AQUEOUS SOLUTIONS AT HIGH PRESSURES AND TEMPERATURES

water was filled into a platinum-iridium cell placed between two opposed tungsten carbide anvils. The water was frozen to -30° C, compressed to the desired pressure and then heated by graphite resistance heaters while the conductance was recorded. *Figure 16* gives isobars of the specific conductance as a function of temperature. The pressures have been calibrated by metal transition points and melting pressure curves of salts and ice. Starting from a low conductance the curves rise while the ice is melting. It is assumed that, after levelling off, that is for example above 500°C for the 98 kb isobar, the curves give the conductance of the fluid water.

The circles showing the earlier shock wave results of Hamann *et al.*⁴² are consistent with the static measurements considering the difficulties and differences of the two methods. This means that water at 1000° C and pressures between 100 and 120 kb has a specific conductance comparable to to the conductance of a concentrated aqueous salt solution at 25°C.

The observed large increase of conductance is mainly due to an increase of the 'ionization product', that is of the product of the activities of hydrogen ions and hydroxyl ions in mole/l. Within the accuracy of this discussion the activities even at these extreme conditions can be relatively well estimated^{45,42} or derived from measurements with shocked salt solutions⁴⁶. The sum of the ion mobilities for hydrogen ions and hydroxyl ions at 670°C



Figure 17. The ionization product K_w of water as a function of density according to Hamann and Linton. • denotes shock wave results of Hamann and Linton, \times static measurements of Holzapfel and Franck⁴⁴.

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and 114 kb for example should be $2090 \pm 500 \text{ cm}^2 \text{ ohm}^{-1} \text{ mole}^{-1}$. Figure 17, which is from Hamann and Linton⁴⁶, gives the ionization product of water derived from static and shock wave experiments as a function of density for several temperatures. At 1000°C and densities between 1.5 and 1.7 g/cm³ the ionization product reaches values of 10^{-2} to $10^{-1} \text{ mole}^2 \text{ 1}^{-2}$. This increase of the product by more than twelve orders of magnitude over the value for standard conditions is not unreasonable if one assumes a constant energy of dissociation and an average reaction volume change for the pressure range between 7 and 10 cm³/mole. It has been suggested that water may become an ionic fluid if compressed to densities higher than about 1.8 g/cm³ at high supercritical temperatures^{46, 47}. This plausible suggestion means that water at these conditions would behave similarly to fused sodium hydroxide. It is indicated in the upper right corner of Figure 17.

Recently conductance measurement with pure fluid ammonia have been made to 600°C and 40 kb using a similar method as in the water experiments⁴⁸. The two broken curves in *Figure 16* give the results. It appears as if the ionization would also be increased substantially by raising temperature and pressure. The ionization product for ammonia at 500°C and 40 kb has been estimated from these conductance data to be 4×10^{-4} mole² 1^{-2} . This would be an increase by about a factor of about 10^{18} over the value for 25°C at saturation pressure⁴⁹.



Figure 18. Specific conductance σ of supercritical caesium and mercury and of supercritical water as a function of reduced density ρ/ρ_c

As a conclusion it may be interesting to compare the electric conductance of dense supercritical water with the conductance of dense gaseous supercritical mercury⁵⁰ and caesium⁵¹. This is done in *Figure 18*. The logarithm of the specific conductance is plotted as a function of reduced density. The actual density is divided by the critical density of each substance. The comparison is made at a reduced temperature of 1.035—slightly above the critical point. For water, however, a second curve for a reduced temperature