

Thermal Monitoring of LiFePO₄ Lithium Battery System for the Use of an Electric Boat with Recharging Based on Photovoltaic Systems.

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Abstract– For the safe operation and control of battery deterioration, thermal monitoring is extremely important. The performance of a battery energy system and its accuracy in State of Charge (SoC) (%) estimation for electric boats and electric vehicles are significantly influenced by ambient temperature. In this study, the battery will operate at extreme temperatures within Titicaca Lake, located at an altitude of 3850 meters above sea level. Therefore, the objective is to study a battery model that considers ambient and extreme temperatures. The cell must operate at critical values with a current of 150 A, a voltage of 3.65 V, and observe thermal aging that affects the capacity due to multiple charges and discharges. The method applied to lithium iron phosphate batteries LiFePO₄ involved wear testing, considering three stress factors (time, temperature, and state of charge (SoC) (%)). Capacity measurements and resistance calculations tracked short-term degradation behaviors. The Shepherd model was established to identify battery aging using a simple but accurate two-step nonlinear regression approach; also, the thermal analysis of the battery cell was conducted to identify the most critical zone of the cell in terms of heat generation. This analysis allowed us to maximize heat dissipation with only a forced air fan mounted on the vital region. For further evaluation of the proposed strategies, a computational fluid dynamics (CFD) model was built in COMSOL Multiphysics® and validated with a thermographic camera to identify the temperature in the cells. The final arrangement of the LiFePO₄ battery is the 16s1p prismatic cell type (16 cells in series and one parallel), with the cell experiencing a temperature variation of approximately 1° C higher than that of the outer parts.

Keywords– LiFePO₄ battery; electric boat; recharging; state of charge; battery thermal management; photovoltaic.

I. INTRODUCTION

Cathodes of LiFePO₄ lithium batteries represent the optimal choice for photovoltaic systems, as well as other renewable energy applications, aiming to achieve energy self-consumption [1]. Certain categories of lithium batteries exhibit specific attributes related to weight, dimensions, safety, and robustness, allowing them to adapt to scenarios where lead-acid batteries are not viable [2].

Lithium iron phosphate LiFePO₄ batteries represent a category of rechargeable lithium-ion batteries designed for

high-power applications [3]. These batteries feature high discharge currents, lack explosiveness, exhibit a significant cycle life, and initially have lower energy density compared to lithium cobalt oxide, but surpass it significantly by the third year due to their greater stability [4].

A. Lithium Iron Phosphate LiFePO₄

In 1996, the University of Texas discovered that phosphate could be used as a cathode material for lithium batteries [5]. This cathode material remains stable even under overload conditions and can tolerate high temperatures without decomposing. The cathode material in lithium iron phosphate batteries is more reliable and safer than other cathode materials such as LiCoO₂ or LiMn₂O₄ batteries [6]. Phosphates have a cell operating temperature range of -30°C to +60°C and a cell packaging temperature range of -50°C to +60°C, which reduces the risk of overheating and fire hazard [7]. This type of battery is made from nanoscale phosphate materials and exhibits low resistance, long lifespan, high capacity to handle heavy loads, improved safety and thermal consistency, no toxic effects, and lower cost. It has a reduced impact on cycle life concerning overcharge and undercharge, although specific energy slightly decreases compared to LiMn₂O₄. It exhibits some negative performance and lifespan properties due to temperature [8]. Lithium-phosphate batteries are capable of providing a specific energy and nominal voltage of approximately 160 mAh/g and 3.40 V, respectively [9]. These characteristics make it straightforward to enhance battery performance. Within lithium ion batteries, according to its chemical composition, LFP (LiFePO₄) is the cathode source, see Figure 1, illustrates the types of cells for design and chemistry [10].

B. Reaction of Lithium Iron Phosphate LiFePO₄ Battery Chemical

A new type of high-performance battery is the lithium iron phosphate LiFePO₄ battery. An electrochemical process underlies its chemical reaction mechanism [11]. It has been extensively utilized in energy storage systems and electric boat because of its benefits, which include high energy density, extended cycle life, and environmental friendliness [12].

The two primary components of the chemical process in a lithium iron phosphate LiFePO_4 battery are the charging reaction and the discharging reaction [13].

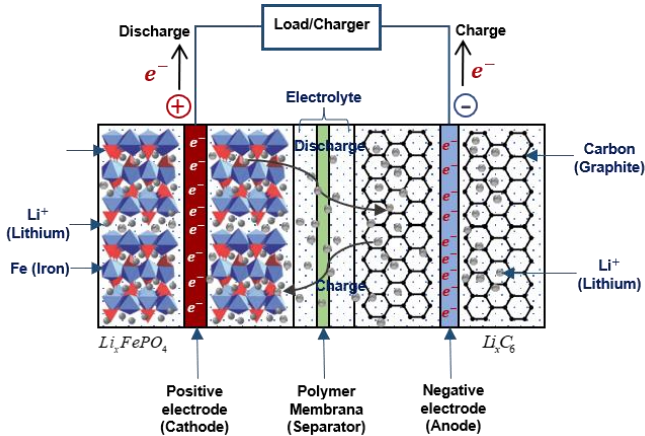
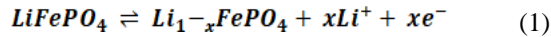


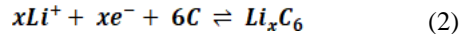
Fig. 1 Charging-discharging process of a LiFePO_4 type battery

In the process of charging, the positive electrode (lithium iron phosphate) undergoes a reaction where it relinquishes oxygen molecules, converting them into iron trioxide (Fe_2O_3), and simultaneously discharges electrons, see Eq. 1. Conversely, the negative electrode (graphite) assimilates electrons and transforms lithium ions into metallic lithium [14], [15]. The process is shown in Figure 1.

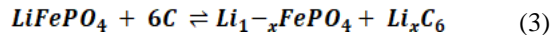
Positive reaction:



Negative reaction:



Total reaction:



LiFePO_4 is the chemical formula of lithium iron phosphate; in the Eq. 1. $\text{Li}_{1-x}\text{FePO}_4$ is a vacancy-containing chemical compound; $x\text{Li}^+$ represents lithium ions that have lost x electrons; xe^- represents the x electrons released; and 6C is the chemical formula of graphite.

When the battery is charged, lithium ions move from the positive electrode to the negative electrode, where they react chemically with graphite and cling to the graphite's surface, forming lithium compounds. At the same time, electrons in the battery travel from the positive to negative electrodes, producing current in the external circuit. As a result, lithium iron phosphate batteries have high energy density, a long cycle life, and are environmentally friendly, making them promising for use in the field of renewable energy.

II. THE OPERATIONAL ENVIRONMENT OF AN ELECTRIC BOAT

2.1 Power System Structure

An autonomous electric boat is being constructed with photovoltaic systems and a special nautical lithium battery that enhances the protection and performance of this electric vessel. It measures 7.5 meters in length from bow to stern, 2.20 meters in width, and is powered by a NAVY 6.0 electric motor with 9.9 HP, 39-60 V dc. It uses a LiFePO_4 lithium-ion battery model E-163 with 8345 Wh and 163 Ah capacity, charged by a CH4200 lithium battery charger of 30A, 220V AC 1440 W. The boat also features a solar charge controller for the E-series 48 V battery, 1600 W, and a photovoltaic system with 6 Monocrystalline RenePV 24 V dc, 270 W, Voltage at Pmax of 31.35V and Current at Pmax of 8.61A.

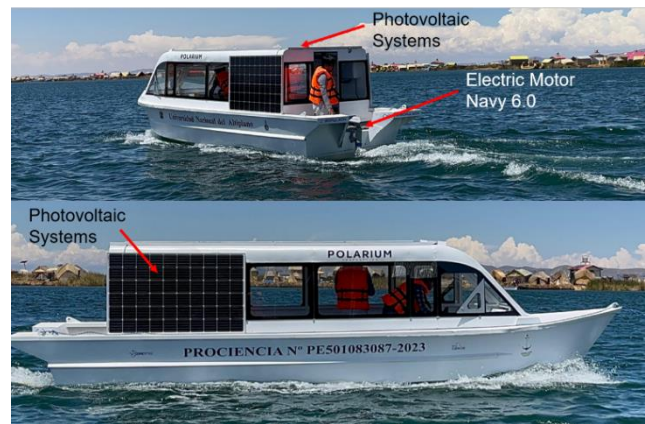


Fig. 2 Here we can see the electric boat in operation with the LiFePO_4 lithium-ion battery model E-163, NAVY 6.0 electric motor, Photovoltaic system, Charged 30A serie CH4200

The choice of the electrical system depends on various factors such as the aquatic environment, navigation conditions, and economic costs. This document considers the basic structure of the hull, vessel, or outer lining that wraps around and waterproofs its framework, forming the hull and the cabin area where the photovoltaic system is mounted. When the electric boat docks, it is powered from the 220 V electrical grid with a transformer capacity of 10 kVA. Single-phase industrial simple charging point, type 1 up to 125 A, SAE J1772/IEC 62196-2, considered within the Battery Electric Vehicle (BEV) technology, which is a fully electric vehicle powered exclusively by a lithium battery.

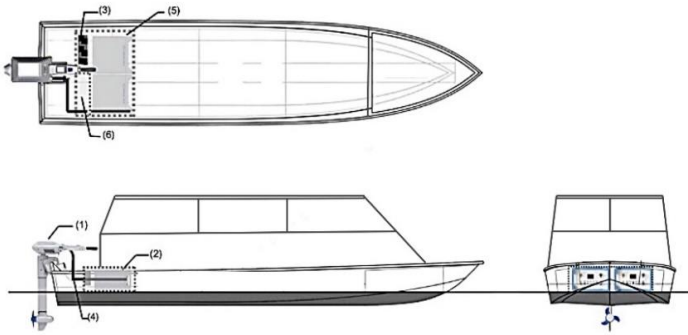


Fig. 3 This is a figure. Schemes follow the same formatting. If there are multiple panels, they should be listed as: (1) NAVY 6.0 electric motor, (2) LiFePO4 lithium-ion battery model E-163, (3) Charged 30 A serie CH4200, 4) Power and data cable (5) Drawer for the system (battery, charger and wiring), (6) secondary drawer for charging cable.

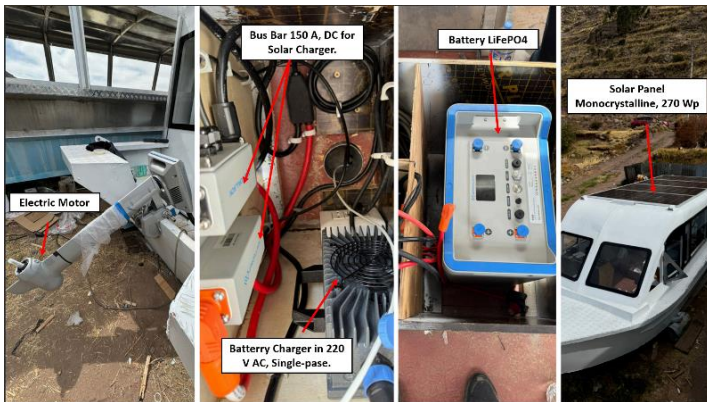


Fig. 4 Parts that make up the propulsion system.

The primary propulsion engine that powers the propeller must adhere to all applicable regulations or standards set out by the maritime inspection organization. These include those pertaining to the marine motor, motor protection class, insulation level, and other technical indications. Furthermore, the motor needs to function properly in environments with mildew, oil, salt, and damp air. To prevent motor blockages due to propeller windings, the motor needs to have a one-minute overload capacity. The motor's speed range need to be sufficient for the propeller's needs. The DC motor has low cost, good durability, and simplicity.

When operating at full speed, the motor's rated power should match the propulsion's rated power. In the Figure 5 displays the motor's primary technical specifications.

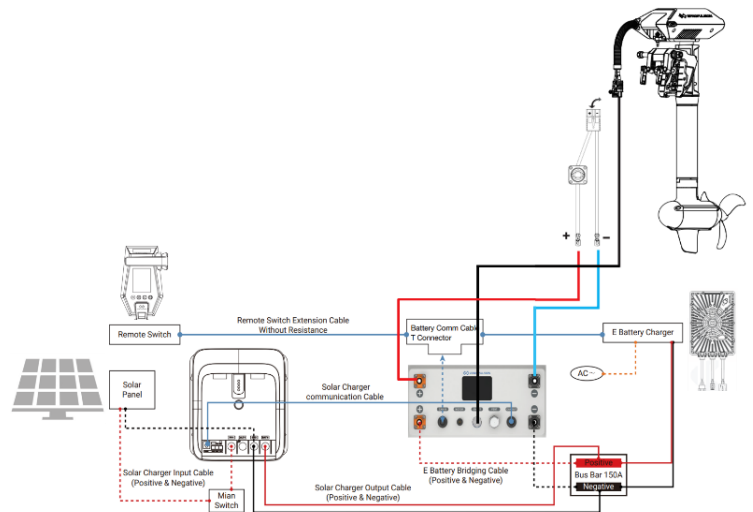


Fig. 5 Components of the propulsion system for the electric boat of the project; solar panel, solar charger, remote switch, battery LiFePO4, motor, battery charger

2.2 Battery Experiment

The capacity of prismatic LiFePO₄ cells typically ranges from 100 to 300 Ah. This capacity range makes them suitable for applications requiring moderate to high energy storage capacity, such as electric vehicles (EVs), energy storage systems (ESS) for residential or commercial use, and industrial applications.

A commercial LiFePO₄ battery with a capacity of 163 Ah, 3.2 V is used in this investigation, is an aluminum-cased lithium iron phosphate prismatic battery that is ideal for power applications such as electric vehicles and boats.

III. RESULT AND ANALYSIS

3.1 Result of Temperature behavior for LiFePO₄ prismatic cells

Heat transfer in fluids in vector form can be described using the concept of heat flux, which is a vector quantity indicating the amount of heat crossing a surface per unit time and area. In vector terms, heat transfer in fluids is expressed as:

$$\vec{q} = -k \cdot \nabla T \quad (4)$$

where; \vec{q} is the heat flux vector, indicating direction and magnitude, k is the thermal conductivity of the fluid, ∇T is the temperature gradient, a vector indicating the direction and rate of temperature change in space.

In the Eq. 4, The negative sign indicates that heat flux always moves from regions of higher temperature to regions of lower temperature, in accordance with the second law of thermodynamics. $k \cdot \nabla T$ represents the amount of heat flowing through a surface per unit area and time, where ∇T defines the direction and magnitude of the temperature gradient.

In the equation 5, this vectorial model is fundamental for formulating and solving heat transfer problems in fluids, especially in applications where the direction and spatial distribution of heat within a fluid system are considered; ρ density, C_p specific heat, \vec{u} Velocity Vector, \vec{q} Heat flux by conduction, Q Heat source, Q_{ted} Thermoelastic effects. (M. Soltani *et al.* 2019). (J. Jaguemont *et al.* 2016).

$$\rho C_p \frac{\partial T}{\partial t} + \rho C_p \vec{u} \cdot \nabla T + \nabla \cdot \vec{q} = Q + Q_{ted} \quad (5)$$

COMSOL Multiphysics® created the 3D-thermal model to determine the cell's thermal behavior. An energy balance equation is used to specify the transient thermal distribution within the cell. This equation formulates the quantity of thermal energy that the cell generates for its surroundings as follows. (M. Soltani *et al.* 2019).

$$m C_p \frac{\partial T}{\partial t} + q_{conv} = k \left[\frac{\partial^2 T}{\partial x^2} + \frac{\partial^2 T}{\partial y^2} + \frac{\partial^2 T}{\partial z^2} \right] + q_g \quad (6)$$

where mass, heat capacity, temperature, thermal conductivity, and heat generation are denoted by the letters m , C_p , T , k , and q_g , respectively. In this paper, the polarization process and the cell's ohmic resistance are used to compute the heat generation of the cell. Additionally, Eq. (7) makes use of the tab domain.

$$q_g = R_{bt} \cdot I^2 + R_1 \cdot I_1^2 + R_2 \cdot I_2^2 \quad (7)$$

wherein, I and R_{bt} represent the current and ohmic resistance of the cell. (J. Jaguemont *et al.* 2016)

$$q_{conv} = hS(T_{amb} - T) \quad (8)$$

where h and S stand for the cell's cross section area and coefficient of heat transfer, respectively. T_{amb} and T also show the battery and the surrounding temperature. A thermal camera clearly shows that the temperature distribution within the cell region is not uniform in Fig. 6.

According to the model described in Eq. 4 and 5, you can see the simulation for the prismatic lithium cells; within the cell, there is a negligible variation in temperature. As would be predicted given that this is the location that is farthest from the cooling surfaces, the highest temperature is reached within the jelly roll, about in the center of the cell. During discharge and charging, the temperature rises; that is, it does not attain a constant state. See the figures 6 to 8.

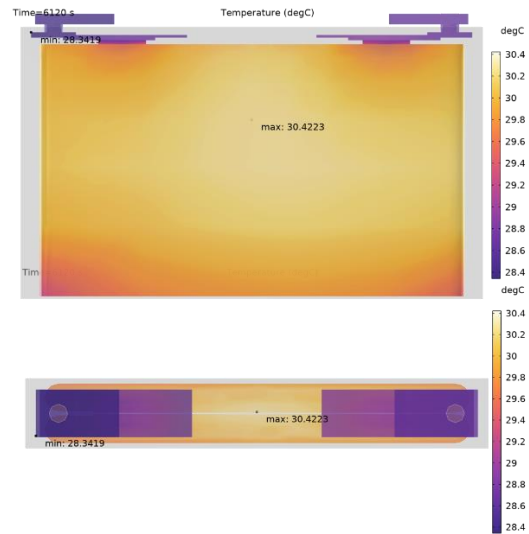


Fig. 6 This is a figure of Temperature Field of prismatic lithium cell, simulated in COMSOL.

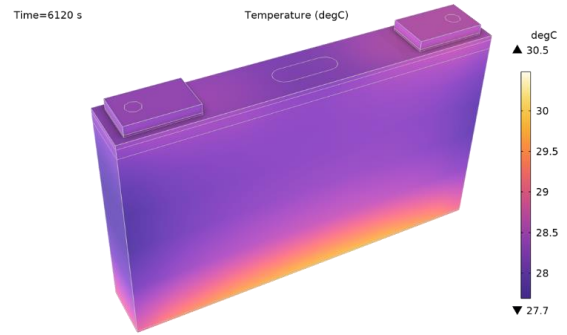


Fig. 7 Surface temperature, simulated in COMSOL

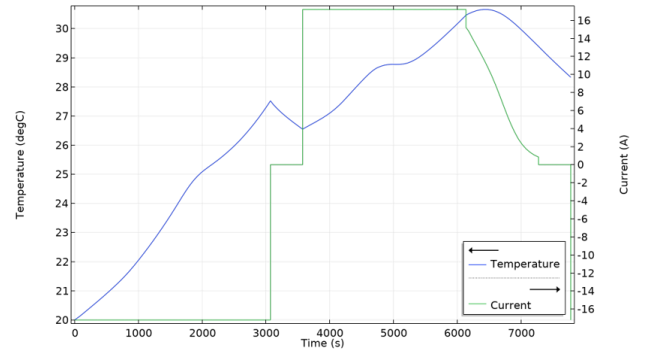


Fig. 8 The temperature increases during discharge and charge.

The significance of the metal foils in cooling the cell is seen by the temperature along the roll layer's cross sections at that location. The temperature displays a local maximum on the negative current collector foil and a local minimum on the positive foil. The cell's outside cooling is the cause of the minimum. Since it is the layer farthest from the cooling tab, the maximum is. At the location of the negative foil, an

identical but opposite effect is observed. The nearest foil that is closest to a tab tends to chill the jelly roll layer's components.

Finally the variance in external temperature is a result of the disparity in thermal conductivity between the metal components and the electrical insulators. The bottom of the cell, where the jelly roll and battery container are in close contact, has the highest temperature. See the Figure 7.

3.2 Result of external temperature behavior for the prismatic LiFePO₄ (16 cells in series and one parallel cells)

The thermographic camera monitors the first charging process of the photovoltaic system and the discharge of the battery to the electric motor, providing information about the temperature distribution and potential short circuits. The automatic detection of hot spots recognizes and automatically reports deviations from the target temperature.

Hot spots in batteries during operation can provide an indication of the areas where the battery will fail. This adverse thermal behavior can quickly spread throughout the battery in a domino effect known as thermal runaway, which can lead to catastrophic failures. Excessive voltage, charge, ambient temperature, or a combination of these factors can trigger thermal runaway. Sealed cells can sometimes explode violently if the safety vents become clogged or fail to function. Lithium-ion batteries are particularly prone to thermal runaway.



Fig. 9 External temperature behavior for the prismatic LiFePO₄ (16 cells in series and one parallel cells).

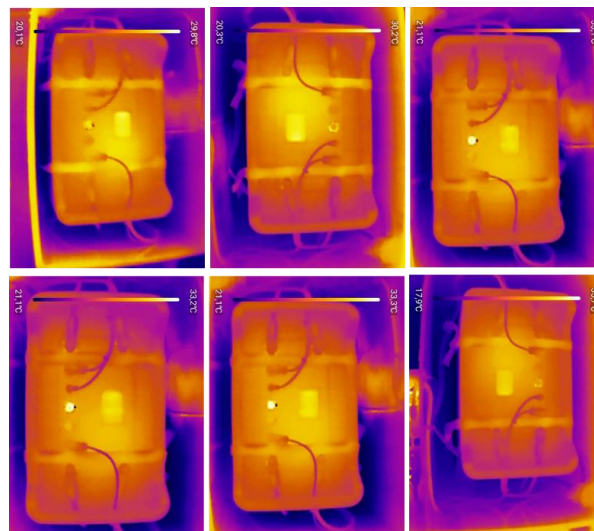


Fig. 10 The temperature increases during discharge and charge 29.8 °C to 35.7° C.

The cell's temperature under natural air cooling is taken into account for the beginning temperatures of 20.1 °C to 35.7 °C at a 53 Amp. discharging rate. However, during charging with the photovoltaic system, the temperature varies between 20 °C at a charging current of 13 A. See the Figures 9 and 10.

IV. CONCLUSIONS

With the proposal to reduce polluting emissions from electric boats within the bay of Lake Titicaca, thereby improving air quality, flora, and fauna, this study presented a hybrid energy system powered by a photovoltaic DC system and conventional AC grid according to the characteristics reflected in current electromobility standards such as SAE J1772, IEC 61851, and GB/T 20234-2. A method for designing parameters of the hybrid energy system based on loads like the 6 kW, 9.9 HP propulsion motor and auxiliary equipment was suggested. The results of the experiments demonstrate that the proposed technique for designing system parameters and control strategy for the hybrid energy system are satisfactory; the energy capacity meets the navigation requirements on Lake Titicaca. The benefits of energy conservation and emission reduction are clearly evident.

Thermal modeling is an effective way to understand the thermal behavior of LiFePO₄ lithium-ion batteries during charging and discharging. With charging and discharging, the internal heat generation of the battery increases, thereby raising its temperature with an uneven distribution and partial degradation of the battery.

Non-destructive evaluation of batteries using thermographic cameras allows for determining the temperature distribution in the lithium-ion battery. The key to solving the safety issue lies in thermal control, including heat generation and internal and external heat transfer. Therefore, a

model was created independently for each cell, capturing the geometric thermal effects on the thermal abuse behavior of the battery based on the charging of the photovoltaic system and discharge to the electric motor.

The analysis of discharge voltage characteristics in the prismatic cell type shows that comparing the rate of voltage changes in the final discharge phase will allow us to detect SoC imbalances. Parameters were obtained that will enable us to develop a damage protection system for the entire cell arrangement for LiFePO₄ batteries in 16S1P configuration.

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