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Erratum

Erratum to "Further studies of the anodic dissolution in sodium chloride electrolyte of aluminium alloys containing tin and gallium" [J. Power Sources 193 (2) (2009) 895–898]

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The publisher regrets that there was an error within Table 1. Please see corrected table below.

The publisher would like to apologise for any inconvenience this may have caused to the authors of this article and readers of the journal.

Table 1The anodic dissolution of the Al alloy, IO, in the small volume cell filled with 2 M NaCl. Current density 170 mA cm⁻². Room temperature except where otherwise stated.

	Experiment	Charge before steep rise in cell voltage ^a (C cm ⁻²)	Comments
1	Fresh Al alloy anode	1090	A 'shroud' formed on alloy anode.
	Fresh electrolyte		·
2	New Al alloy anode	935	The increase in electrolyte pH and the presence of
	Shroud remove with first anode		dispersed solid in the cell, has little influence on anode
	Electrolyte reused		performance.
3	Al alloy anode reused but shroud removed external to cell	1070	Once freed of shroud, the Al alloy anode can continue to
	Fresh electrolyte		be dissolved anodically.
4	Al alloy anode reused but without the shroud being removed	120	The shroud limits anodic dissolution.
	Fresh electrolyte		
5	Fresh Al alloy anode	1090	Increase in temperature does not lead to increased
	Fresh electrolyte		charge density.
	Temperature: 311 K		
6	Fresh Al alloy anode	1585	More material in base of cell as shroud broken up to
	Fresh electrolyte		some extent.
	In ultrasonic bath		

 $^{^{\}rm a}$ Charge density until the cell voltage increases to 3 V.

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