M10 ELECTROLYTE EXCHANGE PROCEDURES FOR 22 AH NICKEL-Cadmium CELGARD CELLS

by

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RAPID ELECTROLYTE EXCHANGE PROCEDURES FOR
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ABSTRACT

State-of-the-health analysis of Nickel-Cadmium aircraft batteries rejected as being unsatisfactory by CF bases has shown that most of the problems encountered can be related to the fact that the KOH concentration in the cells is too low and/or the carbonate concentration is too high.

Exchange of electrolyte restored the performances of the cells but is time consuming, one to two weeks for a battery using the charge-KOH exchange-discharge procedure.

The present technical note describes some of the tests performed and a procedure for rapidly exchanging the electrolyte in 22 AH Nickel-Cadmium Aircraft Celgard cells.

RÉSUMÉ

Une analyse des batteries usagées nickel-cadmium d'aéronef a montré que la baisse de performances observée par les forces armées canadiennes pouvait être attribuée dans une large part à une concentration trop faible en potasse et/ou à une concentration trop élevée en carbonate dans l'électrolyte.

Dans plusieurs cas, un échange d'électrolyte redonna les performances attendues. Toutefois cet échange d'électrolyte prend énormément de temps; une à deux semaines selon la méthode de la charge-échange KOH-décharge de la batterie.

Le présent document offre une nouvelle méthode d'échange de l'électrolyte d'une durée de quelques heures et décrit les tests effectués avec des prototypes. La méthode décrite a été adaptée aux batteries d'aéronef d'une capacité de 22 AH, montées avec des séparateurs "celgard".
# TABLE OF CONTENTS

<table>
<thead>
<tr>
<th>Section</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td>ABSTRACT/RESUME</td>
<td>iii</td>
</tr>
<tr>
<td>TABLE OF CONTENTS</td>
<td>v</td>
</tr>
<tr>
<td>FIGURE CAPTIONS</td>
<td>vii</td>
</tr>
<tr>
<td>INTRODUCTION</td>
<td>1</td>
</tr>
<tr>
<td>BASIC PROCESS</td>
<td>1</td>
</tr>
<tr>
<td>EFFICIENCY OF THE PROCESS</td>
<td>2</td>
</tr>
<tr>
<td>SIX CELL PROTOTYPE</td>
<td>2</td>
</tr>
<tr>
<td>ELECTROLYTE EXCHANGE PROCEDURES FOR 22 AH NICKEL-CADMIUM CELGARD CELLS</td>
<td>3</td>
</tr>
<tr>
<td>Equipment Required</td>
<td>3</td>
</tr>
<tr>
<td>Procedure and Operations</td>
<td>4</td>
</tr>
<tr>
<td>Vacuum Leakage Problems</td>
<td>4</td>
</tr>
<tr>
<td>Turbid Electrolyte</td>
<td>5</td>
</tr>
<tr>
<td>CONCLUSION</td>
<td>5</td>
</tr>
<tr>
<td>REFERENCES</td>
<td>5</td>
</tr>
</tbody>
</table>
LIST OF ILLUSTRATIONS

Figure 1. Exchange of Electrolyte Apparatus: Single Cell Prototype.

Figure 2. Efficiency of Mixing During the First Hour.
# 1 Discharged Cells
# 2 Charged Cells

Figure 3. Exchange of Electrolyte Apparatus: Six Cell Prototype.

Figure 4. Lowering of KOH Concentration in Cells After the Exchange of Electrolyte.
A Water and KOH Exchange Cells
KOH Exchange Cells

Figure 5. Example of Turbidity in Cell Electrolyte.
INTRODUCTION

During recent years, DREO/ECD personnel have conducted state-of-the-health analysis of Nickel-Cadmium cells rejected as being unsatisfactory by CF bases. Experiments have shown that most of the problems encountered result from the KOH concentration in the cells being too low and/or the carbonate concentration being too high (1). Experiments have also shown that after an exchange of electrolyte that brings the KOH and the carbonate concentration within the range suggested in (2), the performance of the cells is restored.

In the past, the cells that were built with a cellophane separator and used for cranking applications lasted less than three years so the cells were often rejected before the need of exchanging the electrolyte appeared. Celgard cells last much longer and it is expected that the electrolyte will need to be exchanged after 3-5 years of field service.

Exchange of electrolyte has always been time consuming (one to two weeks for a battery using the charge-KOH exchange-discharge procedures). DREO/ECD has developed procedures (3,4,5) that reduce this time to about two days. Improvements of the techniques (6) suggest that a complete exchange of electrolyte in a battery can be accomplished within one working day.

The present Technical Note describes some of the tests performed and suggests a procedure for exchanging the electrolyte in 22 AH Nickel-Cadmium Celgard cells. These techniques should not be used for the Nickel-Cadmium cellophane cells still in use in the CF because the application and the release of the vacuum during the process might damage the cellophane separator and promote the formation of internal short-circuits.

BASIC PROCESS

The schematic diagram and details of the required equipment for electrolyte exchange have been covered in (5,6) and will be briefly reviewed for illustrative purposes. Figure 1 shows an exchange of electrolyte apparatus in the simplest configuration. A cell with contaminated electrolyte is mounted on a shaft which can be rotated to position the cell right-side-up or inverted as desired. A modified vent cap and tubing connector make connection to tubing which leads into the bottom region of a volumetric flask referred to as the "Exchange Flask". This flask is linked to a vacuum line. The vacuum is applied and released by a solenoid valve operated by a cycling timer. When the timer causes the vacuum to be applied a quantity of the electrolyte (30-40 ml) from the cell is withdrawn into the exchange flask. It mixes with the much larger quantity of fresh electrolyte causing only small changes in composition. When the vacuum is released on the second
part of the cycle, the electrolyte in the exchange flask flows back into
the cell. This mixes in the cell with the remaining contaminated electrolyte
and the process is repeated. Eventually the electrolyte in the exchange
flask and the cell are of equal concentration. However due to the relative
volumes when the process was started the quality in the cells is now much
higher than it was originally. The major advantage of this process over the
charge-exchange KOH-discharge process is that this process does not tie up,
for extended periods of time, the charge-discharge equipment and it does not
interfere with the daily battery shop duties.

EFFICIENCY OF THE PROCESS

In order to measure the process efficiency and reproducibility from
cell to cell, the electrolyte of four cells in the charged state
and four cells in the discharged state has been changed using the prototype
shown in Figure 1. The initial KOH concentration in the cells was in the
range 29-32% KOH and 500 ml of water was added in the exchange flask each
time the electrolyte in a cell was exchanged. The vacuum mixing cycles were
18 seconds "vacuum on" 12 seconds "vacuum off". The KOH concentration after
24 hours of mixing was considered as 100% mixing. The percent of
mixing with time was calculated according to the formula:

\[
\% \text{ mixing} = \frac{C_t}{C_{24}}
\]

where \( C_t \) = % KOH in the exchange flask at a given
time and

\( C_{24} = \% \text{ KOH after 24 hours of mixing.} \)

The data obtained during the first hour of mixing are presented in
Figure 2. It appears that the mixing in the charged cells took place more
rapidly than in the discharged cells. This is consistent with the fact that
the charged cells have about 30 ml of "free electrolyte" available at the
beginning of mixing and that the plates in the charged state have a more
open structure. The vertical bars represented the range of data obtained
with four different cells. From the height of the bars one can see that the
reproducibility of mixing is also better with the charged cells. After one
hour of mixing, 80% of mixing is completed in the charged cells compared
with 65% with cells in the discharged state. However for safety reasons
during the manipulation of the cells, it is not advised to exchange the
electrolyte while the cells are in the charged state and work has been
pursued with cells in the discharged state.

SIX CELL PROTOTYPE

DREO/ECD has developed a prototype that allows the exchange of
electrolyte in six cells at a time (Figure 3). The prototype has a four
litre exchange flask and has been used to establish the working procedures
to exchange electrolyte in a battery shop.
Water exchange treatments have been conducted in order to try to minimize the difference in KOH concentration from cell to cell prior to the exchange of KOH electrolyte. Six cells containing from 20% to 30% KOH electrolyte were vacuum-cycled for a ten minute period with 2,500 ml of water in the exchange flask. At the end of the 10 minute period, the 2,500 ml of electrolyte was dumped and 2,500 ml of fresh water was added to the exchange flask and mixed for another ten minute period. This treatment reduced the KOH concentration to 5% in each of the six cells. After the water exchange treatment, 3,000 ml of 37% KOH (500 ml/cell) were added to the exchange flask and the cells were vacuum cycled for one hour.

Another set of six cells with from 20% to 30% KOH electrolyte were exempted from the water treatment and were vacuum cycled for one hour with 3,000 ml of 37% KOH in the exchange flask. The KOH concentration was analysed immediately after the treatment and during the following days while both sets of cells were at rest at room temperature.

After the exchange of electrolyte, the cells that received the water treatment showed an average of 28.5 ± 0.7% KOH and the cells without water treatment showed 35.2 ± 0.3% KOH. The difference from cell to cell was more significant with the cells that had received the water treatment. This difference further increased as the cells were left at rest. Figure 4 illustrates the data of the cells with the highest and the lowest KOH concentration for each set of cells. At rest, the cells that received the water treatment decreased in KOH concentration by an average of 1.5% KOH while the cells without the water treatment decreased in KOH concentration by an average of 0.5% KOH. It appears that the water treatment prior to the exchange of KOH electrolyte does not offer any advantages and in fact creates some additional problems in terms of reproducibility from cell to cell. For these reasons the water treatment has been abandoned.

In further experiments, reproducible results were obtained by introducing in the exchange flask 2,400 ml of 32% KOH (400 ml/cell) and vacuum mixing for one hour. The details of the procedure will be explained in the following paragraphs.

ELECTROLYTE EXCHANGE PROCEDURES FOR 22 AH NICKEL-CADMIUM CELGARD CELLS

Equipment Required

- Six cell holder (BBQ type), as illustrated in Figure 3.
- Vacuum facilities: primary vacuum pump (Model No. 0211-V45N-68CX manufactured by Cast or the equivalent) or building vacuum facilities.
- 4,000 ml exchange flask.
- Magnetic stirrer for mixing the electrolytes in the exchange flask.
- 32% KOH solution.
- Solenoid valve (manufactured by ASCO, cat-no. 8329A44N, 117 VAC, orifice 1/8", pipe 1/8").

- Adjustable CAM operated timer switch with 30 second cycle time. (Industrial Timer Corp).

- Tygon tubing with 1/4" bore and 1/16" wall thickness or equivalent.

- Modified vent caps (3,4).

**Procedure and Operations**

Six 22 AH discharged cells are tightly secured in the rotatable cell holder. Each cell has a modified vent cap which is connected via tubing to the 4,000 ml exchange flask. The exchange flask sits on a magnetic stirrer which is used to mix the electrolyte. The exchange flask is filled with 32% KOH (400 ml/cell). The vacuum pump is turned on and the cycling timer is switched on. The suggested cycling mode is 18 seconds "vacuum on" and 12 seconds "vacuum off". Cells are allowed to fill in the upright position for about two minutes. The cell holder is then inverted and the "exchange of electrolyte" operations are left on for at least one hour. At the end of the one hour period, the KOH concentration in each of the cells should be around 30 ±1.0% (tests have been done with cells having a KOH initial concentration of 23 ±1% KOH). As the mixing is not completed after one hour of mixing, a further decrease of about 0.5% KOH is expected. The KOH concentration ends up within the range specified in (2). At the end of the "exchange of electrolyte" treatment the excess of electrolyte in each cell must be emptied prior to charging and the level of electrolyte must be adjusted according to the procedures in (2) at the end of charge.

The volume of KOH electrolyte (400 ml/cell) in the exchange flask has been selected to cover a wide range of cell electrolyte concentrations. The total volume has been selected such that any carbonate concentration in the cells will be reduced or diluted by a factor of about four. For instance, cells with electrolyte in the range 15-30% KOH and carbonate up to 15% should finish after the exchange of electrolyte, with electrolyte in the range suggested in (2).

The suggested exchange procedures require the manipulation of large quantities of KOH electrolyte and it is recommended that safety goggles be worn during the operation. The development of a prototype for the exchange of more than six cells at a time required the use of volumes of electrolyte too large for safe handling in a battery maintenance shop and has been abandoned.

**Vacuum Leakage Problems**

Vacuum leakages can be detected by a constant bubbling in the exchange flask while the vacuum is applied. As some air is left in the cells, bubbling in the exchange flask is normal during the first exchange of electrolyte cycles but should cease. If after two minutes of cycling bubbling persists, a vacuum leakage is indicated. It has been observed that leakages commonly
occurs at the vent cap connection. The wrapping of teflon tape around the vent caps should overcome most of the leakage problems encountered. If leakages persist, the cells responsible for the leakage must be discarded from the exchange treatment and replaced.

Turbid Electrolyte

It is not advised to use this process for cells having "muddy" electrolyte. The electrolyte in each cell should be inspected for turbidity prior to the exchange of electrolyte. Samples of electrolyte may be taken from each cell with a syringe while the battery is in overcharge and gassing, and placed in test tubes for observation. Figure 5 shows examples of electrolyte from cells which had seen extensive use. They are arranged to show the wide range of turbidity which may be found. The two test tubes shown at the right hand end contain pure electrolyte for reference purposes. The two samples at the left are extremely "muddy". Cells having "muddy" electrolyte should be discarded, they will contaminate the electrolyte of the other cells with solid particles during the "exchange of electrolyte" process. It is likely that cells with large quantities of material in the electrolyte, and by inference in the separator materials, would develop short circuits in the near future and should be removed from service.

CONCLUSION

DREO/ECD has developed a prototype that allows the exchange of KOH electrolyte in 22 AH Nickel-Cadmium Celgard Cells within a working day period. The major advantage of this process over the charge-exchange KOH-discharge process is that the new process requires little charge-discharge equipment time and does not interfere with daily battery shop duties. The prototype and the suggested procedures should be evaluated at AMDU in the near future and if found satisfactory should be introduced into CF battery shops that handle 22 AH celgard cells.

REFERENCES

4. K. Feldman. Additional Information to Technical Note 77-8, on Rapid Electrolyte Exchange Technique, DREO Memorandum No. 43/79.
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Nickel-Cadmium Celgard Cells
Nickel-Cadmium Aircraft Batteries
KOH Electrolyte Contamination
Rapid Electrolyte Exchange Technique