

## Electrochemical behavior of hydrogen peroxide at carbosital electrode

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Hydrogen peroxide ( $\text{H}_2\text{O}_2$ ) is neorganic peroxide belonging to the family of oxygenreleasing agents. Hydrogen peroxide and other peroxides are involved in a wide variety of industrial processes. Hydrogen peroxide is used as disinfecting and bleaching agent with applications ranging from the medical, pharmaceutical and paper industry to household washing powder, and they are also attractive reactants for decontamination of chemical warfare agent.

The aim of the research was to investigate the electrochemical behavior of  $\text{H}_2\text{O}_2$  by cathodic voltammetry using carbosital electrode, as indicating electrode. Electrochemical measurements were carried out in the analyzer AVS-1.1 (Volta, St. Petersburg) with a three-electrode scheme by alternating current mode with square wave modulation in potential range  $+1.0 \dots -1.0\text{V}$ ,  $W=1000$  rpm, amplitude  $40\text{mV}$ ,  $\nu=65\text{Hz}$ . CE was used as a working and an auxiliary electrode, and Ag, AgCl /KCl (sat) electrode type EVL-1M4 as a reference electrode.

It was experimentally proved that height of  $\text{H}_2\text{O}_2$  reduction peak decreases and potential of reduction peak is shifted toward more electronegative values with increasing of background electrolyte pH from 1.4 to 4.5. The maximum peak ( $I_p$ ) occurs at a pH approximately 2,2 and at a pH around 4 analytical signal almost disappears. The effect of pH on peak potential ( $E_p$ ) shows the following: when pH value increases in the interval from 2 to 3,  $E_p$  remains almost constant, but  $E_p$  decreases sharply to negative value with pH increasing over 3.5. So, the optimal peak for the analysis ( $E_p=+0.16\text{V}$ ) was obtained at  $\text{pH}\approx 2.2-2.4$  on the background of  $\text{Na}_2\text{SO}_4$  and  $\text{mol L}^{-1}$   $\text{KHSO}_4$ .

The voltammetric study was conducted in solutions with a concentration of  $\text{H}_2\text{O}_2$  from  $1.7 \times 10^{-5}$  to  $10.2 \times 10^{-5} \text{ mol L}^{-1}$ . The peak was obtained at  $E_p= +0.16 \text{ V}$ , whose height was rising proportionally to  $\text{H}_2\text{O}_2$  concentrations increasing. Linear concentration ranges of  $\text{H}_2\text{O}_2$  varied from  $(1.70-10.20) \times 10^{-5} \text{ mol L}^{-1}$ . It is  $I_p = (8,6 \pm 0,3) \times 10^5 \times c$  ( $r = 0.998$ );  $\text{LOD} = 6,16 \cdot 10^{-6} \text{ mol L}^{-1}$ ,  $\text{LOQ}$  is  $2,05 \cdot 10^{-5} \text{ mol L}^{-1}$ . The high sensitivity of this method is accompanied by very good reproducibility. The reproducibility was evaluated from 5 repeated electrochemical signal measurements of model solutions with  $5.10 \times 10^{-5}$ ,  $6.80 \times 10^{-5}$  and  $8.50 \times 10^{-5} \text{ mol L}^{-1}$  concentrations of  $\text{H}_2\text{O}_2$ . The precision of the developed method in terms of the relative standard deviation (RSD) were 4.24 %, 3.27 % and 2.87 %, respectively ( $n = 5$ ,  $P = 0.95$ ).

Thus, the electrochemical behavior of  $\text{H}_2\text{O}_2$  at carbosital electrode was investigated and optimal conditions for its quantitative analysis were established.