metal cocatalysts for photocatalytic hydrogen production becomes a very important topic.

Transition metal phosphides (TMPs) have metallic characteristics and good electrical conductivity [15–19], which are desirable as cocatalysts for photocatalytic hydrogen evolution. As the important members in TMPs, the low cost and high activity of FeP [20,21], Ni₂P [22–25] and CoP [26–29] have been demonstrated as effective cocatalysts towards photocatalytic H₂ evolution. The ternary NiCoP has been reported to exhibit an enhanced electrochemical property than mono-metal phosphides on hydrogen evolution reaction (HER) [30–34]. Since it is widely accepted that an excellent HER electrocatalyst can also promote photocatalytic H₂ evolution as a co-catalyst [35,36], research on NiCoP as a competent co-catalyst has drawn considerable interests [37]. Bi *et al.* [38] adopted one-pot annealing method to fabricate a highly efficient photocatalyst of NiCoP/g-C₃N₄ with an average particle size around 10 nm for hydrogen generation. Qin *et al.* [39] synthesized a NiCoP-based core/shell cocatalyst, which was mixed with g-C₃N₄ by grinding method. Recent progress has provided evidence that the particle size is important to the catalytic activity [40,41]. Generally, reducing the particle size could improve the performance of the photocatalyst due to the enhanced quantum con-

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finement effect, which will shorten the route length of photoexcited charge transmission [42–46]. In order to achieve a better photocatalytic activity, the cocatalyst, i.e., NiCoP, needs to have a tiny particle size as well as uniform distribution on the g-C₃N₄ to avoid impeding light absorption, and couple with g-C₃N₄ to improve the transfer of charge. However, the *in-situ* fabrication of g-C₃N₄ photocatalyst with uniformly distributed tiny NiCoP nanoparticles is still a challenge.

NiCoP/g-C₃N₄ nanohybrids with uniformly distributed NiCoP nanoparticles with a particle size around 5 nm are developed based on an *in-situ* growth strategy. The NiCoP/g-C₃N₄ nanohybrids possess a greatly improved photocatalytic activity than pristine g-C₃N₄ in H₂ evolution due to the enhanced charge separation and transfer, which originate from the small size of NiCoP nanoparticles, uniform distribution of NiCoP on g-C₃N₄, and the coupling between NiCoP and g-C₃N₄. The design, fabrication and in-depth characterization of NiCoP/g-C₃N₄ exhibit the great potential of TMP-based materials as noble-metal-free cocatalysts in photocatalysis for highly efficient hydrogen evolution.

EXPERIMENTAL SECTION

Chemicals

Urea, Ni(Ac)₂·4H₂O, Co(Ac)₂·4H₂O, 1,3-propanediol, isopropanol, sodium hypophosphite hydrate, *N,N*-dimethylformamide (DMF) and methanol were analytical grade and bought from Shanghai Aladdin Ltd. (China).

Synthesis of g-C₃N₄

Bulk g-C₃N₄ was fabricated by the urea precursor *via* a direct thermal condensation process. Urea powder (10 g) was placed in a 50 mL crucible with cover and then thermally treated at 550°C for 2 h under ambient atmosphere with a 5°C min⁻¹ ramp rate. After naturally cooling down, the yellowish agglomerated sample was obtained and carefully ground into fine powders in an agate mortar.

Synthesis of NiCo-OH/g-C₃N₄

The NiCo hydroxides (NiCo-OH/g-C₃N₄) were fabricated by an alcoholothermal method. 200 mg g-C₃N₄ was dispersed in 37.6 mL isopropanol under ultrasonication for 45 min. Meanwhile, certain amounts of Ni(Ac)₂·4H₂O (3.4, 10.2, 17, 23.8, 68 and 102 mg) and Co(Ac)₂·4H₂O (3.4, 10.2, 17, 23.8, 68 and 102 mg), corresponding to the specific loadings of NiCoP on g-C₃N₄ (1, 3, 5, 7, 20 and 30 wt%), were added into 2.4 mL of 1,3-propanediol to

obtain a clear solution. The above solution was dropwise added into the g-C₃N₄ suspension. The obtained mixture was stirred for 2 h, and then moved into a Teflon-lined autoclave, maintained at 160°C for 12 h. After cooling down, the resulting product was washed by deionized water and ethanol through centrifugation, and dried at 80°C to obtain NiCo-OH/g-C₃N₄. Pristine NiCo-OH was synthesized *via* the similar steps without adding g-C₃N₄.

Synthesis of NiCoP/g-C₃N₄

The NiCoP/g-C₃N₄ was fabricated by thermal phosphidation of the as-prepared NiCo-OH/g-C₃N₄. Briefly, 0.1 g NiCo-OH/g-C₃N₄ with different loading amounts of NiCo-OH was mechanically blended with 50 mg of NaH₂PO₂·H₂O using a mortar until a uniform powder was formed. The mixture was then heated in a tube furnace with a ramp rate of 5°C min⁻¹ and kept at 300°C for 2 h under Ar atmosphere. The resulting sample was washed by water and ethanol, and then dried in an oven. The final material was labeled as NiCoP-*x*/C₃N₄ (*x*=1, 3, 5, 7, 20, and 30), where *x* stands for the theoretical weight percentage of NiCoP. The same procedure was also used to obtain pristine NiCoP using 150 mg of NiCo-OH and 350 mg of NaH₂PO₂·H₂O. The 3 wt% of Ni₂P and CoP on g-C₃N₄ were synthesized by the same procedure as that of NiCoP-3/C₃N₄, except for the absence of Co²⁺ or Ni²⁺ in the alcoholothermal reaction, which were denoted as that of Ni₂P-3/C₃N₄ and CoP-3/C₃N₄, respectively. Thermogravimetric analysis (TGA) was conducted to confirm the actual content of cocatalysts on g-C₃N₄ (Fig. S1). TGA results show that the actual weight percentage of NiCoP, N₂P and CoP on g-C₃N₄ is very close to the theoretical value.

Characterization

The X-ray diffraction (XRD) was obtained from a Rigaku SmartLab using Cu K α radiation. X-ray photoelectron spectra (XPS) and ultraviolet photoelectron spectra (UPS) were measured on a Thermo Fisher Scientific ESCALAB 250Xi. Scanning electron microscopy (SEM) imaging was conducted on a FEI Verios 460L. Transmission electron microscopy (TEM) imaging, high resolution TEM (HRTEM) imaging and scanning TEM (STEM) imaging were implemented on a JEOL JEM-3010. UV-Vis diffuse reflectance spectra were examined by a PerkinElmer Lambda 750 with BaSO₄ standard. Photoluminescence (PL) spectrum was recorded at ambient temperature on a Hitachi F-4600 fluorescence spectrophotometer with 365 nm excitation wavelength. TGA was performed under 30% O₂/N₂ at a heating rate of 10 K min⁻¹ on a TA

SDT Q600 analyzer.

Photocatalytic evaluations

The photocatalytic activities for hydrogen generation were assessed in a gastight and evacuation system connected circulation, in which a top-irradiation-type Pyrex cell and a 300 W xenon lamp with 300–780 nm wavelength were used. 100 mg photocatalyst was dispersed in the reactor containing 10 mL methanol and 90 mL water under vigorous stirring. The suspension was degassed by evacuation for 15 min before irradiation. The temperature of the reactor was kept at 20°C by a flowing circular water-cooling system. The amount of evolved hydrogen was analyzed by an on-line gas chromatograph (Tech-comp GC7900, TCD detector and 5A molecular sieve capillary column, Ar as carrier gas). The apparent quantum efficiency (AQE) was measured under the same condition as that of photocatalytic reaction, except using a 420 nm band pass filter. The intensity of monochromatic light on the reactor was $\sim 6.2 \text{ mW cm}^{-2}$ over an area of 19.6 cm^2 . The AQE was calculated using the following Equation (1):

$$\begin{aligned} \text{AQE}(\%) &= \frac{\text{number of reacted electrons}}{\text{number of incident photons}} \times 100 \\ &= \frac{\text{number of evolved H}_2 \text{ molecules} \times 2}{\text{number of incident photons}} \times 100. \end{aligned} \quad (1)$$

Electrochemical analysis

The electrochemical properties were tested on a Chenhua CHI 760E electrochemical workstation by employing a standard three-electrode cell in $0.5 \text{ mol L}^{-1} \text{ Na}_2\text{SO}_4$ solution. The light source was a 300 W Xe lamp equipped with a 400 nm cutoff filter. The reference electrode and counter electrode were Ag/AgCl (saturated KCl) and Pt foil ($1 \times 2 \text{ cm}^2$), respectively. The working electrode for electrochemical measurements was prepared as follow: 1 mg sample was ultrasonically dispersed in DMF and then milled to form homogeneous paste. The paste was uniformly deposited over indium tin oxide (ITO) glass with 2.0 cm^2 of exposed area, and annealed at 300°C under Ar atmosphere.

RESULTS AND DISCUSSION

Fig. 1 displays the XRD results of original $\text{g-C}_3\text{N}_4$ and different $\text{NiCoP-}x/\text{C}_3\text{N}_4$ ($x = 1, 3, 5, 7, 20$ and 30). All the samples have two characteristic diffraction peaks at 13.0° and 27.4° indexed to $\text{g-C}_3\text{N}_4$, which are attributed to the (100) plane with in-plane repeating period of heptazine

Figure 1 XRD results of NiCoP, $\text{g-C}_3\text{N}_4$ and $\text{NiCoP-}x/\text{C}_3\text{N}_4$.

framework and the (002) plane with interlayer stacking of conjugated aromatic structures [47,48]. For the $\text{NiCoP-}x/\text{C}_3\text{N}_4$, when the content of NiCoP is below 20%, the diffraction peaks corresponding to NiCoP cannot be observed due to the small amount and the high dispersity. As the content of NiCoP increases to 20%, obvious diffraction peaks of NiCoP are observed. The diffraction peaks located at 41° and 44.9° could be indexed to (111) and (201) planes of hexagonal NiCoP (JCPDS No. 71-2336), respectively.

The surface chemical composition of the as-fabricated $\text{NiCoP-3/C}_3\text{N}_4$ was probed by XPS. The XPS survey spectrum (Fig. 2a) confirms the existence of Ni, Co, and P. In the C 1s region (Fig. 2b), the peaks located at 284.7, 288.2 and 285.8 eV correspond to the C–C, N–C=N, and C–NH₂ in $\text{g-C}_3\text{N}_4$ [49,50]. The N 1s region of $\text{g-C}_3\text{N}_4$ (Fig. 2c) is deconvoluted into four peaks at 398.6, 399.4, 400.8 and 404.7 eV, attributed to the C–N=C, N–(C)₃, C–N–H and π -excitations of the C=N conjugated systems from $\text{g-C}_3\text{N}_4$, respectively [51,52]. The two peaks of the Ni 2p region (Fig. 2d) located at 853.3 eV (Ni 2p_{3/2}) and 868.5 eV (Ni 2p_{1/2}) imply the presence of partially charged $\text{Ni}^{\delta+}$ ($\delta \approx 0$) in Ni–P compound, and the peaks around 856.4 and 872.7 eV can be ascribed to oxidized Ni^{2+} . The peaks at 862.1 eV (Ni 2p_{3/2}) and 880.0 eV (Ni 2p_{1/2}) are assigned to the satellites [53,54]. In Co 2p region (Fig. 2e), the peak at 778.5 eV is ascribed to Co 2p_{3/2} of partially charged reduced $\text{Co}^{\delta+}$ ($\delta \approx 0$) in the

Figure 2 XPS survey spectrum (a) and C 1s (b), N 1s (c), Ni 2p (d), Co 2p (e) and P 2p (f) spectra of NiCoP-3/C₃N₄.

Co-P compound. Besides, the peaks at 781.5 and 795.4 eV are attributed to Co 2p_{3/2} and Co 2p_{1/2} of the oxidized Co^{2+/3+}, with satellite peaks at 786.3 and 802.3 eV [55,56]. In the P 2p spectrum (Fig. 2f), the peak at 129.7 eV can be ascribed to the reduced P⁰⁻ in the metal phosphides, while the peak at 133.3 eV can be assigned to the phosphorus oxide (denoted as P⁵⁺) due to the exposure to air [57]. Since the XPS signals of Ni 2p and Co 2p in NiCoP-3/C₃N₄ are weak for the low content of NiCoP, the XPS of pristine NiCoP shown in Fig. S2 displays almost the same deconvolution peaks as NiCoP-3/C₃N₄.

The SEM image of NiCoP-3/C₃N₄ is shown in Fig. 3a. The as-prepared NiCoP-3/C₃N₄ aggregates as a highly crumpled structure. The corresponding energy dispersive X-ray spectrometry (EDX) elemental mapping suggests that Ni, Co, and P are evenly spread over the surface of NiCoP-3/C₃N₄ (Fig. S3). The structure and composition of NiCoP-3/C₃N₄ were further characterized by TEM. The TEM image reveals that the bare g-C₃N₄ is composed of irregularly curved layers (Fig. S4). For NiCoP-3/C₃N₄, the tiny nanoparticles are evenly distributed on the g-C₃N₄ nanosheets (Fig. 3b, c). In the synthesis process, the metal ions (Ni²⁺ and Co^{2+/3+}) can be captured by the lone pair electrons of nitrogen in the g-C₃N₄ framework, leading to a uniform distribution [5]. The average particle

size of the NiCoP nanoparticles was calculated as 5 nm (Fig. S5). The HRTEM image (Fig. 3d) of the single NiCoP nanoparticle on g-C₃N₄ clearly presents the (111) plane of NiCoP, with lattice spacing of 0.22 nm [58]. The STEM image and elemental mapping of nanohybrids are demonstrated in Fig. 3e, revealing Ni, Co and P elements are concentrated on NiCoP nanoparticles. The TEM EDX spectrum is given in Fig. S6, suggesting the atomic ratio of Ni:Co:P is around 5:4:5.

The UV-Vis absorption analysis was used to investigate the optical property of g-C₃N₄, NiCoP and NiCoP-*x*/C₃N₄ (Fig. 4a). The colors of the as-prepared NiCoP-*x*/C₃N₄ samples vary from yellow to dark gray (Fig. S7). Besides, NiCoP-*x*/C₃N₄ exhibits remarkably raised absorption in the whole visible spectrum range with increased content of NiCoP, which should be attributed to the inherent absorption of NiCoP nanoparticles. The band gaps of original g-C₃N₄ and NiCoP-3/C₃N₄ calculated from the derived plots of the transformed Kubelka-Munk functions are 2.68 and 2.62 eV, respectively. The almost same value indicates that loading of NiCoP does not cause any obvious change of the intrinsic band gap of the original g-C₃N₄ (Fig. 4b).

In order to further study the band edge positions of NiCoP-3/C₃N₄, Mott-Schottky measurements were car-

Figure 3 (a) SEM, (b, c) TEM, (d) HRTEM images and (e) STEM elemental mapping of NiCoP-3/C₃N₄.

Figure 4 (a) UV-Vis absorption spectrum and (b) the corresponding plots of $(\alpha h\nu)^{1/2}$ versus $h\nu$ of g-C₃N₄ and NiCoP- x /C₃N₄ samples.

Figure 5 Mott-Schottky plots of g-C₃N₄ and NiCoP-3/C₃N₄.

ried out as displayed in Fig. 5. The Mott-Schottky plots of g-C₃N₄ and NiCoP-3/C₃N₄ show positive slopes, corresponding to n-type semiconductors. The flat-band potentials of g-C₃N₄ and NiCoP-3/C₃N₄ can be estimated as -0.76 and -0.84 V vs. the normal hydrogen electrode (NHE, $E_{\text{NHE}} = E_{\text{Ag/AgCl}} + 0.197$ V) [59], which are obtained from the x -axis intercept of the linear region in Mott-Schottky plots. Normally, the flat band potential of n-type semiconductor is more positive about 0.1 or 0.2 V than its conduction band potential [60], so the calculated conduction band bottoms of g-C₃N₄ and NiCoP-3/C₃N₄ from Mott-Schottky plots are determined as -0.96 and -1.04 V vs. NHE, respectively. According to the equation $E_{\text{VB}} = E_{\text{CB}} + E_{\text{g}}$ [61], the E_{VB} of g-C₃N₄ and NiCoP-3/C₃N₄ are calculated as 1.72 and 1.58 V, vs. NHE, respectively. For

comparison, UPS was also employed to measure the valence band of g-C₃N₄ and NiCoP-3/C₃N₄, as given in Fig. S8. The valence band tops of g-C₃N₄ and NiCoP-3/C₃N₄ are determined as 1.84 and 1.68 V, respectively vs. NHE, which are close to the results of Mott-Schottky plots. Obviously, coupling with tiny NiCoP nanoparticles leads to a variation in the band structure of g-C₃N₄. Considering band edge positions of the as-prepared NiCoP-3/C₃N₄, it should be a promising photocatalyst under UV-Vis light.

The light harvesting properties of NiCoP- x /C₃N₄ were investigated for photocatalytic H₂ generation. As shown in Fig. 6, only a trace amount of H₂ could be detected for pristine g-C₃N₄, while NiCoP exhibits no appreciable activity under UV-Vis light irradiation. However, NiCoP

Figure 6 Photocatalytic H₂ generation activities of different samples under UV-Vis light. 100 mg photocatalyst and 10 mL methanol in 90 mL water, $T = 298$ K.

can significantly promote the photocatalytic hydrogen evolution when coupling with g-C₃N₄. The enhancement effect of NiCoP is strongly related to its weight percentage and a volcano-shaped curve can be plotted. The highest photocatalytic H₂ generation rate of $159 \mu\text{mol g}^{-1} \text{h}^{-1}$ is obtained for NiCoP-3/C₃N₄, which is increased by a factor of 2.5 folds compared with CoP-3/C₃N₄ and 35 folds compared with Ni₂P-3/C₃N₄. The AQE is calculated as 4.2% for the optimized NiCoP-3/C₃N₄ photocatalyst under monochromatic 420 nm visible light, which is one of the highest values among the similar TMPs/g-C₃N₄ photocatalytic systems (Table S1). The activity of NiCoP-3/C₃N₄ was compared with the 3 wt% Pt loaded g-C₃N₄ (Pt-3/C₃N₄) (Fig. S9). Although Pt-3/C₃N₄ has a higher photocatalytic activity, NiCoP-3/C₃N₄ has the obvious advantage of low-cost. In addition, the photo-stability of NiCoP-3/C₃N₄ was also evaluated by performing cycling tests. As indicated in Fig. S10, after five cycling tests, the activity had no obvious loss, confirming the sufficient stability of NiCoP-3/C₃N₄ for photocatalytic H₂ evolution.

Under suitable light irradiation, the electrons of g-C₃N₄ are excited from valence band (VB) into conduction band (CB), and holes are left in the VB. The photo-generated charges can migrate to the solid-solution interface of g-C₃N₄, reducing water to hydrogen and oxidizing methanol. However, a rapid recombination of the charges occurs in bare g-C₃N₄ with the absence of cocatalysts. NiCoP is identified as an excellent electron conductor. When coupling with g-C₃N₄, photo-induced electrons in

the CB of g-C₃N₄ will efficiently move to the surface of NiCoP, and react with water to generate hydrogen. At the same time, the holes in the VB of g-C₃N₄ are consumed for oxidizing methanol. The NiCoP nanoparticles can effectively restrain the recombination of photo-induced electron-hole pairs, and thereby facilitate the photocatalytic activity. With the increasing content of NiCoP, the transfer of photo-induced electrons in NiCoP-x/C₃N₄ should be promoted, whereas the light absorbance would be diminished by the light screening effect, and the agglomeration of excess NiCoP also decreases the activity of reaction sites. To balance these two points, NiCoP-3/C₃N₄ was optimized for effective hydrogen evolution (Fig. 7).

PL spectrum is very helpful to explore the electron-hole pairs separation and migration behaviors in semiconductor materials. The emission PL spectra of g-C₃N₄ and NiCoP-x/C₃N₄ were measured at room temperature and shown in Fig. 8. PL signals of all samples are located within 400–600 nm as a result of the surface and internal radiative recombination of photo-induced charges. It is observed that the presence of NiCoP significantly decreases the PL intensity, confirming the role of NiCoP in boosting the charge transfer and suppressing the charge recombination in g-C₃N₄. The high separation efficiency leads to a higher performance of NiCoP-x/C₃N₄. Compared with Ni₂P and CoP cocatalysts, the lower PL intensity indicates NiCoP is more efficient to suppress the charge recombination (Fig. S11).

To reveal the charge-transfer behavior in g-C₃N₄ and NiCoP-3/C₃N₄ photocatalysts, photocurrent responses were finally examined, as shown in Fig. 9a. Pristine g-C₃N₄ shows an obvious current response from potential

Figure 7 Schematic of the separation of photo-generated charge carriers in the photocatalytic H₂ generation for the NiCoP/g-C₃N₄ nanohybrids.

Figure 8 PL spectra of g-C₃N₄ and NiCoP-*x*/C₃N₄.

Figure 9 (a) Photocurrent responses and (b) EIS result of g-C₃N₄ and NiCoP-3/C₃N₄ under ≥ 400 nm light.

of -1.1 V (vs. Ag/AgCl) under light with wavelength ≥ 400 nm. NiCoP-3/C₃N₄ shows a much higher photocurrent density, suggesting the presence of NiCoP significantly enhances the separation and migration efficiency of photo-induced charge carriers. Fig. 9b displays the electrochemical impedance spectroscopy (EIS) of the as-fabricated pristine g-C₃N₄ and NiCoP-3/C₃N₄. The diameter of the semicircle in the Nyquist plots is relative to the charge transfer resistance (R_{ct}). The value of R_{ct} concluded from the EIS of NiCoP-3/C₃N₄ is much smaller than that of pristine g-C₃N₄, reflecting the more rapid electron transfer in NiCoP-3/C₃N₄ photocatalyst. The R_{ct} value of NiCoP-3/C₃N₄ is also much smaller than that of Ni₂P-3/C₃N₄ and CoP-3/C₃N₄ (Fig. S12), supporting the conclusion that NiCoP has a better charge transfer efficiency than Ni₂P and CoP.

CONCLUSIONS

The NiCoP/g-C₃N₄ nanohybrids containing uniformly distributed 5 nm NiCoP nanoparticles coupled with g-C₃N₄ nanosheets were successfully developed *via* an *in-situ* growth strategy. Considering NiCoP as an excellent electron conductor, the photo-induced electrons in g-C₃N₄ may efficiently move to NiCoP, and the recombination of photo-induced charges should be largely suppressed. Accordingly, NiCoP/g-C₃N₄ nanohybrids exhibit greatly enhanced photocatalytic activity than pristine g-C₃N₄. The enhancement effects of NiCoP cocatalyst are relative with NiCoP contents and the optimized NiCoP weight percentage is 3 wt%, leading to a hydrogen production rate of $159 \mu\text{mol g}^{-1} \text{h}^{-1}$ by using UV-Vis light, which is significantly higher than pristine g-C₃N₄, CoP/C₃N₄ and Ni₂P/C₃N₄, respectively. The *in-situ* fabrication of tiny NiCoP nanoparticles coupled with g-C₃N₄ as a highly active photocatalyst in hydrogen production demonstrates an effective strategy for boosting the photocatalytic activity of noble-metal-free cocatalyst.

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Author contributions Ma B fabricated the NiCoP/C₃N₄ nanohybrids and tested the photocatalytic activity; Zhao J fabricated the samples for control experiment; Ge Z did the characterization; Chen Y and Yuan Z designed the experiment and wrote the manuscript.

Conflict of interest The authors declare no conflict of interest.

Supplementary information A listing of supporting data (TGA curves of NiCoP-x/C₃N₄, Ni₂P-3/C₃N₄ and CoP-3/C₃N₄; XPS spectra of Ni 2p, Co 2p and P 2p of pristine NiCoP; SEM elemental mapping of C, N, Ni, Co and P for NiCoP-3/C₃N₄; TEM image of the bare g-C₃N₄; TEM image of NiCoP-3/C₃N₄ and NiCoP particle size distribution on g-C₃N₄; EDX spectrum of the as-prepared NiCoP-3/C₃N₄ in TEM; digital photo of the NiCoP, g-C₃N₄ and NiCoP-x/C₃N₄; UPS of g-C₃N₄ and NiCoP-3/C₃N₄; comparison of photocatalytic H₂ production rate of NiCoP-3/C₃N₄ and Pt-3/C₃N₄ under UV-Vis light; cycling tests of NiCoP-3/C₃N₄ for H₂ evolution; PL spectra of g-C₃N₄, Ni₂P-3/C₃N₄, CoP-3/C₃N₄ and NiCoP-3/C₃N₄; EIS results of Ni₂P-3/C₃N₄, CoP-3/C₃N₄ and NiCoP-3/C₃N₄; comparison of the photocatalytic hydrogen production performance of NiCoP/g-C₃N₄ with the literatures) are available in the online version of the paper.

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负载5 nm磷化钴镍纳米颗粒的石墨相氮化碳高效光催化产氢催化剂

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摘要 本文通过可控醇热反应将5 nm磷化钴镍纳米颗粒负载在石墨相氮化碳上得到一种高效催化产氢催化剂。磷化钴镍是优良电子导体和光催化反应助催化剂。通过原位生长法将微小的磷化钴镍纳米颗粒和氮化碳复合可有效消除磷化钴镍的光遮蔽效应, 同时抑制光生载流子的复合, 提高载流子的迁移率。由此, 磷化钴镍/氮化碳纳米复合物表现出远高于纯氮化碳, 单独的磷化镍或磷化钴/氮化碳复合物的光催化产氢活性。本研究为构筑过渡金属磷化物基光催化剂提供了崭新的思路。