

Formation of Gd-Al Alloy Films by a Molten Salt Electrochemical Process

Concha Caravaca and Guadalupe De Córdoba

CIEMAT, Departamento de Energía/División de Fisión Nuclear/URAA, Avda. Complutense, 22,
Madrid, 28040, Spain

Reprint requests to Dr. C. C.; Fax: +34-91-3466233; E-mail: c.caravaca@ciemat.es

Z. Naturforsch. **63a**, 98 – 106 (2008); received December 1, 2006

Presented at the EUCHEM Conference on Molten Salts and Ionic Liquids, Hammamet, Tunisia, September 16 – 22, 2006.

The electrochemistry of molten LiCl-KCl-GdCl_3 at a reactive Al electrode has been studied at 723 to 823 K. Electrochemical techniques such as cyclic voltammetry and chronopotentiometry have been used in order to identify the intermetallic compounds formed. Cyclic voltammetry showed that, while at an inert W electrode GdCl_3 is reduced to Gd metal in a single step at a potential close to the reduction of the solvent, at an Al electrode a shift towards more positive values occurs. This shift of the cathodic potential indicated a reduction of the activity of Gd in Al with respect to that of W, due to the formation of alloys. The surface characterization of samples formed by both galvanostatic and potentiostatic electrolysis has shown the presence of two intermetallic compounds: GdAl_3 and GdAl_2 . Using open-circuit chronopotentiometry it has been possible to measure the potentials at which these compounds are transformed into each other. The values of these potential plateaus, once transformed into e. f. m. values, allowed to determine the thermodynamic properties of the GdAl_3 intermetallic compound.

Key words: Molten Salts; Gadolinium; Electrolysis; Alloy Formation; Pyrochemical Processes.