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CURRENT GENERATION PROCESSES OF LITHIUM POWER SOURCES WITH CATHODE ON THE BASE OF HYDRATED FORM OF X-RAY AMORPHOUS TITANIUM DIOXIDE

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Current research to improve the electrode materials of lithium power sources (LPS) mainly deal with optimization of their composition, namely purposeful formation of nanoparticle morphology and structure. The perspective base of LPS cathode is electrode material with layered structure that will be stable after lithium insertion processes.

The focus was to explore the current generation processes of lithium power sources based on hydrated form of X-ray titanium dioxide ($TiO_2 \cdot nH_2O$). The five samples of $TiO_2 \cdot nH_2O$, as nanosized particles containing water molecules $n = 2, 1.3, 1.0, 0.9$ and 0.2 , were obtained by controlled $TiCl_4$ hydrochlorid-acid hydrolysis. The electrode materials were studied with testing LPS (electrochemical cell - galvanic pair $Li | 1M$ solution of $LiBF_4$ in γ -butyrolactone | cathode), in which nanoparticles were the base of cathode composition (88 %). To record current generation chemical reactions flow in detail, the cell discharge was carried out in galvanostatic mode at current density $C/250$.

With the help of the method of impedance spectroscopy, it has been shown that the kinetics of lithium ions electrochemical insertion process in the LPS cathode based on $TiO_2 \cdot 2H_2O$ (H_4TiO_4) is characterized mainly by chemisorption of Li^+ to the intercalation degree value $x = 4$, i.e. lithium ions knit electrostatically together with oxygen of each hydroxyl of $Ti(OH)_4$ without formation of lithium titanate phase (Li_4TiO_4). Synthesized $TiO_2 \cdot 1.3H_2O$, titanium oxyhydroxide $TiO(OH)_2$ with layered structure showed the highest absorption capacity at electrochemical insertion of lithium atoms in the material-"master" (specific capacity of LPS into discharge up to $1.5V$ was $2700 mA \cdot h \cdot g^{-1}$). In the first stage of galvanic couples discharge, current generation occurs during Li_2CO_3 formation (data of electron microscopy and the energy X-ray fluorescence spectroscopy) due to the existence of carbonate and carboxil adsorbed grouping on the titanium oxyhydroxide surface followed by Li^+ diffusion in the X-ray structure of $TiO(OH)_2$ and formation of lithium titanate according to reaction: $2Li^+ + H_2TiO_3 \rightarrow Li_2TiO_3 + 2H^+$ (1). The second stage of LPS current generation is carried out $LiOH$ forming, which does not coagulate fully to Li_2O with water formation because of the excess protons presence in current source electrolyte, which freed at

Li_2TiO_3 formation: $LiBF_4 + H_2O \rightarrow LiF + BF_3OH^- + H^+$ (2), $Li^+ + BF_3OH^- \rightarrow LiOH + BF_3$ (3), $BF_3 + LiOH + H^+ \rightarrow LiBF_4 + H_2O$ (4). After exhaustion of excess hydrogen ions, repeated regeneration of electrolyte salt ($LiBF_4$) is terminated and the third stage of current generation takes place due to Li_2O formation: $LiOH + LiOH \rightarrow Li_2O + H_2O$ (5). Chemical reaction 5 immediately and irreversibly occurs when lithium ions are electrochemically inserted in materials $TiO_2 \cdot H_2O$, $TiO_2 \cdot 0.9H_2O$ and $TiO_2 \cdot 0.2H_2O$ with frame structure, providing the value of the LPS specific charge of no more than 900, 800 and $400 mA \cdot h \cdot g^{-1}$.

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EXOMODIFIED FULLERENES AS PROMISING ANTICANCER DRUGS TO THE PHOTODYNAMIC EFFECT

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The creation of a new photosensibilizing anticancer drugs which can be used for photodynamic therapy of tumors occupies the important place in the list of the most current and economically expedient problems of modern Chemistry and Medicine. The photodynamic therapy of tumors is the method of malignancy new growth treatment which in recent years has an active development. That is why, the functionalization of fullerenes allow to get their endo- and exohedral derivatives, which are characterized by a complex of new properties and may be useful in creating such anticancer drugs. Photodynamic activity of exomodified fullerenes in vitro on the cell culture malignant lymphocyte line «Namalva» was determined. The possible application of exomodified fullerenes in the capacity of sensitizers for photodynamic therapy of tumors was shown.

Previously, the mechanism of the inhibitory action of fullerenes was investigated on the example of chain-radical oxidation of organic compounds (benzyl alcohol, hexamethyltriamidophosphate, methyl oleate). It was found that fullerenes inhibit auto-oxidation and initiated oxidation of organic substrates. It was first identify that fullerenes can short-stop the chain of oxidation simultaneously with the participation of peroxy and alkyl radicals.

In 2008 the authors of the project experimentally determined unique and fundamental phenomenon of inversion of antioxidant activity nanoclusters of carbon (derivatives of fullerene

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