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Determination of Ionic Mobilities of Uranium in Aqueous Solutions at 25 °C by Use of Conductivities

G. Marx and W.-D. Wittke

Institut für Anorganische Chemie, Abteilung Radiochemie, der Freien Universität Berlin

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In pure water the limiting ionic conductance of $(\frac{1}{2} UO_2^{2^+})$ was found to be $57 \, \mathrm{cm}^2 \, \Omega^{-1} \, \mathrm{mol}^{-1}$ by use of conductivity measurements of $UO_2(ClO_4)_2 \cdot xH_2O$ in diluted perchloric acid.

In order to optimize nuclear fuel reprocessing, there still is an urgent need for investigating transport processes of the actinides in aqueous solutions. Since for the standard system water ionic mobilities of uranium had already been determined by us from combined transference- and conductance measurements of $UO_2(NO_3)_2 \cdot 6 H_2O$ in diluted nitric acid ¹, the conductivities of $UO_2(ClO_4)_2 \cdot x H_2O$ in perchloric acid solutions were determined in a concen-

Reprint requests to Prof. Dr.-Ing. G. Marx, Forschungsgruppe Radiochemie, WE 1-Institut für Anorganische Chemie, Freie Universität Berlin, Fabeckstraße 34-36, D-1000 Berlin 33.

tration range of $10^{-4} n \le c \le 6 \cdot 10^{-3} n$ now to confirm those results. The concentration of the UO₂-ion constituent was gained from gravimetric determinations by use of 8-hydroxyquinoline, the $p_{\rm H}$ -value of each solution being calculated with relative high accuracy from these data and the analytical concentration of perchlorate, got from gravimetric determination by nitron ².

Using Mac Innes principle ³ the limiting equivalent conductivity of UO₂(ClO₄)₂ could be calculated for the standard state "water" at 25 °C delivering the value

$$A^0_{(1/2\,{
m UO_{\bullet}}({
m ClO_{\bullet}})_{\bullet})} = 124 \pm 2~{
m cm^2}~\Omega^{-1}~{
m mol^{-1}}$$

and the association constant

$$K_{\rm A} = (9800 \pm 500) \, \rm l^2 \, mol^{-2}$$

the limiting ionic conductance of ${\rm ClO_4}^-$ in water ⁴ being $(67.36\pm0.05)~{\rm cm^2}~\Omega^{-1}~{\rm mol^{-1}}$. The corresponding value of the ${\rm UO_2}^{2^+}$ -ion was found to be

$$\begin{array}{l} \lambda_{(1/2\,\mathrm{UO_2}^{2+})}^0 = (57\pm2)~\mathrm{cm^2}~\varOmega^{-1}~\mathrm{mol^{-1}}~\mathrm{or}\\ u_{(\mathrm{UO_2}^{2+})}^0 = (5.9\pm0.2)\cdot10^{-4}~\mathrm{cm^2}~\mathrm{V^{-1}}~\mathrm{s^{-1}} \end{array}$$

respectively.

These values are in excellent accordance with those gained from our previous investigations ¹.

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² J. Loebich, Angew. Chem. 39, 432 [1926].

³ D. A. Mac Innes, J. Amer. Chem. Soc. 47, 1922 [1925].

⁴ R. A. Robinson and R. H. Stokes, Electrolyte Solutions, Butterworths, London 1959, revised 1970.