



# A Closer Look at the Electrical Properties of Al<sub>2</sub>O<sub>3</sub>-Embedded Polypyrrole

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(Received: 14 April 2024

Revised: 1 May 2024

Accepted: 18 June 2024)

## KEYWORDS

PPY, Al<sub>2</sub>O<sub>3</sub>,  
TGA, Semi-conductor, Mott's Hopping

## ABSTRACT:

In the present work, a simple chemistry-based polymerization process was utilized to produce clean and Al<sub>2</sub>O<sub>3</sub> nanoparticle-doped polypyrrole-based samples using three different Al<sub>2</sub>O<sub>3</sub> doping dosages. The prepared samples were characterized using a range of techniques. The X-ray diffraction spectra of the produced samples reveal that they are quasi-crystals. The Fourier transformation infrared spectroscopy of these samples reveals the various required fundamental bands, validating the synthesis of the expected samples. Field emission scanning electron microscopy shows the morphology of the produced materials. The thermal properties of the produced samples were analyzed using thermal gravimetric analysis. Furthermore, the electrical properties of the produced samples were investigated at temperatures ranging from 293 to 393 K. The generated samples' conductivity rises with temperature, suggesting that they are semi-conducting.

## Introduction

Insulators make up most polymers in literature. Polymers can behave like semiconducting materials, according to later findings [1]. Polymers have unique optical, electrical, and thermal properties that make them helpful in many industries. Polypyrrole (PPY) and other conducting polymers are popular due to their non-redox doping [2], thermal and environmental stability, high conductivity, and practicality. As a compound, PPY characteristics vary dramatically. These composites are promising for electrochromic displays [3], temperature and current sensor devices, polymeric batteries, and electromagnetic/radio frequency interface shielding in electronics like computer and cell phone casings [4]. Combining PPY with metal oxides (MOs) improves its mechanical properties, conductivity, and processability. Achieving chemical interactions between PPY and MOs makes making PPY-MO composites difficult [5]. Researchers are still developing MO and PPY composites utilizing various ways due to their increased qualities and possible applications. Conducting polymers have been used as shells around inorganic nanoparticles like Al<sub>2</sub>O<sub>3</sub> to generate nanocomposite materials [6]. Xia and Wang made a

PPY/Al<sub>2</sub>O<sub>3</sub> nanocomposite using ultrasonic irradiation. Schnitzler and Zarbin used sol-gel to synthesize Al<sub>2</sub>O<sub>3</sub> nanoparticles and PPY. The most common cathode choices for lithium. PPY packed with carbonaceous fillers [7], magnetic particles, or dielectric particles combines dielectric/magnetic and electrical properties. In the presence of sodium dodecyl benzenesulfonate, Song and colleagues generated PPY/NiO nanoparticles, nanobelts [8], and nanotubes. Despite multiple studies on PPY/Al<sub>2</sub>O<sub>3</sub> composites, most researchers are currently working on their synthesis and characterization [9]. As far as the author knows, no TGA study has investigated dynamical characteristics for PPY/Al<sub>2</sub>O<sub>3</sub> composites. Thus, a comprehensive study using varied Al<sub>2</sub>O<sub>3</sub> concentrations in PPY and methods is needed [10].

## Experimental Details

### Sample Preparation

Pure PPY and PPY/Al<sub>2</sub>O<sub>3</sub> composite samples were synthesized via chemical oxidative polymerization. Aqueous ferric chloride and pyrrole solutions at 1.25 M and 1 M molar concentrations were prepared [11]. After that, the solutions were refrigerated for two hours. After



that, an ice bath kept the monomer solution between 0 and 4 °C. After dropwise adding the oxidant solution to the monomer solution [12], the polymerization reaction commences quickly and lasts 24 hours. After enough time, the solution filters through filter paper [13]. Rinse the yield with 1 M HCl and acetone until the filtrate is colorless [14]. This yield was air-dried before vacuum-drying. The dried sample was ground into powder with a pestle and mortar. The 5, 10, and 15% weight percentage Al<sub>2</sub>O<sub>3</sub>-doped PPY composites were synthesized similarly [15].

### Experimental Techniques

At a modest scanning rate of 2°/min, the Rigaku Miniflex-II diffractometer produced XRD patterns of MgK $\alpha$  at an angle of 2 $\theta$  (10-80°). XRD measurements determined the sample crystal structure [16]. The powder sample was mixed with dry KBr at a weight ratio of 1:20 for FTIR analysis with Shimadzu IR affinity-1 8000 spectrophotometers. Surface microstructure was examined using the EVO 18 scanning electron microscope [17].

### Results and Discussion

#### X-ray Diffraction (XRD) Analysis

PPY and PPY/Al<sub>2</sub>O<sub>3</sub> composite XRD patterns are displayed in Figure 1. Two identical peaks at 20.36 and 25.01 degrees are seen in the PPY and PPY/Al<sub>2</sub>O<sub>3</sub> composite samples' XRD patterns [18], while a soldier is at 16.10 degrees. The peak strength changes significantly when Al<sub>2</sub>O<sub>3</sub> nanoparticles are introduced to PPY symmetry [19], demonstrating a strong interaction between Al<sub>2</sub>O<sub>3</sub> and PPY. These peaks are caused by quinoid and benzenoid rings in perpendicular and parallel orientations [20]. The processed samples' two peaks indicate a quasi-crystal structure [21].

#### Fourier Transform Infrared (FTIR) Analysis

FTIR spectra of pure and Al<sub>2</sub>O<sub>3</sub>-doped PPY composite samples are shown in Figure 2. In the FTIR spectrum, vibrations between 400 and 2000 cm<sup>-1</sup> are apparent. Bands appear at 512, 808, 1160, 1291, 1483, and 1583 cm<sup>-1</sup>. Para-disubstituted aromatic rings and C-H out-of-plane bending vibration cause 512 and 808 cm<sup>-1</sup> bands. An apparent band at 1291 cm<sup>-1</sup> shows C-N stretching vibrations. Curved planes cause C-H vibration at 1160 cm<sup>-1</sup> [22]. The 1460-1600 cm<sup>-1</sup> bands are likely driven

by non-symmetric C6 ring stretching modes. The peak at 2312 cm<sup>-1</sup> is created by aromatic C-H stretching vibrations, while the band at 2950-3300 cm<sup>-1</sup> is due to aromatic amine N-H stretching [23].

Figure 3(a-d) shows typical FESEM images for pure PPY and PPY/Al<sub>2</sub>O<sub>3</sub> composite samples. FESEM images of current samples show particles with the same morphology as clean and Al<sub>2</sub>O<sub>3</sub>-doped PPY samples [24]. The composite samples FESEM images show no Al<sub>2</sub>O<sub>3</sub> nanoparticles, possibly because they've been integrated into the PPY matrix. Here, Al<sub>2</sub>O<sub>3</sub> nanoparticles are the core and PPY the shell [25].

### Conclusion

PPY and Al<sub>2</sub>O<sub>3</sub>-doped PPY composite samples were synthesized via chemical oxidative polymerization. PPY-based pure and composite samples in the emerald salt form have significantly conducting state peaks of quinoid and benzenoid rings in their FTIR spectra. XRD patterns show that present samples are quasi-crystal. This shows that sample stability increases with Al<sub>2</sub>O<sub>3</sub> nanoparticle doping. FESEM images show particles 1–10  $\mu$ m.

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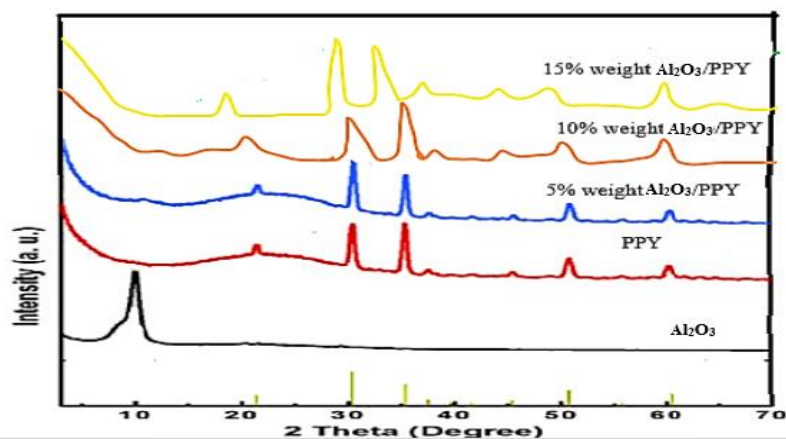


Figure 1: XRD patterns of PPY and PPY/Al<sub>2</sub>O<sub>3</sub> composites

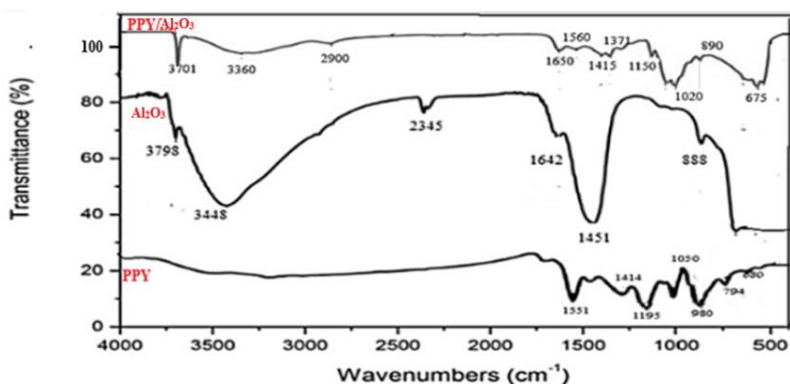


Figure 2: FTIR spectra of PPY and PPY/Al<sub>2</sub>O<sub>3</sub> composites

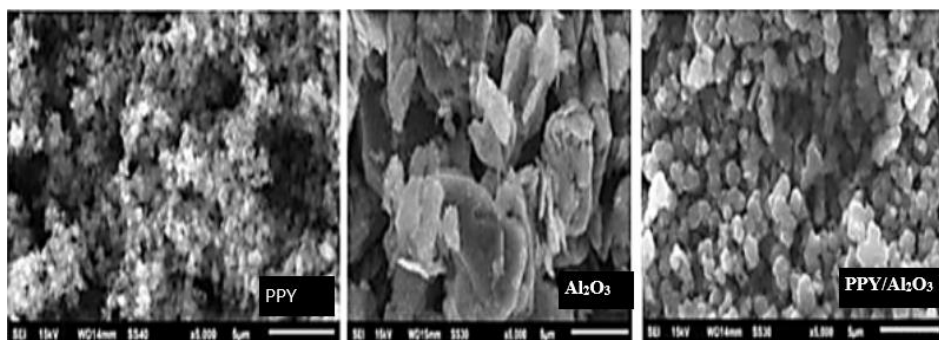


Figure 3: FESEM images of PPY and PPY/Al<sub>2</sub>O<sub>3</sub> composites