

SYNTHESIS AND CHARACTERIZATION OF LiFePO_4 /PPy/CLAY COMPOSITE AS CATHODE MATERIAL FOR Li-ION BATTERY

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ABSTRACT

SYNTHESIS AND CHARACTERIZATION OF LiFePO_4 /PPy/CLAY COMPOSITE AS CATHODE MATERIAL FOR Li-ION BATTERY. Conductor polymers have been used previously as a conducting agent or carbon source in various sol-gel and solid state methods to increase the conductivity of LiFePO_4 . The composite of LiFePO_4 /PPy/Clay were synthesized by mixing LiFePO_4 and clay with a Ppy solution. The characterizations were done by using X-Ray Diffractometer (XRD), Scanning Electron Microscope (SEM) and LCR meter respectively. From XRD analysis it could be deduced that LiFePO_4 has an ordered olivine structure with a Pnma space group. The SEM images provide clear evidence that small crystal particles of LiFePO_4 either coat the surface of the clay particles or lie among them. These LiFePO_4 particles have a small particle size (100–500 nm). With the increasing of LiFePO_4 content, these small crystal particles at the surface or among the clay particles are increased. Increasing the PPy content to increase the conductivity of the material obtained. With the addition of clay from 5 to 10 wt. %, the DC conductivity of the blends is clearly observed to increase up to three orders of magnitude.

Keywords: LiFePO_4 , Clay, Composites, Li-Ion battery

ABSTRAK

SINTESIS DAN KARAKTERISASI KOMPOSIT LiFePO_4 /PPy/CLAY SEBAGAI BAHAN KATODE UNTUK BATTERY Li-ION. Polimer konduktor PPy telah digunakan sebagai bahan penghantar atau sumber karbon dalam berbagai metode sol-gel dan reaksi padatan untuk meningkatkan konduktivitas LiFePO_4 . Komposit LiFePO_4 /PPy/Clay disintesis dengan mencampurkan LiFePO_4 dan tanah liat dengan larutan Ppy. Karakterisasi dilakukan dengan menggunakan X-Ray Diffractometer (XRD), Scanning Electron Microscope (SEM) dan LCR meter. Dari analisis XRD dapat disimpulkan bahwa LiFePO_4 memiliki struktur olivin yang bersesuaian dengan grup ruang Pnma. Gambar SEM memberikan gambaran bahwa partikel kristal dengan ukuran kecil LiFePO_4 melapisi permukaan partikel tanah liat atau berada di antara mereka. Partikel LiFePO_4 ini memiliki ukuran partikel kecil (100-500 nm). Dengan meningkatnya kandungan LiFePO_4 , konduktivitas partikel-partikel kristal kecil di permukaan atau di antara partikel-partikel tanah liat meningkat. Meningkatnya kandungan PPy dapat meningkatkan konduktivitas bahan yang diperoleh. Dengan penambahan tanah liat dari 5 hingga 10 wt. %, konduktivitas DC dari campuran jelas teramati terjadi peningkatan hingga tiga orde.

Kata kunci: LiFePO_4 , Tanah liat, Komposit, Baterai Li-Ion

INTRODUCTION

For their promising potential to be used as power sources for electric vehicles (EVs), so far, Li-ion batteries have captured a large share of the rechargeable battery

market. It is generally recognized that the battery performance strongly depends on the development of electrode materials. Recently, olivine lithium iron

phosphate (LiFePO₄, LFP) has emerged as one of the most promising cathode materials because of its low cost, safety, low toxicity, and high specific capacity of 170 mA h g⁻¹ with a flat discharge–charge potential at 3.45 V vs. Li/Li⁺ owing to the Fe²⁺/Fe³⁺ redox couple. However, one challenging issue regarding LFP cathode is its extremely poor conductivity, which induces lower rate capability. Accordingly, the LFP cathodes cannot satisfy the required power density in Li-ion battery for end- customer needs [1]. However, intrinsic low electrical conductivity and slow Li⁺ ion diffusion directly result in initial capacity loss and poor electrochemical performance, especially at high power applications [2].

Up to the present, many effective approaches have been proposed to solve the above problem. The approaches mainly involve LiFePO₄ coating with fine electronic conductive materials like carbon, reducing particle size, and LiFePO₄ doping with cations. Among the above methods, cations doping at Li-site or Fe-site of LiFePO₄ is very convenient and effective to enhance the electrochemical properties of LiFePO₄. The first-principle calculation on LiMPO₄ (M = Fe, Mn, Co, and Ni) reveals that the lowest Li migration energy for the nonlinear pathway along the [010] channel is curved trajectory between Li sites, and with such one-dimensional pathway, long- range Li conduction may be easily blocked by Li site doping. Therefore, Fe-site doping may be a more suitable strategy to enhance the intrinsic conductivity, and now great efforts have been devoted to investigate the doping effects at Fe-site on the physicochemical and electrochemical properties of LiFePO₄. In addition, an isovalent cation (such as Mg²⁺, Cu²⁺, Co²⁺ and so on) would easily occupy Fe sites of LiFePO₄, it could induce the crystal lattice distortion, provide more space for intercalation/deintercalation of Li, and thus improve the electrochemical properties of LiFePO₄. Although many investigations have been done in order to enhance electrochemical performance of LiFePO₄, it has been rarely reported on isovalent cations co-doping at Fe sites of LiFePO₄ [3].

Concerning the synthetic routes, various methods such as modified solid state reactions, sol gel, microwave synthesis, and hydrothermal reaction have been reported, among which the hydrothermal reaction is recognized as a promising method because of some advantages like simple synthesis process and low energy consumptions compared to high temperature firings and long firing time during the solid state reaction. The hydrothermal synthesis is a method to prepare fine particles, involves dissolution of the reagents in water, precipitation of precursors and conversion of the precursors to target material phase. It is well known that carbon as reducing agent cannot only prevent the formation of Fe³⁺ impurity and the agglomeration of the particles during the preparation of LiFePO₄, but also increases the electronic conductivity [4].

The main limitation with respect to this LiFePO₄ compound is its very poor electronic conductivity which limits the cycling capability. Several methods have been proposed to solve this problem, such as carbon coating or thin metal particle dispersion. But the amount of carbon recover the theoretical capacity (~20 %) and the metal used (copper or silver) are much more expensive than the cathode material itself [5]. Precipitation of precursors and conversion of the precursors to target material phase is well known that carbon as reducing agent cannot only prevent the formation of Fe³⁺ impurity and the agglomeration of the particles during the preparation of LiFePO₄, but also increases the electronic conductivity.

Conducting polymers have been utilized previously as conducting materials or carbon sources in various sol–gel and solid state methods to LiFePO₄. However, those methods coated conducting polymers by post treatment. Pyrrole as a new efficient reducing agent for hydrothermal synthesis of LiFePO₄ powders. We employ the principle of oxidative polymerization of pyrrole to reduce Fe³⁺ to Fe²⁺ and to polymerize pyrrole to polypyrrole (PPy). Therefore, Fe³⁺ oxidized from reactant Fe²⁺ by air can be efficiently reduced back to Fe²⁺ because the hydrothermal solution contains pyrrole. To compare with other hydrothermal syntheses, this is an in situ approach to coat PPy on LiFePO₄ as the hydrothermal synthesis of LiFePO₄, which is more feasible. In addition, the thickness of carbon layer may be controlled by addition of Fe³⁺ as a precursor of LiFePO₄ and an oxidant for the polymerization of pyrrole [6].

Considering simple carbon coating or additive is excluded to obtain perfect rate performance, an electrode consisting of carbon-coated, submicron-sized crystalline LiFePO₄ with 3D porosity is appealing since it can provide fast electronic conduction in the solid phase and ion conduction at reasonable rates in both solid and liquid phases. This has been collaborated by recent work on porous carbon monoliths, showing superior high-rate performance. In view of its facility in terms of synthesis and porous structure, the design is superior to insert submicron sized particles into a porous carbon matrix. Recently, CNTs providing 3D have receive a wide attention due to its most effective in reducing the resistance, and thus improving the electrochemical performance of the composite cathode [7]. Carbon coating does not change its intrinsic conductivity, but it is very effective in enhancing capacity and rate capability. Carbon black and sugar are used as main carbon sources. The electronic conductivity of LiFePO₄/PPy/Clay composite prepared reached 0.1 S/cm [10]. Adding 25% PPy in the starting materials, an improvement in its practical capacity and rate capability was seen. Carbon can control particle growth and enhance electronic conductivity, therefore, higher capacity and better rate capability can be obtained [8].

Clay minerals, a large family of aluminosilicate structures with a range of chemical composition, structure and surface properties, are very reactive materials due to their small grain size, large surface area, adsorption properties and chemical variability. Therefore, they are of considerable recognition in all the physical, chemical and environmental processes in soils and sediments, and frequently control them. The industrial application of clays is enormous and grows with our ability to engineer their properties and use them in the production of composite materials [9]. Zeolites/clays are microporous crystalline hydrated aluminosilicates with a net negative charge due to the presence of aluminum ions in their framework. This negative charge will be balanced by extra framework cations. Clinoptilolite, as the most abundant natural zeolite, has been widely used in research studies and its chemical composition, surface characteristics and ion exchange properties have been reported in many studies [10].

In this research paper, we report on an in-situ synthesis of LiFePO₄/PPy - Clay composite electrode materials aiming to achieve homogenous micro structures and better battery behavior. PPy and Clay will increase the electronic conductivity of LiFePO₄ by means of a different mechanism. The incorporation of LiFePO₄ with zeolites composites in order to achieve three dimensional porous LiFePO₄ architectures also to achieve fast electronic and ion conduction, while keeping an acceptable tap density.

EXPERIMENTAL METHOD

LiFePO₄, LFP, polypyrrole, PPy and clay were purchased from Aldrich with a degree. LFP, PPy and clay were mixed and then heat treated at 70 °C in the oven. The composition of LFP, PPy and clay in the mixtures was varied as follows:

Table 1. The composition of LFP, PPy and clay in the samples

Compound	Code of samples/weight %				
	K1	K2	K3	K4	K5
LFP	90	88.75	82.5	76.25	70
PPy	0	2.5	10	17.5	25
Clay	10	8.75	7.5	6.25	5

Crystalline phases of the samples were identified by XRD at room temperature. Shimadzu XD-610 equipped with a Cu K α source target was utilized to record the XRD pattern of the samples. Diffraction patterns were logged with a goniometer (2 θ) from 10° to 80°. The morphology of the samples was observed by means of SEM on the surface of the samples. JEOL JSM-6510LA equipped with EDS was employed to observe the morphology and elemental analysis of the samples. Electrochemical characterization was carried out for the samples using a HIOKI LCR HiTESTER 3532-50 for EIS.

Impedance spectra were collected in an ambient atmosphere and temperature at an applied voltage of 1 V over a frequency range of 42 Hz – 1 MHz. All the characterization equipments were located at BSBM, PSTBM, BATAN Kawasan Puspipstek Serpong, Tangerang Selatan.

RESULT AND DISCUSSION

Figure 1 represents the XRD patterns of samples, all the peaks are in accordance with the reference LiFePO₄ pattern (PDF No. 40-1499). The peaks which are marked by the appearance of diffraction quite narrow, symmetrical at the 2 θ = 17 °; 20,73 °; 23,14 °; 24,33 °; 26,5 °; 27,45 °; 28,8 °; 29,32 °; 32,11 °; 35,6 °; 36,5 °; 38 °; 42 °; 44 °; 54 ° and 56,93 °. All samples are identified as perfectly single-phase olivine structure indexed to orthorhombic Pnma space group, without any second phase or impurities observed, i.e. the addition of clay does not generate impurities or cause any phase transformation of the samples. LiFePO₄ has an ordered olivine structure with a Pnma space group.

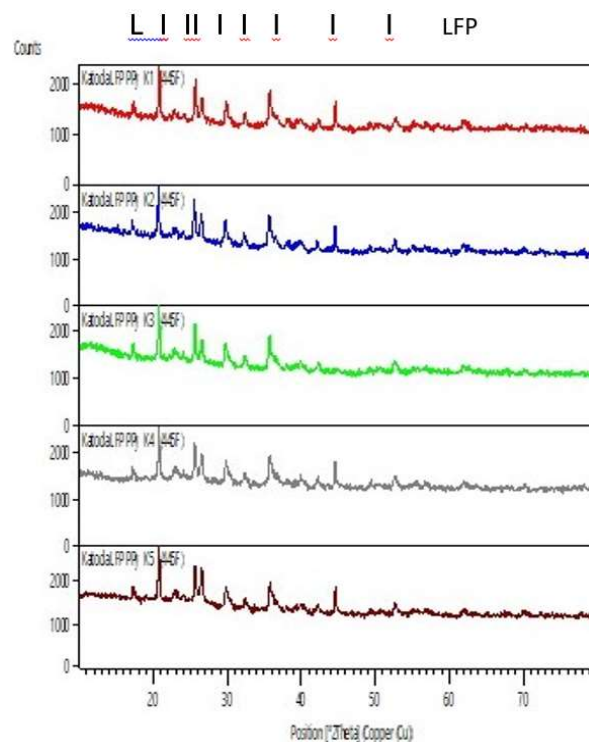


Figure 1. Represents the XRD patterns of samples

The SEM images of composites with different LiFePO₄ contents are shown in Fig. 2. The images provide clear evidence that small crystal particles of LiFePO₄ either coat the surface of the clay particles or lie among them. These LiFePO₄ particles have a small particle size (100–500 nm). The smaller the particles size the better the ionic conductivity will be, because the Li ion can decalate easily in a short pathway, but other variables such as purity and crystallinity must be considered. With

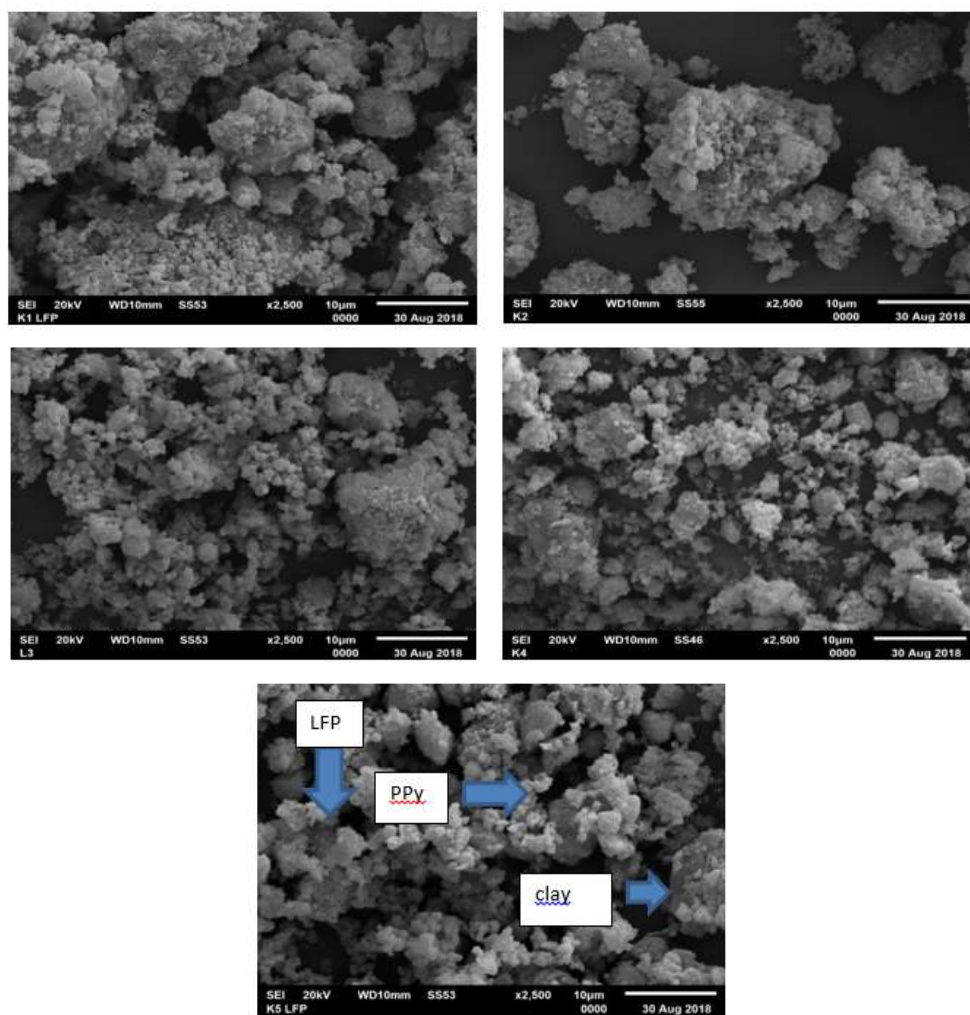


Figure 2. The SEM images of composites with different $\text{LiFePO}_4/\text{PPy}/\text{clay}$ contents.

the increasing of LiFePO_4 content, these small crystal particles at the surface or among the clay particles are increased. Clay providing 3D porosity have most effective in reduced the resistance, and thus improving the electrochemical performance of the composite cathode. An increase in porosity decreases the active surface area, reduces the electrical path into the graphite particles and reduces the accessibility of the lithium ions into the current collector. Due to the agglomeration of the clay during the in-situ synthesis, the surfactant was added to enhance the dispersing of the clay in LFP. Although the pores will accommodate a large volume of electrolyte, they serve as a reaction point during the electrolyte decomposition process [11].

The log-log scale of conductivity measurement on LFP/PPy/Clay is shown in Fig. 3. It can be observed that the conductivity pattern could be divided into two regions, that is, an almost frequency-independent region for low-frequency, and the high-frequency region in which conductivity increases in parallel with the increase in frequency. The linear region of conductivity in the lower frequency characterizes the direct conductivity that is a result of the displacement of charge carriers. At

low to middle frequency the DC phenomenon of this conductivity found of this measurement, meanwhile at frequency of 40 kHz and higher the conductivity followed AC conductivity phenomenon. Increasing PPy content increases the conductivity of the material obtained. With the addition of clay from 5 to 10 wt%, the DC conductivity of the blends is clearly observed to increase up to three orders of magnitude. The incremental increase in conductivity may be ascribed either to the increase in mobile ion concentration or to the increase of free space allowing the ions to move due to the addition of clay. However, DC conductivity is observed to decrease at the clay concentration of 10 wt%. This decrease may be attributed to the dominating role of ion association over free ion formation, which reduces the number of ions for conduction. Consequently, the long-range drift of ions may be one of the sources of ion conduction in the system's [12].

CONCLUSION

The main limitation with respect to this compound is its very poor electronic conductivity which limits the

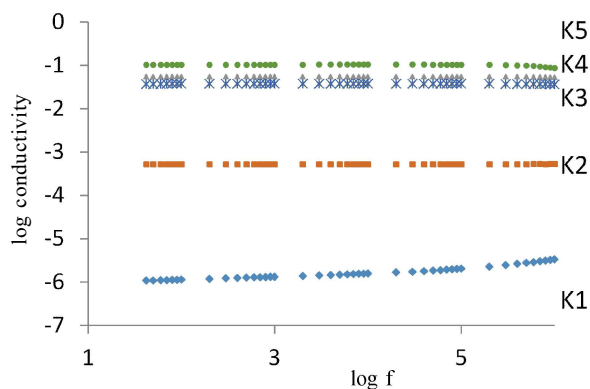


Figure 3. The log-log scale of conductivity measurement on LFP/PPy/Clay.

cycling capability. Several methods have been proposed to solve this problem, such as carbon coating or thin metal particle dispersion. But the amount of carbon recovers the theoretical capacity (~20 %) and the metal used (copper or silver) is much more expensive than the cathode material itself, therefore in this study cheap clay was added to increase the conductivity of LFP.

The XRD patterns of samples, all the peaks are in accordance with the reference LiFePO_4 pattern (PDF No. 40-1499).

The SEM images of composites with different LiFePO_4 contents show the provide clear evidence that small crystal particles of LiFePO_4 either coat the surface of the clay particles or lie among them.

Increasing PPy content increases the conductivity of the material obtained.

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