

FOURTEENTH ANNUAL CONFERENCE

YUCOMAT 2012

Hunguest Hotel Sun Resort Herceg Novi, Montenegro,
September 3–7, 2012
<http://www.mrs-serbia.org.rs>

Programme and The Book of Abstracts

Organised by:
Materials Research Society of Serbia

under the auspices of
Federation of European Material Societies
and
Materials Research Society

Title: THE FOURTEENTH ANNUAL CONFERENCE
YUCOMAT 2012
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Publisher: Materials Research Society of Serbia
Knez Mihailova 35/IV, 11000 Belgrade, Serbia
Phone: +381 11 2185-437; Fax: + 381 11 2185-263
<http://www.mrs-serbia.org.rs>

Editor: Prof. Dr. Dragan P. Uskoković

Technical editor: Aleksandra Stojičić

Cover page: Aleksandra Stojičić and Milica Ševkušić

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Acknowledgment:



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Society**



Printed in: Biro Konto
Sutorina bb, Igalo – Herceg Novi, Montenegro
Phones: +382-31-670123, 670025, E-mail: bkonto@t-com.me
Circulation: 200 copies. The end of printing: August 2012

P.S.B.7.

SYNTHESIS AND CHARACTERIZATION OF $\text{Li}_2\text{FeSiO}_4/\text{C}$ COMPOSITE

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Polyoxyanion compounds, particularly the olivine phosphate LiFePO_4 , are receiving considerable attention as cathodes for rechargeable lithium batteries. Despite its numerous advantages, olivine phosphate severely suffers from poor rate performance due to its inherent conducting properties and limited capacity. More recently, an entirely new class of polyoxyanion cathodes based on the orthosilicates (Li_2MSiO_4 , $\text{M} = \text{Fe, Mn, and Co}$), has been attracting growing interest. Li_2MSiO_4 has two lithium ions per formula unit, suggesting a higher theoretical capacity than phosphate. Lithium iron orthosilicate, $\text{Li}_2\text{FeSiO}_4$, is very important member of orthosilicates family due to its electrochemical stability, cell safety, eco-friendliness, and cost effectiveness. It is proposed as another promising alternative cathode material for the same lattice stabilization effect as in LiFePO_4 through the presence of strong Si–O bond. The lower electronegativity of Si vs. P would result in a lowering of the $\text{Fe}^{2+} \leftrightarrow \text{Fe}^{3+}$ couple and therefore $\text{Li}_2\text{FeSiO}_4$ often possesses a lower electronic band gap and higher electronic conductivity in comparison with LiFePO_4 . Although $\text{Li}_2\text{FeSiO}_4$ is known for several years, it is still a challenge obtaining a phase pure material with desired particle size and good electrochemical characteristics. Here we report citric acid assisted sol-gel method for $\text{Li}_2\text{FeSiO}_4/\text{C}$ composite synthesis. Starting compounds were LiNO_3 , $\text{Fe}(\text{NO}_3)_3$ and $\text{Si}(\text{OC}_2\text{H}_5)_4$ (tetraethyl orthosilicate, abbrev. TEOS). Citric acid was used as a chelating agent. Sol-gel preparation of $\text{Li}_2\text{FeSiO}_4/\text{C}$ powder was conducted via two routes: (i) one starting from water solutions of above mentioned compounds and (ii) other starting from ethanol solutions of the same compounds. Synthesis in alcohol solution proved to be much faster due to fast hydrolysis of TEOS in presence of alcohol and rapid gel formation. Final product obtained from alcohol solution contains a higher percentage of carbon.