

LITHIUM BATTERIES WITH V₂O₅ CATHODIC DEPOLARISANT

G. Taralunga¹, E. M. Rus², D. M. Constantin², Al. Rus²

¹ UNIVERSITY OF AGRICULTURAL SCIENCES AND VETERINARY MEDICINE, 3-5,
Manastur, 3400 Cluj-Napoca

² "BABEŞ-BOLYAI" UNIVERSITY, FACULTY OF CHEMISTRY AND CHEMICAL
ENGINEERING, 11, A. Janos, 3400 Cluj-Napoca, ROUMANIA

Introduction

In recent years, a great attention has been given to the several vanadium oxides which have been proposed as positive electrode materials for use in secondary lithium batteries. Thus, V₂O₅ may be regarded as one of the most promising candidates for cathodic depolarisants, because of its electrochemical behaviour and physical properties, attractive energetic characteristics combined with a good rechargeability [1-4]. In the present work, the lithium intercalation process into V₂O₅ has been studied by cyclic voltammetry and charge-discharge curves.

Experimental

We have prepared V₂O₅ by thermal decomposition of ammonium metavanadate at 500°C in argon atmosphere and characterized by X-ray diffractometry. The electrodes were obtained by pressing V₂O₅ – powder as active material, graphite as electronic conductor and Teflon as binder.

The voltammetrical measurements were done in a three-electrode glass cell at room temperature. The cyclic voltammograms were recorded in 1M LiClO₄ solution in propylene carbonate-dimethyl ethan (PC:DME) mixture (1:1 volumic ratio) as electrolyte, at slow rate (1mV/min).

The electrochemical behaviour of V₂O₅ electrodes has been investigated by charge-discharge curves in galvanostatic regime, at room temperature, using Li/ V₂O₅ button cells. These batteries were discharged at the current density of 1mA/cm², whereas the charge curves were performed at 0.5mA/cm², because the oxidation process unfolds slower [5, 6].

Results and Discussion

The cyclic voltammograms for electrode containing 64.5%V₂O₅, 29% graphite and 6.5% Teflon are showed in figure 1. It can be observed two cathodic peaks (at 3.192 V vs. Li and 3.390V vs. Li), corresponded at lithium ion insertion in crystalline lattice of V₂O₅ and two anodic peaks (at 3.258V vs. Li and 3.446V vs. Li) corresponded to the oxidation of V₂O₅,too. Also, it can be remarked one small anodic peak at 3.194V vs.Li, witch corresponded to the oxidation of intercalation compound Li_x V₂O₅.

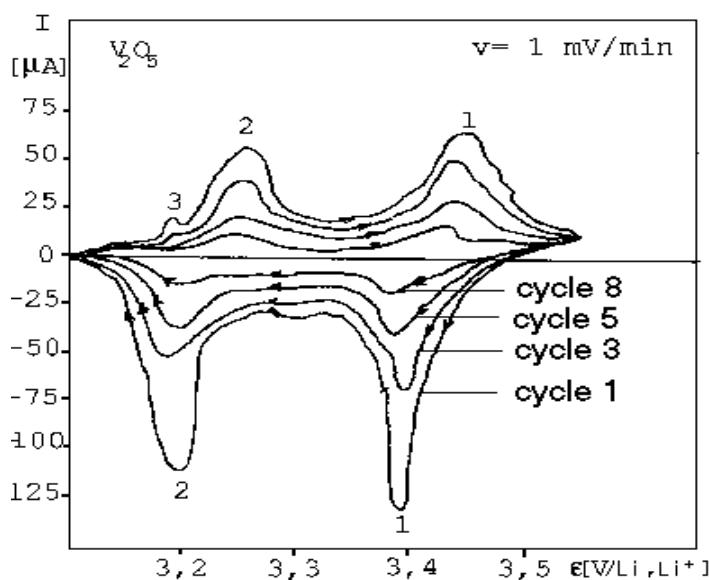


Fig. 1 Cyclic voltammograms of V₂O₅ electrodes.

It was observed one cathodic peak about of 0.75 V vs. Li accordingly to lithium ion insertion in graphite. The cathodic and anodic peaks correspond to the two main plateaus of the charge-discharge curves (see Fig. 2).

From the experimental data, corresponding to the different cycles, discharge capacities C_{disch}, charge capacities C_{ch}, utilization coefficient of active mass and efficiency of batteries have been estimated. The electrochemical characteristics of electrodes with 64.5% V₂O₅, 29% graphite and 6.5% teflon are shown in Table 1.

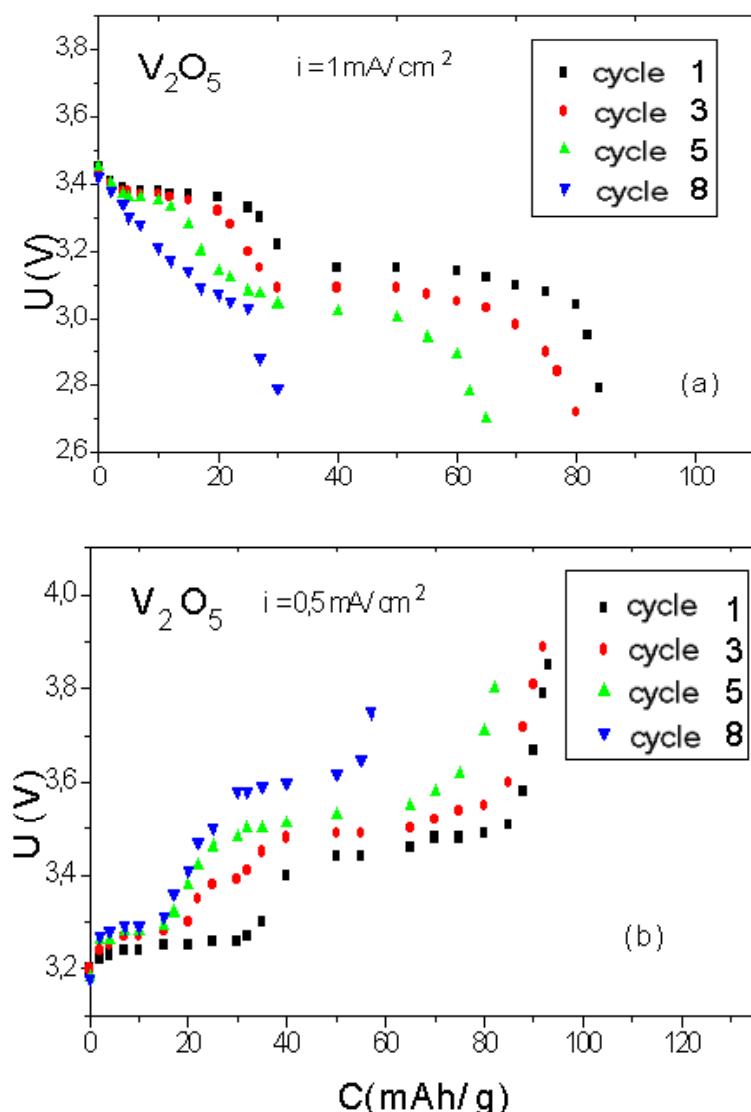


Fig. 2 Performance curves of $\text{Li}/\text{V}_2\text{O}_5$ cell; (a) discharge curves and (b) charge curves.

Table 1 Electrochemical characteristics of V_2O_5

Cycle	C_t [mAh/g]	t_{disch} [h]	C_{disch} [ah/g]	t_{hc} [h]	N_{CH} [ah/g]	F_r [%]	u [%]
1	94.69	20	81.24	45	91.39	88.89	85.57
3	94.69	19	77.18	44	89.37	86.63	81.50
5	94.69	15	60.93	40	81.24	77.88	64.34
8	94.69	8	32.49	30	60.93	53.23	34.31

The reversibility of the electrode processes decreases with the number of charge-discharge cycles, because the oxidation process becomes more difficult in the presence of lithium ions.

Conclusion

From the obtained results it can be establish that V₂O₅ electrodes present good electrochemical characteristics, the intercalation process taking place by diffusion of lithium ions in the crystalline lattice of active mass.

References

1. S. Hub, A. Tranchant, R. Messina, *Electrochim.Acta*, **33**, 997 (1988).
2. K. West, B. Zachau-Christiansen, T. Jacobsen, S. Atlung, *J. Power Sources*, **14**, 235 (1985).
3. B. Garcia, J. Farcy, J.P. Pereira-Ramos, N. Baffier, *J. Electrochem. Soc.*, **144**, 1179 (1997).
4. A. Tranchant, J.M. Blengino, J. Farcy, R. Messina, *J. Electrochem. Soc.*, **139**, 1243 (1992).
5. G. Tarălungă, *Ph. D. Thesis*, "Babeş-Bolyai" University Cluj-Napoca, Romania, 1998.
6. G. Tarălungă, S. Avram, L.D. Bobos, L.Oniciu, *49th Annual Meeting of International Society of Electrochemistry*, Kitakyshu, Japan, 13-18 sept. 1998. Book of Abstracts, p. 654.