Battery AP-B09



Galvanostatic Intermittent Titration Technic (GITT) and its application in analysing of Livion battery



In this application note, GITT method was run on Li-ion battery 2,200 mAh and application of this method on exporting quantitative and qualitative data from battery samples are investigated.





INTRODUCTION

The GITT method consists of two main cycles:

Current Pulse + OCP Cycle no.1

The first cycle include applying galvanostatic charge pulses (positive current) each defined duration, followed by defined relaxation time (OCP), with no current passing through the cell till reach to maximum limit defined potential (depending on the properties of the battery).

Current Pulse + OCP Cycle no.2

Then, discharge negative current will be applied for defined duration, followed by relaxation time (OCP). The discharge pulses are repeated until the lower limit potential is achieved (depending on properties of the battery).

From the potential versus time curve, important information like calculation of the diffusion coefficient and thermodynamics parameters can be obtained.

NOTES:

During imposing charge current, the potential increases rapidly which is related to iR drop of system (consiste all uncompensated resistance of system like charge transfer, cables, etc.). Then it will increase slowly with a constant slope. This happens exactly while imposing discharge current too. It means by imposing negative current, the potential drop fast (related to iR drop) flowing by slowly decreasing in potential value by fixed slope versus time (more details on figure 4).



PARAMETERS

This experiment was performed by OrigaMaster software. The parameters of test are shown in figure 1.

| Galvanostatic Intermittent Titration Technique | | Initialization - Cell configuration | | | |
|--|---|---|---|---|--|
| Current Pulse + OCP Cycle no. 1 | | | Electrodes connected with | OGS/OGF/LDS | |
| 🖂 Galvanostatic Pulse no. 1 | | | Connection cell on | 4 electrodes | |
| E Set Current 1 | 220, mA | | E1 input | No | |
| For Duration | 10, min. | | E2 input | No | |
| Open Circuit Potential no. 1 | | | Temperature sensor | No | |
| 1 Duration | 10, min. | | Settings Instruments | | |
| Exit conditionsCycle no. 1 | | | Delay before disjunction (msec.) | 20 | |
| If Pulse Potentiel is | >, 15000 | | Auto ranging delay (msec.) | 200 | |
| 🗄 If Open Circuit Potential | >, 4200 | | Randwidth limit | No | |
| Current Pulse + OCP Cycle no. 2 | | | | NO | |
| Galvanostatic Pulse no. 2 | | | Stopping criteria | | |
| 🗄 Set Current 2 | -220, mA | | Variables initialization | | |
| I For Duration | 10, min. | | A | | |
| 😑 Open Circuit Potential no. 2 | | l r | | | |
| 1 Duration | 10, min. | | | | |
| Exit conditionsCycle no. 2 | | | -11 | | |
| ⊞ If Pulse Potentiel is | >, 15000 | | Start Start | | |
| If Open Circuit Potential | <, 2800 | 1 | | 111 | |
| 📃 Global parameters | | | | | |
| Meas. period (sec.) | 10, sec. | 1 | | | |
| r record every dE (mV) | 5 | 1 ◀┥ | GITT | | |
| tential range | 5V | | | · · · · | |
| nalog Filter | 1 msec. | | · · · · · · · · · · · · · · · · · · · | 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 | |
| gital Filter | 0 | 1 | · · · · · · · · · · · · · · · · · · · | | |
| pen circuit at end | Yes | 1 | End | | |
| rdinate Y2 | Current | 1 | Lind in the | 111 | |
| tit conditions | | | · · · · · · · · · · · · · · · · · · · | | |
| If Charge variation > | 0, mA.h | | | | |
| Max Total Duration | 0, hour | 1 4 | | | |
| | nostatic Intermittent Titratic rrrent Pulse + OCP Cycle no. Galvanostatic Pulse no. 1 Set Current 1 For Duration Open Circuit Potential no. 1 Duration Exit conditionsCycle no. 1 If Pulse Potentiel is If Open Circuit Potential rrrent Pulse + OCP Cycle no. Galvanostatic Pulse no. 2 Set Current 2 For Duration Open Circuit Potential no. 2 Duration Exit conditionsCycle no. 2 Duration Color Circuit Potential no. 2 Duration Circuit Potential no. 2 Duration Circuit Potential no. 2 Duration Exit conditionsCycle no. 2 If Open Circuit Potential obal parameters Meas. period (sec.) record every dE (mV) tential range lalog Filter open circuit at end dinate Y2 it conditions If Charge variation > Max Total Duration | nostatic Intermittent Titration Technique | Image of the second se | nostatic Intermittent Titration Technique irrent Pulse + OCP Cycle no. 1 Galvanostatic Pulse no. 1 | |

Figure 1: Parameters of GITT method on OrigaMaster software

NOTES:

In the « Start » menu, it is suggested to define 4 electrodes configuration if long cables are used for connecting the battery to potentiostat.



PARAMETERS

According to figure 2 as default settings, the method will start in applying galvanostatic charge pulses 220 mA of current by duration of 10 minutes long, followed by 10 minutes of relaxation time, with no current passing through the cell, from started open circuit potential to 4200 mV.

Then, discharge steps are applied. Each negative current pulses -220 mA are applied for 10 minutes long, followed by 10 minutes of relaxation time. The discharge pulses are repeated until the lower limit of 2800 mV is reached.

The other "Exit condition cycle No. " parameter consist of two main limits. "If pulse potential is" correspond to maximum or minimum value of potential of each pulse during imposing current.

"If circuit potential ..." accords to value of OCP during relaxation times which should not passes the limitations.



Figure 2: Default setting of GITT method

RESULTS AND DISCUSSIONS



Figure 3: The whole curve of GITT at the end of the test

Figure 3 shows the curve results by performing complete cycles of GITT on Li-Ion battery.

The diffusion coefficient of electro analyte inside the battery could be calculated through this curve and the related equation.



RESULTS AND DISCUSSIONS

According to following equation, the diffusion coefficient of the battery sample can be determined:

$$\mathsf{D} = \frac{4}{\pi\tau} \left(\frac{\mathsf{I}_0 V}{zFA}\right)^2 \left(\frac{\Delta Es}{\Delta Et}\right)^2$$

While:

 I_0 is the base current (A)

 τ is the duration of the current pulse (s)

V is the molar volume of battery sample (cm³/mol)

Z number of exchanged electron in the reaction

F faraday constant (96485 C/mol)

A is the surface of electrode/electrolyte (cm²)

 ΔEs is the potential change during the relaxation time correspond to the applied current (V)

 ΔEt is the potential change during imposing current pulse (V)

Figure 4 Shows the ΔEs and ΔEt on the curve of GITT while zooming on one step of pulse imposed and following rest step.



Figure 4: Zoom on pulse steps of GITT curve

INSTRUMENT AND ELECTRODES

| Electrode setup | | | |
|-----------------|--|--|--|
| Sample | Emmerich J127 Li-Ion 3.7V, 14 Wh, 2200 mAh | | |
| Instrument | OrigaFlex OGF500 | | |
| Software | OrigaMaster | | |



Figure 5: OrigaFlex OGF500



Figure 6: Li-Ion battery

NOTES:

As in this application note, a commercial battery was used, some information (like molar volume and surface of electrode/electrolyte) were missed to calculate the qualitative parameters. So, it will just explain how to calculate the desired values.

OrigaLys ElectroChem SAS