### FIFTEENTH ANNUAL CONFERENCE

# **YUCOMAT 2013**

Hunguest Hotel Sun Resort Herceg Novi, Montenegro, September 2-6, 2013 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 FIFTEENTH ANNUAL CONFERENCE

YUCOMAT 2013

Programme and The Book of Abstracts

**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

Editors: Prof. Dr. Dragan P. Uskoković and Prof. Dr. Velimir Radmilović

Technical editor: Aleksandra Stojičić

Cover page: Aleksandra Stojičić and Milica Ševkušić

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#### **Acknowledgments:**



**Printed in:** Biro Konto

Sutorina bb, Igalo – Herceg Novi, Montenegro

Phones: +382-31-670123, 670025, E-mail: bkonto@t-com.me Circulation: 220 copies. The end of printing: August 2013

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P.S.A.12.

### PREPARATION AND CHARACTERIZATION Pt CATALYST ON Ru DOPED TIN OXIDE SUPPORT FOR OXYGEN REDUCTION

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Platinum nanocatalyst on Ru doped tin oxide support (Ru-SnO<sub>2</sub>) was synthesized and characterized as a catalyst for oxygen reduction reaction in 0.5 mol dm $^{-3}$  HClO<sub>4</sub> solution, at 25°C. Ru doped tin oxide support was synthesized by sol-gel procedure. Synthesized support was characterized by BET (Brunauer, Emmett, Teller), X-ray diffraction, HRTEM (high resolution transmission electron microscopy) and cyclic voltammetry techniques. Specific surface area of the support determined from nitrogen adsorption/desorption isothermal curves was 141 m $^2$  g $^{-1}$ . XRD spectra contained mainly SnO<sub>2</sub> belonging peaks. The corresponding crystallite size for Ru-SnO<sub>2</sub>, determined by Scherrer's equation was 4nm.

Platinum nanocatalyst at Ru-SnO<sub>2</sub> support was synthesized by borohydride reduction method and characterized by X-ray and TEM techniques. Homogenous Pt nanoparticles distribution over the support, without pronounced particle agglomeration was confirmed. The average Pt particle size was 5.3 nm. Electrochemically active surface area of the catalyst was determined from adsorption/desorption charge of hydrogen atoms, after double layer charge correction, taking into account the reference value of 210  $\mu$ C cm<sup>-2</sup> for full monolayer coverage with adsorbed hydrogen species.

The oxygen reduction reaction at Pt/Ru-SnO<sub>2</sub> catalyst was studied by cyclic voltammetry and linear sweep voltammetry at rotating disc electrode. Two different Tafel slope were observed: one close to 60 mV dec<sup>-1</sup> in low current density region, and other close to 120 mV dec<sup>-1</sup> at high current densities region, as it was already reported in literature for oxygen reduction at pure polycrystalline Pt, as well as at Pt nanoparticles in acid solutions. The specific activities, expressed in terms of kinetic current densities per electrochemically active surface area, as well as per mass of Pt loaded, at the constant potential of practical interest, were compared to carbon supported (Vulcan XC-72) catalyst. Pt/Ru-SnO<sub>2</sub> catalyst exhibited similar catalytic activity for oxygen reduction reaction like carbon supported one. The durability of the catalysts was evaluated by repetitive cycling up to 1.4 V vs RHE. Better stability of Pt/Ru-SnO<sub>2</sub> catalyst compared to Pt on carbon support was confirmed by determination of the loss of platinum electrochemically active surface area after potential cycling tests.