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Experimental Analysis of Thermal Runaway in 18650 Cylindrical Li-Ion Cells Using an Accelerating Rate Calorimeter

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Abstract: In this work, commercial 18650 lithium-ion cells with LiMn_2O_4 , LiFePO_4 , and $\text{Li}(\text{Ni}_{0.33}\text{Mn}_{0.33}\text{Co}_{0.33})\text{O}_2$ cathodes were exposed to external heating in an accelerating rate calorimeter (es-ARC, Thermal Hazard Technology (THT), Bletchley, UK), to investigate the thermal behavior under abuse conditions. New procedures for measuring the external and internal pressure change of cells were developed. The external pressure was measured utilizing a gas-tight cylinder inside the calorimeter chamber, in order to detect the venting of the cells. For internal pressure measurements, a pressure line connected to a pressure transducer was directly inserted into the cell. During the thermal runaway experiments, three stages (low rate, medium rate, and high rate reactions) were observed. Both the pressure and temperature change indicated different stages of exothermic reactions, which produced gases or/and heat. The onset temperature of the thermal runaway was estimated according to the temperature and pressure changes. Moreover, the different activation energies for the exothermic reactions could be derived from Arrhenius plots.

Keywords: Li-ion cell; thermal runaway; accelerating rate calorimeter (ARC); pressure change

1. Introduction

To eradicate the dependence on petroleum and to reduce CO_2 emissions, the best near-term solutions for vehicles are electric (EV) and hybrid electric vehicles (HEVs). However, in order for them to extensively penetrate the market, one of the most urgent requirements is to develop lithium-ion cells and batteries, which are safe and reliable, even at higher temperatures. Several exothermic chemical reactions can occur inside a cell while the temperature rises. This may generate heat that accumulates inside the cell and accelerates the chemical reaction between the cell components, if the heat transfer to the surroundings is not sufficient. In this case, thermal runaway can occur, resulting in leaks, smoke, gas venting, flames, etc., which may lead to a fire or explosion.

To describe the thermal runaway, the main exothermic chemical reactions have to be identified. According to [1–3], the thermal runaway process can be described by four main steps:

- (1) At $T > T_1$, the solid-electrolyte interface (SEI) decomposes in an exothermic reaction;
- (2) At $T > T_2$, an exothermic reaction between the intercalated Li-ions and the electrolyte starts;
- (3) At $T > T_3$, an exothermic reaction between the positive material and the electrolyte takes place with the evolution of oxygen inside the cell;
- (4) At $T > T_4$, the electrolyte decomposes.

A literature overview is given in Table 1. Abraham et al. [3] presented ARC measurements on 18650 lithium-ion cells to decipher the sequence of events leading to thermal runaway. At 84 °C, self-heating of the cell started. The gases generated in the cell were analyzed by gas chromatography/mass spectroscopy (GC/MS). Jhu et al. [4,5], Chen et al. [6], and Lu et al. [7] studied thermal runaway characteristics (initial exothermic temperature, self-heating rate, pressure rise rate, maximum temperature, and pressure) of Li-ion cells with different state of charge (SOC) with vent sizing package 2 (VSP2). The cathode materials of their tested commercial batteries were LiCoO₂, Li(Ni_{1/3}Co_{1/3}Mn_{1/3})O₂, and LiFePO₄. For all cathode materials, high SOC levels aggravated the exothermic reactions. They also compared the self-heating behaviors of cells with different voltages by the heat of reaction (ΔH), which was the product of mass of a cell, the specific heat capacity of the cell, and the rise of temperature under adiabatic conditions [4–6,8]. In addition, the activation energy and the frequency factor were calculated with the Arrhenius equation. Fleischhammer et al. [8] showed the differences in the safety behavior between un-aged and aged 18650 Li-ion cells by ARC. The cells that were cycled at a low temperature were proven to be the most hazardous, in comparison to the un-aged cells and cells cycled at a high current. Hence, lithium plating leads to a significant increase of heat formation and the ageing history has a strong influence on the safety of the cells. Mendoza-Hernandez et al. [9] used accelerating rate calorimeter (ARC) to study the thermal runaway behavior of LiCoO₂ (LCO) and LiMn₂O₄ cathode materials at different SOCs. The cell with the LiMn₂O₄ cathode material was found to be more thermally stable than the cell with LiCoO₂. In a recent report of the Sandia National Laboratories [10], the effects of cell size on the thermal runaway energetics of the cylindrical cells from 3 to 50 Ah of both LiFePO₄ (LFP) and LiNi_{0.80}Co_{0.15}Al_{0.05}O₂ (NCA) chemistries were studied. The results showed that for both LFP and NCA cells, the normalized heating rate (W/Ah) increases roughly linearly for cells from 3 to 38 Ah, while the normalized total heat released (kJ/Ah) is relatively constant over that cell size range. Selman et al. [11] also used ARC to study the thermal runaway behavior of LiCoO₂ cathode materials at different voltages. The fully charged Li-ion cell had the lowest onset self-heating temperature.

Golubkov et al. [12] ran thermal runaway tests on batteries of LiFePO₄ (LFP), Li(Ni_{0.45}Mn_{0.45}Co_{0.10})O₂ (NMC), and blended LCO/NMC cathode materials by using a pressure-tight and airtight reactor, which was able to collect gases. CO, CO₂, and H₂ represented about 80% of the produced gases, and the others were CH₄ and C₂H₄. The onset temperatures and amount of released gas of LFP, NMC, and LCO/NMC were 195 °C and 50 mmol, 170 °C and 150 mmol, 150 °C and 270 mmol, respectively.

In this work, four kinds of experimental methods were applied to measure the surface temperature of new unaged commercial 18650 cells, the external pressure change due to the released gases, and the internal pressure change in re-sealed cells. For the internal pressure test, a hole was drilled via the tab welding and the housing of the cell, a pressure line was then introduced, and finally, the cell was resealed using epoxy resin. Before each test, the cells were charged to SOC 100. Because this study was focused on the elaboration of the correlation between temperature and pressure using different testing methods, only unaged cells and SOC 100 were considered. The correlation between the pressure and temperature behavior of lithium ion cells can characterize the degree of violence of thermal runaway [4], and is very helpful in defining the threshold, the stages of thermal runaway, and the different reactions which occur during thermal runaway.

Table 1. Literature overview on accelerating rate calorimetry measurements of thermal runaway behavior by cylindrical lithium-ion cells. SOC: state of charge; NCA: lithium nickel cobalt aluminum oxide; LCO: lithium cobalt oxide; NMC: lithium nickel manganese cobalt oxide; LFP: LiFePO₄; LMO: LiMn₂O₄; ARC: accelerating rate calorimeter; VSP: vent sizing package; and BTPV: battery testing pressure vessel.

Author	Cell Type					Calorimeter Type	SOC in % or Voltage in V	Onset Temperature of Self-Heating in °C	Maximum Temperature T_{max} in °C	Heat of Reaction ΔH in kJ	T_{vent} in °C	Maximum pressure P_{max} in Bar
	Manufacturer/Name	Capacity in Ah	Dimensions	Weight in g	Materials (Cathode/Anode)							
Abraham [3]	Quallion LLC, Sylmar, CA, USA/-	1	Cylindrical 18650	-	NCA/graphite	ARC	100	84	-	-	124	-
Jhu [4]	-	2.6	Cylindrical 18650	44.9	LCO/-	VSP2	100	131.5	708.8	18.9	-	18.6
	-	2	Cylindrical 18650	41.7	NMC/-		100	175.4	665.6	14.9	-	11.3
Jhu [5]	Sanyo Electric Co. Ltd., Osaka, Japan/UR18650F	2.6	Cylindrical 18650	45.5	LCO/-	VSP2	4.2	131.3	708.7	19.2	-	13.5
	Sony Energy Devices Corporation, Fukushima, Japan/SE US18650GR			44.5			3.7	172.8	402.6	7.7	3.6	
	Samsung SDI Co Ltd, Yongin, Korea/ICR 18650-26D			44.8			4.2	125.2	654.3	17.9	24.5	
				44.8			3.7	154.8	517.2	12.3	3.1	
	LG Chem Ltd, Seoul, Korea/LGDB218650			46.5			4.2	116.5	683.2	18.6	18.1	
			3.7	152.8	454.8	10.2	4.0					
				4.2	142.5	691.9	18.4	15.8				
				3.7	168.0	407.1	8.0	3.3				
Chen [6]	Sony Energy Device, s. Jhu [5]/SE US18650GR	2.6	Cylindrical 18650	46.5	LCO/graphite	VSP2	0	-	-	-	200	-
							30	175	241	2.5	175	2.9
							50	174	441	10.2	175	4.0
							80	140	638	19.0	175	5.9
							100	122	660	20.6	160	16
Lu [7]	-		Cylindrical 18650	38.15	LFP/-	VSP2	3.3	200.5	217.75	0.47	-	17.2
				32.84			3.6	199.94	243.23	1.02	-	17.89
				44.8			3.7	160.45	280.10	3.91	34.34	
				44.89								LCO/-
Fleischhammer [8]	-	1.5	Cylindrical 18650	-	NCM/LMO carbon/graphite	ARC	Un-aged		-	-	-	-
							97.9					
							High-rate aged					
Low temp. aged		42.1										

Table 1. Cont.

Author	Cell Type				Calorimeter Type	SOC in % or Voltage in V	Onset Temperature of Self-Heating in °C	Maximum Temperature T_{max} in °C	Heat of Reaction ΔH in kJ	T_{Vent} in °C	Maximum pressure P_{max} in Bar	
	Manufacturer/Name	Capacity in Ah	Dimensions	Weight in g								Materials (Cathode/Anode)
Mendoza-Hernandez [9]	-	0.8	Cylindrical 18650	-	LCO/graphite	ARC	0	165.3	-	-	-	-
							25	159.8				
							50	155.1				
							75	140.1				
							100	124.9				
	120	116.5										
	-	0.72	Cylindrical 18650	-	LMO/graphite	ARC	0	-	-	-	-	-
							25	109.9				
							50	104.9				
							75	104.9				
100							104.9					
120	99.8											
Orendorff [10]	-	2.6	Cylindrical	-	LFP	ARC BTPV	-	175	-	19.4	-	0.16
		3.4			NCA			160		31.1		0.45
		20			LFP			235		150		1.3
		27			NCA			110		254		4.1
		30			LFP			250		272		-
		38			LFP			233-246		289		1.8
Selman [11]	Sony Energy Device, s. Jhu [5]/US18650		Cylindrical 18650	-	LCO/carbon	ARC	2.8 4.06	144 104	-	-	-	-
This work	-	1.65	Cylindrical 18650	43.4	LMO	ARC	100	91	303	7.8	-	
		1.1		39.7	LFP			90	259	7.3		
		2.2		41.7	NMC			91	731	24.9		

2. Experimental

For commercial 18650 lithium-ion cells with LiMn_2O_4 (LMO), LiFePO_4 (LFP), and $\text{Li}(\text{Ni}_{0.33}\text{Mn}_{0.33}\text{Co}_{0.33})\text{O}_2$ (NMC) cathodes, four different thermal runaway testing methods have been applied in an accelerating rate calorimeter (es-ARC, Thermal Hazard Technology (THT), Bletchley, UK). The calorimeter chamber with a diameter of 10 cm and a height of 10 cm has one heater and one thermocouple located in the lid and in the bottom; and two heaters and thermocouples (all type N) in the side wall. In order to ensure comparable starting conditions, all tested cells were cycled three times with the constant current constant voltage (CCCV) method, which means charging at a constant current with a rate of 1C, followed by charging with the maximal voltage until the current decreases to C/10. Afterwards, the cells were discharged with a 1C rate until the minimal voltage was reached (see Table 2). Finally, before the ARC test, the cells were charged to SOC 100. Following this, the tested cell was fixed on the lid of the calorimeter, as shown in Figure 1. The calorimeter temperature is controlled by a main or so-called bomb thermocouple attached to the surface of the cell. The heaters work until reaching the required temperatures regarding the measurement conditions and modes. An external steel casing serves as protection.

Table 2. Specifications of 18650 Li-ion cells with different cathode materials.

Cathode Material	LiMn_2O_4	LiFePO_4	$\text{Li}(\text{Ni}_{0.33}\text{Mn}_{0.33}\text{Co}_{0.33})\text{O}_2$
$U_{\min}-U_{\max}$ in V	2.5–4.2	2.0–3.6	2.75–4.2
Nominal capacity in mAh	1650	1100	2200
Cycle life	1000	>1000	>300
Weight in g	43.4	39.7	41.7
Operating temperature in °C	−20–60	−30–60	−20–60

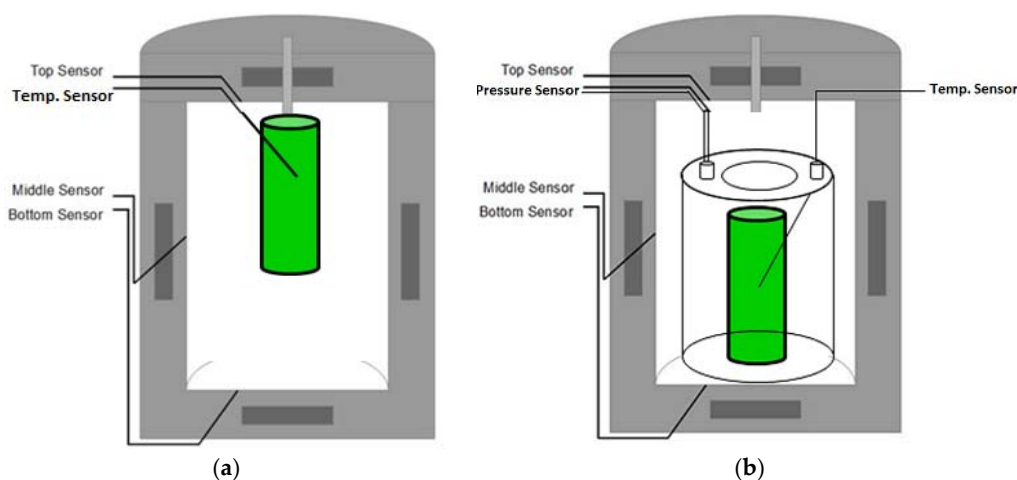


Figure 1. Setup of the cell in the ARC: (a) normal and internal pressure measurement; and (b) external pressure measurement.

2.1. First Method (Only Temperature Measurement OT)

In the first method, only the temperature and temperature rate were recorded (Figure 1a). The standard heat-wait-seek (HWS) method was applied for all measurements. This method allows an investigation of the thermal abuse of stored cells by excessive external heating, while they are neither charged nor discharged. The procedure and the recorded data are shown in Figures 2 and 3, respectively. The HWS method starts at 25 °C in the heat mode, by heating up the cell in small temperature steps of 5° C (see Figure 3). At the end of each temperature step, the wait mode is activated for 15 min, to reach thermal equilibrium. After reaching thermal equilibrium, the system enters Seek Mode, which monitors the temperature rate and presents two possible modes—exotherm

mode or heat mode. The decision is related to the selected Temperature Rate Sensitivity. If the measured temperature rate is larger than this value, the system goes into Exotherm Mode. This mode provides an adiabatic environment, which means that no heat exchange between the sample and the surrounding space exists, so that the heat energy of the reactions can be monitored by the measured temperature. On the other hand, if the temperature rate is smaller than the value, the system goes back into Heat Mode. If the temperature exceeds the given final temperature T_f (250 °C), the heaters completely shut down and cool down the system by introducing pressurized air into the calorimeter chamber.

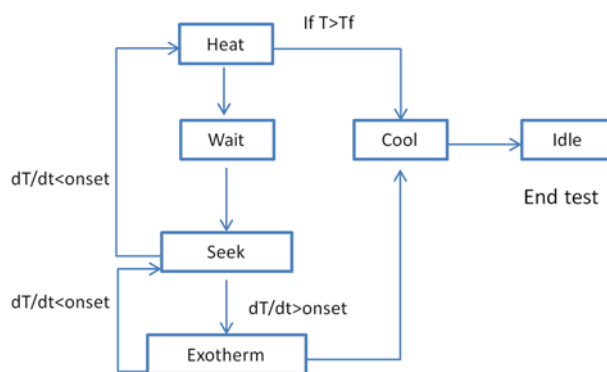


Figure 2. Process of the heat-wait-seek (HWS) method.

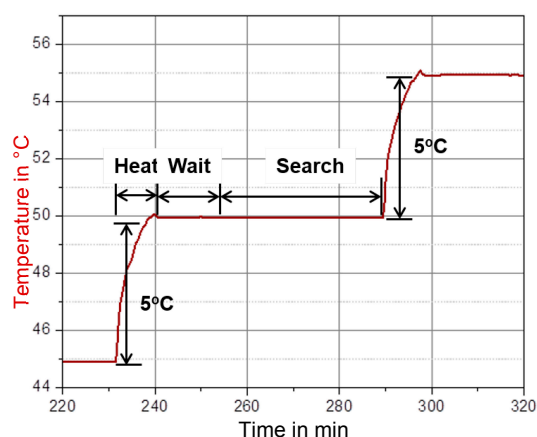


Figure 3. Example of cell surface temperature curve during HWS experiment in an ARC.

2.2. Second Method (External Pressure Measurement EP)

For the second method, an additional cylinder chamber was used in external pressure measurements (see Figure 1b). The thermocouple was introduced through a well-sealed hole and a pressure line connected with a pressure sensor was screwed onto the cylinder chamber. It was critical to screw every part of the cylinder chamber extremely tightly, to prevent leaks. Similar to the adiabatic environment, the system is only capable of monitoring the external pressure change if it is under an air-tight condition.

2.3. Third Method (Internal Pressure Measurement IP)

In contrast to the set-up of the second method, where the capillary leading to the pressure sensor was connected to the cylinder, in the third method, the capillary was directly introduced into the functional cell for internal pressure measurements (Figure 4). Therefore, a suitable point, which allows the capillary to pass through without disturbing the functionality of the cell, had to be found with the help of X-ray tomography images.

X-ray tomography was conducted in the X-ray Imaging Laboratory at the Karlsruhe Institute of Technology (KIT), Institute for Applied Materials-Applied Materials Physics (IAM-AWP). X-ray micro-tomography scans were performed on the cell using a Phoenix v/tome/x s system (Phoenix X-ray GE Sensing & Inspection Technologies GmbH, Wunstorf, Germany). The battery was rotated 360° around its long axis, whilst 1600 projections were captured with the microfocus tube operating at 200 kV and 370 (230) μ A. The sample and detector were positioned in projection magnification, providing an effective voxel size of 75 (48) μ m for the high resolution imaging. Tomography scans of the full cell were obtained, and a subsequent 3D analysis of the reconstructed image data was performed using VG studio max software 3.0, Volume Graphics Company, Heidelberg, Germany. For instance, Figure 4 shows a 3D rendering of the tomographic dataset for the 18650 cell with an LMO cathode. The spiral layer structure of the cell and the hollow central section are clearly revealed, as well as the safety valve on the top. It was easy to draw the conclusion that the best position for the pressure line is at the bottom center. The reasons for this are:

- (1) The hollow is big enough for inserting a 1/16-inch diameter capillary.
- (2) Inside the battery, the hollow section is located exactly in the center; nothing will be damaged when drilling a hole through the anode tab welding.

Then, the cells were resealed with epoxy resin. The described preparation procedure was conducted in an Argon-filled glovebox, because the cell components are very sensitive to moisture and oxygen.

Furthermore, the cells were cycled three times, and discharged completely to introduce the capillary. After resealing, the cells were cycled three times, and finally charged to SOC 100. This shows that, even if the anode tab was damaged by the drilling, the cell was still functional after this procedure.

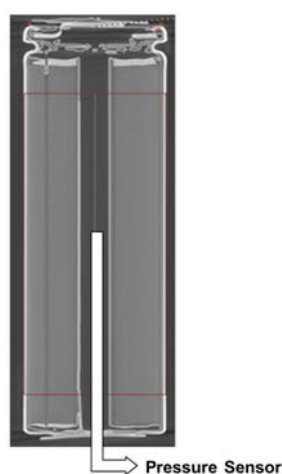


Figure 4. Setup for internal pressure measurement.

2.4. Fourth Method (Ramp Heating Method (Hotbox Test))

For the “ramp heating” method, the cells have been set up in the same way as described before for the HWS tests. However, instead of a stepwise heating of the cells, they were heated up continuously from 30 °C to 250 °C in the ARC, at a constant rate of 5 °C/min. This mimics a hot box test. For this purpose, the measurements were operated in “ramp mode”, with a high sensitivity value of 6 °C/min to prevent automatic switching into the “exotherm mode”. The system was used under adiabatic conditions, as in the other tests.

2.5. Cells Preparation

Before the thermal runaway tests, the capacity of the cells was determined by three full cycles, and finally, they were fully charged to state of charge 100% (SOC 100), because this is the most critical

condition for thermal runaway according to [7]. The method of charge was CC/CV. This means that the cell is initially charged at a constant current at a 1C rate (CC) until the maximal voltage, and is then charged at a constant voltage (CV) with a decreasing current, until the current reaches 10% of its starting value.

3. Results and Discussion

3.1. Results for 18650 Cells with LiMn_2O_4 Cathode by Using Different Methods

In the following, the first three test methods are exemplarily demonstrated for the cell with the LMO cathode in the temperature range of 25 °C to 250 °C. Figure 5 compares the time evolution of the measured surface temperatures for the three different methods with three identical cells at SOC 100. From this figure, it is easy to distinguish the heat-wait-see periods from the exotherm periods. At temperatures below about 80 °C, the temperature curve increased with steps of 5 °C. Then, a first exothermic reaction can be observed at 91 °C, where the self-heating rate is higher than 0.02 °C/min. Following this, the system switched back to heat mode at 100 °C, and the HWS mode was active until 110 °C. Finally, the temperature continuously increased in exotherm mode, until thermal runaway occurred.

Therefore, three stages were observed in the thermal runaway tests, which, according to the literature [1–3], were assigned according to:

- (1) At about 90 °C, the SEI decomposed.
- (2) At about 110 °C, an exothermic reaction between the embedded Lithium ions and electrolyte started, which resulted in the reduction of the electrolyte at the negative electrode.
- (3) At about 200 °C, an exothermic reaction between the active material and electrolyte at the positive electrode took place.

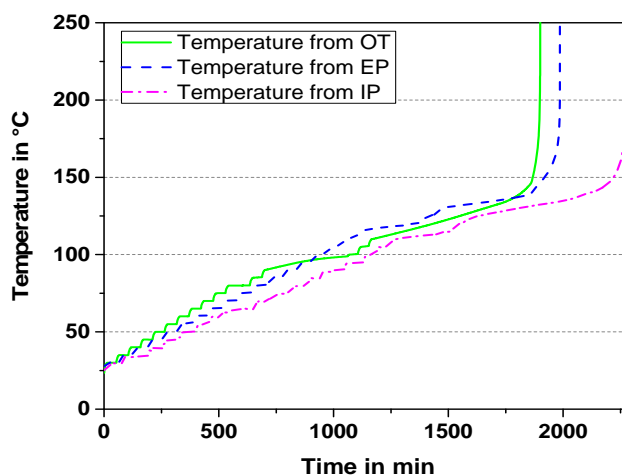


Figure 5. Comparison of temperature versus time curves measured with different methods.

When comparing the three curves, the time period in the HWS mode before switching into exotherm mode, increased from the only temperature via the external up to the internal pressure measurements. The reason for this is that, for the external pressure tests, the system needs to heat up the cylinder, and that for the internal pressure tests, some heat loss occurs via the metallic pressure line. Even though the three curves do not perfectly overlap, the tendencies are the same. Figure 6, which shows the temperature rate vs. the temperature, gives us a more specific view. From this figure, the three stages can be distinguished more easily. In the temperature region from 90 °C to 130 °C, the temperature rate is lower than 0.1 °C/min. From 130 °C to 200 °C, the energy was released with a medium rate which was lower than 25 °C/min. Above 225 °C, the exothermic reactions were violent,

with a temperature rate over 100 °C/min, and could reach over 1000 °C/min. These results indicated the degree and the destructive development of the exothermal reactions.

In Figure 7, the HWS method without pressure measurements (see Figure 7a) is compared to the ramp heating method (Figure 7b). From the latter, less information could be extracted: it can be seen that the temperature increased linearly at the beginning, as shown in Figure 7b, and that the inflection point is visible at about 150 °C, which could also be observed in the curve of temperature rate vs. temperature (see Figure 6). Comparatively, the HWS Mode is relatively sensitive and reveals the entire process of the thermal runaway.

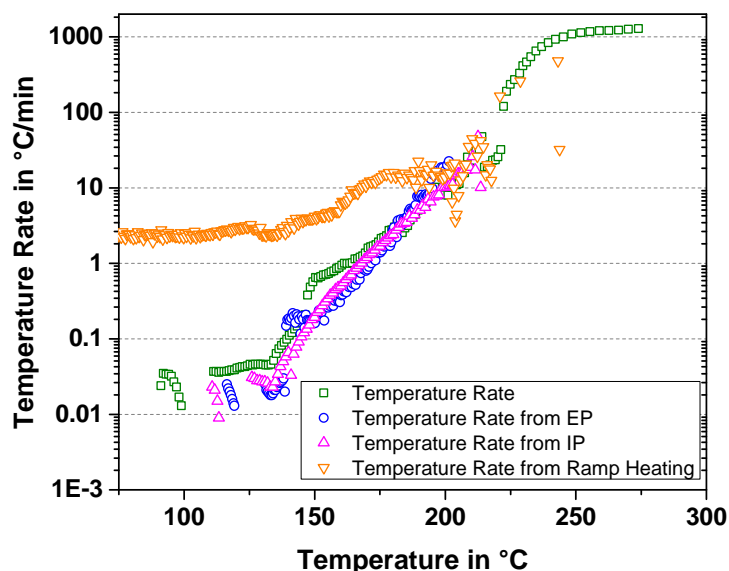


Figure 6. Comparison of temperature rate versus temperature curves measured with different methods.

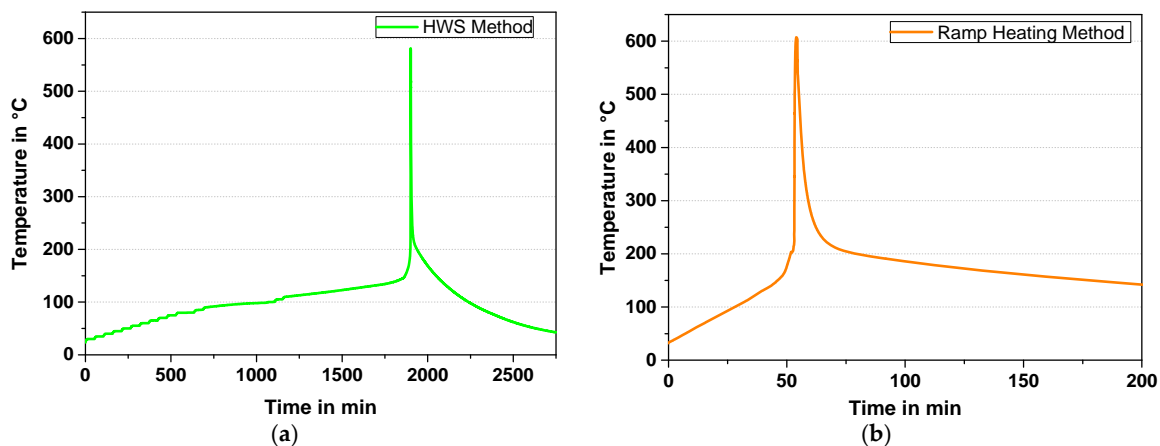


Figure 7. Comparison of temperature versus time curves: (a) HWS Method; and (b) ramp heating method.

In Figure 8, the pressure vs. time curves for the external and internal pressure tests are compared. In the external pressure curve (blue line) at about 130 °C, a small pressure rise can be observed, which marks the beginning of air leaking out of the cell. At the end of stage 2, it could be deduced that the battery had opened and gases were released. The external pressure increased continuously and dramatically until 21 bar at stage 3, matching the temperature change very well. From the internal pressure curve (magenta line), we can see that the internal pressure was increasing steadily during stage 1 and 2. In the transition from stage 2 to 3, the cell vented and gases were released, which was

indicated by a rapid internal pressure drop. The external pressure measurements and internal pressure measurements were different measurements that were operated separately, they do not represent parallel measurements on the same cell. Therefore, the time when the venting starts is different.

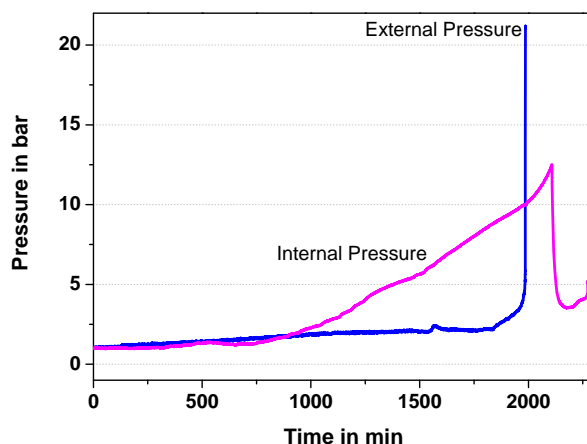


Figure 8. Comparison of pressure curves vs. time with different methods.

Finally, in order to correlate the temperature and pressure measurements, the pressure values measured from the external and the internal pressure tests are plotted vs. the temperature, as can be seen in Figure 9. For both the external and internal pressure curves, the inflexion occurred at 130 °C, where the cells vented produced gases to reduce the internal pressure. At the same time, it caused an increase of the external pressure. The additional red line represents the expansion of the air in the additional cylinder chamber with increasing temperature according to the ideal gas law. As long as the cell is not venting, the ideal gas law should be a good approximation. For the external pressure measurement, the difference between the measured blue line and the red line can be regarded as the amount of gas that is released from the leaking cell. As the temperature increased, the difference became larger, indicating that more gases were produced. There was a gas leak from 130 °C and produced gases were released into the cylinder, due to exceeding the threshold pressure of the safety valve. At 200 °C, which was also the start temperature of stage 3, the difference was significant. For the internal pressure measurement, starting at 80 °C, the difference continuously increased, indicating the beginning of gas production and SEI layer decomposition. After cooling, which is not shown here, the pressure fell below 10 bar again, which is the upper limit for the validity of the ideal gas law. Thus, according to Golubkov et al. [12], the ideal gas law equation can be used in order to calculate the amount of gas produced:

$$n_0 = (p_0 \cdot V_0) / (R \cdot T_0) \quad (1)$$

$$n_1 = (p_1 \cdot V_0) / (R \cdot T_1) \quad (2)$$

where n_0 is the original air amount in the cylinder, p_0 is the pressure of the original air, T_0 represents the temperature of the original air, V_0 represents the volume of the cylinder, and R is the gas constant, taken as $8.316 \text{ J} \cdot \text{K}^{-1} \cdot \text{mol}^{-1}$. Similarly, n_1 represents the actual amount of gas measured during the measurements, p_1 is the pressure of the actual gases, and T_1 represents the temperature of the actual gases. From the parameters, depth: 72 mm, outer diameter: 82 mm, and thickness: 5 mm, the volume is calculated: $2.73 \times 10^{-4} \text{ m}^3$. At the beginning of the measurement, the temperature and pressure in the cylinder were 27.83 °C (300.98 K) and 1.11 bar ($1.11 \times 10^5 \text{ Pa}$), which gives $n_0 = 12.05 \text{ mmol}$. At the end of the measurement, the thermocouple and pressure sensor detected that the sample temperature was 35.0 °C (308.15 K) and the pressure in the cylinder was 8.31 bar ($8.31 \times 10^5 \text{ Pa}$), producing $n_1 = 88.53 \text{ mmol}$. Thus, the amount of gas produced during thermal runaway is (88.53–12.05) mmol, which is 76.48 mmol.

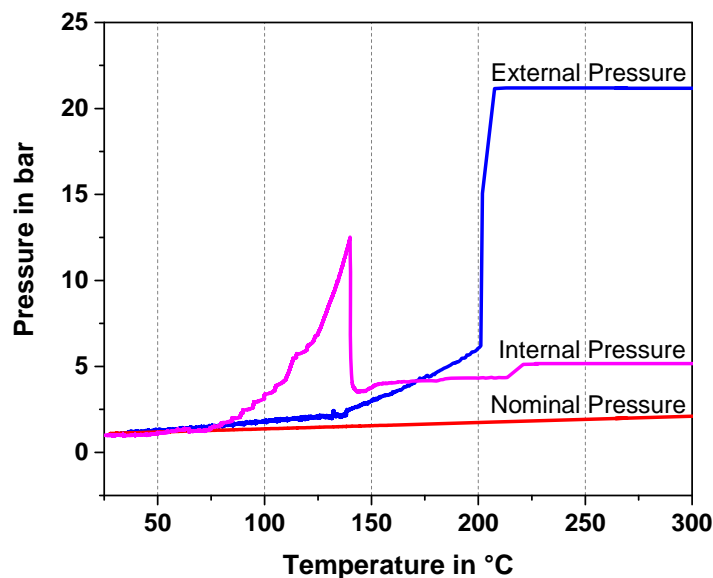


Figure 9. Comparison of pressure vs. temperature curves with different methods.

A similar procedure was described by Golubkov et al. [12] and the amount of gas produced was calculated from the measured external pressure. The cell with the LMO cathode released 76.5 mmol gas during thermal runaway. Golubkov et al. reported that 18650 cells with LCO/NMC cathode material released 265 mmol of gas, and those with pure NMC released 149 mmol. This comparison indicates a better exothermic stability of the LMO cathode.

3.2. Results for 18650 Cells with Different Cathode Materials

In the market, there are many kinds of batteries, varying from geometry to material. The most common structures of cathode materials are spinels, layered oxides, and olivines. Not only their electrical performance, but also their thermal properties, are quite different. Thus, every kind of these three active materials should be investigated and compared in “thermal runaway” studies [13]. The second part of this work focuses on a comparison of their thermal behavior in cylindrical 18650 cells. Their specifications are listed in Table 2. LMO, LFP, and NMC are chosen as examples for spinel, olivine, and layered structures, respectively, in this work.

Figure 10 shows the thermal behavior of 18650 cells during “thermal runaway” measurements. In all tests, the cells were heated between 30 °C and 250 °C with 5 °C increments in the search for self-heating at the sensitivity threshold of 0.01 °C/min. In the plot temperature vs. time, all of the onset temperatures of the exothermal reactions were similar, at about 90 °C. A comparison of the maximum temperatures gives the highest value of 731 °C for the NMC cathode, and a value of only 259 °C for the LFP cathode. One reason for this is the difference between the capacities, i.e., the nominal capacity of NMC (see Table 2) is twice that of LFP. Because the nominal capacity of LMO lies in between, its maximum temperature is 581 °C. In terms of the length of heating time which was required to activate the exothermal reactions, NMC and LFP took significantly longer than LMO.

In the temperature rate vs. temperature plot shown in Figure 11, the three different stages of thermal runaway mentioned before can be observed and distinguished. The LFP cell never entered stage 3, with temperature rates staying below 3 °C/min. In contrast, the LMO cell reached a maximum rate of 1285 °C/min and the NMC cell reached 7580 °C/min. In the temperature region between 80 °C and 130 °C, the SEI decomposed. The temperature rate of LFP was the lowest among these three materials, while NMC and LMO were in the same range. The much lower maximum temperature rate of LFP can not only be attributed to the lower capacity, but also to the much higher thermal stability of this nanoscale material.

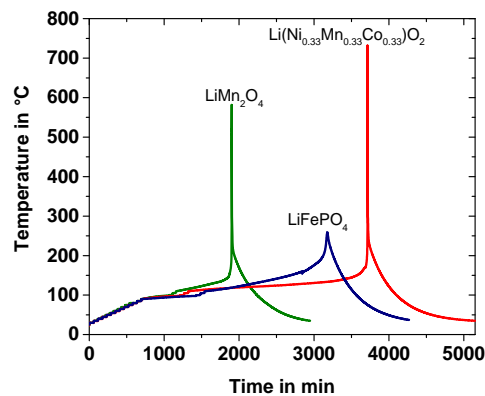


Figure 10. Temperature versus time curves for 18650 cells during HWS tests in an ARC with different cathode materials.

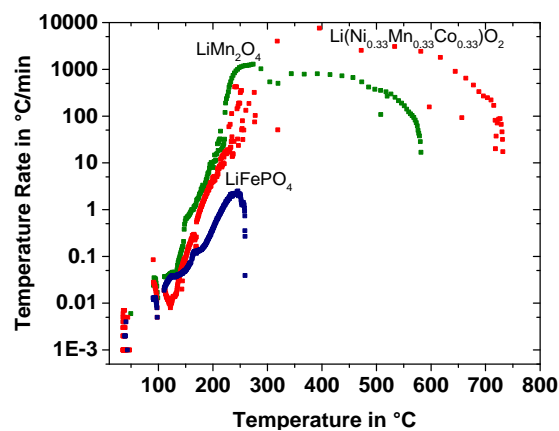


Figure 11. Temperature rate versus temperature curves for 18650 cells during HWS tests in an ARC with different cathode materials.

According to the simplified equation for calculating the thermokinetic parameters proposed by Jhu et al. [4,5], which is based on the Arrhenius law, the temperature rate or self-heating rate can be expressed as follows:

$$\ln(dT/dt) \approx \ln(\Delta T_{ad} \cdot A) - E_a/(k_B \cdot T) \quad (3)$$

When the ARC is operating under adiabatic conditions in exotherm mode, there is no heat loss to the surroundings. Thus, we can consider that the temperature rise is only caused by the self-heating reactions in the cell and the mass specific heat of the reaction (J/g) for each stage can be calculated with the following equation:

$$\Delta H/m_{cell} = c_p \cdot \Delta T_{ad} \quad (4)$$

where A is the pre-exponential factor (min^{-1}); ΔT_{ad} is the adiabatic temperature rise (K), i.e., the difference between the initial exothermic temperature T_0 and the maximum temperature T_{max} ; E_a is the activation energy (eV); k_B is Boltzmann's constant ($8.62 \times 10^{-5} \text{ eV} \cdot \text{K}^{-1}$); m_{cell} is the cell mass (g); and c_p is the heat capacity ($\text{J/g} \cdot \text{K}$). Thus, by plotting the natural logarithm of the self-heating rate versus the inverse of temperature, as shown in Figure 12, the activation energy and frequency factor of the different cells can be calculated. The data obtained from this Arrhenius plot are listed in Table 3. Li-ion cells with an LFP cathode have the lowest T_{max} and $(dT/dt)_{max}$, while cells with the NMC cathode show the highest T_{max} and $(dT/dt)_{max}$. For calculating the heat of the reaction using Equation (4), the heat capacity was measured as described in Schuster et al. [14]. The listed heat of the reaction is the sum of the heats of the reactions of the different stages. The heats of the reactions are in good agreement with the results from the literature [2,15] as are the activation energies [5].

The next step could be to use the derived thermokinetic parameters as input parameters for our coupled electrochemical thermal model [16]. This model is based on the porous electrode theory of Newman and Tiedemann [17] and has been extended with a simple combustion model emerging from reaction kinetics, including various types of heat sources based on an Arrhenius law.

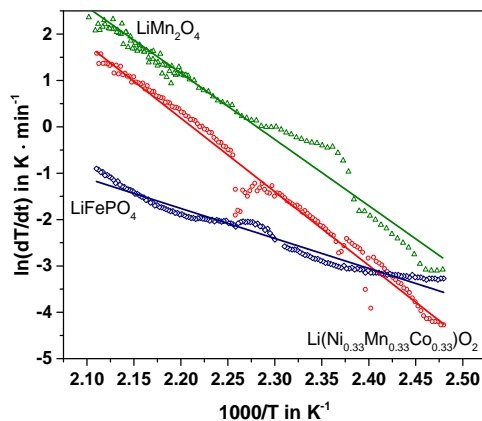


Figure 12. Natural logarithm of the self-heating rate versus inverse of temperature curves for 18650 cells with different cathode materials. The three lines are linear fits according to the Arrhenius law.

Table 3. Thermal runaway test results of 18650 Li-ion cells with different cathode materials.

Cathode Material	LiMn ₂ O ₄	LiFePO ₄	Li(Ni _{0.33} Mn _{0.33} Co _{0.33})O ₂
Onset temperature of self-heating in °C	91	90	91
T_{\max} in °C	581	259	731
$(dT/dt)_{\max}$ in °C/min	1285	3	7577
c_p at 60 °C SOC 100 in J/g·K	0.83	1.19	0.95
E_a in eV	1.24	0.56	1.37
Reaction heat in J/g	391	184	597
Reaction heat in J/g	350–640 [2,15]	260 [15]	600 [15]

4. Conclusions

Three stages have been observed in thermal runaway tests in 18650 Li-ion cells with an LMO cathode by correlating temperature and pressure curves. The first stage began with the production of gas at about 90 °C. The reaction in the first stage was likely to be the SEI layer decomposition. The onset temperature of thermal runaway, which means the temperature at which the self-heating reaction starts, thus leading to thermal runaway, was about 110 °C. The amount of gas produced was 76.48 mmol in thermal runaway. In future studies, it could be interesting to confirm these stages and analyze the gas produced and the material components after thermal runaway, using post-mortem analysis by opening the cells that were frozen at the different stages. Furthermore, the effects of different SOCs and of ageing could be studied. In addition, the developed pressure tests could be applied to cycling in the ARC under adiabatic conditions, in order to study the effect of different charging and discharging rates.

For 18650 Li-ion cells with different cathode materials, the thermal runaway tests confirmed that cells with an LFP cathode exhibited a good stability under thermal abuse, while cells with an NMC cathode presented a poor temperature tolerance at high temperatures.

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