

**Elucidation of Electrochemical Characteristics of Room-Temperature Li/FeS<sub>2</sub> Batteries**

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Pyrite (Iron disulfide) has been demonstrated to be very useful material for the development of electrodes used in lithium batteries. The abundance, non-toxicity, and high theoretical specific energy of the Li/FeS<sub>2</sub> couple (1273 Wh/kg based on 4e<sup>-</sup>/FeS<sub>2</sub>) make the natural mineral, pyrite, a promising cathode material for lithium batteries [1].

Experimental lithium secondary cells with pyrite electrodes have presented excellent reversible specific energy density at high temperatures (400~450°C) or specific capacity at moderate temperatures (625 mAh/g at 135°C) [2]. However, lithium cells with pyrite electrodes tested at room-temperature have shown inadequate characteristics and poor cycling properties [3-5].

In this work, we have studied electrochemical properties of synthetic and natural pyrite composite electrodes at room temperature. FeS<sub>2</sub> composite electrodes (comprised of FeS<sub>2</sub>, PVdF-HFP and carbon black) were characterized in a lithium cell at room temperature. The electrolyte used was LiN(CF<sub>3</sub>SO<sub>2</sub>)<sub>2</sub> in a solution of tetraglyme and 1,3-dioxolane (1 mol/L). The FeS<sub>2</sub> electrodes were characterized by SEM, XRD and electrochemical methods. For galvanostatic discharge/charge at a current rate of C/10 and between voltage limits of 2.6 and 1.1V, its discharge capacity up to 750mAh/g at the first cycle and about 350mAh/g at the 15<sup>th</sup> cycle are shown in Figure 1. The variation of interfacial resistances with storage time is also shown in Figure 2.

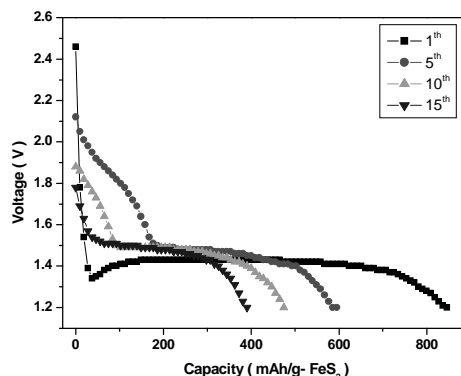


Figure 1. Discharge curves of the Li/FeS<sub>2</sub>.

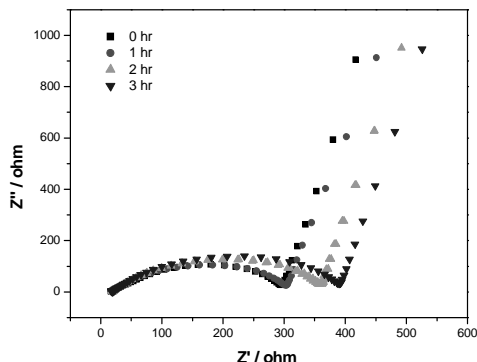


Figure 2. Interfacial resistances of Li/FeS<sub>2</sub> cell vs. storage time at RT.

**References**

- [1] G. Ardel, D. Golodnitsky, K. Freedman, E. Peled, G.B. Appetecchi, P. Romagnoli, B. Scrosati, J. Power Sources, 110 (2002) 152-162.
- [2] Yang Shao-Horn, Steve Osmialowski, Quinn C. Horn, J. Electrochem. Soc., 149 (12) A1547-A1555.
- [3] D. Golodnitsky, E. Peled, Electrochim. Acta, 45 (1999) 335-350.
- [4] E. Strauss, D. Golodnitsky, E. Peled, Electrochim. Acta, 45 (2000) 1519-1525.
- [5] E. Peled, D. Golodnitsky, E. Strauss, J. Lang, Y. Lavi, Electrochim. Acta, 43 (1998) 1593-1599.