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Parameters of electrolysis of zinc sulfate solutions

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Abstract

The purpose of the work was to calculate the exchange currents, the total differential capacity according to the previously developed equations for the electrolysis of sulfate solutions of zinc in the presence of various surfactants on a solid electrode. Electrochemical studies were carried out on a sulfate electrolyte containing 0.005, 0.0125 and 0.025 mol/l ZnSO₄ in a background 0.5 mol/l solution of Na₂SO₄. Some studies were performed using electrolytes of composition: 0.25 mol/l ZnSO₄; 0.25 and 0.75 mol/l ZnSO₄ + 18 and 54 g/l H₂SO₄. In the electrochemical cell flocculants were dosed in the form of an aqueous solution with a concentration of 2.5 g/l in an amount of 25-50 mg/l. The dosage of flocculants corresponded to their consumption in the hydrometallurgical cycle. Lignosulfonate was added in an amount of 80 mg/l. Galvanostatic studies and the removal of polarization curves in a dynamic regime were carried out on the potentiostat *PotentiostatP-30Jcom*. *Elins Electrochemical Instruments»*, using a three-electrode cell. The working electrode (cathode) is made of ZOA zinc with the area of 0.35 cm², the auxiliary (anode) is made of platinum plate with the area of 0.20 cm², the reference electrode is chlorine silver.

The exchange current was calculated from galvanostatic measurements, building the Tafel semi logarithmic dependences of the polarization change on the logarithm of the current density at the initial instant of time. In this case, the calculations were based on a section of an inclined straight line with a correlation coefficient (R^2) greater than 0.99. In the region of low deviations in the potential from the equilibrium calculation of the exchange currents, it was estimated from micro-polarization measurements in the overvoltage region of less than 5÷15 mV. The total polarization capacitance was calculated by the equation, assigning a linearly varying voltage to the electrode at a sweep rate of 100 mV/s and recording the dependence of the current on the electrode potential in the initial fractions of the seconds of the process.

As a result of the work performed, it was shown that additives of flocculants reduce the total polarization capacity in the entire range of zinc content in the background electrolyte of sodium sulfate, in contrast to the additions of lignosulfonate due to the higher molecular weight of the flocculants. At the same time, molecules of organic substances are relatively large in size, and their adsorption leads to an increase in the distance between the capacitor plates in the double layer and thereby increases the polarization capacity.

It is noted that the exchange current and the total polarization capacitance are increases substantially under conditions of intense mixing. The calculations of the currents of exchange by different methods have shown in most cases a high convergence. Using the example of electrolysis of zinc sulfate solutions with lignosulfonate additives, it was shown that the exchange current decreases with increasing surfactant of addition, and with increasing concentration of zinc, the exchange current and total polarization capacity are increases.

The performed investigations was made it possible to obtain new data of the exchange currents and the polarization capacity of zinc electrolysis on a solid electrode in the presence of various surfactants.

References

- [1] A.V. Kolesnikov. Studies of reasons for the effective use of lignosulfonate in the electrolysis of zinc. *Butlerov Communications.* **2014**. Vol.40. No.12. P.110-116. ROI: jbc-02/14-40-12-110
- [2] A.V. Kolesnikov. Investigation of the effect of surfactant in the inversion-voltammetric method analysis of metals. *Butlerov Communications*. 2016. Vol.47. No.7. P.93-96. DOI: 10.37952/ROI-jbc-01/16-47-7-93
- [3] A.V. Kolesnikov. Studies of the discharge of zinc from the background solution of sodium sulfate in the presence of lignosulfonate. *Butlerov Communications*. 2017. Vol.49. No.1. P.128-135. DOI: 10.37952/ROI-jbc-01/17-49-1-128

Kazan. The Republic of Tatarstan. Russia. _____ © Butlerov Communications. 2017. Vol.51. No.8. _____

Full Paper

- [4] A.V. Kolesnikov, K.V. Semenov. Studies of kinetic parameters of the zinc sulfate solution in the presence of an electrochemical system lignosulfonate. Butlerov Communications. 2016. Vol.47. No.7. P.70-73. DOI: 10.37952/ROI-jbc-01/16-47-7-70
- [5] A.V. Kolesnikov. Influence of flocculants on the electroreduction of zinc from sulfate solutions. Bulletin of SSTU "Chemistry and Chemical Technologies". 2014. No.3 (76). P.47-52. (russian)
- [6] A.V. Kolesnikov, K.V. Semenov. Change in the parameters of electrochemical processes in the presence of lignosulfonate. Materials 8 of the international scientific and practical conference "Fundamental and Applied Sciences Today" on May 10-11, 2016. NorthCharleston, USA. Vol.1. P.115-118. (russian)
- [7] A.V. Kolesnikov, K.V. Semenov. Electrolysis of zinc from sulfate acidic and neutral solutions in the presence of lignosulfonate. Actual Problems of the Humanities and Natural Sciences, 2016. Vol.1. No.4 (87). P.57-60. (russian)
- [8] A.V. Kolesnikov. Cathodic and anodic processes in solutions of zinc sulfate in the presence of surfaceactive substances. News of universities "Chemistry and Chemical Technology". 2016. Vol.59. Iss.1. P.53-57. (russian)
- [9] A.V. Kolesnikov. Reduction of copper by metallic zinc in aqueous solutions in the presence of highmolecular surfactants. Condensed Media and Interface Boundaries. 2016. Vol.18. No.1. P.46-54. (russian)
- [10] I.V. Minin, N.D. Solovyova. Kinetics of electrically reducing zinc from sulfate electrolyte in the presence of surfactant additives. Herald of the SSTU. Chemistry and Chemical Technology. 2013. No.1 (69). P.58-62. (russian)
- [11] A.V. Kolesnikov, L.A. Kazanbaev, P.A. Kozlov. Effect of organic substances on the processes of carburization and electrolysis of zinc. Non-ferrous Metals. 2006. No.8. P.24-28. (russian)
- [12] A.I. Biryukov, A.P. Tronov, V.S. Myhametianov. Features corrosion-electrochemical behavior of steels with different chromium content in the highly acidic sulfate media. Butlerov Communications. 2016. Vol.46. No.5. P.61-66. DOI: 10.37952/ROI-jbc-01/16-46-5-61
- A.V. Kolesnikov. The electroreduction are investigated zinc from the background solution of sodium [13] sulfate in the presence of cationic and anionic flocculants. Butlerov Communications. 2017. Vol.49. No.2. P.130-136. DOI: 10.37952/ROI-jbc-01/17-49-2-130
- [14] F.I. Danilov, V.F. Voloshin, M.A. Loshkarev. Effect of adsorption of caproic acid on the kinetics of the discharge of Cd^{2+} . II. Determination of the kinetic parameters of electroreduction of cadmium from sulfate electrolyte. In the Collection. Adsorption and a Double Electric Layer in Electrochemistry. Moscow: Nauka. 1972. P.193-197. (russian)
- [15] P. Etkins. Physical Chemistry. Vol.2. Moscow: Mir. 1980. 584p. (russian)
- [16] A.L. Rotinyan, K.I. Tikhonov, I.A. Shoshina. Theoretical electrochemistry /edited by A.L. Rotinyan. Leningrad: Chemistry. 1981. 422p. (russian)
- [17] V.V. Skorchelletti, Theoretical electrochemistry, Ed. 4th corr. and add. *Leningrad: Chemistry*. 1974. 568p. (russian)