2008

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Recommended Citation

Pen-Cheng Wang, Yaping Dan, Luke A. Somers, Alan G. MacDiarmid, and A.T. Charlie Johnson, "Effect of Thermal Treatments on the Transduction Behaviors of Conductometric Hydrogen Gas Sensors Integrated with HCl-Doped Polyaniline Nanofibers", . January 2008.

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2008. Pen-Cheng Wang, Yaping Dan, Luke A. Somers, Alan G. MacDiarmid, and A.T. Charlie Johnson. Mater. Res. Soc. Symp. Proc. Vol. 1042, Paper #1042-S06-06.
URL: http://www.mrs.org/s_mrs/index.asp

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Abstract

We present the effect of thermal treatments on the transduction behaviors of HCl-doped polyaniline (PANI) nanofibers integrated in conductometric devices upon exposure to 1% H₂ (carried by N₂). After drying in N₂ at 25°C for 12 hours, devices showed a ~10% decrease in electrical resistance upon exposure to 1% H₂. However, devices subject to 12-hour drying in N₂ at 25°C followed by further thermal treatments in N₂ at 100°C, 164°C or 200°C for 30 minutes showed different transduction behaviors. Specifically, devices subject to thermal treatments at 100°C and 164°C showed a decrease in electrical resistance by ~7% and <0.5%, respectively. More interestingly, the device subject to thermal treatment at 200°C showed a transduction behavior with opposite polarity, i.e. a ~5% increase in electrical resistance upon exposure to 1% H₂. SEM, FTIR and TGA were employed to investigate the effect of thermal treatments on the morphology and chemical characteristics of HCl-doped polyaniline nanofibers. The results indicated that the change in the devices' interesting transduction behaviors might be related to the thermal treatment effects on the HCl-doped PANI nanofibers in (i) removal of adsorbed water, and (ii) crosslinking and/or degradation of polymer backbones.

Keywords

Heat treatment, Bacteriophages, Crosslinking, Dewatering, Doping (additives), Electric currents, Electric resistance, Fourier transform infrared spectroscopy, Hydrogen, Hydrogen storage, Nanofibers, Polyaniline, Adsorbed water, Chemical characteristic, Conductometric, Doped polyaniline, Electrical resistances, FTIR, Hydrogen gas sensors, Polymer backbones, SEM, Thermal treatment

Comments

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Effect of Thermal Treatments on the Transduction Behaviors of Conductometric Hydrogen Gas Sensors Integrated with HCl-Doped Polyaniline Nanofibers

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ABSTRACT

We present the effect of thermal treatments on the transduction behaviors of HCl-doped polyaniline (PANI) nanofibers integrated in conductometric devices upon exposure to 1% H₂ (carried by N₂). After drying in N₂ at 25°C for 12 hours, devices showed a ~10% decrease in electrical resistance upon exposure to 1% H₂. However, devices subject to 12-hour drying in N₂ at 25°C followed by further thermal treatments in N₂ at 100°C, 164°C or 200°C for 30 minutes showed different transduction behaviors. Specifically, devices subject to thermal treatments at 100°C and 164°C showed a decrease in electrical resistance by ~7% and <0.5%, respectively. More interestingly, the device subject to the thermal treatment at 200°C showed a transduction behavior with opposite polarity, i.e. a ~5% increase in electrical resistance upon exposure to 1% H₂. SEM, FTIR and TGA were employed to investigate the effect of thermal treatments on the morphology and chemical characteristics of HCl-doped polyaniline nanofibers. The results indicated that the change in the devices' interesting transduction behaviors might be related to the thermal treatment effects on the HCl-doped PANI nanofibers in (i) removal of adsorbed water, and (ii) crosslinking and/or degradation of polymer backbones.

INTRODUCTION

As one of the main themes in the development of hydrogen economy, hydrogen storage has become a very dynamic research area. Conducting polymers have been reported to be promising materials for this application. [1, 2]. However, later studies on hydrogen storage using conducting polymers gave controversial results, showing either much lower H₂ uptake or no H₂ uptake at all [3-5]. It is believed that this controversy could be related to the ambiguous and somewhat inconsistent materials processing procedures used, as the properties of conducting polymers (e.g. oxidation states, doping levels etc) can be affected by many subtle processing variables [6]. Thus, it is of particular interest to further investigate the effects of those processing variables on the interactions between H₂ and conducting polymers.

While hydrogen storage using conducting polymers is under active investigation, conductometric devices integrated with conducting polymers for H₂ sensing have also been demonstrated [7, 8]. The studies on the interactions between H₂ and conducting polymers using conductometric sensors showed that conducting polymers subject to different experimental conditions could result in different transduction behaviors [7, 8], which suggests that these conductometric gas sensors may be used to acquire information for the research on exploiting conducting polymers for hydrogen storage application.

Since the controversy in hydrogen storage using PANI-based materials might be caused by the different thermal treatments on HCl-treated PANI [1, 2, 5], the effect of thermal treatments on the transduction behaviors of HCl-doped PANI nanofibers integrated in conductometric devices upon exposure to 1% H₂ (carried by N₂) at 25°C was investigated in this work. In addition, SEM, TGA and FITR were also used to study the morphology and chemical characteristics of HCl-doped PANI subject to thermal treatments.

EXPERIMENT DETAILS

HCl-doped PANI nanofibers were synthesized by template-free oxidative chemical polymerization in a single-phase acidic aqueous medium [9-11]. Briefly, 10 ml of 0.02 M (NH₄)₂S₂O₈ in 1.0 M HCl_(aq) was quickly poured into a beaker containing 10 ml of 0.02 M aniline in 1.0 M HCl_(aq) under magnetic stirring. The mixed solution in the beaker was magnetically stirred for 5 minutes, after which the magnetic stirring was stopped and the polymerization was allowed to proceed in the beaker for 2 hours. After the 2-hour reaction, the reaction mixture was transferred to a dialysis tubing (Spectra/Por, 12-14,000 MW cutoff). The reaction mixture was dialyzed against 4-L deionized water for 20 hours. The dialysis bath was changed 6 times during the course of the 20-hour dialysis. The dialyzed aqueous dispersion of PANI nanofibers was then transferred to a storage vial for future use.

A home-built, sealed apparatus used to thermally treat HCl-doped PANI nanofiber samples deposited onto glass slides in a controlled atmosphere of flowing nitrogen gas (99.99%) is shown in Figure 1. For computer-controlled conductometric measurements (accuracy of current measurements: ± 100 pA), the same heater chamber was used, with samples of PANI nanofibers deposited onto a glass slide with pre-fabricated, addressable gold electrode lines (~ 2 cm $\times$ 5 μ m; gap between the electrode lines: 500 μ m). By modulating the gas-flow parameters, N₂ or 1% H₂ carried by N₂ was controllably delivered into the sealed gas-flow system to investigate the electrical response of HCl-doped PANI nanofibers subject to various thermal treatments.

Figure 1. Schematic illustration of the home-built apparatus used for performing thermal treatments or conductometric measurements on PANI samples deposited on glass slides (not drawn to scale).

DISCUSSION

Conductometric Measurements

The conductometric device used in this study was composed of two adjacent parallel gold electrode lines bridged by deposited HCl-doped PANI nanofibers (diameter of the sample spot: slightly $>500\text{ }\mu\text{m}$). After the conductometric device was assembled into the experimental set-up (see Figure 1), the device was dried in N_2 at 25°C for 12 hours. The conductometric response of the devices subject to the different thermal treatment conditions and then exposed to several cycles of 1% H_2 is shown in Figure 2. In general, after drying in N_2 at 25°C for 12 hours, devices showed a $\sim 10\%$ decrease (when the device transduction pattern was stabilized; same for the following discussion) in electrical resistance upon exposure to 1% H_2 . However, devices subject to 12-hour drying in N_2 at 25°C and further thermal treatments in N_2 at 100°C , 164°C or 200°C for 30 minutes showed different transduction behaviors. Specifically, devices subject to thermal treatments at 100°C and 164°C showed a decrease in electrical resistance by $\sim 7\%$ and $<0.5\%$, respectively. More interestingly, the device subject to the thermal treatment at 200°C showed a transduction behavior with opposite polarity, i.e. a $\sim 5\%$ increase in electrical resistance upon exposure to 1% H_2 . The above results indicate that the transduction behaviors of these devices can be altered by thermal treatments, and that the transduction behavior is a function of the thermal treatment temperature.

Figure 2. Conductometric response of the device subject to (A) 12-hr drying at 25°C in N_2 with no further thermal treatment, and 12-hr drying at 25°C in N_2 with a further 30-min thermal treatment in N_2 at (B) 100°C , (C) 164°C , or (D) 200°C . For (A), R_0 is the stabilized resistance after 12-hr drying at 25°C . For (B), (C) or (D), R_0 is the stabilized resistance after 12-hr drying at 25°C followed by a 30-min thermal treatment in N_2 at 100°C , 164°C , or 200°C .

SEM

SEM images of the HCl-doped PANI nanofiber samples subject to 12-hour drying in N₂ at 25°C with no further thermal treatment or with a further thermal treatment at 100°C, 164°C or 200°C in N₂ for 30 minutes are shown in Figure 3. Although the general morphology (nanofiber mat) was not significantly changed by the employed thermal treatments, it can be seen from the figure that, upon thermal treatments, inter-fiber fusion occurred in certain areas with higher density of nanofibers.

Figure 3. SEM images of HCl-doped PANI nanofibers subject to (A) 12-hr drying in N₂ at 25°C with no further thermal treatment, and 12-hr drying in N₂ at 25°C followed by a further 30-min thermal treatment in N₂ at (B) 100°C, (C) 164°C or (D) 200°C.

FTIR and TGA

Figure 4 shows the FTIR spectra of the HCl-doped PANI nanofiber samples, each subject to 12-hour drying in N₂ at 25°C with no further thermal treatment, or with a further thermal

treatment at 100°C, 164°C or 200°C in N₂ for 30 minutes. The sample subject to 12-hour drying in N₂ at 25°C with no further thermal treatment gave the FTIR spectrum characteristic of HCl-doped polyaniline (e.g. 1585 cm⁻¹ – stretching of quinoid unit; 1486 cm⁻¹ – stretching of benzenoid unit; 1295 cm⁻¹ – aromatic C-N stretching; 1115 cm⁻¹ – a mode of quinoid unit; 759 cm⁻¹, 698 cm⁻¹ and 544 cm⁻¹ – peaks associated with substituted benzene rings) [12, 13]. The sample subject to 12-hour drying in N₂ at 25°C followed by a 30-minute thermal treatment in N₂ at 100°C still gave the FTIR spectrum with the peaks characteristic of HCl-doped PANI, but the relative intensities of those peaks changed. Our TGA study (data not shown) indicated that a major weight loss occurred when HCl-doped polyaniline was heated from room temperature to 100°C in N₂, which is attributed to the loss of adsorbed water [14]. As the loss of adsorbed water would not significantly affect the chemical structures of polymer backbones, the change in the spectral features might be related to thermal crosslinking of polymer backbones without major degradation [15]. However, HCl-doped PANI samples subject to thermal treatments at 164°C or 200°C in N₂ for 30 minutes gave dramatically different FTIR spectra. The above results indicate that HCl-doped PANI samples subject to the thermal treatments at 164°C or 200°C in N₂ for 30 minutes underwent more dramatic change in their chemical structures and/or composition. The overall FTIR study also indicates that the different transduction behaviors observed in the conductometric measurements involving H₂ and HCl-doped PANI nanofibers subject to various thermal treatments are related to the change in the chemical structure and/or composition of the HCl-doped PANI nanofiber samples.

Figure 4. FTIR spectra of HCl-doped PANI nanofibers subject to (A) 12-hr drying in N₂ at 25°C with no further thermal treatment, and 12-hr drying in N₂ at 25°C followed by a further 30-min thermal treatment in N₂ at (B) 100°C, (C) 164°C or (D) 200°C.

CONCLUSIONS

In this study, we have shown that conductometric devices integrated with HCl-doped PANI nanofibers subject to different thermal treatments can exhibit different transduction

behaviors when exposed to 1% H₂ carried by N₂. FTIR spectra indicate that the chemical structure and/or composition of the HCl-doped PANI samples can be dramatically altered by some of the employed thermal treatments. The study also suggests that the controversy in hydrogen storage studies involving conducting polymers may be related to the ambiguous thermal treatments on HCl-treated PANI samples. In addition to investigating the interactions between H₂ and conducting polymers, the conductometric devices used in this study may be exploited to help systematically screening and identifying the active chemical species based on conducting polymers for hydrogen storage application. Although the interaction mechanisms between H₂ and the HCl-doped polyaniline nanofibers used in our devices were not investigated in the present work, the interesting transduction behaviors presented in this paper indeed indicate that they deserve to be studied in more detail in the future.

ACKNOWLEDGMENTS

The authors wish to acknowledge the supports from DOE (PCW and AGM; grant number: DE-FC36-05GO15075), ARO (PCW and ATCJ; grant number: W911NF-06-1-0462), NIRT (YD; grant number: ECS-0303981) and JSTO-DTRA (LAS).

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