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**Fabrication of Ceramic Components  
for Use in Plutonium Electrorefining**

LOS ALAMOS NATIONAL LABORATORY



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by

S. D. Stoddard  
D. E. Nuckolls



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## FABRICATION OF CERAMIC COMPONENTS FOR USE IN PLUTONIUM ELECTROREFINING

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### ABSTRACT

A method of fabricating ceramic cells and associated components for use in the electrolytic refining or electrowinning of plutonium metal is described. Details for making such ware of impervious alumina and magnesia are described. References on the development of processes for the electrochemical preparation of up to 3.5 kg per batch of plutonium and the ceramics required for such processes are also given.

### 1. INTRODUCTION

Although the electrochemical preparation of Pu metal was reported over 20 years ago by Kolodney<sup>1</sup> of this Laboratory, the feasibility of a process for producing the metal in relatively large quantities was not reported until 1958 by Leary et al.<sup>2,3</sup> Batch processes for production of up to 3.5 kg of metal were reported by Mullins, Leary, Morgan, and Maraman<sup>4,5</sup> in the open literature in 1963. Subsequent reports have dealt with improved processes and purities attained therein.<sup>6,7,8</sup>

During some of this period, workers at Hanford Atomic Products Laboratory (HAPCO),<sup>9</sup> Argonne National Laboratory,<sup>10</sup> and the Dow Chemical Company at Rocky Flats<sup>11</sup> have also reported electrochemical processing of Pu. The melt container and associated components have evolved from metals, namely Ta, to ceramics, the fabrication of which is the subject of this paper.

The fabrication of ceramic components for the HAPCO electrowinning and batch electrorefining process wherein the ceramic body is composed of 4 mole %  $Mg_2TiO_4$  and 96 mole % MgO has been described by Rosenfels.<sup>12</sup> Work with magnesia components at the

Dow Chemical Rocky Flats Plant has been reported by Kessel et al.<sup>13</sup>

At this Laboratory the first ceramic cells were made of aluminum oxide ceramics utilizing metallic Ta stirrers. The alumina cells were found satisfactory in all-chloride electrolytes. It was seen that by substituting MgO for  $Al_2O_3$ , the impurity content of the product metal could be cut from 40 ppm of Al to less than 5 ppm of Mg. Incidental to the lowering of the total impurity content by this substitution, it was found that  $PuF_3$  or  $PuF_4$  could be used in place of the uncommon and hygroscopic  $PuCl_3$  electrolyte. Mixed fluoride-chloride electrolytes were found unsatisfactory when alumina cells were used because the Al concentration of the Pu increased from 40 to 450 ppm. Consequently, all ceramic components, i.e., anode container, electrolyte container, stirrer, and protective anode sheath, were thenceforth slip cast of impervious magnesia ( $MgO + 3$  wt %  $Y_2O_3$ ).

Figure 1 illustrates various ceramic components produced at LASL for use in refining processes.



Fig. 1. LASL produced ceramic components for refining processes.

## 2. MATERIALS

### Properties Required

Ceramic accessories for electrorefining Pu metal must have the following properties.

1. The cell must contain the low viscosity melt (NaCl, KCl, + 2 mole % PuCl<sub>3</sub>, PuF<sub>3</sub>, or PuF<sub>4</sub>) employed at temperatures up to approximately 850°C.

2. The cell and components should have high thermodynamic stability to minimize the possibility of reaction between the container and the molten salts or molten Pu, or both.

3. The cell and components must have adequate thermal shock resistance. The cells are heated at 50°C/h to 740°C and held 1 h at temperature before the temperature is raised to 840°C after which they are furnace cooled.

4. The material must be mechanically strong to withstand thermal and mechanical shock. The stirrer revolves at up to 1000 rpm, imposing high tensile stresses, even though it revolves in a low viscosity medium.

5. The cells and components should be commercially available, fabricable by established techniques, and reasonably inexpensive. Although the cell is for one-time use, the stirrers and anode sheaths are used repeatedly. An average of seven cycles is realized with the stirrers, and 10 to 12 cycles with the anode rod sheaths.

Figure 2 illustrates the latest electrorefining process assembly, predecessors of which are completely described elsewhere.<sup>14</sup> Figure 3 shows the ceramic components in greater detail.

The choices of container materials for the molten electrolytes and molten Pu are rather limited. Both corrosion resistance and corrosion products are of consideration in the selection; i.e., Al cannot be easily removed from the product metal where Mg is easily removed by vacuum melting. From a consideration of their free energies of formation, the following oxides could be used in order of increasing stability (-ΔF, K cal/g-atom oxygen at 1000°K): SiO<sub>2</sub> (84), Al<sub>2</sub>O<sub>3</sub> (109), Pu<sub>2</sub>O<sub>3</sub> (109), MgO (119), Y<sub>2</sub>O<sub>3</sub> (119), ThO<sub>2</sub> (124), and CaO (127). Owing to their availability within this Laboratory,<sup>15,16</sup> first Al<sub>2</sub>O<sub>3</sub> and then MgO were enlisted. Although technology is available for the production of impervious Y<sub>2</sub>O<sub>3</sub> ware and CaO ware,<sup>17</sup> cells have not been

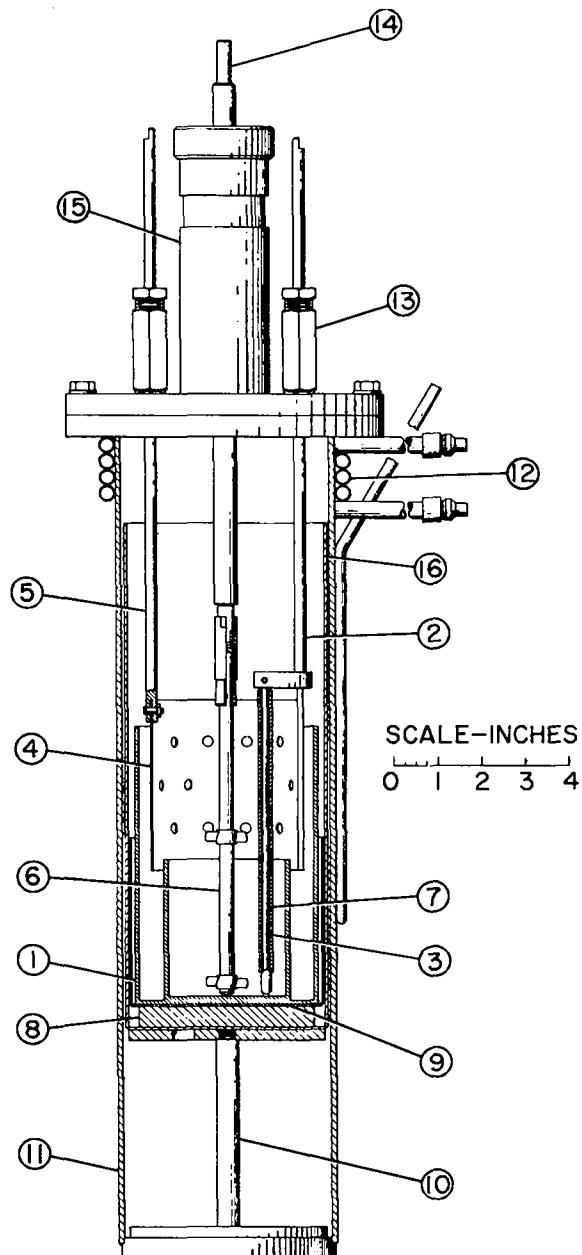


Fig. 2. Electrorefining process assembly.

1. Alumina or magnesia cell
2. Nickel anode lead
3. Tungsten anode rod
4. Tungsten cathode tube
5. Tantalum cathode lead
6. Alumina or magnesia stirrer assembly
7. Magnesia anode sheath
8. Magnesia disc 4-1/8-in. diam x 3/8-in. thick
9. Tantalum safety crucible
10. Stainless steel pedestal
11. Stainless steel tube
12. Copper cooling coils
13. Teflon lined electrode seals
14. Stainless steel stirring shaft
15. Stirrer gland
16. Stainless steel can

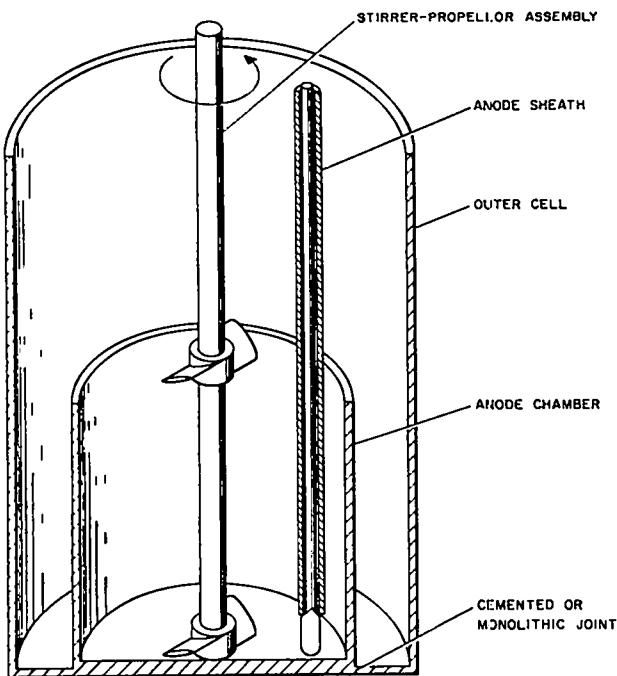


Fig. 3. Ceramic components for electrorefining.

fabricated of either material.  $\text{La}_2\text{O}_3$  together with the referenced series of oxides is considered a good possibility with respect to favorable free energy of formation, but it, like the  $\text{CaO}$ , is hygroscopic, making it difficult to store for production applications. HAPCO also evaluated  $\text{MgAl}_2\text{O}_4$ ,  $\text{Al}_2\text{O}_3$ ,  $\text{BeO}$ ,  $\text{ZrO}_2$ , and porous  $\text{MgO}$ , as applied to electro-winning, with varying results.<sup>18</sup>

The impervious  $\text{Al}_2\text{O}_3$  and  $\text{MgO}$  ware tested at LASL was fabricated with no additives other than incidental impurities in the as-received material and with grain-growth inhibitors or sintering aids such as  $\text{MgO}$  (in the former) and  $\text{Y}_2\text{O}_3$  (up to 3 wt %) in each. A wide variety of commercially available fused  $\text{MgO}$  powders were evaluated and found equally satisfactory in resistance to corrosion. It is interesting that cells made of fused raw material containing as much as 1000 ppm B did not contribute measurable B (<0.5 ppm) to the refined metal.

### 3. FABRICATION PROCEDURES

#### A. $\text{MgO}-\text{Y}_2\text{O}_3$ Preparation - General

The starting material used in this process is fused  $\text{MgO}$  containing less than 1% total impurities, the major contaminants being  $\text{CaO}$  and  $\text{SiO}_2$ . The particle size distribution is 100% -200 mesh, 40% +325 mesh, and 60% -325 mesh. This material is reduced

in particle size by ball milling for 42 h at 55 rpm and 1°C. A 1-gal mill charge includes 2000 g of  $\text{MgO}$ , 600 ml of water, 3185 g of 1-in.-diam balls, and 1015 g of 3/4-in.-diam balls. High density alumina balls are used in porcelain mills. Chemical analyses of the milled  $\text{MgO}$  revealed that only about 50 ppm Al were picked up during milling.

After this initial milling, 500 additional g of the  $\text{MgO}$  are added together with 3% by weight of 99.9% pure, -325 mesh  $\text{Y}_2\text{O}_3$ , and the milling is continued for 2 h. The extra  $\text{MgO}$  is added to achieve the desired grain size distribution to minimize firing shrinkage. Andreason pipette data indicated that the average particle size was approximately  $2\mu$  although it was not certain that the suspensions were completely deagglomerated.

After being milled for 44 h the slip is deflocculated through the addition of Formamide\* (1 ml per 100 g of slip). The viscosity of the slip is approximately 100 cP as measured with a Brookfield Syncro-Electric Viscometer Model RVT at 10 rpm using a No. 1 spindle. The pH of the slip is between 10 and 11, thereby contributing to long mold life.

The mechanism of deflocculation of this slip is not fully understood. It is suspected that a controlled amount of hydration is necessary to produce a slip having optimum viscosity. The weight loss of samples upon firing indicates that approximately 16%  $\text{Mg}(\text{OH})_2$  is present in the cast ware.

The casting operation is carried out in conventional plaster molds made from three parts water and four parts plaster. The rate of casting is relatively slow; approximately 15 min are required to form a 1/16-in.-thick wall. The casting rate decreases appreciably as the wall thickness increases beyond 1/16 in. As a consequence it is often difficult to cast thick-walled or solid pieces. Another problem sometimes encountered is caused by the gradual rise of temperature of the slip during casting. The rate of hydroxide formation increases as the slip warms, and the 30% volume expansion accompanying the reaction may cause the casting to expand in the mold. This problem is not serious except when very thick sections are

\*A product of Matheson, Norwood, and Bell, Norwood, Ohio.

being cast. In this case it is advantageous to cast in chilled molds, thereby reducing the rate at which the slip warms. The slip is kept chilled (approximately 20°C) between casts.

The slip has excellent drain characteristics permitting sharp interior corners to be formed. The drying shrinkage of the cast ware is very low, and it is thus advisable to incorporate a slight taper into one-piece molds. The pieces release from the mold quite easily, and no mold release agent is needed.

The cast pieces are dried at about 35°C. They may then be stored in the open air for indefinite periods before firing.

#### B. $\text{Al}_2\text{O}_3$ Preparation - General

The starting material used in this process is fused  $\text{Al}_2\text{O}_3$  containing less than 0.2% total impurities, the major contaminants being Si, Fe, Cr, and Zn. The average particle size is 1.7 $\mu$  measured with a Fisher Subsieve analyzer. This material is reduced in particle size by ball milling for 30 h at 55°C. A 1-gal mill charge includes 2000 g of fused  $\text{Al}_2\text{O}_3$ , 3.29 g Marex\* deflocculant, 2.64 g dry sodium meta-silicate, 3185 g of 1-in.-diam balls and 1015 g of 3/4-in.-diam balls. High density  $\text{Al}_2\text{O}_3$  balls are used in porcelain mills. After 28 h of milling, 40 g (2 wt %) of  $\text{Y}_2\text{O}_3$  is added and milling is continued 2 hr for a total milling time of 30 h. After the slip is discharged from the mill, approximately 7 ml Darvan No. 7\*\* together with 1 ml formaldehyde is added to the batch. The formaldehyde is used to prolong the shelf life of the slip, while the Darvan acts as a deflocculant. The viscosity of the slip at this stage is 250 cP as measured with a Brookfield Syncro-Electric Viscometer Model RVT at 10 rpm using a No. 1 spindle. The pH of the slip is between 8 and 9, thereby contributing to long mold life.

The casting operation is carried out in the same manner as that employed for the magnesium oxide except that there is no need to chill the mold. The casting rate is essentially identical to that for the magnesia.

\*Marex, an ammonium alginate from Kelco Company.

\*\*Darvan No. 7, a polyelectrolyte from R. T. Vanderbilt, Inc.

#### C. Fabrication Details

##### 1. $\text{MgO-2 wt \% Y}_2\text{O}_3$ Cells (LASL Dwg. No. 26Y-72325-8, Rev. A, Part 75\*).

The outer cell and the anode chamber are cast separately in conventional plaster molds and after removal from the mold are dried separately as described above. They are then nested together and fired on prefired, coarse-mix (21-1), dry-pressed  $\text{MgO}$  setter plates from which they are separated by a 1/16-in.-thick layer of fused -20 +50 mesh (U.S. Standard series)  $\text{MgO}$  grain. This setting is fired in natural gas-air fueled kilns to 1775°C and held at peak temperature for 20 to 24 h.

During the firing of pieces prepared by this method it is important to use a low heating rate until all of the  $\text{Mg}(\text{OH})_2$  has been decomposed. This treatment yields relatively translucent ware which has no open porosity, exhibits approximately 11% firing shrinkage, and has a density ranging from 94 to 97% of theoretical density.

It is felt that the  $\text{Y}_2\text{O}_3$  addition and the long soak at this relatively high temperature are responsible for the successful bond achieved between the cells.

##### 2. $\text{Al}_2\text{O}_3$ Cells (LASL Dwg. No. 26Y-72325-8, Rev. A, Part 75).

The  $\text{Al}_2\text{O}_3$  cells are cast and fired in air in nearly the same manner as the  $\text{MgO}$ . Although also fired on  $\text{MgO}$  setter plates, the cells are separated from the setters in this case by a 1/32-in.-thick layer of -60 mesh fused  $\text{Al}_2\text{O}_3$  grain. The setting is fired to 1750°C in the same type of kiln and held at peak temperature for 20 h. The fired body has no open porosity, a density of 93 to 94% of theoretical, and a fired shrinkage of approximately 10-1/2%.

##### 3. Stirrers (LASL Dwg. No. 26Y-72500).

The stirrers are solid cast rather than drain cast, employing the same casting slip as the cells, and are assembled from three separate components. The shaft, the propellers, and the straightening weight are cast separately and prefired to 1250-1300°C in an oxidizing atmosphere. After prefiring, the propellers are cemented in position, and the shaft weight and shaft are grooved and cemented together as indicated in Fig. 4. The  $\text{MgO}$  casting

\*This drawing and all others referenced herein are available from the Clearinghouse for Federal Scientific and Technical Information.

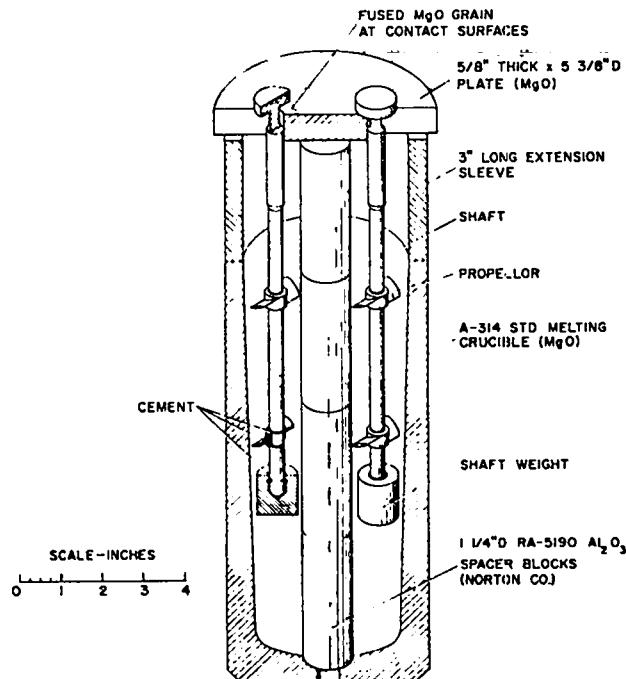


Fig. 4. Setup for firing stirrer assemblies.

slip is employed as the cement for both joints. The cemented assembly is then hung from the flanged end through a notched, coarse-grain MgO plate, separated at the flange-plate interface by a layer of fused MgO grain as shown in Fig. 4.

This setup is then fired with the same schedule as the refining cells. Following firing and "furnace cooling," the weight and flange are cut off utilizing a conventional cut-off saw and the enlarged shaft section is ground to 0.312-in. diam concentric with the balance of the as-cast shaft.

Stirrers of both the MgO and Al<sub>2</sub>O<sub>3</sub> compositions have been produced in the manner described. In cementing the alumina propellers to the shafts, the Al<sub>2</sub>O<sub>3</sub> slip replaces the MgO slip used for the magnesia assemblies.

#### 4. Anode Sheaths (LASL Dwg. No. 26Y-75970 A-1 and 26Y-72495, Detail 42).

The anode sheaths have been fabricated of the MgO composition only as it was in full production before the advent of a protective sheath on the anode rod. The slip used for these pieces is different from that described in Section 3A in that no MgO, only Y<sub>2</sub>O<sub>3</sub>, is added after the 44 h milling time. The elimination of the coarse fraction increases drying shrinkage and promotes removal from the mold.

The sheaths are drain cast in conventional plaster molds with a flange at the mouth of the mold to facilitate hanging during the firing process (see Fig. 4). As was the case with the stirrers, it was noted at once that the required straightness could be achieved only by hanging the ware. The flange is cut off using a conventional wet-cutting diamond saw after firing.

#### 4. DISCUSSION

##### A. Cementing Slip-Cast Cells

Although the process described for joining the outer cell and anode chamber created no problem at LASL, it was soon discovered that for other fabricators this was a severe problem. The mode of failure noted in commercially supplied cell assemblies is failure of the bond between the two nested cups. If the as-received cells pass the RF bond test\* (as devised by personnel at the Dow Chemical Rocky Flats Plant) wherein a dowel is used as a lever to pry against the inner cell with an applied load of 3 lb, they often fail on initial heatup of the assembly. If the cell does not break loose under this treatment, the bond is satisfactory. Other investigators, in addition to using casting slip for stickup, also evaluated some proprietary commercial cements. These, too, have thus far proved unreliable. It is postulated that the principle reason for the success enjoyed here is the high maturing temperature and the long soaking times routinely utilized. Apparently the body is sufficiently plastic under these maturing conditions to allow the two walls to join together and form an essentially monolithic wall. The reluctance of some commercial suppliers to utilize the Y<sub>2</sub>O<sub>3</sub> addition (because of its relatively high cost) also is felt to contribute to their problems by lowering the sintering temperature of MgO base bodies. Unlike the HAPO process described by Rosenfels,<sup>12</sup> the LAMEX process<sup>6</sup> utilizes a cemented cell, making a reliable joint an absolute necessity.

##### B. Stirrer Fabrication

The stirrers used in the refining process must be straight because of their relatively high

\*See Detail A, LASL Dwg. No. 26Y-72325, Part 75.

rotational speed. They must also be free of voids at the juncture of the shaft and stirrer to prevent Pu from entering the voids and ultimately causing fracture because its thermal expansion is greater than that of the MgO.

The presence of voids in the joint is a function of the care and skills employed in that step of the operation. Although this has been reported elsewhere<sup>11</sup> to have been a problem, no difficulties have been experienced here.

Several techniques were evaluated for setting the stirrer assemblies to minimize warpage during firing. A high degree of success (approximately 95%) has been reached with the setup illustrated in Fig. 4. Others report 60% loss in fabrication of stirrers by the slip casting method.

The weight was found necessary to ensure straightness during firing. The weight of the assembled stirrer itself is insufficient to overcome internal stresses induced during fabrication. Likewise it was found that reducing the shaft diameter near the brazing collar was beneficial toward this end.

Early in the processing development, slight warpage in the shaft length was obviated by building up the last 2 inches of the shaft with oxyacetylene flame-applied (Metco Type "P" gun\*) W and subsequently grinding this built-up layer so that it was generally concentric with the major part of the shaft-stirrer assembly. A stirrer embodying this technique is illustrated in Fig. 1.

## 5. SUMMARY

Fabrication methods have been developed to produce cells, sheaths, and stirrers of monolithic, impervious, high purity  $Al_2O_3$  and MgO ceramics to satisfy the rigorous requirements for the electrorefining of Pu.

Although other fabrication methods and compositions have been employed at other installations, the slip-cast magnesia components have been found most satisfactory for use at this Laboratory. The cell is used only once, but the stirrers have been used an average of seven cycles, and the sheaths 10 to 12 cycles, without failure.

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\*A product of Metallizing Engineering Company.