TOWARDS TOTAL ELIMINATION OF THE ANODE EFFECT
AND PFC EMISSIONS?

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Abstract

In connection with the electrolytic production of aluminum the anodic reactions have been studied on carbon microelectrodes by voltammetry and chronoamperometry. Anode gases have been analyzed by gas chromatography on-line during controlled-potential electrolysis in a laboratory-scale aluminum reduction cell. When the voltage exceeds a critical value (about 3 V vs Al/Al³⁺), the cell current drops precipitously. This is attributed to the formation of a highly resistive film on the surface of the anode. The existence of this putative film was shown to be strictly potential dependent - the film could be formed and removed at will by regulation of applied potential. The rate of PFC generation was found to vary with the magnitude of the anodic overpotential. The anode effect can be averted by reducing cell current so as to prevent voltage from exceeding a critical value.
INTRODUCTION

In the primary production of aluminum by the Hall-Heroult process, a cell malfunction known as the anode effect results in the generation of the perfluorocarbons (PFCs) CF$_3$ and C$_2$F$_6$ [1]. Owing to the high global warming potentials of these gases [2] and the fact that, in the U.S., aluminum smelting is the number one point source of PFC emissions, the Environmental Protection Agency and the primary aluminum producers have established the Voluntary Aluminum Industrial Partnership (VAIP) with the goal of substantially reducing PFC emissions [3]. To gain a better understanding of the mechanism of PFC generation, VAIP has undertaken two projects: (1) measurements of PFC emissions from industrial smelters [4], and (2) basic research into the attendant electrochemistry. The results of the latter project have been reported recently and are summarized here for the convenience of the reader [5].

Previous work in this laboratory had shown that, in conformity with industrial data, PFCs are generated only when the cell goes on anode effect [6]. Furthermore, time-resolved data taken by Kimmerle et al. on industrial cells [7] and by Nissen and Sadoway on laboratory-scale cells [6] confirm that by far the highest PFC levels are attained during the first several minutes after the cell goes on anode effect. Clearly, any plan to reduce emissions needs to address this fact. Accordingly, the present investigation was directed at the underlying kinetics of PFC generation, more specifically the nature of the reactions at the electrode. The fundamental electrochemistry was studied by cyclic voltammetry and chronocamperometry. Subsequently, controlled-potential electrolysis with analysis of anode gas was performed. It was found that when the voltage exceeds a critical value (about 3 V vs Al/Al$^3+$), the cell current drops precipitously. This is attributed to the formation of a highly resistive film on the surface of the anode. The existence of this putative film was shown to be strictly potential dependent, i.e., the film could be formed and removed at will by regulation of applied potential. The rate of PFC generation was found to vary with the magnitude of the anodic overpotential. In controlled current electrolysis it was found that the anode effect can be completely averted by reducing cell current so as to prevent voltage from exceeding a critical value. There is reason to believe that the results can be applied in an industrial setting.

EXPERIMENTAL

The cell design was largely identical to that used previously in this laboratory [6,8]. Electrochemical measurements were made with a potentiostat (Solartron Electrochemical Interface, model 1287, Allentown, PA) controlled by a personal computer running Corrware (Scribner Associates, Southern Pines, NC). The working electrode was made of either a graphite or a glassy carbon rod, about 2 to 3 mm in diameter, shrouded by BN, so that about 10 mm in length was exposed to the melt and electrochemically active. The molybdenum crucible served as the counter electrode. The reference electrode was Al/AlF$_3$ based upon the design patented by Sadoway [9,10].

For the analysis of anode gas during the electrolysis, a tubular anode was used [8]. The outer vertical surface is insulated with BN; hence, only the inside walls (8 mm in dia., 10 mm height) contact the electrolyte. Anode gas was drawn through an alumina tube from the chamber formed above the melt “inside” the electrode. Gas analysis was performed by gas chromatograph (Model M200 Dual Gas Analyzer, MTI Analytical Instruments, Fremont, CA). The electrolyte was prepared from reagent-grade chemicals: AlF$_3$ (98%), CaF$_2$ (99%) and Na$_3$AlF$_6$ (97%) dried at 500°C under Ar flow for about 12 hours and pre-melted before being introduced into the reduction cell. Prior to cell testing, melts were subjected to pre-
electrolysis to rid them of impurities. All experiments were conducted at a temperature of 975°C.

RESULTS AND DISCUSSION

CYCLIC VOLTAMMETRY

Cyclic voltammetry was performed on melts doped with alumina. Because the technique is conducted with a microelectrode and relies upon depletion of the electroactive species on a time scale comparable to that of the voltage sweep, it is necessary to fix the alumina concentration at a value much lower than is typically encountered in industrial operations. In this set of experiments, alumina levels ranged from 0.01 wt % to 0.60 wt %. The solvent was 84 wt % Na₃AlF₆ - 11 wt % AlF₃ - 5 wt % CaF₂. Figure 1 shows the behavior of a melt containing 0.01 wt % Al₂O₃ on a graphite electrode at a sweep rate of 100 mV s⁻¹. On the forward (anodic) sweep, peaks are evident at 1.8 V, 3.4 V, and 4.3 V. Repeatedly, on the reverse scan nothing measurable was found in the way of a cathodic peak, indicating that the electrode process is highly irreversible. Actually, the voltammogram of the reverse scan is rather similar to that of forward scan. We attribute the rise in current at 1.8 V to the discharge of oxide species, since the potential is not extreme enough to support the discharge of fluoride: the lowest potential for PFC evolution is 2.5 V for CF₄. We suspect that the peaks at 3.4 V and 4.3 V are related to the discharge of fluoride.

![Cyclic voltammogram](image)

**Fig. 1** Cyclic voltammogram of 0.01 wt% Al₂O₃ melt.

Graphite electrode, v = 100 mV s⁻¹.

Figure 4 is the voltammogram of a melt containing 0.25 wt % alumina. The potential was swept over an exceptionally wide range – out to 14.0 V. As above, the current rises with potential up to about 3.0 V at which point there is a distinct decrease in current. What is surprising is that beyond 3.0 V, the current remains low, all the way out to 14.0 V. With the exception of a tiny peak at 4.0 V there is no evidence of faradaic activity on a par with that observed below 3.0 V. On the reverse sweep, there is no cathodic activity (as above), and the anodic peak at 3.0 V is retracted with some offset to more cathodic potentials (~2.8 V).
Figure 4 Cyclic voltammogram of 0.25 wt% \( \text{Al}_2\text{O}_3 \) melt. Graphite electrode, \( v = 500 \text{ mV s}^{-1} \).

Figure 5 Cyclic voltammetry of various melts. Forward and reverse scans depicted separately. Graphite electrode, \( v = 100 \text{ mV s}^{-1} \).

Figure 5 shows the effect of alumina concentration in a set of voltammograms all taken at the same sweep rate of 100 mV s\(^{-1}\). The current of the first wave starting at 1.2 V increases with alumina concentration which is consistent with the assignment of this peak to the discharge of oxide. The elementary process is in fact the oxidation of \( \text{O}^{2-} \) which is assumed to be present.
as part of an oxyfluoro-anionic species. At all alumina concentrations the current falls at intermediate potentials (3.0 - 4.0 V) and remains low even as potential increases beyond 4.0 V. There appears to be some correlation between the potential at which the current drops off (we call this the critical potential, EC) and the alumina concentration: EC rises with alumina concentration. However, on the reverse sweep, the potential at which current rises (we call this the recovery potential, ERC) shows no dependence on alumina concentration.

CHRONOAMPEROMETRY

Chronoamperometry involves stepping the potential on the working electrode from rest potential value and measuring the current between the working and counter electrodes. Figure 6 shows chronoamperometric traces taken at various potentials for a sample containing 0.2 wt % alumina. A trend similar to that seen in the sweep voltammetry is observed, i.e., current increases with potential up to about 3.0 V and then falls abruptly at more extreme potentials. This is summarized in Figure 7 which plots the average current sampled from the chronoamperometric traces as a function of applied potential. On the same plot are the data from linear sweep voltammetry. While there are modest differences between the behaviors of graphite and glassy carbon electrodes, for each material the two techniques give identical results. This, together with the fact that the linear sweep voltammogram does not depend on sweep rate, indicates that the kinetics of the electrode process must be controlled by an interfacial step and not by mass transfer.

Fig. 6 Stepped - potential chronoamperometry of 0.2 wt% Al₂O₃ melt. Graphite electrode.
GAS ANALYSIS DURING CONTROLLED-POTENTIAL ELECTROLYSIS

The close correspondence between voltammograms measured by swept-potential and stepped-potential techniques indicates that the electrode reaction rate is dependent mainly upon the electrode potential; hence, controlled-potential electrolysis combined with chemical analysis of the gas produced at the anode should prove to be insightful. Such experiments were conducted using a tubular anode in a cell described previously [8]. Simultaneously, the anode gas was analyzed before, during, and after application of the potential. Figure 8 displays the data for a potential step of 4.0 V in a melt containing alumina at a concentration of 0.1 wt %. Trace (a) shows the potential step and the current response, while (b) and (c) show how the concentration of CO, CO₂, CF₄, and C₂F₆ vary with time in the anode chamber. Apart from carbon monoxide, which is always expected to be present at the temperature of the experiment (975°C) due to the Boudouard reaction,

\[
    \text{CO}_2 + \text{C} \rightarrow 2 \text{CO}, \quad (1)
\]

all other gases were detected only when current passed through the cell. Furthermore, their concentrations fell to zero when current ceased to flow through the cell. We assume, therefore, that the change of gas composition during electrolysis is due to faradaic processes occurring in the cell. In another words, the change in composition should be governed by the rate of the electrode reaction that generates gases. Measurements were carried out for different potential steps up to 14.0 V. In all experiments no peaks other than those of N₂, O₂, CO, CO₂, CF₄ and C₂F₆ were detected.
Figure 8 Current response and anode gas composition during potential-controlled electrolysis ($C_{\text{AlO}} < 0.1\%$).

Figure 9 summarizes the results of many such experiments by plotting the average values of current and gas concentration as functions of applied potential. The melt had an estimated alumina level of $< 0.1\%$ Al$_2$O$_3$ and had been treated by pre-electrolysis. In this figure, $C_{\text{CO}}$ is the concentration of CO corrected by subtraction of the nonfaradaic baseline value. Over the entire range of potential, CO is the biggest component of the anode gas. Indeed, the shape of the CO concentration history mimics that of the cell current. At low potentials ($< 2.8$ V) CO$_2$ levels are comparable to those of CO ($C_{\text{CO}} / C_{\text{CO}_2} \approx 4$); at higher potentials CO$_2$ levels drop significantly. CF$_4$ and C$_2$F$_6$ were detected at potentials exceeding 2.8 V. At high
potentials (>5.0 V), the concentration of CF₄ is comparable to that of CO (C₃O / C₄CF₄ □ 2.5), and the concentration of C₂F₆ is much lower than that of CF₄ (C₃CF₄ / C₄C₂F₆ □ 50).

![Graphs](image)

**Fig. 9** Potential dependence of current and anode gas composition during potential-controlled electrolysis (C₃ADS < 0.1%).

**ANODIC REACTIONS AND THE ANODE EFFECT**

The results of the anode gas analysis clearly indicate that the first faradaic process, occurring at potentials exceeding 1.2 V, is the oxidation of oxide with the formation of CO and CO₂. At more extreme potentials (>2.8 V) we observe the oxidation of fluoride with the formation of CF₄ as expressed by several rather weak peaks. The relationship between the oxide and fluoride peak heights is in stark contrast to the concentrations of the two species. In all our experiments the oxide (Al₂O₃) concentration was very low (<0.25 wt %): the melts were
essentially fluorides doped with a tiny amount of alumina. In spite of this, the fluoride peaks are small relative to the oxide peak. One explanation is that at potentials capable of supporting PFC evolution the anode is covered by a film that acts as an electrical insulator. On the basis of this work alone not much can be said about the chemistry of the film. However, in an analogous system, molten KF-HF, Imoto et al. [11] and Bai et al. [12] have suggested that the formation of a “CF” film will inhibit the electrode process and contribute to anode effect. Our results are consistent with the formation of an insulating film at intermediate potentials which, in turn, inhibits the electrode process at more extreme potentials.

Further support can be found in the following.
(1) In some of our experiments the anode was vertically disposed (the inside wall of a graphite tube) which made it impossible to accumulate gas on its surface, yet the cell went on anode effect.
(2) The recovery potential, ERC, does not depend on the oxide concentration. Thus, we expect the resistant film to be a fluorocarbon compound – formed when the discharge of fluoride ion occurs and sustained only at high potentials.

![Graph](image)

**Fig. 12** Cyclic voltammogram of 0.40 wt% $A_2O_3$ melt. Graphite electrode, $v = 500$ mV s$^{-1}$. 
Figure 12 is a set of cyclic voltammograms displaying the key effect of the electrode potential on the anode effect. The voltammogram (a) is fully retraceable when the reverse potential is lower than the critical potential. The anode effect occurs when the potential exceeds 3.4 V (b), and the electrode becomes faradically active again when the potential is swept back to potentials lower than 2.8 V. It is as though the shift to less anodic potential promotes active stripping of the putative resistive film, thereby rendering the electrode electrochemically active. Perhaps the hysteresis seen in (b) is evidence of activation barriers associated with film formation on the anodic sweep and with film removal on the return sweep.

![Figure 12](image12.png)

Figure 13 compares the current responses to two different potential waveforms. In both cases, the electrode was held at 4 potentials, E₁, E₂, E₃, and E₄, for ten seconds each, and the current was monitored. E₁ and E₄ were set to the rest potential of the electrode, 0.85 V, and E₂ was set to 2.0 V. The only difference between the two waveforms is in the value of E₃: 3.0 V in one case and 3.5 V in the other. These potentials were chosen to straddle the critical potential, EC. The current response for both waveforms is identical at E₁, E₂, and E₄. There is a pronounced difference at E₃. Figure 14 shows this effect even more vividly. In this case the potential
waveform is repeated: $E_1 = 0.85 \text{ V}$, $E_2 = 2.0 \text{ V}$, $E_3 = 5.0 \text{ V}$, and $E_4 = 3.0 \text{ V}$. It is clear that when the electrode potential is set at $5.0 \text{ V}$ an insulating barrier blocks current flow. The effect is reversed as soon as the potential drops to $3.0 \text{ V}$, which is below the critical potential. These current responses are reproducible for many cycles.

![Graph showing current response of square potential wave](image)

*Fig. 14 Current response of square potential wave.*

$C_{\text{Al}_2\text{O}_3} = 0.15 \text{ wt\%}$, graphite electrode.

**POSSIBLE WAY TO REDUCE THE AE AND PFC GENERATION**

Once the electrode goes on anode effect, the current will fall quickly, and the electrode will follow the polarization curve shown in Figures 4, 5, and 7. In the high-potential region the current rises slightly with increasing potential, and the fraction of PFC in the anode gas increases. This tendency can be seen in Figures 9 and 10. The results of measurements made in aluminum smelters [7] and in laboratory-scale cells [6] indicate that the most intense period of PFC emission is the first moments of high voltage immediately following the onset of the anode effect.

Accordingly, we propose the following without any regard for instant practicality: reduction in cell current so as to prevent voltage from rising to values extreme enough to cause the electrode to go on anode effect. Experiments performed in our laboratory have shown that it is not necessary to shunt the entire cell current in order to prevent anode effect. In fact, as cell voltage rises to near critical values, only a modest reduction in cell current (on the order of several per cent) is needed to stave off anode effect. During this time, conventional measures of intervention can be initiated. Data from industrial smelters indicate that the concept should be adaptable to commercial cells.

Clearly, the ability to avert anode effect translates into total elimination of PFC emissions.
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