Suppression of Zinc Dendrites in Zinc Electrode Power Cells

Addition of one of various tetraalkyl quaternary ammonium salts, to the alkaline zincate electrolyte of a cell, prevents formation of zinc dendrites during charging of the zinc electrode (see table); the electrode's capacity is not impaired. Elimination of dendrites prolongs the cell's life.

The fact that 0.2 M Br⁻ (as KBr) is completely ineffective in preventing initiation of dendrites implies that the influence of the tetraalkyls is due to the adsorbed cation. The effect of the tetrapentyl cation on both initiation and propagation is exactly analogous to that of the same concentration of lead (see Tech Brief 70-10153). It is concluded that at the same
concentration the tetrapentyl is as effective as lead at -100-mV overpotential. Addition of soluble lead salts (acetate or nitrate) confirmed the results reported in Tech Brief 70-10153 (see fig.). At a lead concentration of $10^{-4} M$, dendrites are eliminated provided that the zinc overpotential does not exceed -140 mV.

It is proposed that adsorption of either lead ions or tetraalkyl cations interferes with the initiation of zinc dendrites because of the adsorption at certain preferential sites and of the slowing of the normally fast surface-diffusion step. The net result of these effects is formation of many very small grains rather than of fewer large grains that normally precedes initiation of dendrites.

Minimum concentrations of quaternary ammonium salts for complete suppression of zinc dendrites from an electrolyte of 10$M$ zincate-10% KOH at 30°C and -100 mV. At higher concentrations the electrodeposit’s morphology is always compact.

<table>
<thead>
<tr>
<th>Quaternary salt</th>
<th>Minimum concentration, $M$</th>
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<tbody>
<tr>
<td>$(C_2H_5)4NBr$</td>
<td>$2 \times 10^{-1}$</td>
</tr>
<tr>
<td>$(C_3H_7)4NBR$</td>
<td>$2 \times 10^{-2}$</td>
</tr>
<tr>
<td>$(C_4H_9)4NBr$</td>
<td>$2 \times 10^{-3}$</td>
</tr>
<tr>
<td>$(C_5H_{11})4NI$</td>
<td>$2 \times 10^{-4}$</td>
</tr>
</tbody>
</table>

Note:
Requests for further information may be directed to:
Technology Utilization Officer
Headquarters
National Aeronautics and Space Administration
Washington, D.C. 20546
Reference: TSP70-10434

Patent status:
Inquiries about obtaining rights for the commercial use of this invention may be made to NASA, Code GP, Washington, D.C. 20546.
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