

PANI/NaTaO₃ composite photocatalyst for enhanced hydrogen generation under UV light irradiation

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A PANI/NaTaO₃ composite was successfully synthesized by an oxidative polymerization of aniline monomer in hydrochloric acid solution containing sodium tantalate. NaTaO₃ at a monoclinic structure was produced via hydrothermal method. The photocatalytic activities of the unmodified NaTaO₃ and PANI/NaTaO₃ were evaluated for hydrogen generation from an aqueous HCOOH solution and under UV light irradiation. The results showed that the evolution rate of H₂ increased significantly when NaTaO₃ was modified with PANI. The enhancement of the photocatalytic activity of PANI/NaTaO₃ composite was ascribed to the effective charge transfer and separation between NaTaO₃ and PANI, which reduced their recombination. This indicates that PANI modification of tantalate photocatalysts may open up a new way to prepare highly efficient catalytic materials for H₂ generation.

Keywords: polyaniline, NaTaO₃, composite photocatalysts, hydrogen generation.

INTRODUCTION

In the past few decades, a variety of research studies have been performed to develop heterogeneous photocatalysis for the removal toxic organic and inorganic contaminants from aqueous and gas phase and water splitting into hydrogen and oxygen. A wide group of photocatalytic materials which showed high activity for H₂ evolution is tantalates^{1–10}. In 1998 Kato and Kudo reported that alkali tantalates such as LiTaO₃, NaTaO₃ and KTaO₃ are efficient photocatalysts for water decomposition into hydrogen and oxygen in the UV region¹. Thereafter, different tantalate catalysts (Na₂Ta₂O₆, K₂Ta₂O₆, CaTa₂O₆, SrTa₂O₆, BaTa₂O₆, Rb₄Ta₆O₁₇, Sr₅Ta₄O₁₅, Ba₅Ta₄O₁₅ and K₃Ta₃B₂O₁₂) have been studied in the reaction of photocatalytic hydrogen production^{4–7}. Moreover, for improvement of the photocatalytic activity of tantalates many different strategies such as metal deposition, doping of metal or non-metal ions or mixing with another oxide have been investigated^{8–10}.

Conducting polymer composites have received significant attention due to their wide applications in various fields such as charge storage materials, catalysts, solar cells, immunodiagnostic assays and electrorheological ER fluids¹¹. Here, many composites of polyaniline (PANI), polypyrrole (Ppy), polyphenylene vinylene (PPV) and polythiophene (PTh) with inorganic compounds (LiMnO₂, Pt, Cu, SiO₂, WO₃, Ta₂O₅, MnO₂, TiO₂) have been studied^{11–12}. Moreover, both unmodified conducting polymers and conducting polymer nanocomposites can be used as catalysts in the photocatalytic processes. For example, Kandiel et al.¹³ reported that TiO₂ modified with Pt-polypyrrole and synthesized via “in situ” simultaneous reduction of Pt(IV) and the oxidative polymerization of Ppy monomers showed enhanced photocatalytic H₂ production activity in respect to Pt-loaded TiO₂. Zhang et al.¹⁴ studied photocatalytic H₂ production from water with Na₂S and Na₂SO₃ as sacrificial reagents and under vis-light irradiation in the presence of PANI/PdS-CdS catalysts. It had been stated that the PANI/PdS-CdS showed high photocatalytic activity for H₂ evolution despite the absence of Pt co-catalyst. Furthermore, the

anti-photocorrosion performance of PdS-CdS was also significantly enhanced. Moreover, the PPy/CdS composite materials were investigated to split water under visible-light irradiation¹⁵. The results showed that for PPy/CdS composite the rate of H₂ evolution was 4.4 times higher than that of unmodified CdS.

In the current study, we report the synthesis of a PANI/NaTaO₃ composite photocatalyst via oxidative polymerization of aniline monomer in the presence of NaTaO₃. The photocatalytic reaction was carried out in the presence of formic acid as an electron donor and under UV light irradiation. It has been stated that the as-prepared PANI/NaTaO₃ exhibited higher activity in the reaction of photocatalytic H₂ generation in comparison to the pristine NaTaO₃.

EXPERIMENTAL SECTION

Preparation of NaTaO₃

Sodium tantalate (NaTaO₃) was prepared by hydrothermal method using Ta₂O₅ (purity 99.99%, Aldrich) and NaOH (purity 98%, Sigma-Aldrich) as precursors. In the first step of the NaTaO₃ synthesis, 4 g of Ta₂O₅ was added to 50 cm³ of aqueous solution of 8M NaOH. Then the solution was sonicated at the room temperature for 1 h. Thereafter, the mixture was poured into Teflon lined stainless steel with a capacity of 70 cm³. The hydrothermal reaction was carried out at 200°C for 12 h. The final product was collected by filtration, washed with distilled water and dried at 100°C for 48 h.

Preparation of PANI/NaTaO₃ composite

PANI/NaTaO₃ composite was prepared as follows: 2 g of NaTaO₃ was dispersed in the solution that contains 8 ml H₂O and 2 ml HCl. Then aniline monomer (0.1 ml) was added to the above suspension with constant mixing for 1 h. After that the reactor vessel was transferred to an ice bath and the APS ((NH₄)₂S₂O₈, 8 cm³, 0.01 mol/dm³) was added by dropwise. The obtained mixture was stirred for 2 h. Finally, the sediment was filtrated, washed with water and then dried at 100°C for 48 h.

Experimental procedure and techniques

The morphology of the prepared samples was investigated by high-resolution transmission electron microscopy (HRTEM, Tecnai F30 with a field emission gun operating at 200 kV). The phase composition of the samples was determined using X-ray diffraction analysis (XPERT PRO Philips diffractometer, $\text{CuK}\alpha$ radiation). UV-vis diffuse reflectance spectra were recorded using a UV-vis spectrometer (Jasco, Japan). The FTIR spectra were carried out on a Nicolet 308 spectrometer. Photoluminescence (PL) properties at 270 nm excitation of studied materials were studied by using F-7000 Spectrofluorometer (Hitachi).

Photocatalytic reactions

The photocatalytic activities of the obtained NaTaO_3 and PANI/ NaTaO_3 composite in the reaction of H_2 evolution in the presence of formic acid were investigated. The reactions were carried out under atmospheric pressure in a closed system with inner-irradiation-type reactor. A medium pressure mercury lamp (150 W) was used as a light source. In a typical photocatalytic experiment, 300 mg of catalyst was dispersed in 700 cm^3 of 0.1 M aqueous solution of formic acid. Afterwards, the suspension was mixed with a magnetic stirrer for 60 min, and during this time argon was passed through the reaction mixture to remove the dissolved oxygen. Finally, the solution was irradiated for 120 min without argon purging. The evolved hydrogen was collected and analysed by a gas chromatography (Thermal Conductivity Detector (TCD), Ar as a gas carrier, model: Chrome5).

RESULTS AND DISCUSSION

Figure 1 shows the X-ray diffraction patterns of pure PANI, NaTaO_3 and PANI/ NaTaO_3 composite. XRD pattern of PANI shows a weak and broad diffraction peaks at 2θ value of 20° and 25° , corresponding to the periodicity parallel to the polymer chains of PANI¹⁶. The sample obtained after the hydrothermal reaction of NaOH and Ta_2O_5 is composed of sodium tantalate phase at a monoclinic structure (NaTaO_3 , JCPDS card

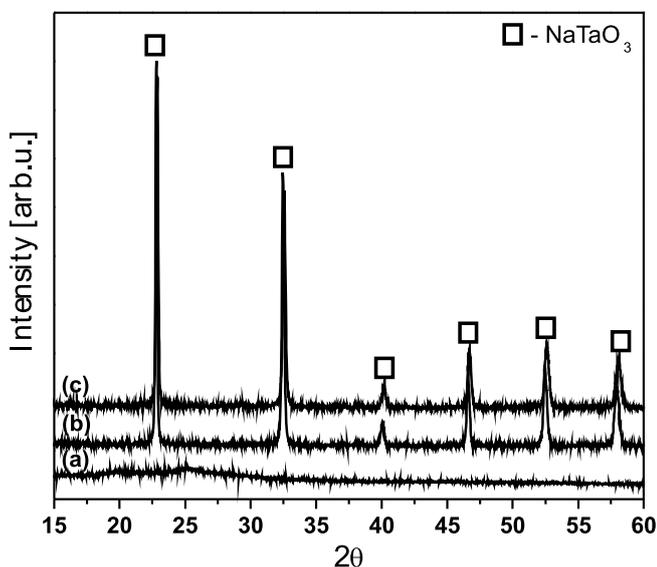


Figure 1. A comparison of X-ray diffraction patterns of samples: (a) PANI, (b) NaTaO_3 and (c) PANI/ NaTaO_3

no 74-2479). For PANI/ NaTaO_3 composite, only the diffraction peaks of NaTaO_3 phase have been detected. The absence of reflections for PANI could be ascribed to the weak crystallinity and small doping content of PANI in composite¹⁶⁻¹⁷.

Figure 2 shows the TEM images of the NaTaO_3 and PANI/ NaTaO_3 composite at different magnifications.

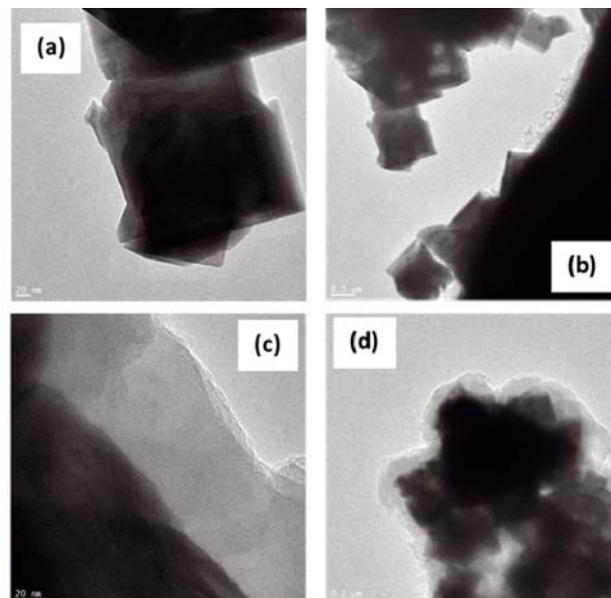


Figure 2. TEM images of samples NaTaO_3 (a–b) and PANI/ NaTaO_3 (c–d)

From TEM images (a–b) it is clearly seen that NaTaO_3 shows the well crystallized cubic morphology. From Figure 2 (c–d), it is clearly seen that PANI/ NaTaO_3 is composed of two phases (c–d) and polyaniline formed a layer on the surface of NaTaO_3 . The thickness of the PANI layer deposited on NaTaO_3 is equal about 80 nm. Moreover, the agglomeration of NaTaO_3 particles in the PANI composite was noted (see Fig. 2, image d).

The FTIR transmission spectra of the produced PANI, NaTaO_3 and PANI/ NaTaO_3 composite are shown in Figure 3. The main characteristic bands of pure PANI are assigned as follows: the peaks at 1573 cm^{-1} and 1498 cm^{-1} are attributed to the C=N and C=C stretch-

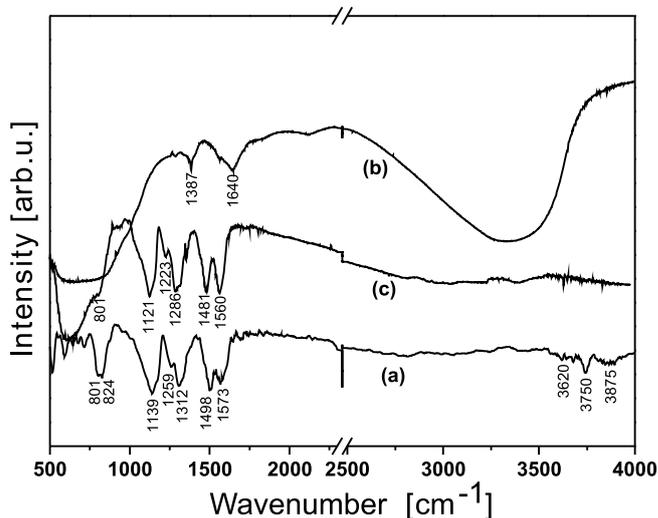


Figure 3. FTIR/DRS spectra of (a) PANI, (b) NaTaO_3 and (c) PANI/ NaTaO_3

ing modes of the quinonoid and benzenoid rings, the bands at 1259 cm^{-1} and 1255 cm^{-1} are assigned to C-N stretching mode for benzenoid units, while the peak at 798 cm^{-1} is associated with C-C and C-H for benzenoid units¹⁷⁻²⁰. Otherwise, the broad and sharp peaks in the range of $3419\text{--}3508\text{ cm}^{-1}$ are attributed to N-H stretching vibration¹⁸. The FTIR spectrum of NaTaO_3 shows bands at 1387 cm^{-1} , 1630 cm^{-1} , a broad band in the region of $2600\text{--}3600\text{ cm}^{-1}$ and a broad band in the range of $500\text{--}1200\text{ cm}^{-1}$. A broad and strong peak in the range of $2600\text{--}3600\text{ cm}^{-1}$ and a peak at 1630 cm^{-1} are assigned to the bending of O-H-O and stretching of O-H modes due to the presence of physically adsorbed water on the surface of the samples²¹. The bands between $500\text{--}1200\text{ cm}^{-1}$ and at 1387 cm^{-1} correspond to the Ta-O and Na-O vibrations, respectively²². The FTIR spectrum of PANI/ NaTaO_3 composite shows seven peaks at about 783 cm^{-1} , 1144 cm^{-1} , 1310 cm^{-1} , 1500 cm^{-1} , 1588 cm^{-1} , 3230 cm^{-1} and 3450 cm^{-1} . All those peaks are characteristic bands of PANI and confirmed the presence of PANI in the synthesized composite. Moreover, for PANI/ NaTaO_3 composite all characteristic peaks of PANI shifted to the lower wavenumber in comparison to the pure PANI. This effect can be explained by the strong interaction between PANI and NaTaO_3 in the composite.

TGA curves of PANI and PANI/ NaTaO_3 composite are shown in Figure 4. For both samples two steps of weight loss are observed. The first weight loss ($25\text{--}120^\circ\text{C}$) of about 12% for PANI and 3% for PANI/ NaTaO_3 is due to the removal of water adsorbed on the surface of the samples. The second significant weight reduction ($330\text{--}600^\circ\text{C}$) is attributed to the polymer backbone decomposition²³. Moreover, according to the data from TG analysis, it was found that the content of PANI in PANI/ NaTaO_3 composite is about 30 wt%.

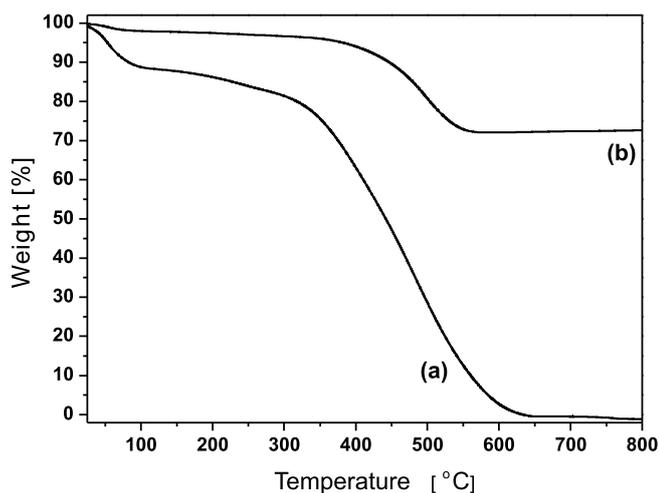


Figure 4. TGA profiles of PANI (a) and PANI/ NaTaO_3 (b) samples

Figure 5 shows a comparison of UV-vis absorbance spectra of PANI, NaTaO_3 and PANI/ NaTaO_3 composite. The spectrum of PANI shows two strong peaks. The peak at about 320 nm is assigned to the $\pi\text{-}\pi^*$ transition of benzene rings and the peak at about 630 nm is attributed to the transition of the quinoid rings in long PANI chains²⁰. For pure NaTaO_3 a single absorption edge at around 300 nm , indicating the band gap energy of 4.11 eV

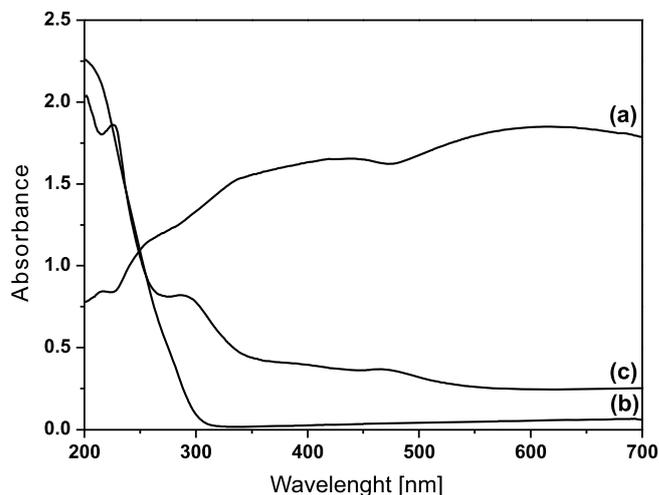


Figure 5. DR-UV-vis spectra of (a) PANI, (b) NaTaO_3 and (c) PANI/ NaTaO_3

can be observed. The estimated band gap of unmodified NaTaO_3 is similar to that described in literature^{24, 25}. Moreover, it can be noted that the characteristic bands of both polyaniline and NaTaO_3 have been observed for PANI/ NaTaO_3 composite. Here, a clear agreement with XRD, TEM and FTIR data is demonstrated.

The photocatalytic H_2 production studies under UV light irradiation were evaluated over unmodified NaTaO_3 and PANI/ NaTaO_3 composite using HCOOH as an electron donor. The control reactions without HCOOH have been also investigated. Here, a trace amount of H_2 was observed in these reaction conditions for both unmodified and modified NaTaO_3 . The results of hydrogen evolution are presented in Figure 6. Here, one can notice that PANI/ NaTaO_3 exhibits higher photocatalytic activity than unmodified NaTaO_3 . It indicates that polyaniline plays a crucial role in the photocatalytic reaction. Also, the reaction of photocatalytic H_2 evolution during the first 120 min of UV light irradiation can be described by zero order kinetics. The rate constants of H_2 evolution are about $87\text{ }\mu\text{mol/h}$ and $163\text{ }\mu\text{mol/h}$ for NaTaO_3 and PANI/ NaTaO_3 , respectively. It is known that the result of a photocatalytic reaction depends mainly on separation, mobility and the lifetime of photogenerated electrons and holes. Here, the photoluminescence (PL) analysis was used to study the efficiency of electron-hole

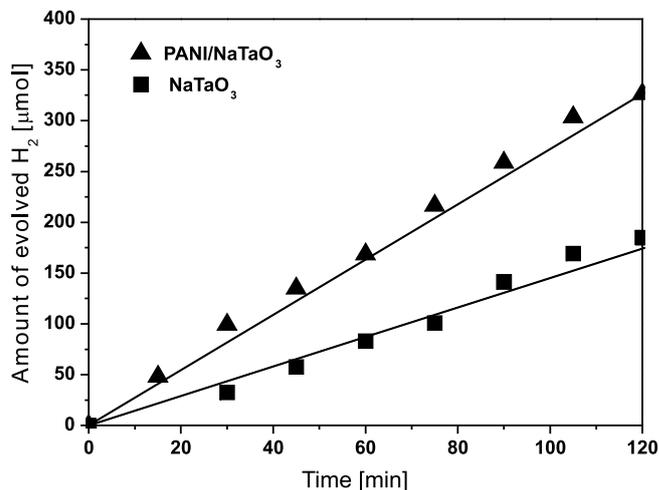


Figure 6. Photocatalytic H_2 evolution over NaTaO_3 and PANI/ NaTaO_3 catalysts

carriers separation and transfer in semiconductors^{26–28} and to understand why the introduction of PANI leads to the enhance the photoactivity of NaTaO₃. Figure 7 presents the PL emission spectra of PANI, NaTaO₃ and PANI/NaTaO₃ composite. PANI shows a broad spectrum with low PL intensity. The PL response for NaTaO₃ exhibits a strong and broad emission band in the range of 370–500 nm with a maximum at about 420 nm^{29–30}. Moreover, it is found that PL spectrum of the PANI/NaTaO₃ composite in the same range of wavelength is much lower than that of pure NaTaO₃. This results indicated that the recombination of photogenerated electrons and holes was effectively inhibited in the PANI/NaTaO₃ composite^{26–30}. It suggests that the higher photocatalytic activity of PANI/NaTaO₃ can be explained by efficient separation of electrons and holes between those two phases, reducing their recombination.

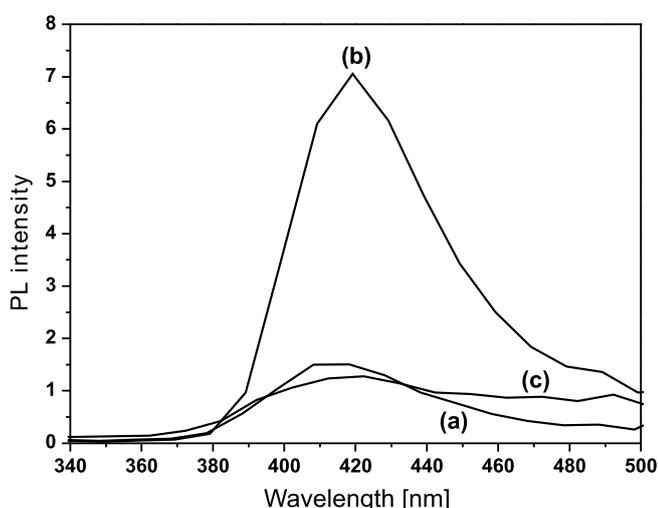


Figure 7. Photoluminescence spectra (a) PANI, (b) NaTaO₃ and (c) PANI/NaTaO₃

CONCLUSION

In summary, PANI/NaTaO₃ composite was successfully prepared via oxidative polymerization of aniline monomer in the presence of NaTaO₃. A detailed analysis (TEM, XRD, FTIR, TGA and DR-UV-vis) indicated that PANI formed the layer on the surface of NaTaO₃. The photocatalytic properties of the synthesized samples were evaluated for the reaction of photocatalytic H₂ production. It was found that, PANI/NaTaO₃ composite exhibited significantly higher photocatalytic activity in respect to the unmodified NaTaO₃. The improvement of photocatalytic performance of PANI/NaTaO₃ composite was attributed to the efficient photogenerated electron-hole transfer and separation.

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