CHARACTERISTICS OF VANADIUM DOPED AND BAMBOO ACTIVATED CARBON COATED LIFePO₄ AND ITS PERFORMANCE FOR LITHIUM ION BATTERY CATHODE

Nofrijon Sofyan^{1,2}, Subkhan Alfaruq¹, Anne Zulfia^{1,2}, and Achmad Subhan³

¹Department of Metallurgical and Materials Engineering, Faculty of Engineering, Universitas Indonesia, Depok 16424, Indonesia

²Tropical Renewable Energy Center, Faculty of Engineering, Universitas Indonesia, Depok 16424, Indonesia

³Research Center for Physics - LIPI, PUSPITEK, Tangerang, Banten 15310, Indonesia

E-mail: nofrijon.sofyan@ui.ac.id

Received: 21 Maret 2018; revised: 22 Maret 2018; accepted: 11 April 2018

ABSTRACT

CHARACTERISTICS OF VANADIUM DOPED AND BAMBOO ACTIVATED CARBON COATED LIFePO₄ AND ITS PERFORMANCE FOR LITHIUM ION BATTERY CATHODE. Vanadium doped and bamboo activated carbon coated lithium iron phosphate (LiFePO₄) used for lithium ion battery cathode has been successfully prepared. Lithium iron phosphate was prepared through a wet chemical method followed by a hydrothermal process from the starting materials of LiOH, NH₄H₂PO₄, and FeSO₄.7H₂O. The dopant variations of 0 wt.%, 3 wt.%, 5 wt.%, and 7 wt.% of vanadium and a fixed 3 wt.% of bamboo activated carbon were carried out via a solid-state reaction process each by using NH₄VO₃ as a source of vanadium and carbon pyrolyzed from bamboo tree, respectively. The characterization was carried out using X-ray Diffraction (XRD) for the phase formed and its crystal structure, Scanning Electron Microscope (SEM) for the surface morphology, Electrochemical Impedance Spectroscopy (EIS) for the conductivity, and battery analyzer for the performance of lithium ion battery cathode. The XRD results show that the phase formed has an olivine based structure with an orthorhombic space group. Morphology examination revealed that the particle agglomeration decreased with the increasing level of vanadium concentrations. Conductivity test showed that the impedance of solid electrolyte interface decreased with the increase of vanadium concentration indicated by increasing conductivity of 1.25×10^{-5} S/cm, 2.02×10^{-5} S/cm, 4.37×10^{-5} S/cm, and 5.69×10^{-5} S/cm, each for 0 wt.%, 3 wt.%, 5 wt.%, and 7 wt.% vanadium, respectively. Vanadium doping and bamboo activated carbon coating are promising candidate for improving lithium ion battery cathode as the initial charge and discharge capacity at 0.5C for LiFePO₄/C at 7 wt.% vanadium is in the range of 8.0 mAh/g.

Keywords: Activated carbon, Bamboo, LiFePO₄, Lithium ion battery

ABSTRAK

SIFAT-SIFAT LIFEPO4 YANG DIDOPING VANADIUM DAN BERLAPIS KARBON AKTIF DARI BAMBU SERTA KINERJANYA SEBAGAI KATODE BATERAI ION LITIUM. Dalam penelitian ini, bahan aktif litium besi fosfat (LiFePO4) yang didoping vanadium dan berlapiskan karbon yang dipirolisis dari bambu untuk katode baterai ion litium telah berhasil dibuat. Bahan LiFePO4 disiapkan melalui metode kimia basah yang diikuti proses hidrotermal dari bahan awal LiOH, NH₄H₂PO₄, dan FeSO₄.7H₂O. Penambahan variasi 0 %berat, 3 %berat, 5 %berat, dan 7 %berat vanadium dan 3 %berat karbon dilakukan melalui proses reaksi padatan masing-masing menggunakan reagen NH₄VO₃ sebagai sumber vanadium dan arang yang dipirolisis dari bambu. Proses karakterisasi dilakukan menggunakan Difraksi Sinar-X (XRD) untuk melihat fase yang terbentuk dan strukturnya, Mikroskop Elektron (SEM) untuk melihat morfologi permukaan, Spektroskopi Impedansi Elektrokimia (EIS) untuk menguji konduktivitas, dan penguji baterai untuk melihat aktivitas dan kinerja bahan sebagai katode baterai ion litium. Hasil pengujian XRD memperlihatkan bahwa fase yang terbentuk memiliki struktur dasar olivin dengan grup ruang ortorombik. Pengujian morfologi memperlihatkan bahwa aglomerasi partikel semakin berkurang dengan naiknya konsentrasi vanadium. Pengujian konduktivitas memperlihatkan bahwa impedansi antarmuka padat-elektrolit semakin berkurang dengan naiknya konsentrasi vanadium yang dibuktikan naiknya konduktivitas sekitar 1,25 x 10⁻⁵ S/cm; 2,020 x 10⁻⁵ S/cm; 4,371 x 10⁻⁵ S/cm; dan 5,69 x 10⁻⁵ S/cm, masing-masing untuk 0 %berat, 3 %berat, 5 %berat, dan 7 %berat vanadium. Penambahan vanadium dan pelapisan dengan karbon dari hasil pirolisis bambu ini memiliki potensi untuk meningkatkan kinerja katode baterai ion litium dimana kapasitas charge dan discharge awal dari bahan pada 0,5C mampu mencapai 8,0 mAh/g untuk LiFePO₄/C pada 7 %berat vanadium.

Kata kunci : Karbon aktif, Bambu, Litium besi fosfat, Baterai litium

INTRODUCTION

With the cumulative problems of global warming due to the use of non-renewable fossil based fuels, many efforts have been made to replace this non-renewable fossil fuels with other sustainable and green energy sources such as solar, wind, and hydroelectric power (Mathew et al. 2014). Unfortunately, these sources are uncontrollable and intermittent in nature, which then brings a great research interest in material development for energy storage (Y. D. Zhang et al. 2015). Lithium Ion Battery (LIB) is one of the devices developed for this purpose.

The electrochemical performance cathode materials is very important determining the property of LIBs, however, the current cost is still nearly twice as much as the anode that make the research about the developments and optimizations of the cathode active materials have never ended (Y. D. Zhang et al. 2015). LiFePO₄ is the most promising cathode material because of its excellent cyclic stability, non-toxic, environmental friendliness, low cost, and high theoretical capacity of 170 mAh/g at relatively high potential (~ 3.45 V vs. Li/Li⁺) (Qin, Ma, and Wang 2014). However, LiFePO₄ usually shows a poor rate capability because of its low intrinsic electronic conductivity (~10⁻⁹ S/cm) and low diffusion coefficient (Lei et al. 2015).

Carbon coating is the most promising method to improve the electronic conductivity on the surface of LiFePO₄ particles (Li et al. 2017). It has been shown that the presence of activated carbon inside these composite electrodes improves the electronic conductivity and the performance of the materials at high current density of the LiFePO₄ (Böckenfeld et al. 2012). Lithium ion diffusion can be improved by supervalent metal ion doping or substitution at Fe site of LiFePO₄ (Su et al. 2017). Vanadium substitution can significantly improve the of electrochemical performance LiFePO₄ cathode material for lithium ion battery (Chen et al. 2015). This is true because of V doping can create a vacancy and lattice disorder in the crystal structure of LiFePO₄, therefore Li ion can move easier between cathode materials (Chiang et al. 2012).

There are many types of bamboo found in Indonesia, one of which is a clumping bamboo called bambusa balcooa (Bystriakova et al. 2013). Many research have been performed to increase the added value of this bamboo such as using it as cheap carbon source for activated carbon (Tsubota et al. 2018) and biochar for lithium-sulfur battery (Gu et al. 2015). They found that bamboo derived carbon has the potential to be used in many applications due to its simple process and abundant source in nature.

In this work, activated carbon pyrolyzed from bamboo is used as a carbon source applied as active material coating in lithium ion battery cathode. At the same time, the influence of vanadium doping coated by this bamboo activated carbon on the electrochemical behavior of LiFePO₄ cathode material in the forms of composite LiFe_{1-x}V_xPO₄/C, where $0 \le x \ge 0.07$ has been investigated. The results obtained from the investigation are provided and discussed in detail.

MATERIALS AND METHODS

Preparation of Bamboo Activated Carbon

Old bamboo (*Bambusa balcooa*) stems with an average diameter of 10 cm were cut and chopped into small pieces and dried in an oven at 105 °C for 24 hours. The dried bamboo pieces were then heated in a furnace at 400 °C for 4 hours to form bamboo charcoals. The steps in producing this bamboo charcoal can be seen in Figure 1. The bamboo charcoals were then crushed to form charcoal powder in a ball mill and sieved into fine carbon fragments.



Figure 1. The steps in producing bamboo charcoal. Left to right is bamboo tree, pieces of bamboo internodes, chopped and dried bamboo, and bamboo charcoal

The fine carbon fragments were further activated through a chemical activation by soaking the fine bamboo carbon fragments in a closed system using 30% phosphoric acid (H₃PO₄, E-Merck) solution with impregnation ratios of 4 : 1 (w/w) for 24 hours. The sample was then dried in an oven at 105 °C for 20 hours. The dried sample was then physically activated through a heating process at 500 °C before being neutralized using NaOH and successively rinsed with distilled water until the filtrate reached a stable pH of 7. Finally, the moisture was removed by drying the bamboo activated carbon at 105 °C for another 20 hours and ready for characterization.

Preparation of LiFePO₄

LiFePO₄ was firstly synthesized via hydrothermal method using precursors of NH₄H₂PO₄, LiOH, and FeSO₄.7H₂O, all of which analytical grade reagents from E-Merck. The precursors were taken as starting materials with a molar ratio of 1 : 2 : 1. The precursors were dissolved in deionized water and were mixed using a magnetic stirrer. The resulting solution was placed in a Teflon-lined stainless steel autoclave and the hydrothermal reaction was performed at 170 °C for 20 hours. The obtained product was cooled to room temperature and washed with distilled water until the pH reached 7 before being dried at 105 °C to obtain greenish color of LiFePO₄ powder.

Composite LiFe_{1-x}V_xPO₄/C Preparation

Composite LiFe_{1-x}V_xPO₄/C was fabricated through a solid-state reaction by firstly preparing the as-synthesized LiFePO₄ powders from the previous step with variations of 0 wt.%, 3 wt.%, 5 wt.%, and 7 wt.% ammonium metavanadate (NH₄VO₃, Sigma Aldrich) and a fix amount of 3 wt.% bamboo activated carbon. LiFePO₄ powder was mixed with NH₄VO₃ in a ball mill for 20 minutes. The next, bamboo activated carbon was added into the chamber and the milling was continued until a homogeneous mixture was obtained. The ratio of LiFePO₄, activated carbon, and NH_4VO_3 was (97-x) : 3 : x, where x = 0 wt.%, 3 wt.%, 5 wt.%, and 7 wt.%. The mixture was then sintered at 800 °C in a nitrogen atmosphere for 4 hours. Finally, the product was cooled down to room temperature and ready for the characterization.

Battery Fabrication and Testing

The LiFe_{1-x}V_xPO₄/C composite form previous process was mixed with poly(vinylidene fluoride) (PVDF, MTI) in N-methyl pyrrolidone (NMP, MTI) solvent to obtain active material. Active material was coated onto an aluminum sheet (MTI) using doctor blade and heated at 80 °C. Battery fabrication was prepared in the

form of a coin in the glove box on the split test cell using Li metal anode for half-cell using LiPF $_6$ as an electrolyte. The conductivity was tested through an electrochemical impedance spectroscopy, whereas the battery performance was measured using a battery analyzer.

Measurement Apparatus and Condition

X-Ray Diffraction (XRD, ARL OPTX-2050) was carried out using Cu K α radiation (λ = 1.5406 Å) at 40 kV and 35 mA on powder specimen to characterize the structural properties of the materials. Scanning Electron Microscope (SEM, FEI Quanta-650) equipped with energy dispersive X-ray spectroscopy (EDX) was performed at 20 kV to characterize surface morphology and elemental compositions. Electrochemical Impedance Spectroscopy (EIS, Hioki LCR 3532-50) was used to measure the conductivity of the active materials, whereas the battery performance was measured using a battery analyzer (MTI BST8-3 Battery Analyzer). All measurements were performed under ambient condition with no control on humidity and pressure.

RESULTS AND DISCUSSION

Diffraction Patterns

X-ray diffraction was carried out to confirm the phase and crystal structure of the sample and the result is given in Figure 2. As seen in Figure 2, the activated carbon from bamboo has formed indicated by dominant peaks at 2θ 26.61°, 43.45°, 46.32°, 54.81°, and 56.68° corresponding to (111), (100), (110), (222), and (211) orthorhombic crystal planes, respectively. In addition to these peaks, however, there are some other peaks formed at 20 15.13, 23.186, 24.16, and 31.64 corresponding to $(\overline{2}02)$, $(\overline{1}13)$, $(\overline{2}13)$, and (021) monoclinic crystal planes of diphenyl acetylene, respectively. This result shows that pyrolyzing the bamboo at 400 °C and activated it of up to 500 °C has not transformed the cellulose to carbon completely. Nonetheless, quantification of the two phases revealed that more than 10% of the cellulose have been transformed to carbon. Because of that, the result was then used for further testing.

Using Scherrer's formula (Cullity 1978), it can be shown that the crystallite size of the carbon is around 35 nm. This suggests that the crystallinity of the bamboo activated carbon is relatively high. Further, the X-ray diffraction patterns of LiFe_{1-x}V_xPO₄/C composite are given in Figure 3. As seen in Figure 3, all of diffraction patterns of the samples are in agreement with that of LiFePO₄ reference with a structure crystal indexed to olivine structure of Pnma orthorhombic (JCPDS No.083-2092).

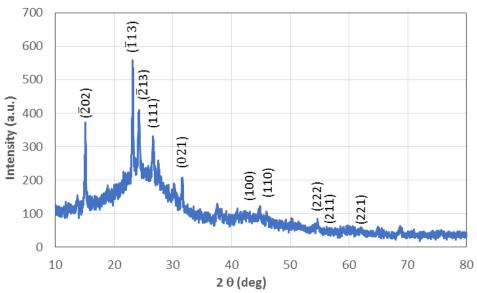


Figure 2. X-ray diffraction pattern of bamboo activated carbon pyrolyzed at 400 °C

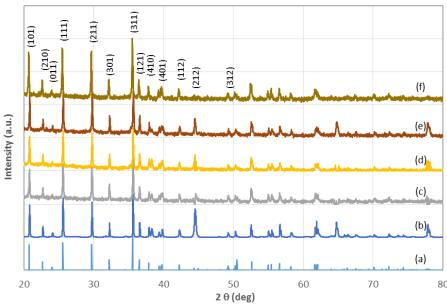


Figure 3. X-ray diffraction pattern of (a) reference, (b) commercial LiFePO₄, (c) as-synthesized LiFePO₄, (d) LiFePO₄ 3 wt.% vanadium, (e) LiFePO₄ 5 wt.% vanadium, and (f) LiFePO₄ 7 wt.% vanadium

No other peak appears for crystallized carbon because of the small amount and the carbon contained is dominated of amorphous phase of acetylene or not enough crystallized phase exists to be detected by XRD (Long et al. 2015). The carbon in this composite does not affect the structure of LiFePO₄. Shifting of the peaks with increase of vanadium concentration, can be observed. This can be however. understood since would be there lattice mismatch due to vanadium insertion into the crystal structure of LiFePO₄. The shifting of the lattice parameters implies that vanadium doping has been successfully occupied the sites of Fe²

in the LiFePO $_4$ structure to form solid solution of LiFe $_{1-x}V_xPO_4$

Morphology and Particle Distribution

SEM characterization was carried out in order to evaluate the effect of vanadium doped on the morphology and particle size distribution of LiFe_{1-x}V_xPO₄. According to the SEM images in Figure 4, it can be observed that LiFePO₄ powders are mainly fine particles between 0.5 μ m and 2 μ m sizes. Some of pure LiFePO₄ particles form agglomeration structure with the hexagonal look shaped particles.

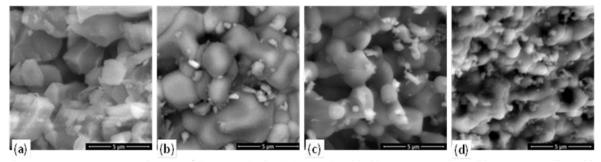


Figure 4. Secondary electron images of the as-synthesized LiFePO₄/C with (a) 0 wt.% vanadium, (b) 3 wt.% vanadium, (c) 5 wt.% vanadium, and (d) 7 wt.% vanadium. Bar scale is 5 μm

The LiFePO₄ with 3 wt.% vanadium shows decrease in agglomeration and particle changes to be more spherical in shape. For the vanadium 5 wt.% and 7 wt.% doped LiFePO₄, the morphology shows spherical shaped particles and less agglomerate. It is worth to note that the particle size gets smaller with the increase of vanadium concentration. This result is agreement with the finding from others (Y. Zhang et al. 2012). This also implies that vanadium doping has a great effect on the structure of LiFePO₄ particles and expected to

be beneficial to the enhancement of LiFePO₄ electronic conductivity (Y. Zhang et al. 2012).

For qualitative purpose, Figure 5 shows Energy Dispersive X-Ray Spectroscopy (EDS) results from the as-prepared sample and the one containing vanadium. As seen in Figure 5a, the sample has the basic elements of C, Fe, P, and O, which indicated the presence of LiFePO₄ compound in the structure coated by carbon, whereas the presence of vanadium shown in Figure 5b, 5c, and 5d reveal that the vanadium is already distributed in LiFePO₄.

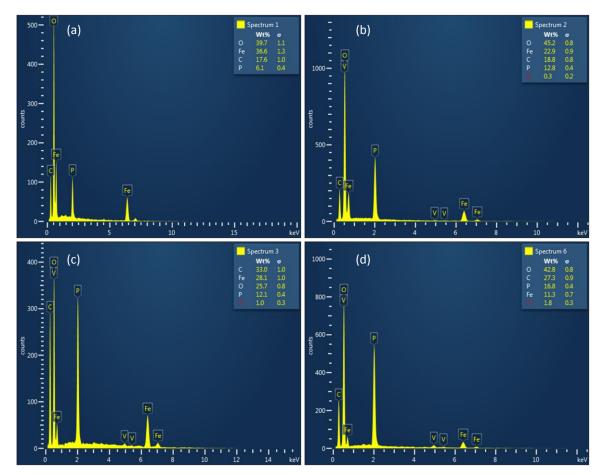


Figure 5. Energy dispersive X-ray spectroscopy of the sample showing the basic elements of C, Fe, P, and O for 0 wt.% vanadium (a) C, Fe, P, O, and V for 3 wt.%, (b) 5 wt.%, (c) 7 wt.%, and (d) vanadium

Conductivity and Battery Performance

Electrochemical Impedance Spectroscopy (EIS) measurements were carried out to explore the effect of vanadium doping and bamboo activated carbon coating on the electrochemical performance of the LiFePO₄ cathode. The results are given in Figure 6 in the forms of Nyquist plot. As seen in Figure 6, the values of the cross point of the semicircle on the horizontal axes represent the charge transfer resistance. The straight line represents the Warburg impedance, which is associated with the lithium ion diffusion in the inner of LiFePO₄ particles. In this instance, charge transfer resistance has a close relationship between composition and conductivity of the interface. Pure LiFePO₄ has very low conductivity (~10⁻⁹ S/cm) (Chung, Bloking, and Chiang 2002). As shown in Figure 6, bamboo activated carbon coated only LiFePO₄ (LFP 0% V) has increased in the conductivity of about 1.25 x 10⁻⁵ S/cm. This indicates that bamboo activated carbon coating can decrease the resistance and increase the conductivity of LiFePO₄. Doping with vanadium even increases the conductivity of LiFePO₄. As the vanadium levels increases, each for 3 wt.%, 5 wt.%, and 7 wt.%, the conductivity increases 2.02 x 10⁻⁵ S/cm, 4.37 x 10⁻⁵ S/cm, and 5.69 x 10⁻⁵ S/cm, respectively. The vanadium doped sample shows superior electrochemical performance than the undoped sample due to higher conductivity and better electrode kinetics related to the synergistic effect of V3+ in the lattice (Pietrzak et al. 2013).

Cyclic voltammetry and charge/discharge measurements were performed for the two active materials only, i.e. LiFePO₄/C at 3 wt.% and 7 wt.% vanadium to represent the active materials with the lowest and the highest conductivity, respectively. The graphs are given in Figure 7. As seen in Figure 7, cyclic voltammetry curves shown at the top have reduction-oxidation peaks at 3.70 volts and 3.10 volts for 3 wt.% vanadium (top left) and at 3.60 volts and 3.20 volts for 7 wt.% vanadium (top right), respectively. The charge/discharge testing was performed at the potential range of 2.5 V until 4.2 V at two capacities, i.e. 0.5C and 4C. The obtained curves are shown at the bottom in Figure 7 for the LiFePO₄/C at 3 wt.% vanadium (bottom left) and 7 wt.% vanadium (bottom right). The initial charge and discharge capacity at 0.5C for LiFePO₄/C at 7 wt.% vanadium (8.0 mAh/g) is higher than that of 3 wt.% vanadium (4.2 mAh/g). At 4C however, the charge and discharge capacity of 7 wt.% vanadium drops to around 4.0 mAh/g, whereas for LiFePO₄/C at 3 wt.% vanadium drops to around 1.0 mAh/g. This can be understood since at high vanadium concentration, at low cycle it can accommodate high capacity. At high cycle, however, the uneven presence of vanadium and carbon and the presence of pores might start forming a bulk layer that blocks and decreases the electrical conductivity and thus the capacity.

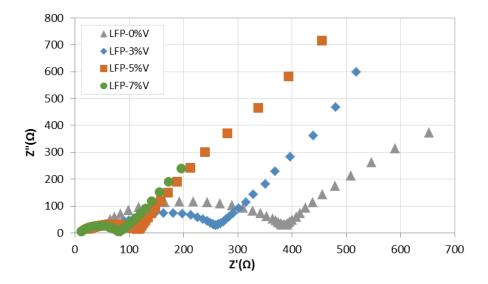


Figure 6. Nyquist plots of LiFePO₄/C (grey triangle), 3 wt.% (blue diamond), 5 wt.% (orange square), and 7 wt.% (green circle) vanadium

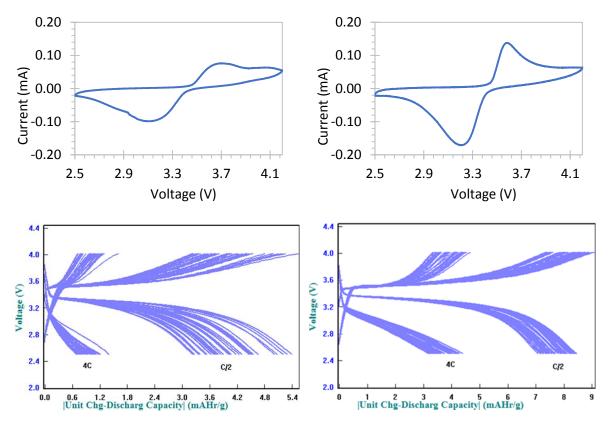


Figure 7. Cyclic voltammetry (top) and charge discharge (bottom) testing results of 3 wt.% (left) and 7 wt.% (right) vanadium

This current result is promising in terms of improving the conductivity of LiFePO₄ for lithium ion battery cathode using cheap natural resources of activated carbon. In comparison to other works, Bhardwaj et al. (2007) have used bamboo carbon for lithium ion battery anode instead of using it to improve the conductivity of LiFePO₄ cathode with a capacity of 130 mAh/g. Gu et al. (2015) have also used bamboo biochar for lithium-sulfur batteries with a capacity of up to 550 mAh/g. Their results can be understood since they used the bamboo carbon for different purposes. Nonetheless, compared to our previous results on the use of activated carbon pyrolyzed from rice husk (Sofyan, Sekaringtyas, et al. 2018) and coconut shells (Sofyan, Rachmawati, et al. 2018) for the same purpose on improving the conductivity of LiFePO₄, the performance of this current result is much better.

CONCLUSION

In this study, activation process of bamboo charcoal was successful using H_3PO_4 reagent and produce a good crystallinity of activated carbon that can increase the conductivity of LiFePO $_4$ from 10^{-9} S/cm to 1.25×10^{-5} S/cm. Doping with vanadium even increases the conductivity of LiFePO $_4$ in which for 3 wt.%, 5 wt.%, and 7 wt.% vanadium, the conductivity increases 1.246×10^{-5} S/cm,

 2.020×10^{-5} S/cm, 4.371×10^{-5} S/cm, and 5.688×10^{-5} S/cm, respectively. At 0.5C, the initial charge and discharge capacity for LiFePO₄/C at 7 wt.% vanadium is 8 mAh/g. This value is higher than that of 3 wt.% vanadium (4.2 mAh/g). At 4C however, the charge and discharge capacity of 7 wt.% vanadium drops to around 4 mAh/g, whereas for LiFePO₄/C at 3 wt.% vanadium drops to around 1 mAh/g.

ACKNOWLEDGMENT

The authors would like to express their gratitude for the funding from the Directorate of Research and Community Services (DRPM), Universitas Indonesia, through Hibah PITTA No. 822/UN2.R3.1/HKP. 05.00/2017.

REFERENCES

Bhardwaj, S., M. Sharon, T. Ishihara, S. Jayabhaye, R. Afre, T. Soga, and M. Sharon. 2007. "Carbon Material from Natural Sources as an Anode in Lithium Secondary Battery." *Carbon Letters* 8 (4): 285–91.

Böckenfeld, N., T. Placke, M. Winter, S. Passerini, and A. Balducci. 2012. "The Influence of Activated Carbon on The Performance of Lithium Iron Phosphate Based Electrodes." *Electrochimica Acta*

- 76: 130–36. doi:10.1016/j.electacta.2012.04.152.
- Bystriakova, N., V. Kapos, I. Lysenko, and C. M. A. Stapleton. 2013. "Distribution and Conservation Status of Forest Bamboo Biodiversity in The Asia-Pacific Region." Biodiversity and Conservation 12: 1833—
- Chen, M., L.-L. Shao, H.-B. Yang, T.-Z. Ren, G. Du, and Z.-Y. Yuan. 2015. "Vanadium-Doping of LiFePO4/Carbon Composite Cathode Materials Synthesized with Organo Phosphorus Source." *Electrochimica Acta* 167: 278–86. doi:10.1016/j.electacta.2015. 03.185.
- Chiang, C., H.-C. Su, P.-J. Wu, H.-J. Liu, C.-W. Hu, N. Sharma, V. K. Peterson, H-W. Hsieh, Y-F. Lin, W. C. Chou, C-H. Lee, J-F. Lee, and B-Y. Shew. 2012. "Vanadium Substitution of LiFePO4 Cathode Materials to Enhance the Capacity of LiFePO4-Based Lithium Ion Batteries." *The Journal of Physical Chemistry C* 116: 24424–29. doi:10.1021/jp307047w.
- Chung, S. Y., J.T. Bloking, and Y. M Chiang. 2002. "Electronically Conductive Phospho-Olivines as Lithium Storage Electrodes." *Nature Materials* 1: 123–28. doi:10.1038/nmat732.
- Cullity, B. D. 1978. *Elements of X-Ray Diffraction*. 2nded. Massachusetts: Addison-Wesley.
- Gu, X., Y. Wang, C. Lai, J. Qiu, S. Li, Y. Hou, W. Martens, N. Mahmood, and S. Zhang. 2015. "Microporous Bamboo Biochar for Lithium-Sulfur Batteries." *Nano Research* 8 (1): 129–39. doi:10.1007/s12274-014-0601-1.
- Lei, X., H. Zhang, Y. Chen, W. Wang, Y. Ye, C. Zheng, P. Deng, and Z. Shi. 2015. "A Three-Dimensional LiFePO4/Carbon Nano-Tubes/Graphene Composite as a Cathode Material for Lithium-Ion Batteries with Superior High-Rate Performance." *Journal of Alloys and Compounds* 626: 280–86.
 - doi:10.1016/j.jallcom.2014.09.169.
- Li, L., L. Wu, F. Wu, S. Song, X. Zhang, C. Fu, D. Yuan, and Y. Xiang. 2017. "Review—Recent Research Progress in Surface Modification of LiFePO4 Cathode Materials." *Journal of The Electro-Chemical Society* 164 (9): A2138–50. doi:10.1149/2.1571709jes.
- Long, Y.-F., J. Su, X.-R. Cui, X.-Y. Lv, and Y.-X. Wen. 2015. "Enhanced Rate Performance of LiFePO4/C by Co-Doping Titanium and Vanadium." *Solid State Science* 48: 104–11.doi:10.1016/j.solidstatesciences.2015.0 7.011.
- Mathew, V., S. Kim, J. Kang, J. Gim, J. Song, J. P. Baboo, W. Park, D. Ahn, J. Han, L. Gu,

- Y. Wang, Y.-S. Hu, Y.-K. Sun, and J. Kim. 2014. "Amorphous Iron Phosphate: Potential Host for Various Charge Carrier Ions." *NPG Asia Materials* 6: e138. doi:10.1038/am.2014.98.
- Pietrzak, T. K., M. Wasiucionek, I. Gorzkowska, J. L. Nowiński, and J. E. Garbarczyk. 2013. "Novel Vanadium-Doped Olivine-Like Nanomaterials with High Electronic Conductivity." *Solid State Ionics* 251: 40–46. doi:10.1016/j.ssi.2013.02.012.
- Qin, G., Q. Ma, and C. Wang. 2014. "A Porous C/LiFePO4/Multiwalled Carbon Nano-Tubes Cathode Material for Lithium Ion Batteries." *Electrochimica Acta* 115: 407–15. doi:10.1016/j.electacta.2013.10.177.
- Sofyan, N., A. D. Rachmawati, A. Zulfia, and A. Subhan. 2018. "Pyrolysis of Activated Carbon from Coconut Shell and Its Characteristic in The LiFePO4/V/C Composite for Lithium Ion Battery Cathode." In IOP Conference Series: Earth and Environmental Science. Vol. 105. doi:10.1088/1755-1315/105/1/012017.
- Sofyan, N., P. Sekaringtyas, A. Zulfia, and A. Subhan. 2018. "Use of Carbon Pyrolyzed LiFePO4/V/C Rice Husk in Composite and Its Performance for Lithium Ion Battery Cathode." In IOP Conference Series: Earth and Environmental Science. Vol. 105. doi:10.1088/1755-1315/105/1/012023.
- Su, W., K. Xu, G. Zhong, Z. Wei, C. Wang, and Y. Meng. 2017. "Enhanced Electrochemical Performance of LiFePO4 as Cathode for Lithium Ion Battery by Codoping with Titanium and Nitrogen." International Journal of Electrochemical Science 12: 6930–39. doi:10.20964/2017.08.13.
- Tsubota, T., K. Ishimoto, S. Kumagai, S. Kamimura, and T. Ohno. 2018. "Cascade Use of Bamboo as Raw Material for Several High Value Products: Production of Xylo-Oligosaccharide and Activated Carbon for EDLC Electrode from Bamboo." Journal of Porous Materials 25: 1–9. doi:10.1007/s10934-018-0567-6.
- Zhang, Y. D., Y. Li, X. H. Xia, X. L. Wang, C. D. Gu, and J. P. Tu. 2015. "High-Energy Cathode Materials for Li-lon Batteries: A Review of Recent Developments." Science China Technology Sciences 58 (11): 1809–28. doi:10.1007/s11431-015-5933-x.
- Zhang, Y., Q.-Y. Huoc, P.-P. Du, L.-Z. Wang, A.-Q. Zhang, Y.-H. Song, Y. Lv, and G.-Y. Li. 2012. "Advances in New Cathode Material LiFePO4 for Lithium-Ion Batteries." *Synthetic Metals* 162 (13): 1315–26. doi:10.1016/j.synthmet.2012.04.025.