



Article Coupled Electrochemical-Thermal Simulations and Validation of Minichannel Cold-Plate Water-Cooled Prismatic 20 Ah LiFePO₄ Battery

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Abstract: This paper discusses the quantitative validation carried out on a prismatic 20 Ah LiFePO₄ battery sandwiched between two minichannel cold-plates with distributed flow having a single U-turn. A two-way coupled electrochemical-thermal simulations are performed at different discharge rates (1–4 C) and coolant inlet temperatures (15–35 °C). The predicted battery voltage response at room temperature (22 °C) and the performance of the Battery Thermal Management System (BTMS) in terms of the battery surface temperatures (maximum temperature, T_{max} and temperature difference, ΔT) have been analyzed. Additionally, temperature variation at ten different locations on the battery surface is studied during the discharge process. The predicted temperatures are compared with the measured data and found to be in close agreement. Differences between the predicted and measured temperatures are attributed to the assumption of uniform heat generation by the Li-ion model (P2D), the accuracy of electrochemical property input data, and the accuracy of the measuring tools used. Overall, it is suggested that the Li-ion model can be used to design the efficient BTMS at the cell level.

Keywords: Li-ion battery; BTMS; minichannel cold-plates; C-rates; coolant temperature; COM-SOL software

1. Introduction and Literature

Global issues such as air pollution and the energy crisis have led to the rapid development of electric vehicles (EVs) in the past decade and a half, and Li-ion batteries (LIBs) are more widely used in EVs than other battery types (Pb-acid, Ni-Cd, Ni-MH). However, the challenges faced with LIBs include the battery cycle life, power, adaptability, and fast charging, which are primarily dependent on the process parameters and operating temperature. To design the suitable BTMS, it is crucial to know the perfect operating conditions for the batteries to work in a highly efficient manner. In recent times, it was shown that the performance of LIBs is affected by the operating temperatures, and hence, an optimum temperature range of 15 to 35 °C was suggested [1]. At these temperatures, the LIBs are at their maximum capacity and there are minimal degradation effects. For instance, an experimental test showed a 22.5% increase in discharge time for a constant current discharge at 25 °C compared to 50 °C [2]. Different types of Li-ion batteries are used in EVs and they require different thermal cooling designs for an effective thermal management. Additionally, designing an appropriate battery thermal management system (BTMS) design requires a detailed understanding of heat generation, transportation, and



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Copyright: © 2021 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https:// creativecommons.org/licenses/by/ 4.0/). dissipation at a cell level. The paragraphs below discuss the researchers' recent experimental and theoretical works on the heat generation rates and thermal responses of different LIBs operated at different process and cooling conditions.

The heat generation rate by a stack of four Li-ion titanate (10 Ah, 2.3 V) batteries measured at different discharge rates (1–4 C) and ambient temperatures (28–42 °C) shows that the discharge C-rate has a more significant effect than the ambient temperature [3]. The irreversible and reversible heat generations at different discharge rates (1–5 C) of the 10 Ah LiFePO₄ battery revealed that the former is relatively more stable than the latter [4]. The measured heat generation rate of 8 Ah LiMn₂O₄ battery during charge–discharge cycles taking into account the effect of operating load, SOC, temperature, and aging have shown that the joule heating rises with a decrease in the battery operating temperature, while the heat of reaction does not differ notably [5]. The heat generation rates obtained from overpotential and entropic heats with negligible heat of mixing in a pouch type 45 Ah LiFePO₄ battery have shown that the heat generation is higher during discharge than the charge, and hence an effective cooling system is necessary to prevent thermal runaway at C-rates [6]. The electrochemical-thermal model predicted that the voltage and temperature of LiFePO₄ battery were compared with the measured data and found that the reaction heat contributes more, followed by the contact resistance heat and the Joule heat [7]. However, the net heat generation rate increases with the decrease in battery ambient temperature. The measured heat generation rate of a prismatic 20 Ah LiFePO₄ cell at different ambient conditions (-10 to 40 °C) and discharge C-rates (0.25–3 C) have shown that the heat generation increases with the decrease in operating temperature [8]. Furthermore, the heat generation rate rises first, then a plateau region, and then further increases with depth-of-discharge (DOD). The heat generation rate of a cylindrical 2.6 Ah LiFePO₄ cell was predicted at different C-rates based on the measured battery surface temperature and surface heat flux and using a simple energy balance [9]. The heat generation rates of 19.5 Ah LiFePO₄ cell have shown that the overpotential and voltage difference methods predicted similar results at all SOCs and C-rates [10]. The heat generation rate of prismatic 40 Ah Li-ion NMC cells measured under adiabatic and constant ambient test conditions showed an overall exothermic behavior with the adiabatic test, but an exothermic behavior during discharge and endothermic behavior during charge with ambient test [11].

After understanding heat generation, it is important to understand its effect on the battery in terms of temperature response. The measured heat generation rates of a battery module of cylindrical cells (18650) of Li-ion NiMnC cathode were used in the thermallumped treatment of each cell [12], and the effects of discharge/charge rate, liquid flow rate, and heat exchange contact surface area were analyzed. A coupled electrochemical-thermal model developed using COMSOL software for a 4 Ah Li-ion NCA/graphite battery has shown that the maximum temperature is in the proximity of tabs and positive current collector and is due to the high ohmic heat generation [13]. Additionally, the absence of tabs resulted in a more uniformity of battery temperature at the bottom. A review of the battery models and thermal management methods has suggested that the Peukert's equation can be used instead of the more complex electrochemical model [14]. An analytical model based on predicted temperature distribution in 1.5 Ah Li-ion NMC (INR18650-15L) batteries subjected to charge-discharge cycles with a rest period has shown a significant temperature difference between the core and surface at the end of 50 cycles [15]. A numerical study on the response of a commercial cylindrical LiFePO₄/C battery using a 2D axisymmetric electrochemical-thermal model at different discharge C-rates and controlled ambient conditions has shown a good match between the predicted and measured discharge voltage and battery surface temperatures [16]. The thermal behavior and evolution of electrochemical reactions in a 20 Ah LiFePO₄ prismatic battery with the ambient temperature at 25 °C were studied using single and multi-cell approaches and at different discharge C-rates [17]. The predicted voltage and thermal responses were compared with the measurements and it was stated that the cooling system is indispensable to maintain the battery surface temperature within the permissible range.

Now, it is important to understand how to control the temperatures achieved after charge/discharge and which method to use in order to reduce the battery temperatures. A review on the performance of battery pack systems with different liquid coolants (water/oil/liquid-metal/nano-fluid) and coolant configurations (internal/external) was discussed [18]. A review presented on the effect of ambient temperatures and cooling technologies for battery thermal management stated that water-cooling had been widely researched and used in BTMS, and cold-plates are used where there is a space constraint [19]. The study on the effectiveness of air and liquid cooling of Li-ion battery packs by comparing the overall heat transfer coefficient of the respective designs has concluded that water-cooling is three times more effective than air for the same rise in battery surface temperature [20]. Single prismatic 20 Ah LiFePO₄ cells sandwiched between cold-plates with a single channel and multiple turns were experimentally studied at different C-rates (1-4 C) and coolant inlet temperatures (5-35 °C), and it was found that the maximum battery surface temperature increases with the increase in C-rate as well as coolant inlet temperature [21,22]. A drive cycle test with a battery pack of three 10 Ah LiFePO₄ cells in series and sandwiched between cold-plates was conducted at different C-rates (1-4 C) and coolant inlet temperatures (10–40 °C) and found the maximum battery surface temperature below the optimum range at all C-rates [23]. A battery pack of 30 cylindrical cells (18650 Li-NCA/C) was simulated using a coupled electrochemical-thermal model and found that the surface temperatures doubled when the coolant flow rate reduced to half [24]. A conjugate heat transfer model developed and used for a pouch type Li-ion battery pack with coolant channels between the batteries has concluded that the coolant flow rate significantly affects the maximum surface temperature at higher C-rates than at lower C-rates. However, the authors assumed an initial temperature profile at the solid-liquid interface rather than calculating the heat generation rate during the discharge process [25]. A simulation study performed on the effectiveness of liquid cooling of 55 Ah Li-ion battery with different configurations (1 \times 4 to 4 \times 4 channels), flow rates (0.2 to 1 L/min), and flow direction has shown increased cooling performance with the increase in the number of channels as well as with all inlets and outlets on one side of the battery [26]. A simulation study performed on cylindrical Li-ion batteries (18650) with coolant pipe placed between the spiral fins has shown a significant drop in maximum battery temperature and temperature difference across the battery by varying the fin height [27]. A 3D thermal model study on a battery module of 7 Ah Li-ion batteries sandwiched between aluminum cold-plates and discharged at a 5 C discharge rate has revealed that the coolant flow direction towards the electrode showed the best performance, and the coolant mass flow rate has a significant effect on the battery temperatures up to a particular value only [28]. The electrochemicalthermal models developed for prismatic 20 Ah LiFePO₄ cells using COMSOL Multiphysics software have shown that the multi-layer approach-based model predictions are in very good agreement with the experiments than the single layer approach [29]. A review comparing uncoupled and coupled battery models stated that the lack of precise experimental data of several parameters in real-time conditions and simplified assumptions such as constant and uniform properties made the electrochemical model (P2D) qualitative [30]. Therefore, the predicted peak temperature and the temperature difference cannot match the measured data. However, the P2D model parameters were taken from the literature as well as physical testing of the battery electrodes. The authors suggested that the model could be scaled up to simulate a battery pack.

It can be noted from the above-detailed literature survey on the experimental and theoretical research works carried out that there are interesting numerical works based on liquid cooling of Li-ion batteries. Nevertheless, the validation is carried out chiefly in terms of voltage profile and maximum temperature of the battery. The models developed also need to be tested based on temperature variation with time at different locations on the battery surface, apart from average battery temperature and heat flux. The primary motive of this research is to bridge this gap. In the current research, a prismatic LiFePO₄ battery with a nominal capacity of 20 Ah sandwiched between cold-plates with multiple channels containing water flow (i.e., distributed flow) is being considered to investigate the battery thermal response at different discharge C-rates and coolant inlet temperatures. The temperature variation with discharge time is plotted at 10 different locations across the principal surface of the battery and is validated with experimental works to test the reliability of the model parameters and also the model itself. A transient, coupled electrochemical-thermal model has been developed using COMSOL Multiphysics software. The electrical and thermal responses predicted in terms of the voltage-time response, battery surface heating, peak temperatures (hot spots), and temperature difference across the battery have been analyzed and validated with the measured data.

2. Experimental Test Bench for Battery with Cold-Plates

As part of a detailed study and quantitative validation of the model predictions, experiments were conducted for voltage and temperature responses by the Battery Thermal Management System (BTMS) during the charge-discharge cycle. A test bench set-up developed for EVs at the University of Waterloo, Canada (as shown in Figure 1) consisting of a Battery Management System (BMS) for charge/discharge process and Thermal Management System (TMS) for heat removal from the prismatic 20 Ah LiFePO₄ battery was utilized along with the measured temperature data at selected locations across the principal surface of the battery. Different tools used include a Lambda ZUP20-40-800 for low-voltage supply, a TDI Dynaload 232 50-150-800 for load (current), a MotoTron controller to monitor the voltage, current, and temperature. A separate monitor for LabVIEW and Keithley tools with accessories were also used. Sensors (T-type and K-type) were used for the temperature measurements on the battery principal surfaces, tabs, cold-plates, and at minichannel inlet and outlet. Heat flux sensors were also used. A flow meter was installed at the inlet of the cold-plate(s). The complete assembly (battery and the cold-plates) was sandwiched using 12.7 mm thick polycarbonate sheets, as shown in Figure 1. Spacers with equal size of the battery are provided to change the distance between polycarbonate and cold-plate for the inlet/outlet fittings. For the whole test, charging was carried out at constant current constant voltage (CCCV) and discharging was carried out at constant current. To ensure proper insulation of the test set-up from its surrounding atmosphere, a compression rig was created consisting of two transparent 12.7 mm thick polycarbonate sheets enclosing the battery with cold-plates. Bolts were used to tighten the assembly. For further details on the conduct of experiments, see the thesis reports published by Waterloo University [31,32].



Figure 1. (a) Battery with cold-plates and apparatus (b) schematic of line diagram.

3. Mathematical and Numerical Modelling

The overpotential heat, entropy change heat, and enthalpy of mixing heat are the three primary sources that contribute to the heat generation in Li-ion batteries. However, the heat of mixing is insignificant and can be ignored. In general, the total heat generation is attributed to the overpotential (ohmic and polarization) heat which depends on temperature and current, and the entropy change heat, which depends on state-of-charge (SOC). It has been shown in the literature that considering just entropic and overpotential heats is enough to produce sufficiently accurate results which closely agree with the measured data. An electrochemical-thermal model referred to as the Li-ion battery model is developed and coupled with a 3D conjugate heat transfer model and numerically solved using the COMSOL Multiphysics software. The Li-ion battery model is the most accurate one among the different theoretical models available in the literature. It solves various PDEs and requires many input parameters. However, these parameters are specific to batteries and operating temperatures, and therefore it is practically impossible to measure at every possible condition that the EVs and HEVs operate. Therefore, the use of numerical modelling and simulations tools are very handy and must be adopted to design an efficient Thermal Management System (TMS) once the essential battery parameters are known and/or supplied by the manufacturer. The coupling of the battery model and thermal model (Figure 2) is performed to update the average battery temperature and heat generation. The following subsections discuss the governing equations of each of the models considered in the current research.



Figure 2. Coupling of battery and thermal models.

3.1. Li-ion (P2D) Battery Model

A Li-ion model considers various well-known theories, such as Ohm's law, porous electrode, mass and charge transfer in electrolyte and solid phases, and intercalation/deintercalation kinetics. Therefore, various partial differential equations (PDEs) are required to study these phenomena. The electrochemical process parameters required in the Li-ion battery model include the diffusion coefficient of lithium in the solution and solid phases, the ionic electrical conductivity in the solution phase, the open circuit potential, the entropy coefficient, and the reaction rate [4]. The Li-ion battery model provides four parameters namely, the solid phase potential (ϕ_s), electrolyte phase potential (ϕ_e), the concentration of Li-ion in the solid phase (C_s) and concentration of Li-ion in electrolyte phase (C_e) by solving two conservation of charge and two conservation of species equations in the solution and solid phases. Butler Volmer kinetics is used to determine reaction rate and also to couple the above equations. The governing equations are not given here for brevity, but they can be found in the literature [4,33].

3.2. Conjugate Heat Transfer Model

This 3D model is used for the cold-plates with water as coolant, as part of Thermal Management System (TMS). The value of heat generation obtained from the Li-ion model is passed on to the cold-plates through mating surfaces, which are further diffused and dissipated by the coolant flow through the channels. A fixed flow rate is supplied through individual channels, and the flow is laminar for the channel shape and cross-section used. Thus, the 3D conjugate heat transfer model solves the following general transient, incompressible fluid flow, and energy equations;

- Mass conservation equation:

$$\nabla . V = 0 \tag{1}$$

- Momentum conservation equation:

$$\frac{\partial(\rho V)}{\partial t} + V.\nabla(\rho V) = -\nabla p + \nabla.(\mu \nabla V)$$
⁽²⁾

where *V* is velocity vector, *p* is pressure, ρ and μ are density and viscosity of coolant, respectively.

- Energy conservation equation:

$$\frac{\partial(\rho C_p T)}{\partial t} + V \cdot \nabla(\rho C_p T) = \nabla \cdot (k \nabla T) + \dot{q}_h^{\prime\prime\prime}$$
(3)

where *k* is thermal conductivity, C_p is specific heat, $\dot{q}_h^{\prime\prime\prime} (= Q_h / \forall)$ is volumetric heat generation rate in the battery, and \forall is battery volume.

The instantaneous average heat generation obtained from the electrochemical (P2D) model is diffused into the cold-plates in the thickness direction and is then carried away by the liquid coolant flowing through the channels. It should be noted here that heat transfer takes place in the battery and cold-plates (as solid domains) as well as in the channels (as a fluid domain). The fluid flow equations are solved using an algebraic multigrid solver, and PARDISO solver is used for heat transfer. The backward differentiation formula (BDF) method is used for time stepping because of its stability and versatility. Furthermore, it can be noted that a stop condition is used to terminate the simulation when the discharge voltage falls below 2.0 V. At high discharge rates, it is observed that the voltage reaches the stop condition before the theoretical discharge time. Hence, there is a difference in actual and theoretical discharge times at 2 C and above.

3.3. Geometry and Mesh

The computational domain (geometry model) consists of a commercial prismatic LiFePO₄ battery sandwiched between two cold-plates, as shown in Figure 3. The height and width of the battery and the cold-plates are 227 mm and 157 mm, respectively. The thicknesses of the cold-plates and battery are 1.7 mm and 7.25 mm, respectively. The crosssection of the channels is hexagonal, and the width is 2 mm. To reduce the computational cost and due to the plane symmetry, only half of the geometry in the thickness direction (i.e., one cold-plate and half of the battery) is considered as the computational domain for the numerical simulations. A tetrahedral mesh is used for the battery and cold-plate, while the hexahedral mesh is used for the tabs. The maximum and minimum element sizes in the fluid domain are 0.625 mm and 0.25 mm, respectively. Additionally, four layers are used in the near-wall region to resolve boundary flow effects. The mesh elements in the fluid domain and its surrounding solid domains is shown in Figure 4. The present electrochemical (P2D) model requires the material property data of anode (graphite), cathode (LiFePO₄), and electrolyte (carbonate), and it does not require the property data of a separator. Additionally, the P2D model does account for the tabs. Thus, the thermal model consists of the active battery material and tabs. It can be noted that the tabs are

heated by the diffusion of heat generated in the battery. The joule heating in the tabs is ignored in the present simulations.



Figure 3. (a) Exploded view of battery and cold-plates, and (b) cold-plate with minichannels.



Figure 4. Mesh size variation in all domains (right) with a zoomed view of fluid domain (left).

3.4. Grid Independence Study

A grid independence study has been performed by comparing the maximum temperature on the battery surface and at different discharge C-rates with mesh counts of 3.5, 4.2, and 5.4 million, and the results are shown in Table 1. The mesh refinement is carried out in the fluid domain and resolves the boundary layer effects near the channel wall surfaces. It can be noted that the difference in maximum temperature with 4.2 and 5.4 million mesh counts is insignificant. However, the fine mesh with 5.4 million elements is considered for all case studies because of better convergence, but at the cost of the slight increase in the computational time.

	Maximum Surface Temperature (°C)			
Discharge C-Rate	3.5 Million Mesh Elements	4.2 Million Mesh Elements	5.4 Million Mesh Elements	
1 C	17.00	16.75	16.74	
2 C	19.90	19.56	19.50	
3 C	23.82	22.67	22.65	

 Table 1. Battery temperature vs. mesh element count.

4. Results and Discussion

In order to test the effectiveness of the minichannel cold-plate design, which is a commercially used thermal management design for prismatic Li-ion batteries, two-way coupled electrochemical-thermal simulations are performed at different discharge rates (1–4 C) and coolant inlet temperatures (15–35 °C). The predicted battery voltage and thermal responses have been analyzed. Furthermore, the predicted temperatures at selected locations on the battery surface during the discharge process are compared with the measured data. The following sections briefly discuss the effects of coolant inlet temperature and discharge C-rate on the battery thermal response in terms of the heat generation and dissipation rates, the maximum surface temperature, and the temperature difference across the battery, along with key observations, which would help in better understanding the overall behavior of the BTMS.

4.1. Battery Electrical Response

Table 2 lists the input data used in the Li-ion model, popularly known as the pseudo-2D (P2D) model available in COMSOL Multiphysics software exclusively for Li-ion batteries. Figures 5 and 6 show the Li-ion model simulated result of battery voltage variation in consideration at different C-rates and at an atmospheric temperature of 22 °C. It can be noted that the battery voltage drops faster as the C-rate increases, and the battery discharge time is reduced significantly from its theoretical time (Figure 6). However, the initial drop in voltage at different C-rates is not observed with the 1D electrochemical battery model in contrast with the measured data, and the same can be attributed to the underlying assumptions such as no spatial variation in battery heat generation, constant property data, and a single layer consisting of cathode, anode, and separator is considered in spite of the battery having multi-layers. It can be noted here that a stop condition of 2.0 V is set in the Li-ion model, and hence the discharge process terminates before the theoretical time of the corresponding battery load (current) used. Figure 7 shows that the Li-ion model predicted the power dissipation density of the 20 Ah LiFePO₄ battery at different C-rates and at room temperature of 22 °C. It can be observed that the battery heat generation rate increases linearly during the discharge process. However, it increases slowly at a low C-rate and rapidly at a high C-rate.

Material Property (Unit)	Positive Electrode	Negative Electrode				
Solid phase lithium diffusivity (m ² /s)	$1 imes 10^{-13}$	$3.9 imes10^{-14}$				
Particle radius (m)	$12.5 imes 10^{-6}$	$8 imes 10^{-6}$				
Electrolyte phase volume fraction	0.444	0.357				
Electrode phase volume fraction	0.297	0.471				
Filler phase volume fraction	0.259	0.172				
Max solid phase concentration (mol/m ³)	22,806	31,370				
Initial concentration of active material (mol/m ³)	14,870	14,870				
Solid phase conductivity (S/m)	3.8	100				
Thickness (µm)	183	117				
Other data						
Initial electrolyte salt concentration (mol/m	1000					
Bruggeman coefficient	1					
Thickness of separator (µm)	52					

Table 2. Li-ion model input data for 20 Ah LiFePO₄ [34].



Figure 5. Predicted voltage–discharge capacity variation in 20 Ah LiFePO₄ at different C-rates by Li-ion model.



Figure 6. Predicted and measured discharge voltage–time response at different C-rates of 20 Ah LiFePO₄.



Figure 7. Predicted power density variation with time at different C-rates of 20 Ah LiFePO₄ by Li-ion model.

4.2. Battery Thermal Response

The thermophysical property data for the 20 Ah LiFePO₄ battery and cold-plate materials used in the simulations are given in Table 3. It can be noted that the property data given in Tables 2 and 3 are taken from the published experimental research works and compiled in the references [33,34]. Figure 8 shows the battery surface temperatures (maximum temperature, T_{max} and temperature difference, ΔT) at the end of the discharge process with coolant inlet at 25 °C and at different C-rates. It is to be noted that the coolant inlet is on the left (anode side) and the exit is on the right (cathode side). The coolant flow rate through the individual channels considered in the present study is 150 mL/min, and it flows from the bottom left inlet on the anode side towards the bottom right exit of the cathode side with a single U-turn of individual channels. As the coolant flows inlet to the exit, water absorbs the heat energy and heats up, and hence the heat absorption capacity decreases in the flow direction. Therefore, it is obvious that battery temperatures would be higher on the cathode side and towards the coolant exit. The tabs on the battery heat up due to heat diffusion from the battery only, as the current flow through the tabs is not considered in the present study. Therefore, the cathode side tab temperature is less than battery surface temperature (near the coolant exit) in contrast to the reality wherein the current flow through the tabs results in higher temperatures due to additional internal heating. The difference in predicted values of T_{max} and ΔT can be observed from the colour contrast, as the upper temperature limit in the legend bar is fixed with respect to the highest C-rate. For instance, at 1 C discharge rate and 25 °C coolant inlet, the values of T_{max} and ΔT are 26.44 °C and 1.44 °C, respectively, while they are 36.88 °C and 11.88 °C at 4 C discharge rate and 25 °C coolant. Therefore, it is clear from Figure 8 that the T_{max} and ΔT (non-uniformity) increase with an increase in discharge rate and this is attributed to the increased heat generation due to an increase in load (current) on the battery.

Material/Property	Density (kg/m³)	Thermal Conductivity (W/m-K)	Specific Heat (J/kg-K)	
Active battery	2092	18.2	678	
Positive tab	2719	202.0	871	
Negative tab	8978	387.6	381	
Cold-plate	2702	237.0	903	

Table 3. Thermophysical property data of 20 Ah LiFePO₄ [34,35].

Ten different locations (a to j) on the battery surface are selected to analyze the rate of rise in temperatures and the maximum temperature across the battery (see Figure 9). Anode side (a, d, f), cathode side (c, e, h), and middle (b, g, i, j) are the three sets of locations. The rise in temperature during the constant current discharge procedure at various C-rates is seen in Figure 10. The discharge process is considered in three phases in order to better understand the battery surface heating rate owing to heat generation: the beginning of discharge, the middle of discharge, and the end of discharge. It can be observed from Figure 10 that the temperatures increased continuously during the discharge process (all three phases) at all C-rates, with a very rapid increase in phase 1. However, at a low C-rate (1 C), the rate of rise in temperatures is distinct in all three phases with a plateau in phase 2, while without a plateau at high C-rates (2-4 C), which confirms that the heat generation rate is higher than the heat dissipation by the coolant and hence a sharp rise in temperatures at higher C-rates. Additionally, at the end of the discharge process, the maximum surface temperatures on the anode side (a, d, f) are lower than the temperatures of the cathode side (c, e, h) and are consistent at all C-rates. Moreover, the maximum temperatures are towards the coolant exit.



Figure 8. Predicted battery surface temperatures with minichannel cold-plate (water inlet at 25 °C), and at different discharge rates; (a) 1 C (b) 2 C (c) 3 C (d) 4 C.

Figures 11–13 show the comparison of predicted and measured rise in temperatures at five different locations (f, g, h, i, j) during the discharge process at different C-rates (2–4 C) and coolant inlet temperatures (15–35 $^{\circ}$ C). The above five locations are away from the tabs and are merely selected for a quantitative comparison with the measured temperature data. The temperatures measured at a low C-rate (2 C), shown in Figure 11, the rate of rise is rapid in phase 1, and reduced in phase 2, resulting in a plateau, and then increased in phase 3. The predicted maximum temperatures by the Li-ion model at the end of the discharge process are in close agreement with the measured values, except at low coolant inlet at 15 °C and 2 C (Figure 11a). However, the rate of rise in temperatures is not as much as that which is found with the measured data. With the increase in C-rates (3-4 C) and at all coolant inlet temperatures (Figures 12 and 13), the plateau in phase 2 vanished in the measured data (solid curves), and the same has been predicted by the present Li-ion model (dotted curves) as well. At 4 C discharge rate and at all coolant inlet temperatures (Figure 13), the rise in temperature is rapid and continuous in all three phases of the discharge process, as shown by the measurements. The Li-ion model also predicted the trend, except that it over-predicted the temperatures across the battery surface. Table 4 provides the complete data on measured and predicted maximum temperatures and the temperature difference across the battery surface at all discharge rates (1-4 C)and coolant inlet temperatures (15–35 °C) undertaken in the present study. Based on Table 4, the following quantitative observations can be drawn to show how effective the Li-ion model is to consider for the BTMS.

- 1. The maximum temperature (T_{max}) and the temperature difference (ΔT) across the battery surface rose with an increase in discharge C-rate at the given coolant inlet temperature $(T_{w, in})$. At medium coolant inlet (25 °C), and at all C-rates, the predicted values of T_{max} and ΔT are in good agreement with the measured values.
- 2. At low coolant inlet (15 °C), the Li-ion model under-predicted the values of T_{max} and ΔT at 1 C and 2 C. However, at high coolant inlet (35 °C), the model slightly over-predicted the values T_{max} and ΔT at 4 C in comparison to the measured values.
- 3. Furthermore, the percentage difference between the predicted and measured values of maximum surface temperature at all C-rates are within 2.6% at high coolant inlet ($35 \degree$ C), within 4.8% at medium coolant inlet ($25 \degree$ C), and within 14.6% at low coolant inlet ($15 \degree$ C).

Water Inlet T (°C)	Discharge C-Rate	Maximum Battery Surface Temperature, T _{max} (°C)		Battery Surface Temperature Difference, ΔT	
		Experiments	Li-ion Model	Experiments	Li-ion Model
15	1 C	19.6	16.74	4.0	1.74
15	2 C	22.9	20.46	7.3	5.46
15	3 C	25.4	25.49	9.8	10.49
15	4 C	28.0	31.14	12.4	16.14
25	1 C	27.6	26.44	3.1	1.44
25	2 C	30.4	29.3	5.9	4.3
25	3 C	32.6	33.02	8.1	8.02
25	4 C	35.2	36.88	10.7	11.88
35	1 C	35.9	36.13	2.4	1.13
35	2 C	38.2	38.15	4.7	3.15
35	3 C	40.3	40.17	6.8	5.17
35	4 C	42.3	43.38	8.8	8.38

Table 4. Predicted vs. measured temperatures of a 20 Ah LiFePO₄ battery with cold-plates.



Figure 9. Ten different locations on battery surface.



Figure 10. Predicted surface temperature variation at 10 different locations with water inlet at 25 °C and at different discharge rates; (a) 1 C (b) 2 C (c) 3 C (d) 4 C.



Figure 11. Predicted (dotted curves) and measured (solid curves) surface temperatures at 5 different locations at 2 C discharge rate and at different coolant inlet temperatures; (a) 15 °C (b) 25 °C (c) 35 °C.



Figure 12. Predicted (dotted curves) and measured (solid curves) surface temperatures at 5 different locations at 3 C discharge rate and at different coolant inlet temperatures; (**a**) 15 °C (**b**) 25 °C (**c**) 35 °C.



Figure 13. Predicted (dotted curves) and measured (solid curves) surface temperatures at 5 different locations at 4 C discharge rate and at different coolant inlet temperatures; (**a**) 15 °C (**b**) 25 °C (**c**) 35 °C.

Therefore, the above results clearly indicated that the performance of the current coolant design with distributed minichannel cold-plate water-cooling is effective for medium to high coolant inlet (25–35 °C) but not at low coolant inlet (15 °C). This could be due to a higher heat generation rate at low coolant inlet temperature. Interestingly, it is observed that at a given C-rate, T_{max} increased linearly with an increase in $T_{w,in}$. Additionally, at a given coolant temperature ($T_{w,in}$) the temperature difference (ΔT) increased with increase in discharge C-rate and is due to increased heat generation rate with increase in C-rates. However, ΔT decreased with increasing $T_{w,in}$ at all C-rates. Furthermore, increasing C-rates resulted in more non-uniformity in battery temperatures. Additionally, at all coolant inlet T_{win} , the predicted temperature difference is less than 5 °C at 1 C and 2 C discharge rates, and is about 8–16 °C at 3 C and 4 C discharge rates. Thus, it can be concluded that the temperature uniformity across the battery surface primarily depends on the discharge C-rate and the coolant inlet temperature has a secondary effect, whereas the maximum temperature depends both on C-rate and coolant inlet temperature. Additionally, the present thermal management system (TMS) with distributed minichannel cold-plates is more suitable for 1–2 C discharge rates and coolant inlet temperatures between 25–35 °C, because the values of T_{max} and ΔT lie within the desired optimum working range (T_{max} between 25–40 °C and $\Delta T < 5$ °C).

5. Summary

In COMSOL software, a two-way coupled transient electrochemical-thermal model for a 20 Ah LiFePO₄ battery sandwiched between two minichannel cold-plates with the distributed fluid flow has been created. The P2D Li-ion model calculates the instantaneous heat generation rate at the average battery temperature and feeds it into the 3D thermal model for the battery and cold-plates with fluid flow. Initially, the Li-ion model is evaluated at various C-rates (1–4 C) and room temperature (22 °C), and found that battery discharge capacity declines as C-rates rises. Following that, the effectiveness of the current BTMS is investigated in terms of heat generation rate, maximum temperature (T_{max}), and temperature difference (ΔT) across the battery surface at various discharge C-rates and water-coolant inlet temperatures ($T_{w,in}$: 15–35 °C). The predicted battery surface heating rate and end temperatures are also compared to the measured data at 10 different locations. The following are derived from the current research;

- 1. Both T_{max} and ΔT increase with an increase in the C-rate, and the rise in T_{max} is high at low $T_{w,in}$ in contrast to high $T_{w,in}$, and is attributed to more heat generation at low $T_{w,in}$.
- 2. T_{max} increases linearly with an increase in $T_{w,in}$. However, ΔT is more (i.e., non-uniformity) at low $T_{w,in}$ than at high $T_{w,in}$.
- 3. Predicted temperatures are in close agreement with the measured data at 3 C discharge rate. However, at low C-rates (1–2 C) the model slightly under-predicted the temperatures and slightly over-predicted at high C-rate (4 C).
- 4. The current BTMS shows best performance meeting the desired operating range $(T_{max} : 25-40 \text{ °C}, \Delta T < 5^{\circ}\text{C})$ at nominal discharge capacity (1 C) and at all coolant temperatures studied.

Based on the above observations, the coupled electrochemical-thermal model predicted the effects of coolant inlet temperatures and discharge C-rates on T_{max} and ΔT across the battery surface as well as the battery surface heating rate, and are in close agreement with the measured data. The differences in the model predictions and measurements can be attributed to the assumption of homogenous reactions in the P2D model and uniform heat generation across the battery. Additionally, the Joule heating in the tabs is not considered in the present model. Furthermore, the constant input data pertaining to the electrochemical phenomena may also account for the percentage differences between the predicted and measured data. It can be concluded that the Li-ion model (P2D) available in COMSOL software can be used to design the (BTMS) at the cell level. However, to improve the battery model prediction accuracy, it is recommended to use the multi-layer approach instead of a single-layer approach, and a 3D electrochemical model with temperature spatial dependent electrochemical and thermal property data.

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