

Recent Advances in Cathode Materials for Sodium-Ion Batteries: Structure–Property Relationships and Performance Optimization Strategies

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Abstract. Sodium-ion batteries show great potential for large-scale applications due to abundant sodium resources and low cost. The performance of cathode materials is crucial for determining the energy density and cycling stability of these systems. This paper reviews three representative cathode materials, discussing their structural features and electrochemical behavior, with a comparative analysis of capacity, rate performance, and cycling life. The comparison reveals that while each material demonstrates specific advantages in storage mechanism, structural stability, and cost, challenges such as capacity fading and limited rate capability remain. Finally, research progress is summarized, and future directions including structural optimization, interface engineering, and cost reduction are outlined to guide the development of advanced sodium-ion cathodes.

Keywords: Sodium-ion batteries, Cathode material, Layered transition metal oxides, Polyanionic compounds, Prussian blue compound Introduction

1.Introduction

The global energy structure is undergoing profound transformation, with renewable sources such as solar and wind power expanding rapidly but exhibiting inherent intermittency and vulnerability to climatic fluctuations, which creates instability in power output and challenges for grid reliability. To mitigate these issues, energy storage technologies play a critical role by balancing supply and demand, enabling peak shaving and valley filling, and enhancing system flexibility. Among the various storage approaches, electrochemical energy storage has attracted particular attention due to its high energy density, rapid response, and deployment flexibility, with lithium-ion batteries (LIBs) being the most mature and widely applied in electronics, electric vehicles, and power grids[1]. However, the uneven distribution and high cost of lithium resources, combined with environmental concerns from mining, constrain their long-term scalability [2,3]. This has driven research and industry to seek resource-abundant and cost-effective alternatives, among which sodium-ion batteries (SIBs) are emerging as a promising candidate due to the chemical similarity of sodium to lithium and its global abundance. Nevertheless, sodium's larger ionic radius results in slower diffusion kinetics and incompatibility with many existing LIB electrode materials, creating both challenges and opportunities for material innovation. In particular, cathode materials largely

determine the energy density, while anode materials strongly influence cycle life, making the development of high-capacity, long-life cathodes central to advancing practical SIB applications [4]. This paper therefore reviews the performance of three representative cathode materials, evaluates their respective advantages and limitations, and highlights future research directions, aiming to provide both a theoretical foundation and practical guidance for accelerating the commercialization of sodium-ion batteries in large-scale energy storage systems.

2. Methodology

This review adopts a structured research design to ensure the reliability and comprehensiveness of the analysis. Literature was retrieved primarily from the China National Knowledge Infrastructure (CNKI) database, with the keywords “cathode materials for sodium-ion batteries” and a time span from 2020 to 2025. To maintain academic rigor, only peer-reviewed journal articles and master’s theses were included, while patents were excluded. The collected literature was examined using a combination of qualitative, classification, and comparative analytical methods. Specifically, qualitative analysis was employed to describe the crystal structure characteristics, ion migration mechanisms, and electrochemical behaviors of different cathode materials. The materials were then classified by structural type and chemical composition into layered transition metal oxides, polyanionic compounds, and Prussian blue analogues, and the advantages and limitations of each category were systematically discussed. Finally, a comparative analysis was carried out to highlight the similarities and differences among these materials in terms of energy density, cycling stability, and ion transport rates, thereby providing a comprehensive and multi-perspective reference for the optimization and future design of high-performance sodium-ion battery cathodes. Ethical considerations were not applicable in this study, as the research is based entirely on previously published literature without the involvement of human participants or animal experiments.

3. Results

Sodium-ion battery cathode materials continue to encounter challenges related to cycle life, ionic/electronic conductivity, and rate performance. Given the diverse structural characteristics and degradation mechanisms across different material systems, targeted optimization strategies must be formulated to improve their overall electrochemical performance. Due to the differences in structural characteristics and failure mechanisms of different material systems, corresponding optimization strategies need to be formulated based on specific structural characteristics and failure mechanisms to improve their comprehensive electrochemical properties.

Layered oxides represent one of the most extensively studied cathode materials for sodium-ion batteries, with the general formula Na_xMO_2 (M = transition metals such as Ni, Mn, or Co). Depending on the Na^+ arrangement and oxygen stacking sequence, the structures are typically classified into O3, P2, and P3 types. O3-type materials deliver high initial capacities but suffer from phase transitions during cycling, resulting in poor long-term stability. P2-type compounds provide wide Na^+ diffusion channels and excellent rate performance, but their structural instability under high voltage limits practical application[5,6]. P3-type structures share features of both O3 and P2 but have been less investigated. Overall, layered oxides offer high capacity and tunable structures, though they face challenges in cycling stability and air sensitivity, necessitating further optimization and surface modification. Figure 1 shows the structural diagram of P2 type (P63/mmc) and O3 type (R-3m) materials.

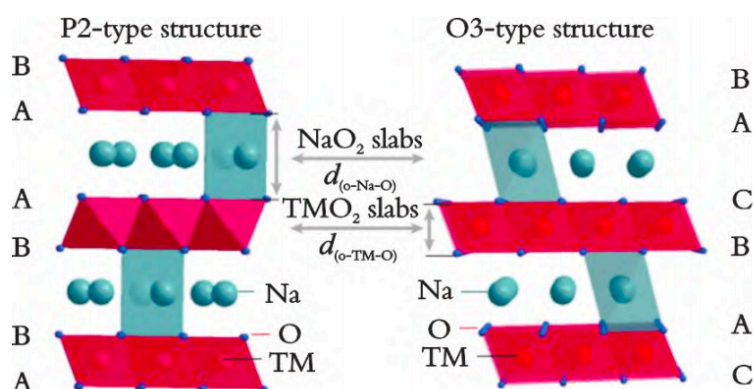


Figure 1. Structural schematic of P2-type (P63/mmc) and O3-type (R-3m) materials[7]

Polyanionic compounds, typically composed of stable three-dimensional frameworks formed by PO_4 , SO_4 , or SiO_4 anions, are represented by materials such as $\text{Na}_3\text{V}_2(\text{PO}_4)_3$ and $\text{Na}_2\text{Fe}_2(\text{SO}_4)_3$. Their primary advantages lie in robust structural stability, high operating voltages, and excellent thermal safety, which contribute to long cycle life and reliable safety performance[4]. However, their specific capacities are generally lower than those of layered oxides, and poor electronic conductivity remains a major limitation, often requiring carbon coating or cation doping to enhance electrochemical performance. In summary, polyanionic compounds hold great promise for large-scale energy storage, though further improvements in energy density are essential. The crystal structure of polyanion-type compounds is shown in Figure 2 to illustrate their typical coordination framework [8].

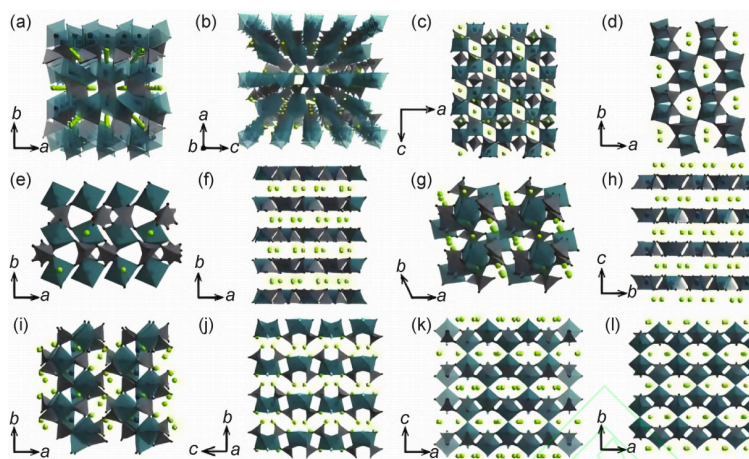


Figure 2. Crystal structure of representative polyanion-type compounds

(a) NaFePO_4 -olivine; (b) NaFePO_4 -maricite; (c) NASICON-type $\text{Na}_3\text{V}_2(\text{PO}_4)_3$; (d) NaFe_2PO_7 ; (e) NaFeP_2O_7 ; (f) $\text{Na}_2\text{CoP}_2\text{O}_7$; (g) $\text{Na}_2\text{CoP}_2\text{O}_7$; (h) $\text{Na}_2\text{CoP}_2\text{O}_7$; (i) $\text{Na}_4\text{Fe}_3(\text{PO}_4)_2\text{P}_2\text{O}_7$; (j) $\text{Na}_2\text{FePO}_4\text{F}$; (k) $\text{Na}_3\text{V}_2(\text{PO}_4)_2\text{F}_3$; (l) $\text{Na}_3\text{V}_2(\text{PO}_4)_2\text{O}_2\text{F}$ [8]

Prussian Blue Analogues (PBAs) possess an open cubic framework formed by $\text{M}-\text{C}\equiv\text{N}-\text{Fe}$ linkages, which provide large Na^+ diffusion channels, leading to excellent rate capability and cost advantages[9-11]. Moreover, PBAs can be synthesized under mild conditions, making them suitable for large-scale production. However, issues such as structural water and vacancy defects often impair electrochemical stability and capacity retention. Recent advances in synthesis techniques,

defect control, and water content regulation have significantly improved their performance. Nevertheless, long-term cycling stability and energy density remain critical challenges that need to be addressed for practical applications. The representative framework of Prussian-blue materials is depicted in figure3 [12].

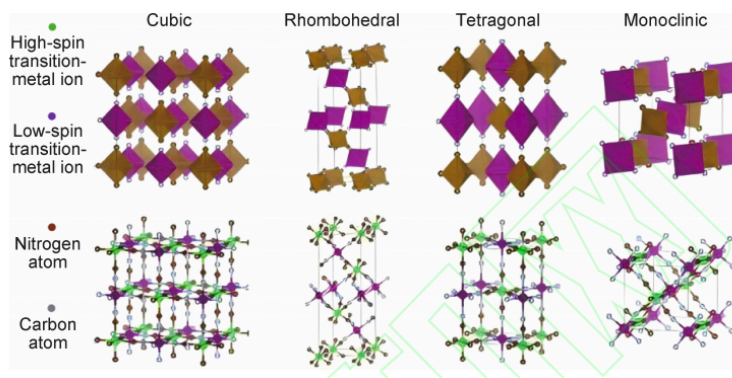


Figure 3. Crystal structures of Prussian-blue materials

4. Conclusion

4.1 Summary of Key Findings

Layered transition metal oxides, polyanionic compounds, and Prussian blue analogues each exhibit distinct advantages and limitations as sodium-ion battery cathode materials. Layered oxides provide relatively high capacity and voltage but suffer from phase transitions, Na^+ loss, and structural collapse during cycling. Polyanionic compounds offer high structural stability and long cycle life but are limited by low ionic and electronic conductivity, resulting in poor rate performance. Prussian blue analogues feature simple synthesis, low cost, and fast ion diffusion but are prone to crystalline water incorporation and Fe–CN bond breakage, leading to severe capacity decay.

4.2 Comparison with Previous Studies

Sodium Layered Transition Metal Oxides. Layered oxides such as $\text{NaNi}_x\text{Co}_y\text{Mn}_z\text{O}_2$ ($x+y+z=1$) have been widely investigated for their high capacity (160–200 mAh/g) and operating voltage (3.0–4.2 V vs. Na^+/Na). For example, $\text{NaNi}_{0.5}\text{Mn}_{0.5}\text{O}_2$ delivers ~160 mAh/g with 80% retention after 500 cycles. However, phase transitions to rock-salt or spinel phases, particularly above 4.2 V, accelerate capacity fading (>30% within 100 cycles). Moreover, Na^+ extraction induces interlayer shrinkage, triggering particle cracking and electrolyte corrosion, which further accelerates performance degradation.

Polyanionic Compounds. $\text{Na}_3\text{V}_2(\text{PO}_4)_3$ (NVP) exemplifies polyanionic cathodes with a rigid three-dimensional framework, offering outstanding stability (>90% retention over 10,000 cycles) and an operating voltage of ~3.4 V. With a theoretical capacity of 117 mAh/g, NVP typically delivers ~100 mAh/g, making it suitable for grid-scale applications requiring long cycle life. Nonetheless, the insulating nature of polyanionic groups results in low electronic conductivity ($\sim 10^{-10}$ S/cm) and limited Na^+ diffusivity ($\sim 10^{-14}$ cm²/s), yielding poor rate capability (1C discharge capacity ~60% of 0.1C).

Prussian Blue Analogues. Prussian blue analogues (PBAs) are attractive for low-cost and fast-charge applications due to facile aqueous co-precipitation synthesis, inexpensive precursors

(ferrocyanide salts <10 USD/kg), and open three-dimensional frameworks (Na^+ diffusivity up to 10^{-9} cm^2/s). For instance, $\text{Na}_2\text{Fe}[\text{Fe}(\text{CN})_6]$ achieves ~ 150 mAh/g and supports rapid charging within 15 minutes (3C). However, crystalline water incorporated during synthesis occupies Na^+ sites, causing an initial capacity loss exceeding 20%. Furthermore, Fe–CN bond breakage during cycling destabilizes the framework, leading to poor cycle life, with capacity retention below 60% after 100 cycles.

4.3 Implications of the Findings

This study systematically analyzes the performance bottlenecks and optimization strategies for sodium-ion battery cathode materials. The findings provide theoretical guidance and practical insights for improving electrochemical performance, offering valuable reference for advancing the large-scale application of sodium-ion batteries.

4.4 Limitations of the Study

This study focused on summarizing material categories and optimization strategies but did not propose specific candidate materials for immediate application.

4.5 Suggestions for Future Research

Future research should integrate structural regulation and performance balancing to develop more robust composite cathodes. Advanced characterization techniques are also needed to elucidate failure mechanisms, while efforts toward green, low-cost synthesis routes will be critical to accelerate commercialization of sodium-ion batteries.

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