## Design of Electrode / Electrolyte Interphase for Rechargeable Magnesium Batteries

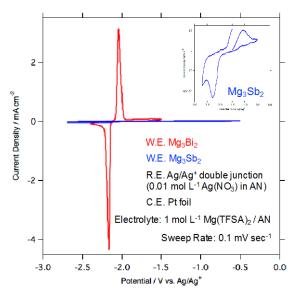
## M. Matsui

Department of Chemical Science and Engineering, Kobe University E-mail: matsui @godzilla.kobe-u.ac.jp

Rechargeable magnesium batteries are expected as potential beyond lithium ion systems, because of their high theoretical capacity densities owing to the divalent magnesium ion. One of the biggest challenges of the magnesium-based negative electrode is passivation of the electrode surface. We have been working on a magnesium-based intermetallic compound Mg<sub>3</sub>Bi<sub>2</sub> Zintl phase as a potential anode active material for the magnesium-ion batteries, because the Mg<sub>3</sub>Bi<sub>2</sub> anode has relatively high volumetric capacity of 1906 mAh cm<sup>-3</sup> and is compatible with conventional ionic electrolyte solutions such as magnesium bis(trifluoromethane

sulfonyl amide) (Mg(TFSA)<sub>2</sub>) dissolved in acetonitrile (AN) solution.

Here we synthesized the Mg<sub>3</sub>Bi<sub>2</sub> and the Mg<sub>3</sub>Sb<sub>2</sub> powders via conventional solid state process. The Mg<sub>3</sub>Bi<sub>2</sub> electrode shows excellent reversibility as reported [1]. On the other hand, the Mg<sub>3</sub>Sb<sub>2</sub> shows poor reversibility as shown in Figure 1. We think the reversibility of the intermetallic anode represents the "SURFACE and BULK INSTABILITY", because the Mg3Bi2 powder shows hydrolysis in ambient atmosphere, while the Mg<sub>3</sub>Sb<sub>2</sub> is extremely stable even with excess H<sub>2</sub>O. We also investigated the compatibility of the Mg<sub>3</sub>Bi<sub>2</sub> with various electrolyte solutions. The Mg<sub>3</sub>Bi<sub>2</sub> shows good reversibility in ether-based and mononitril solvents containing Mg(TFSA)<sub>2</sub>.



**Figure 1:** Electrochemical activity of Mg<sub>3</sub>Bi<sub>2</sub> (red) and Mg<sub>3</sub>Sb<sub>2</sub> (blue and inset) in Mg(TFSA)<sub>2</sub>-AN solution

## References

[1] Timothy Arthur, Masaki Matsui, and Nikhilendra Singh, *Electrochemistry Communications*, **16**, 103-106 (2012)