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Entropy-Driven Mechanisms in P2-Type Layered Oxide Cathodes for Sodium-Ion Batteries: New Insights from First-Principles and Electrochemical Analysis

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Arianna Massaro,^{a,b,} Silvia Porporato,^{b,c} Miriam Botros,^{d,*} Alessandro Piovano,^{b,c,*} Hamideh Darjazi,^{b,c} David Stenzel,^d Giuseppina Meligrana,^{b,c} Ana B. Muñoz-García,^{b,e} Ben Breitung,^d Michele Pavone,^{a,b} and Claudio Gerbaldi^{b,c}*

^a Department of Chemical Sciences, University of Naples Federico II, Via Cintia 21, Napoli 80126, Italy

^b National Reference Centre for Electrochemical Energy Storage (GISEL), INSTM, Via Giusti 9, Firenze 50121, Italy

^c GAME Lab, Department of Applied Science and Technology (DISAT), Politecnico di Torino, Corso Duca degli Abruzzi 24, Torino 10129, Italy

^d Institute of Nanotechnology, Karlsruhe Institute of Technology (KIT), Hermann-von-Helmholtz Platz 1, 76344 Eggenstein-Leopoldshafen, Germany

^e Department of Physics “E. Pancini”, University of Naples Federico II, Via Cintia 21, Napoli 80126, Italy

***corresponding authors:** *Arianna Massaro* (arianna.massaro@unina.it), *Miriam Botros* (miriam.botros@kit.edu), *Alessandro Piovano* (alessandro_piovano@polito.it)



Abstract

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P2-type Na_xMO_2 layered oxides ($x < 1$) are highly promising cathodes for Na-ion batteries (NIBs) but suffer from phase transitions, transition-metal (TM) migration, and structural distortions that limit cycling stability. Here, we combine first-principles modeling and electrochemical measurements to elucidate how configurational entropy governs their structural and electronic response. By comparing low-, medium-, and high-entropy compositions, we show that higher configurational entropy mitigates TM-centered octahedral distortions, suppresses shear-type deformations associated with P2 \rightarrow O2 layer gliding, and distributes redox activity across multiple cations (Ni, Co, Fe), avoiding local over-oxidation. Defect-formation analyses reveal that high-entropy mixing significantly discourages out-of-layer TM migration, reducing TM/ Na_{vac} antisite formation and stabilizing the layered framework upon deep desodiation. Consistently, medium- and high-entropy materials exhibit superior capacity retention and structural reversibility compared to the low-entropy analogue, with further performance enhancement when using room-temperature ionic-liquid (RTIL)-based NaFSI-Pyr₁₄FSI electrolyte, which mitigates Mn dissolution and accounts for enhanced efficiency upon cycling. These findings demonstrate that configurational entropy is a powerful design parameter for achieving robust, high-performance P2-type layered cathodes and provide clear guidelines for entropy-assisted materials engineering in next-generation NIBs.

Keywords

High-Entropy Oxide; Na-ion Battery; P2-type Cathode; DFT Calculation; Ionic Liquid.



1. Introduction

Driven by the demand for sustainable and cost-effective storage technologies, secondary Na-ion batteries (NIBs) have re-emerged as promising alternative to Li-ion batteries (LIBs), particularly in sectors where material abundance and low cost are essential, such as grid-level/residential stationary energy storage, light mobility, backup power systems, low-cost electronics and Internet of Things (IoT).¹ Owing to the natural abundance and low cost of sodium, NIBs offer a viable pathway toward greener electrochemical technologies.² In NIBs, the cathode material is the primary factor dictating the achievable energy density, structural stability, and cycling lifetime, as most degradation mechanisms and voltage-limiting processes originate from the layered cathode framework. As such, its design is central to the development of next-generation Na-based storage systems.

Among the various cathode families, layered transition-metal oxides with the general formula Na_xTMO_2 (TM = transition metal) have attracted considerable attention due to their high theoretical capacities and favorable Na^+ transport properties.³ According to the Delmas notation, these materials are categorized into O3, P3, P2, and O2 structures depending on the stacking of the TMO_2 layers and the coordination environment of Na^+ ions.⁴ In particular, P2-type oxides, where Na^+ ions reside in trigonal prismatic sites, enable faster diffusion kinetics compared to O3-type counterparts, making them appealing for high-rate applications.⁵ However, their structural reversibility remains a challenge: repeated Na^+ extraction at high voltages (> 4.2 V) can induce P2 \rightarrow O2 transitions *via* layer gliding or TM migration, leading to capacity fading and voltage hysteresis.^{6,7} Furthermore, Mn-based P2 oxides, though cost-effective and environmentally benign, suffer from Jahn-Teller distortions associated with Mn^{3+} formation at low potentials, causing lattice strain and structural degradation.^{8–10} Strategies to mitigate these issues include compositional tailoring, partial TM substitution, and the introduction of structural disorder to enhance robustness and suppress phase transitions.^{11–14}

Recently, the high-entropy concept, originally proposed for multicomponent alloys,¹⁵ has emerged as a promising route to design structurally and chemically resilient electrode materials.¹⁶ By incorporating multiple metal species into a single-phase lattice, a large configurational entropy term can stabilize metastable phases, promote homogeneous cation distribution, and trigger beneficial “cocktail effects”.^{17,18} Both O3- and P2-type high-entropy layered oxides have demonstrated enhanced cycling stability and rate capability compared to



their low-entropy analogues.^{19,20} Nevertheless, the fundamental mechanisms by which configurational entropy influences structure, electronic behavior, and electrochemical stability remain insufficiently understood.

In this work, we address this knowledge gap by combining first-principles calculations and electrochemical characterization to elucidate the role of configurational entropy in the stabilization of P2-Na_xMO₂ cathodes (where M = Mn, Ni, Co, Ti, Mg, Al, Fe). Configurational entropy at the metal site can be easily calculated from the compound's stoichiometry as:

$$S = -R \sum x_i \ln(x_i) \quad (1)$$

Accordingly, compositions are classified as low-, medium-, or high-entropy when the corresponding configurational entropy (S) is lower than $1.0R$, lies in the $1.0R$ - $1.5R$ range, or exceeds $1.5R$, respectively. Herein, three representative compositions with increasing entropy are investigated as model systems. Through DFT and hybrid-DFT analyses, we examine the local structural distortions, electronic structure evolution, and defect formation tendencies upon Na extraction. These computational insights are directly correlated with experimental evidence from X-ray diffraction (XRD), cyclic voltammetry (CV), and galvanostatic cycling, enabling a comprehensive understanding of how entropy governs the structural and electrochemical responses of P2-type cathodes. Consistently, when experimentally tested in lab-scale Na-based electrochemical cells, medium- and high-entropy oxide materials (hereafter referred to as MEO and HEO) exhibit superior capacity retention and structural reversibility compared to the low-entropy analogue (LEO). Further performance enhancement is attained when cycling the MEO and HEO cathode materials using a room-temperature ionic liquid (RTIL)-based electrolyte (*i.e.*, the sodium bis(fluorosulfonyl)imide salt (NaFSI) dissolved in N-butyl-N-methylpyrrolidinium bis(fluorosulfonyl)imide (Pyr₁₄FSI)), suggesting a pivotal role of RTIL solvents in suppressing Mn dissolution and promoting the formation of a denser and more stable solid electrolyte interphase (SEI). Although the reported electrochemical performance does not yet exceed that of state-of-the-art cathodes, this work represents a substantive conceptual advance by delivering new fundamental understanding of entropy-driven stabilization in NIB cathodes, and by articulating clear design principles for next-generation materials. Rather than introducing new compositions, the study reveals previously unexplored mechanistic insights, demonstrating how configurational entropy can be deliberately harnessed to stabilize functional cathode frameworks and guide the rational development of high-performance NIB systems.



2. Experimental Section

2.1. Materials synthesis

The P2-type cathode materials $\text{Na}_{0.68}\text{Mn}_{0.68}\text{Ni}_{0.32}\text{O}_2$, $\text{Na}_{0.68}\text{Mn}_{0.44}\text{Ni}_{0.19}\text{Co}_{0.25}\text{Ti}_{0.09}\text{Mg}_{0.03}\text{O}_2$, and $\text{Na}_{0.68}\text{Mn}_{0.44}\text{Ni}_{0.19}\text{Co}_{0.19}\text{Ti}_{0.09}\text{Mg}_{0.03}\text{Al}_{0.05}\text{Fe}_{0.01}\text{O}_2$, respectively low-entropy, medium-entropy and high-entropy (namely, LEO, MEO and HEO) were synthesized *via* solid-state reactions. In the first step, stoichiometric amounts of the precursors Na_2CO_3 (Acros Organics, 99.95%), MgO (abcr GmbH, 99.5%), Al_2O_3 (Alfa Aesar, 99.5%), TiO_2 (Alfa Aesar, 99.5%), Mn_3O_4 (abcr GmbH, 97%), Fe_3O_4 (Alfa Aesar, 97%), Co_3O_4 (Alfa Aesar, 99%) and NiO (Alfa Aesar, 99%) were thoroughly mixed by ball milling at 400 rpm for 2 h and compacted into pellets under 5 tons. The pellets were then heated to 500 °C for 2 hours and calcined at 900 °C for 12 hours under air. The final product was obtained by naturally cooling to room temperature and then kept inside an Ar-filled glovebox ($\text{O}_2 < 0.1$ ppm, $\text{H}_2\text{O} < 0.1$ ppm) to prevent further air exposure.

2.2. Computational details and structural models

The series of P2-type layered oxide cathodes, LEO, MEO and HEO, was modeled using three $8 \times 4 \times 1$ supercells of $\text{Na}_{0.68}\text{MO}_2$ parent material containing 236 atoms within the $P6_3/mmc$ hexagonal space group. Mixed metal occupancy at the 2a Wyckoff site (*i.e.*, the M one) introduces configurational disorder, which was simulated *via* the special quasi-random structure (SQS) approach, as implemented in the *mcsqs* module of the Alloy Theoretic Automated Toolkit (ATAT) package.^{21,22} For each material, SQS optimization was performed by matching pair and triplet correlation functions up to a maximum triplet radius of 5.0 Å. The final structures exhibit correlation mismatches below ~ 0.03 for all targeted clusters, demonstrating a close approximation to a fully random alloy. Na atoms have been placed in edge and face sites with Na(e)/Na(f) ratio being close to 2. To simulate the cathode charge, we model the P2- Na_xMO_2 series by varying the sodium content with $x_{\text{Na}} = 0.68, 0.50, 0.30, 0.12$, with desodiation upon charge representing different States of Charge (SoC). Spin-polarized density functional theory (DFT) calculations have been performed with the DFT+U Hubbard-like correction scheme to overcome the large self-interaction error (SIE) that affects DFT when applied to mid-

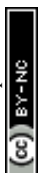


to-late first row TM oxides with unpaired tightly localized electrons in *d* orbitals. Projector-augmented wave (PAW) potentials and plane wave (PW) basis sets have been used, as implemented in the Vienna Ab-initio Simulation Package (VASP) code (version 6.4.2).^{23–25} For all the calculations, the following PAW potentials have been considered: Na_pv [Be]2p⁶3s¹; Mn [Ar]3d⁷; Ni [Ar]3d¹⁰; Co [Ar]3d⁹; Ti [Ar]3d⁴; Mg [Ne]3s²; Al [Ne]3s²3p¹; Fe [Ar]3d⁸; O[He]2s²2p⁴. The Perdew-Burke-Ernzerhof (PBE) exchange-correlation functional has been employed with $U_{\text{eff}} = 4.0$ eV parameter for Mn, Ni, Co, and Fe atoms. This choice of a common effective U value has been validated against hybrid DFT in similar systems.²⁶ The D3-BJ dispersion correction was also added to account for van der Waals (vdW) interactions that play a crucial role in layered structures.^{27–29} A kinetic cut-off energy of 600 eV and Γ -point only sampling have been used; these values ensure converged energies within 3 meV/f.u. with respect to the PW basis set size and Brillouin zone sampling, respectively. For all the self-consistent field calculations, the convergence threshold for total energy has been set to 10^{-5} eV. Lattice constants and atomic positions have been fully relaxed until the maximum forces acting on each atom were below 0.03 eV/Å. Single-point calculations have been carried out at the hybrid DFT level of theory with the range-separated Heyd-Scuseria-Ernzerhof density functional with a screening parameter μ set to 0.2 \AA^{-1} (*i.e.*, HSE06), and analysis reported in terms of projected density of states (PDOS) and net magnetization on TM atoms.³⁰

2.3. Structural and electrochemical characterization

The crystalline structure of the samples was analyzed by X-ray diffraction (XRD), using a Xpert3 MRD Panalytical diffractometer, equipped with Cu K α radiation source. The XRD patterns were collected in Bragg-Brentano geometry in the 2θ range of 30° - 80° . Rietveld refinement was performed with FullProf Suite software.

For electrochemical tests, electrodes were prepared with a composition of 70 wt.% of active material, 20 wt.% of C-65 (from Imerys) as conductive carbon additive, and 10 wt.% polyvinylidene fluoride (PVdF, Solef 5130 from Solvay) as binder. The three components were mixed in dry conditions and then dispersed in N-methyl pyrrolidone (NMP, from Sigma-Aldrich), forming a homogeneous slurry (solid content ≈ 30 %) that was then cast onto the Al current collector (from MSE Supplies) using a doctor-blade (indicative thickness of 200 μm of wet film). NMP solvent was removed by evaporation at ambient temperature and further drying

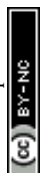


under dynamic vacuum at 120 °C for 24 h prior to be transferred into an Ar-filled glovebox (MBraun, O₂ and H₂O levels < 1 ppm). The thickness reduces to about 50 μm, with an active mass loading in electrodes of ca. 3 mg cm⁻². Na-metal cells were assembled within ECC-Std test cells (EL-Cell GmbH), using metallic Na (from Sigma-Aldrich) as counter electrode, a Whatman GF/A glass wool disk as the separator, and as electrolyte either NaClO₄ 1 M in propylene carbonate (from Solvionic) or NaFSI in Pyr₁₄FSI (molar ratio of 1:4, from Solvionic). Cyclic voltammetry (CV) was performed on a VMP3 workstation (BioLogic Instruments) using Swagelok-type 3-electrode cells in the range from 1.9 to 4.2 V vs. Na⁺/Na with a scan rate of 0.1 mV s⁻¹. Galvanostatic cycling tests were carried out at ambient laboratory temperature (≈ 25 °C) with an ARBIN BT2000 battery tester, from 2.0 to 4.2 V vs. Na⁺/Na, setting the current densities on the basis of an expected capacity of 100 mAh g⁻¹. For both CV and galvanostatic cycling test the cut-off voltage was set to 4.2 V vs. Na⁺/Na to avoid any irreversible phase transitions, as already reported in previous studies for analogous materials at very high voltages.³¹ In this way, we could compare the redox behavior of the three materials under the same sodiation/desodiation conditions; as a matter of fact, the behavior of the three materials is expected to greatly diverge at high voltages, as shown in the sodiation/voltage profile computed with DFT (see Figure S2 in SI)). The reproducibility of the electrochemical measurements was verified by testing additional cells under identical conditions. Therefore, representative results are reported in the manuscript.

3. Results and Discussion

The key properties of LEO, MEO and HEO are hereby addressed in terms of their physicochemical and electrochemical characteristics, with the primary aim of elucidating the role of configurational entropy in determining their stability, performance, and potentiality for future applications in NIBs.

In the following, we provide an in-depth analysis on the structural and electronic features of LEO, MEO and HEO, as well as their evolution upon charging, as revealed by first-principles calculations. Furthermore, we assess their electrochemical performance in laboratory-scale NIB cells employing both standard electrolytes based on organic solvents (NaClO₄ 1M in PC) and RTIL-based electrolytes (NaFSI in Pyr₁₄FSI 1:4). Our goal is to emphasize the influence of



mixing entropy on the structural stability of layered oxide cathodes, and to showcase promising implementations using innovative electrolytes for high-performance energy storage devices.

Figure 1a displays the XRD patterns of the three as-synthesized samples, the diffraction peaks of which are all assigned as reflections of $P6_3/mmc$ hexagonal space group.^{32,33} The results of Rietveld refinement (perfectly aligned with previous works on these materials)¹⁶ are reported in Figure S1 and Table S1, pointing out a slight shrinkage of the cell volume with increasing entropy (from 80.8 to 79.9 Å³ for LEO and HEO, respectively). The first-principles characterization shows a good agreement with XRD data, the lattice constants being optimized at PBE+U-D3BJ level of theory within <1% error deviation from experiments (as shown in Figure 1b). To account for mixing entropy and reliably model homogeneously distributed arrangements, the Na/vacancy and metal atoms orderings are incorporated into the crystalline lattice using the SQSgen code, which leads to the geometry solution corresponding to the maximum configurational entropy (*i.e.*, special quasi-random structures, SQS).³⁴ The SQS results for the three systems within the MO₂-layers can be visualized in Figure 1c.

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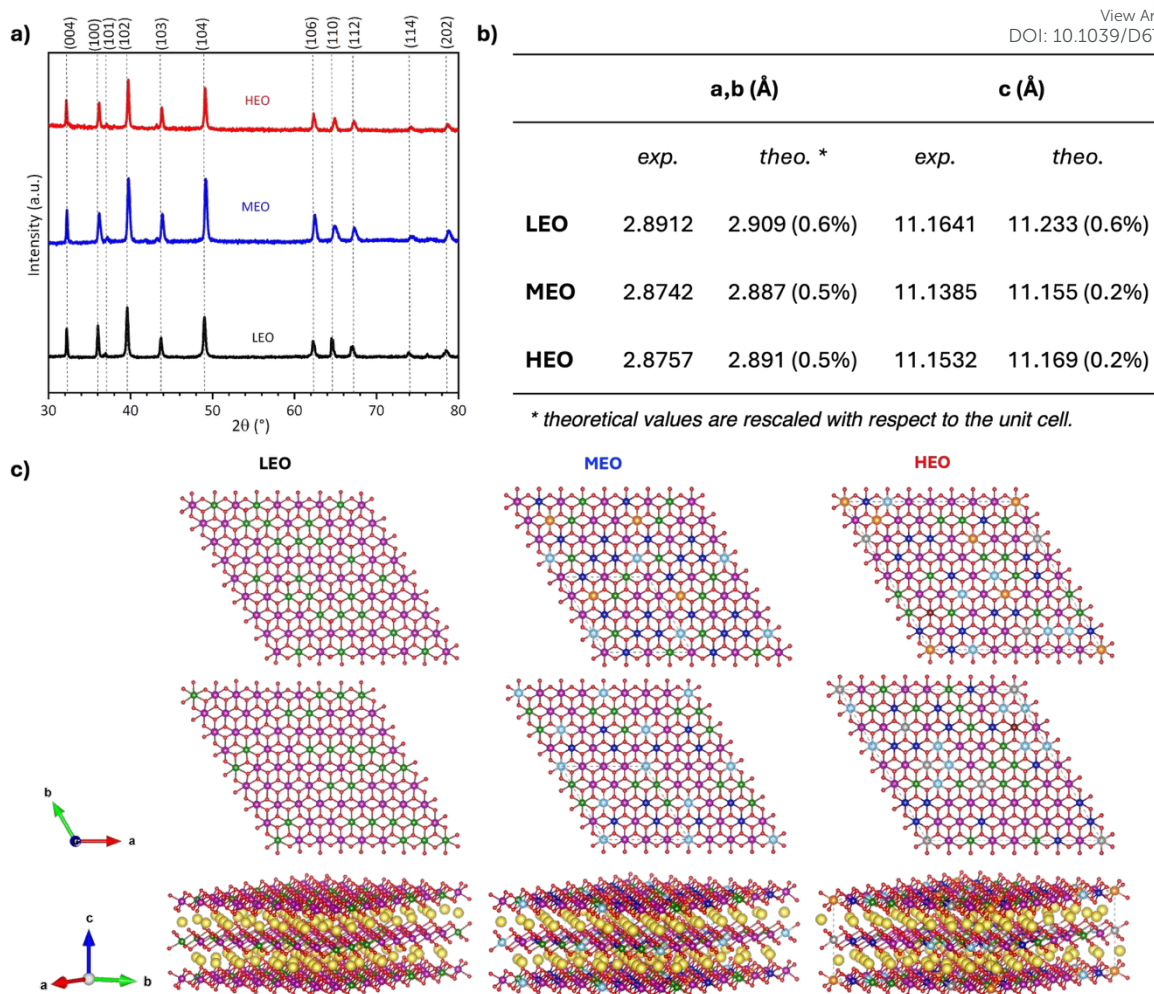


Figure 1. **a)** XRD diffractograms of the three $P2-Na_xMO_2$ samples, labelled as LEO, MEO and HEO, with all the reflections indexed according to $P6_3/mmc$ hexagonal space group. **b)** Lattice parameters of the three samples as obtained from Rietveld refinement and PBE+U-D3BJ level of theory. Percentage error deviation is reported in parentheses. **c)** Structural theoretical models for $P2-Na_xMO_2$ compounds: **(top)** random distribution of metal atoms within the MO_2 -layers; **(bottom)** 3D view of the hexagonal cells in the $P6_3/mmc$ space group. Atoms are represented as spheres. Colour code: Na, yellow; Mn, violet; Ni, green; Co, blue; Ti, cyan; Mg, orange; Al, grey; Fe, brown; O, red.

3.1. Charge compensation mechanisms regulated by configurational entropy

Once the model structure was established, theoretical investigation has been carried out to address the electrochemical behavior of the explored cathodes, namely the evolution of their electronic structure at decreasing sodiation, mimicking the charge/discharge process (the voltage/sodiation correlation is shown in Figure S2, whereas Table S2 reports the corresponding



lattice parameters). By analyzing the magnetization trend, we could derive the oxidation states of each transition metal site at different sodium contents, thus assigning the involved redox couples at each voltage step. It is noteworthy that such compositionally diverse oxides containing multiple metal atoms with intricate and non-trivial spin configurations call for a proper treatment of the strongly localized *d*-electrons. Accordingly, we have performed hybrid DFT calculations with the HSE06 functional, which ensures a reliable description of the spin states across all the investigated systems (see PDOS in Figure S3 of SI). As reported in Figure 2, Mn is predominantly found in mixed Mn³⁺/Mn⁴⁺ oxidation states across all systems, becoming fully oxidized to Mn⁴⁺ at $x_{\text{Na}} = 0.30$. The few Mn³⁺ at high sodium content could act as Jahn-Teller (JT) active centers, and thus may be responsible for structural distortion upon cycling. This aspect will be further highlighted in the following discussion (see Section 3.3). On average, however, Mn oxidation can be considered a minor contribution in all systems (see Figure S4 in SI). In contrast, Ni exhibits a clear and progressive oxidation from Ni²⁺ to Ni³⁺, and further to Ni⁴⁺ in MEO and HEO compositions at the highest degree of desodiation. Again, the presence of JT-active Ni³⁺ species can result in substantial octahedral distortions, as clarified below. Similarly, Co undergoes sequential oxidation from Co³⁺ to Co⁴⁺ in both MEO and HEO at $x_{\text{Na}} = 0.30$. Finally, in HEO, Fe also participates in the redox process through a Fe³⁺ → Fe⁴⁺ oxidation upon sodium removal. Overall, these results demonstrate that Ni, Co and Fe are the primary redox active species responsible for charge compensation during desodiation. Because multiple species share the redox burden, the systems avoid local over-oxidation and evenly distribute not only structural but also electronic changes. This facilitates redox processes and enables more stable electrochemical response of higher-entropy compounds (MEO and HEO) compared to LEO.

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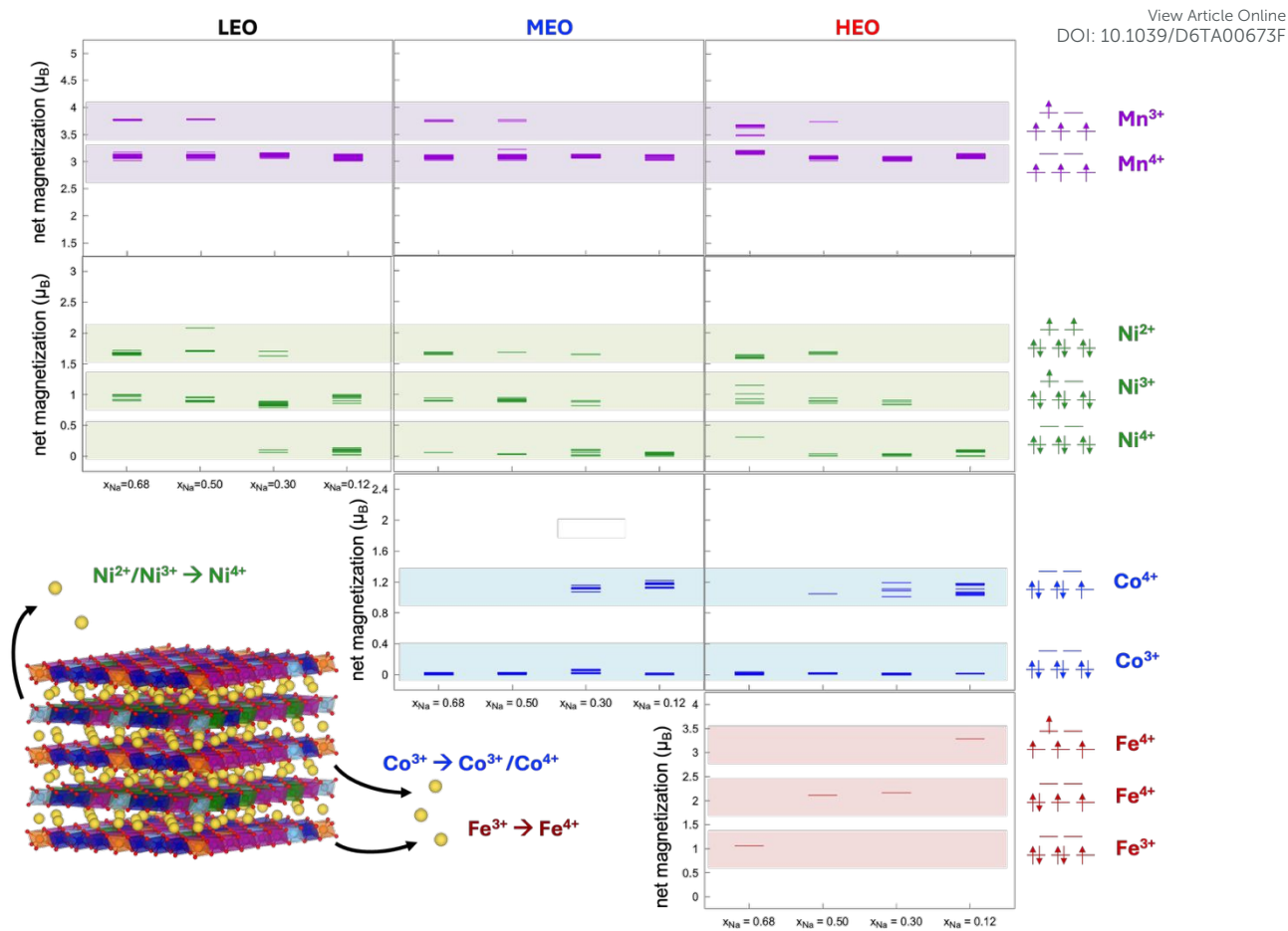


Figure 2. Electronic structure analysis: net magnetization of Mn, Ni, Co and Fe atoms in LEO, MEO and HEO as function of sodiation degree computed at HSE06 level of theory. Electronic configurations are declared. Inset: pictorial representation of a $P2\text{-Na}_x\text{MO}_2$ lattice with the redox couples involved in the charge compensation upon Na^+ extraction. Color code as in Figure 1, Na yellow. Polyhedral visualization is adopted for the MO_6 units.

The theoretical prediction of the redox active species responsible for charge compensation upon desodiation in the three materials is well aligned with the cyclic voltammetry (CV) profiles experimentally measured in Na-based cells employing a standard electrolyte (*e.g.*, NaClO_4 1M in PC). As visible in Figure 3 (panels a, b and c), all samples exhibit a first oxidation peak (at 2.0 V *vs.* Na^+/Na for LEO, and at 2.2 V for MEO and HEO) corresponding to the partial conversion of Mn^{3+} to Mn^{4+} ,³⁵ which is consistent with the detection of Mn^{4+} species from DFT even at high Na content ($x_{\text{Na}} = 0.68$ corresponds to ca. 2.5 V *vs.* Na^+/Na according to the capacity-voltage profiles in Figure S2).



At higher potentials (*i.e.*, increasing desodiation degree), the CV profile of LEO remarkably differs from the other two (Figure 3a *vs.* 3b and 3c): the Ni²⁺/Ni³⁺ transition at 3.0 V is barely visible,³⁶ whereas the rest of the profile is dominated by very intense and sharp peaks due to reversible yet significant structural changes. As reported by Islam *et al.*, *in situ* synchrotron XRD analysis on analogous materials revealed progressive phase transitions from P2 (*P6₃/mmc*) to P'2 (*Cmcm*) and finally to OP4 (*P6₃*) taking place in this voltage range,³⁵ which were classified by Yan *et al.* as Na⁺ ions/vacancies ordering processes.³⁷ On the other hand, the CV profiles of MEO and HEO (Figure 3b and 3c, respectively) suggest a more extended redox participation, due to the presence of many redox active species, partially overlapping with each other in the voltage range from 3 to 4 V *vs.* Na⁺/Na. In particular, both samples display a sharp oxidation peak at 3.0 V due to Ni²⁺/Ni³⁺ transition,³⁶ which is followed in the case of MEO by a broad oxidation signal centered at 3.25 V, compatible with Co³⁺/Co⁴⁺ transition,³⁸ such signal is possibly present also in HEO profile, but in that case it is overlapping with a prominent oxidation peak at 3.35 V assigned to Fe³⁺/Fe⁴⁺ transition. All these transitions are in fair agreement with the predicted oxidation from the computational results reported in Figure 2, taking into account the correlation between sodiation degree and voltage shown in Figure S2. The CV profiles of MEO and HEO also exhibit an oxidation peak at 3.6 V that is reasonably associated to Na⁺ ions/vacancies ordering,³⁷ and both profiles conclude at 4.2 V with an oxidative current that corresponds to the incipient Ni³⁺/Ni⁴⁺ transition; as a matter of fact, DFT results reveal the presence of Ni⁴⁺ at $x_{\text{Na}} = 0.30$, but the whole oxidation peak cannot be properly observed in the profiles of Figure 3b and 3c as it closely lies at the cut-off voltage of the CV measurements.³⁶

Beyond metal-specific contributions to charge compensation, configurational entropy itself clearly modulates the electrochemical response. Indeed, the CV profiles of LEO and MEO/HEO differ not only in the presence of multiple redox peaks associated with different transition metals, but also in the shape of structure-related signals attributed to Na/vacancy ordering. This indicates that mixing entropy can indirectly tune structural stabilization and evolution during cycling, as examined in detail in the following sections.



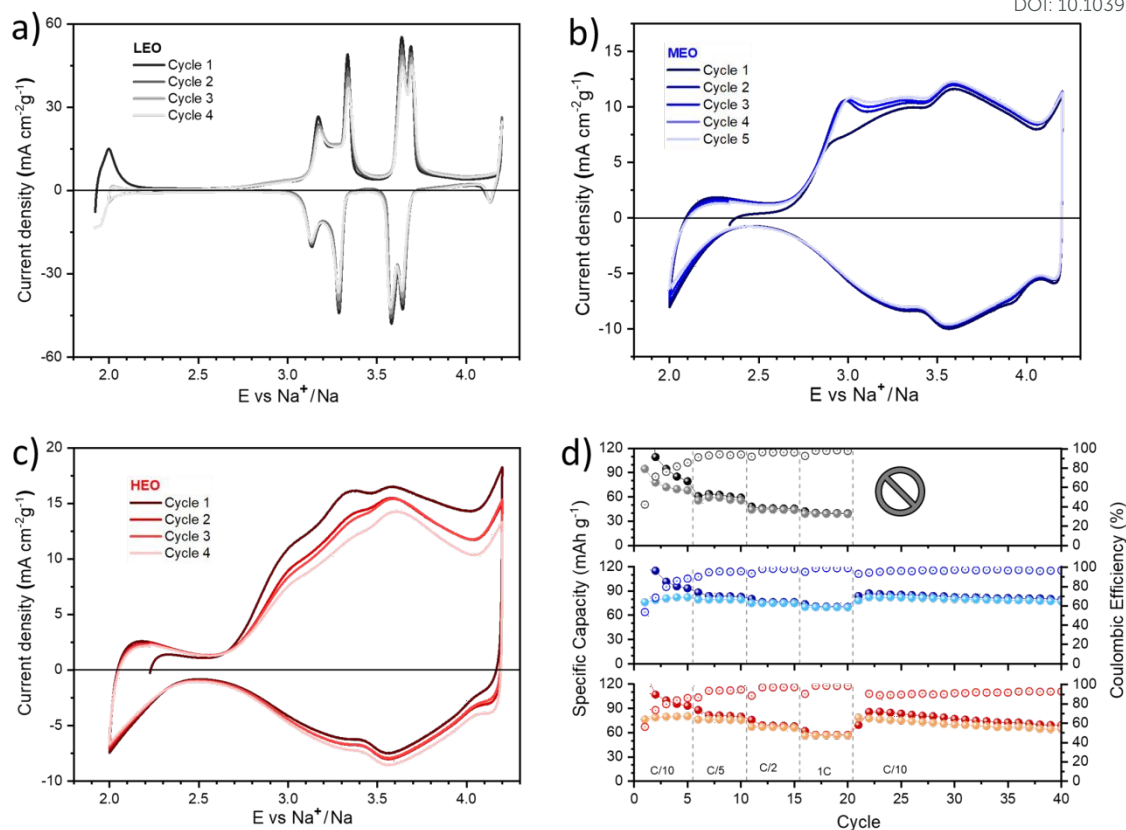


Figure 3. Cyclic voltammetry curves of **a)** LEO, **b)** MEO, and **c)** HEO with standard electrolyte (NaClO_4 1M in PC). **d)** Specific capacity and Coulombic Efficiency of Na-metal cells based on LEO, MEO, and HEO with standard electrolyte, upon galvanostatic cycling at different C-rates (representative voltage profiles vs. specific capacity are reported for each sample in Figure S5).

3.2. Electrochemical performance controlled by configurational entropy

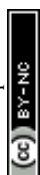
When testing $\text{P2-Na}_x\text{MO}_2$ samples for galvanostatic charge/discharge cycling in combination with standard liquid electrolyte (NaClO_4 1M in PC), it is well evident that the cell with LEO fails after 20 cycles (top panel in Figure 3d): after the rate capability test, upon resetting the current to C/10 regime, the cell gets stuck in charging mode without reaching the targeted voltage because all the electric current is consumed in parasitic reactions, such as Mn dissolution. Indeed, even if not expressly quantified in this work, this is a well-known issue of Mn-containing cathodes for NIBs, typically exacerbated at higher states of charge.³⁹ On the



other hand, MEO and HEO (middle and bottom panels in Figure 3d, respectively) demonstrate to completely recover the initial capacity (ca. 80 mAh g⁻¹) after the rate capability test and stably sustain charge/discharge cycles up to the end of the testing protocol (although with a rather limited Coulombic Efficiency (CE), viz. 97 and 93% for MEO and HEO, respectively). However, also MEO and HEO, tested in these conditions, suffer from high initial polarization, resulting in a CE lower than 90% during the first 6 cycles.

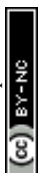
Clearly, managing large compositional diversity arising from metal mixing introduces complexity into the system, which can make any prediction or understanding of electrochemical behavior more difficult. Notwithstanding the beneficial effects that high entropy can induce by delocalizing the electronic response, individual metal species may also promote undesired processes.⁴⁰ In principle, a metal ion can migrate into the sodium layer and occupy a vacant Na site, either created during cycling or intrinsically present in Na-deficient formulations.⁴¹ The resulting metal-Na_{vac} antisite defect disrupts the ideal alternating Na-layer/M-layer stacking, leading to partially mixed cation layers.⁴² Such defects can be detrimental for the battery performance by causing capacity loss, voltage hysteresis and structural degradation: (i) when sitting in Na sites, the metal ion physically blocks the diffusion channels and lowers the ionic conductivity; (ii) M-rich local environments along Na⁺ pathways can hinder subsequent Na⁺ re-insertion, leading to irreversible capacity loss and voltage hysteresis; (iii) metal ion migration can also trigger layer gliding (*e.g.*, P-to-O transitions) and generate local strain that promote structural transformations, micro-cracking and long-term instability; (iv) migrated metal ions can diffuse themselves into the electrolyte and promote detrimental ion dissolution. For all these reasons, the tendency of a P2-Na_xMO₂ cathode to release metals from their layers can be considered as an early step toward degradation processes. To clarify this aspect, we focus on the metal ion out-of-layer migration in our investigated P2-Na_xMO₂ cathodes, so as to unveil whether the increasing entropy may prevent it, thus inducing major stability toward degradation.

The migration of a metal ion into a sodium vacant site is simulated as the formation of a metal/Na_{vac} antisite defect under low-sodiation conditions ($x_{\text{Na}} = 0.30$ and 0.12). We focus on Mn and Ni as the most abundant metals in MEO and HEO formulations and the only species present in LEO. Figure 4 depicts the energy landscape associated to the process: the overall energetics of antisite formation becomes more favorable at lower sodium content ($\Delta E \sim -0.7$ to 4.6 eV *vs.* -5.3 to 1.6 eV at $x_{\text{Na}} = 0.30$ and 0.12 , respectively), reflecting the stronger



interactions established to compensate for the increased number of Na⁺ vacancies. We should point out that not only negative values of ΔE , but also slightly positive energy variations would still be accessible under the battery operating conditions (*i.e.*, under an applied voltage upon charge), indicating that most of these defects are likely to occur. Across the series, LEO easily accommodates the antisite defects, with the Mn being particularly stabilized at $x_{\text{Na}} = 0.12$ due to a 4-fold coordination with the oxygen atom exposed from the opposite MO₂-layer. A similar stabilization occurs for the Mn antisite defect in MEO already at $x_{\text{Na}} = 0.30$, which is even more pronounced in the 5-fold coordination attained at $x_{\text{Na}} = 0.12$. Together with the low-energy Ni antisite defects, MEO clearly allows easy metal out-of-layer migrations, which raises concerns regarding its electrochemical durability. However, the Ni antisite detected at $x_{\text{Na}} = 0.12$ retains its Ni⁴⁺ oxidation state, suggesting that the foreseen capacity loss may remain limited. Conversely, the HEO composition offers a noticeably different behavior, with ΔE values shifted upward compared to LEO and MEO. At $x_{\text{Na}} = 0.30$, Mn antisite formation is highly unfavorable, whereas Ni antisites remain energetically accessible. Further desodiation leads to a mostly unchanged picture for Ni antisite, while Mn antisite formation is slightly stabilized owing to a 5-fold coordinated intermediate. To sum up, the low-entropy composition seems highly prone to out-of-layer metal migration, with thermodynamically convenient formation of Mn²⁺/Na_{vac} antisite occurring already at $x_{\text{Na}} = 0.30$. This finding can be fairly regarded as an easy tendency of LEO to undergo Mn dissolution upon cycling, which could be a feasible reason for the cell failure denoted in Figure 3d. Even though the energetics of Mn²⁺/Na_{vac} antisites in MEO seems to suggest accessible intermediates at $x_{\text{Na}} = 0.30$, the established geometries in this case are sensibly different. The higher coordination attained by the migrated Mn²⁺ allows to pin the oxide slabs, reinforce the crystalline structure and thus prevent layers gliding, which is the main driver in the P-to-O phase transition. By enabling the formation of migrated yet peculiar Mn²⁺ sites that act as lattice pillars, the medium-entropy composition seems able to limit structure collapse, as confirmed by the stable capacity showed upon cycling (Figure 3d). Analogously, the steady performance of HEO perfectly aligns with the largely unlikely formation of Mn²⁺/Na_{vac} at the same SoC predicted by DFT, suggesting a mitigated tendency for Mn dissolution and a consequent enhancement of the electrochemical durability with increasing configurational entropy in the cathode formulation. It is worth mentioning that in the antisite defect calculations, the oxidation state of the transition metal was not imposed *a priori*; instead, the electronic structure was determined self-consistently within the DFT framework, allowing

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the metal charge state associated with migration into a Na vacancy to emerge naturally from the calculation.

Here, the indirect influence of configurational entropy becomes even more pronounced: despite involving the same metals (Mn and Ni), the three compositions exhibit markedly different behaviors due to their highly diverse local chemical environments. Out-of-layer migration of Mn and Ni into Na vacancies is strongly modulated by configurational entropy, with increasing metal mixing progressively disfavoring the formation of TM/Na_{vac} antisite defects. This highlights the role of entropy in promoting structural robustness by suppressing detrimental cation migrations, which can lead to Mn dissolution and cathode degradation.

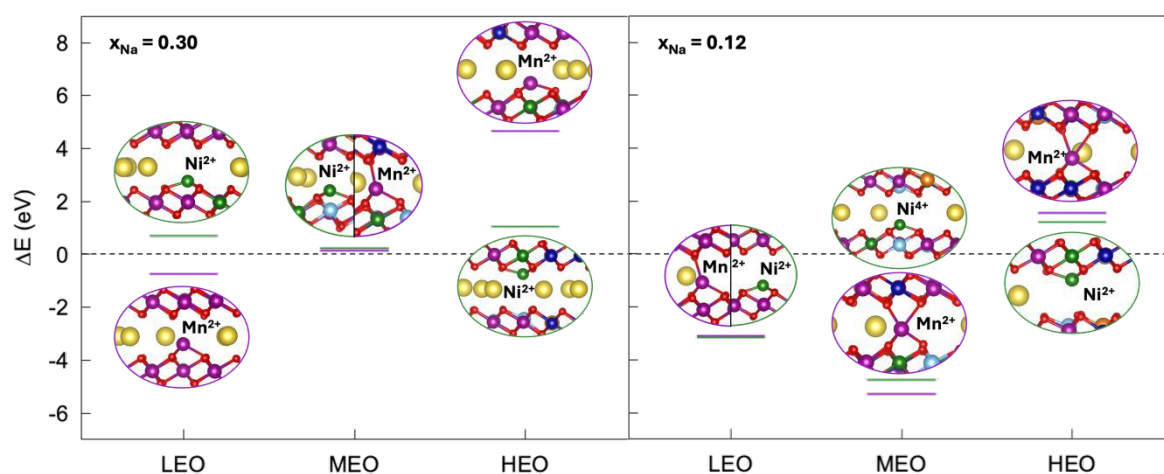


Figure 4. Mn and Ni out-of-layer migration, i.e., M/Na_{vac} antisite defects. ΔE is the formation energy of each defect, calculated as the energy difference between the defective and the pristine structure, plotted for LEO, MEO and HEO at (left) $x_{Na} = 0.30$ and (right) $x_{Na} = 0.12$. Insets: structural detail of the optimized configurations. Oxidation states of migrated Mn/Ni are declared. Color code as in Figure 1.

3.3. Structural effects of configurational entropy

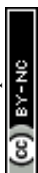
With the scenario depicted so far, we aim at further elucidating the electrochemical performance, stability and durability of our cathode systems. The use of RTIL-based electrolytes is known to beneficially mitigate the detrimental Mn dissolution,⁴³ by avoiding the formation of the convenient solvation shell around Mn²⁺ that in the case of carbonate solvents



can compete with and even exceed the solvation of Na^+ ions.⁴⁴ Therefore, we employ NaFSI in Pyr₁₄FSI (1:4) as electrolyte in combination with the three cathode materials and perform additional galvanostatic cycling tests. The performance of LEO under these conditions considerably improves, with an almost stable capacity of 90 mAh g⁻¹ in the first 5 cycles at C/10 (with a CE of 97 %) and a good capacity retention after the rate capability test (around 70 mAh g⁻¹). The capacity strongly reduces at higher currents (60, 28, and barely 8 mAh g⁻¹ at C/5, C/2, and 1C, respectively). Also, in the case of the RTIL-based electrolyte, much better performances are achieved by MEO and HEO (Figure 5b and 5c, respectively). HEO, in particular, displays an initial capacity of 83 mAh g⁻¹ at C/10 with a CE above 99% starting from the fourth cycle. However, a decrease of the capacity is observed upon increasing current (64, 52 and 42 mAh g⁻¹ at C/5, C/2 and 1C, respectively), a decrease even more pronounced than in the case of standard electrolyte (Figure 3c) reasonably due to the higher viscosity of RTILs compared to carbonates. HEO completely retains the initial capacity upon resetting the current regime to C/10, maintaining a stable performance in terms of capacity and CE (> 99 %) up to the end of the cycling test (final capacity at the 40th cycle of 76 mAh g⁻¹).

The different response of the three materials in the galvanostatic cycling test is reflected also in the *post-mortem* XRD patterns, collected by disassembling the cells in the glovebox after 40 charge/discharge cycles. The XRD pattern of HEO (Figure 5f) is perfectly maintained, demonstrating the high structural stability of the material that reversibly sustains Na^+ insertion/extraction. In the pattern of MEO (Figure 5e), barely noticeable shifts of some peaks reveal minor changes in the lattice parameters; in particular, the left-shifted peaks related to lattice parameter *c* (such as (004) and (106)) and the right-shifted peaks related to *a* and *b* parameters (such as (100), (110) and (112)) indicate a minimal anisotropic cell distortion, with *c*-axis elongation and basal-plane contraction. On the contrary, the pattern of LEO (Figure 5d) reveals a general decrease in the crystallinity of the material and the coexistence of multiple phases (as previously discussed for the redox processes observed in CV profile in Figure 3a); for instance, the OP4 phase is well evident with (004) reflection at 31.5°.⁴⁵

Overall, these results confirm the beneficial effect of RTILs as electrolyte solvents with the investigated cathode materials in Na-based cells, especially at low current densities, where longer time usually allows the parasitic reactions to become more relevant and even compete with the main operating redox processes (at high currents this issue is negligible because of the very high rate of the main processes). With CE normally decreasing as current density



decreases, the improved performance under low current regime clearly indicates a key role of the RTIL-based electrolyte: the highly electrochemically stable and non-volatile RTIL-based electrolyte can efficiently suppress the unwanted side reactions by acting on several levels. First, they discourage ion dissolution and formation of intermediate species that typically feed parasitic reactions, including self-decomposition, which is largely reduced thanks to their high chemical stability. The promotion of a denser and more stable SEI induced by RTILs is also desirable to physically screen and protect the electrode surface and thus to block long-term side reactions. The distinct solvent properties of RTILs can also reduce the solubility of by-products into the electrolyte, confining them on the electrode surface.

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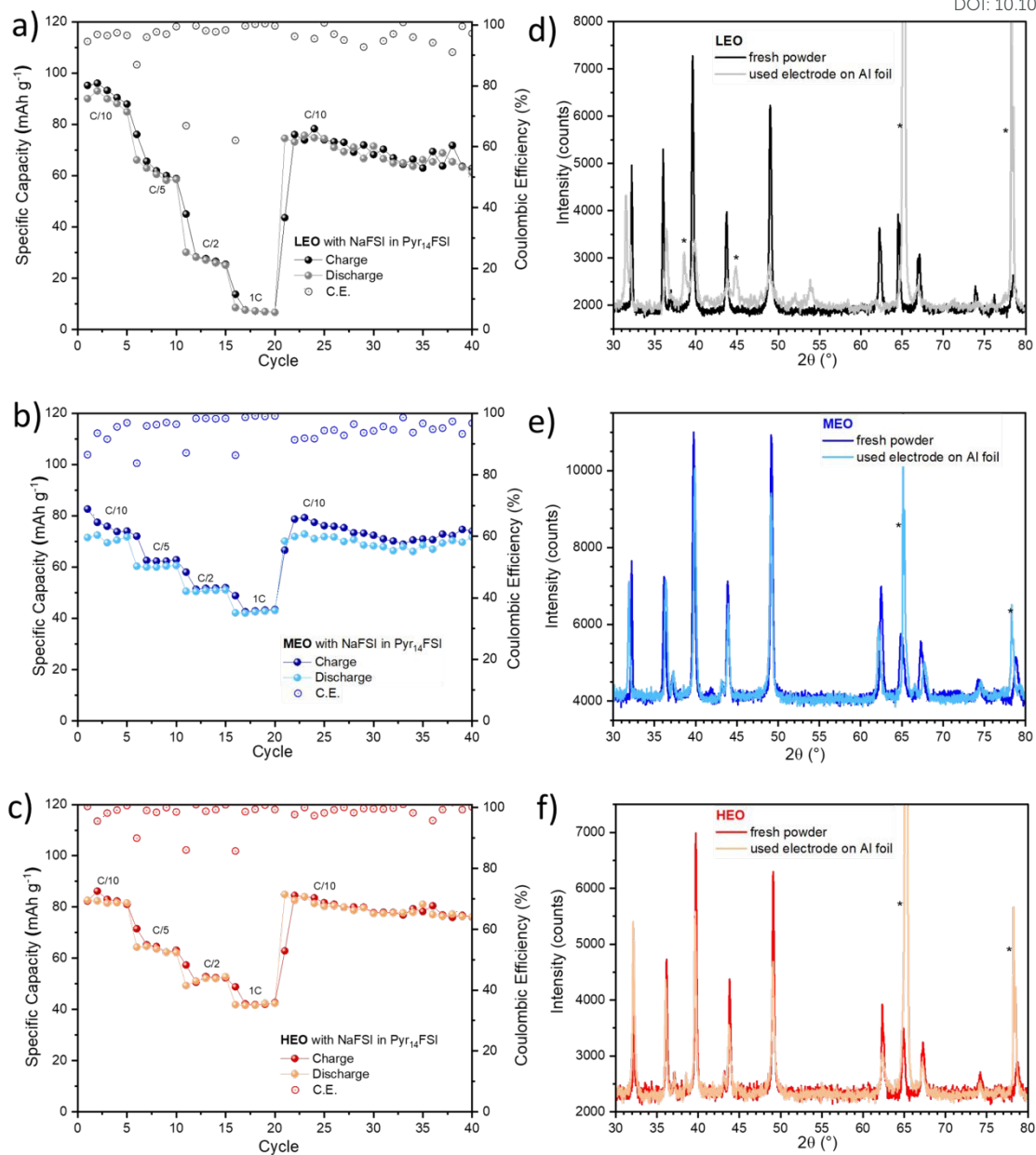
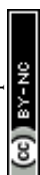


Figure 5. Specific capacity and Coulombic Efficiency of Na-metal cells containing **a)** LEO, **b)** MEO, and **c)** HEO in combination with RTIL-based electrolyte (NaFSI in Pyr₁₄FSI 1:4), upon galvanostatic cycling at different C-rates (representative voltage profiles vs. specific capacity are reported in Figure S6). XRD patterns of **d)** LEO, **e)** MEO, and **f)** HEO, comparing the fresh powder and the electrode after galvanostatic cycling with RTIL-based electrolyte (the reflections due to the Al foil current collector are marked with an asterisk).



Provided the mitigated Mn dissolution owing to RTIL-based electrolyte, we need to understand the enhanced structural retention featured by the higher-entropy compounds. The Na-O coordination illustrated in Figure S7 of SI suggests that Na⁺ ordering is almost constant among the three compounds across the explored sodiation range. On the other side, the pair distribution functions, *i.e.*, PDFs, for M-O coordination are largely affected by desodiation (see Figure S8 in SI). Therefore, we dissect the contribution of each metal sublattice to the overall structural evolution upon charge, that is the analysis of distortion modes within any MO₆ octahedron. Indeed, Na⁺ extraction from the layered crystal structure upon cathode charging can induce variations in both cell volume and geometry, potentially affecting phase stability and mechanical integrity. Whether such distortions arise primarily from oxidation of the central M atom or depletion of the Na layer, different metals may exhibit distinct distortion behaviors. The degree of octahedral tilting is evaluated through variations in both M-O bond lengths and O-M-O bond angles within the MO₆ octahedron at different sodiation levels. Following the definitions by Nagle-Cocco and Dutton, the variations of bond lengths and bond angles accounting for the octahedral tilting can be quantified in terms of the six VanVleck modes (Q1, ..., Q6). By extracting these quantities with the VanVleckCalculator for each M-centered octahedron from our PBE+U-D3BJ minimum-energy structures, we could calculate the average distortion magnitude (ρ , that contains the information on bond length variations) and the fraction parameter (η , which relates to the angular motions) as function of desodiation (as shown from the equations displayed on the right-hand side of Figure 6).^{46,47} In LEO (left panels), Ni-based octahedra exhibit increasing ρ values up to $x_{\text{Na}} = 0.30$, followed by a sharp decrease at $x_{\text{Na}} = 0.12$, while η increases steadily upon desodiation (the PDF, reported in Figure S8, shows significant shifts to lower Ni-O distances with decreasing sodium content, from ~ 2.05 to ~ 1.85 Å). The Mn sublattice contributes minimally to the overall distortion (the PDF in Figure S8 is rather constant upon desodiation, the highest peak lying at ~ 1.90 Å). In MEO (middle panels), the inclusion of Co, Ti and Mg atoms, yielding a medium-entropy composition, does not significantly alter the octahedral distortions associated with bond lengths (as also mirrored in the PDF of Figure S8, with the bond length being kept constant in the range 1.8-2.1 Å), but it modifies the angular displacements. Similarly to LEO, Ni-centered octahedra dominate the ρ trend, with negligible contributions from the other metals. However, Mg and Ti markedly impact the fraction parameter, reducing the contribution of shear-type deformations ($\eta \ll 1$). A similar behavior is observed for HEO (right panels), where Ti-based octahedra limit the angular shear motions and Ni-based ones drive the increasing ρ trend upon charge even



after incorporation of Al and Fe atoms. Because the η trend correlates with the angular motion of the oxide ligands, it serves as an indicator of materials' propensity for layer gliding. As demonstrated by Langella *et al.*,⁶ the P-to-O phase transition is governed by shear-like deformations; thus, lower η values indicate a reduced tendency for phase transformation. Accordingly, increasing configurational entropy seems to enhance cathode phase stability, with MEO and HEO compositions showing lower shear-like structural distortions compared to LEO (see black dashed lines in the three panels). In particular, Ti-centered octahedra play a critical structural role, substantially mitigating distortion effects and promoting structural retention upon desodiation. This analysis aligns well with the XRD patterns of the samples collected before and after the galvanostatic cycling test previously discussed, where HEO and MEO (Figure 5f and 5e, respectively) feature a highly retained structure compared to LEO (Figure 5d).

Also in this case, the functional role of a given metal element appears to be influenced by configurational entropy. For example, the mitigation of shear deformations (reflected by a decrease in η values with increasing SoC) ascribed to Ti sublattice is more pronounced in MEO than in HEO. This suggests that the enhanced structural retention in the HEO composition can be ascribed not only to a metal-specific effect, but also to an additional synergic contribution of entropy-driven lattice stabilization.

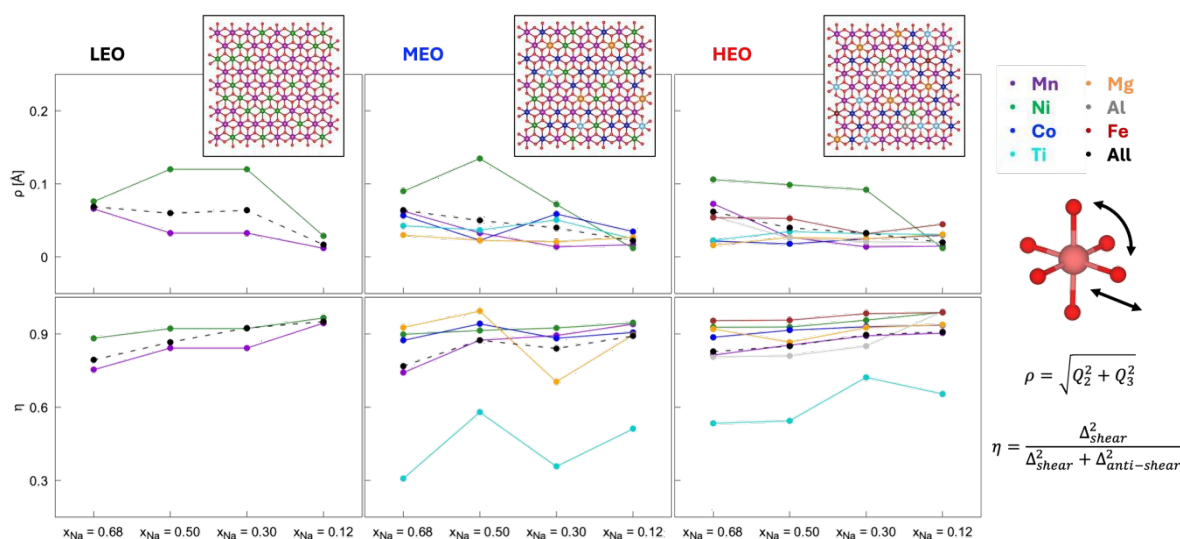


Figure 6. Structural distortion analysis: magnitude of distortion, ρ , and shear fraction parameter, η , calculated for (from left to right) LEO, MEO, and HEO, with the VanVleckCalculator. Equations are reported on the right side of the figure. Insets: structural



detail of the random distribution of metal atoms within the *ab* lattice plane. Color code Mn purple, Ni green, Co blue, Ti cyan, Mg orange, Al gray, Fe brown, O red. Atoms are represented as spheres.

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4. Conclusions

In this work, we demonstrated that combining first-principles modeling and electrochemical analysis can lead to a comprehensive knowledge on how configurational entropy governs the behavior of P2-type Na_xMO_2 layered oxides cathode materials for Na-ion batteries.

At the purpose, three representative compositions with increasing entropy, *i.e.*, $\text{Na}_{0.68}\text{Mn}_{0.68}\text{Ni}_{0.32}\text{O}_2$ (LEO), $\text{Na}_{0.68}\text{Mn}_{0.44}\text{Ni}_{0.19}\text{Co}_{0.25}\text{Ti}_{0.09}\text{Mg}_{0.03}\text{O}_2$ (MEO), and $\text{Na}_{0.68}\text{Mn}_{0.44}\text{Ni}_{0.19}\text{Co}_{0.19}\text{Ti}_{0.09}\text{Mg}_{0.03}\text{Al}_{0.05}\text{Fe}_{0.01}\text{O}_2$ (HEO), have been synthesized *via* solid-state reactions and fully characterized by means of XRD measurements, DFT calculations, cyclic voltammetry and galvanostatic cycling in Na-based cells assembled with both standard NaClO_4 -PC and NaFSI-Pyr₁₄FSI RTIL-based electrolytes.

From our outcomes, configurational entropy is shown to be a key driver on several levels:

- (i) as unveiled from HSE06-resolved electronic structures and CV profiles, increasing entropy can improve the electrochemical response of the cathode material by enabling charge compensation across multiple redox-active species (*i.e.*, $\text{Co}^{3+}/\text{Co}^{4+}$ and $\text{Fe}^{3+}/\text{Fe}^{4+}$ in addition to $\text{Ni}^{2+}/\text{Ni}^{3+}/\text{Ni}^{4+}$), which avoid local over-oxidation, prevent phase transitions (observed instead in the CV of LEO and confirmed by *post-mortem* XRD analysis), reduce voltage hysteresis, thus improving cycling reversibility upon subsequent charging/discharging.
- (ii) DFT results on out-of-layer metal migration suggest that increasing entropy can effectively discourage Mn dissolution (*e.g.*, unfavorable formation of $\text{Mn}^{2+}/\text{Na}_{\text{vac}}$ antisite defects), which is instead highlighted as the main cause of cell failure for the low-entropy formulation. While LEO suffers from rapid degradation with conventional carbonate electrolytes, MEO and HEO materials deliver superior cycling stability and capacity retention. In particular, the best performance is obtained by HEO in combination with the RTIL-based NaFSI-Pyr₁₄FSI electrolyte (92% of capacity retention at the 40th cycle with respect to the initial capacity of 83 mAh g^{-1}), which is consistent with a mitigation of Mn dissolution and with the formation of a dense and stable SEI.



(iii) as evinced from *post-mortem* XRD measurements, increasing entropy can greatly improve structural retention upon cycling. By dissecting each metal contribution to the overall structural evolution upon charge, we elucidate the role of metal mixing in mitigating octahedral distortions occurring upon desodiation and retaining the shear-type deformations that typically drive layer gliding and thus the P-to-O phase transitions.

Overall, configurational entropy emerges as an effective design lever for stabilizing P2-type layered cathodes. This study focuses on a specific set of materials and further efforts would be required to fully assess the transferability of these results, as for example through a systematic exploration of the complete compositional space *via* stepwise elemental incorporation. However, several robust trends already emerge and provide a solid foundation for transferable design guidelines. By implementing broader metal mixing, it is possible to attenuate Jahn-Teller distortions and any octahedral tilting, suppress shear-type lattice deformations linked to P-to-O layer gliding, and distribute charge compensation across multiple redox-active cations. Such collective effects stabilize the host framework and reduce the driving force for detrimental metal ion migration. Beneficial effects are highlighted as metal-specific and may therefore be achieved through appropriate metal doping strategies. For example, Co and Fe act as redox-active elements that promote redox delocalization and stabilize the electronic response, while Ti serves as an electrochemically inert species that suppresses shear-induced structural distortions. However, these mechanisms appear to be further enhanced in the presence of high configurational entropy. Whether exerting direct or indirect effects on the structural and electronic responses to the electrode functioning, mixing entropy engineering can enable robust and durable Na-ion battery performance. These insights provide clear guidelines for the development of next-generation high-entropy layered oxides for sustainable energy storage.

Supporting Information

The Supporting Information is available free of charge and contains:

- XRD pattern and Rietveld refinements
- Lattice parameters obtained from DFT+U calculations
- PDOS and average magnetization obtained from hybrid DFT calculations
- Representative voltage profiles at different C-rate upon galvanostatic charge/discharge cycles
- PDF of Na-O and M-O distances.



Authors Contribution

A.M.: conceptualization, coordination, theoretical modelling and DFT simulations, data analysis and interpretation, manuscript drafting and writing. S.P.: electrochemical measurements, data analysis, figure preparation. M.B.: materials synthesis, conceptualization, coordination, manuscript drafting and writing. A.P.: conceptualization, coordination, electrochemical measurements, data analysis and interpretation, manuscript drafting and writing. H.D.: XRD measurements, data analysis, figure preparation. D.S.: materials synthesis, data analysis and interpretation. G.M.: XRD measurements, data analysis and interpretation. A.B.M.G.: theoretical modelling and DFT simulations, data analysis and interpretation. B.B.: materials synthesis, data analysis and interpretation, manuscript writing. M.P.: theoretical modelling and DFT simulations, data analysis and interpretation, manuscript writing. C.G.: electrochemical measurements, data analysis and interpretation, manuscript writing.

Authors Information

Corresponding Authors

- **Arianna Massaro** - Department of Chemical Science, University of Naples Federico II, Compl. Univ. Monte Sant'Angelo, Via Cintia 21, Naples 80126, Italy; National Reference Centre for Electrochemical Energy Storage (GISEL) - INSTM, Via Giusti 9, Firenze 50121, Italy; ORCID <https://orcid.org/0000-0003-2950-6745>; e-mail: arianna.massaro@unina.it
- **Miriam Botros** - Institute of Nanotechnology, Karlsruhe Institute of Technology (KIT), Hermann-von-Helmholtz Platz 1, 76344 Eggenstein-Leopoldshafen, Germany; ORCID <https://orcid.org/0000-0003-1547-8149>; e-mail: miriam.botros@kit.edu
- **Alessandro Piovano** - GAME Lab, Department of Applied Science and Technology (DISAT), Politecnico di Torino, Corso Duca degli Abruzzi 24, Torino 10129, Italy; National Reference Centre for Electrochemical Energy Storage (GISEL) - INSTM, Via Giusti 9, Firenze 50121, Italy; ORCID <https://orcid.org/0000-0002-5784-6897>; e-mail: alessandro_piovano@polito.it

Authors

- **Silvia Porporato** - GAME Lab, Department of Applied Science and Technology (DISAT), Politecnico di Torino, Corso Duca degli Abruzzi 24, Torino 10129, Italy;



National Reference Centre for Electrochemical Energy Storage (GISEL) - INSTM, Via Giusti 9, Firenze 50121, Italy; ORCID <https://orcid.org/0009-0009-7599-2087>

- **Hamideh Darjazi** - GAME Lab, Department of Applied Science and Technology (DISAT), Politecnico di Torino, Corso Duca degli Abruzzi 24, Torino 10129, Italy; National Reference Centre for Electrochemical Energy Storage (GISEL) - INSTM, Via Giusti 9, Firenze 50121, Italy; ORCID <https://orcid.org/0000-0002-3255-3887>
- **David Stenzel** - Institute of Nanotechnology, Karlsruhe Institute of Technology (KIT), Hermann-von- Helmholtz Platz 1, 76344 Eggenstein-Leopoldshafen, Germany
- **Giuseppina Meligrana** - GAME Lab, Department of Applied Science and Technology (DISAT), Politecnico di Torino, Corso Duca degli Abruzzi 24, Torino 10129, Italy; National Reference Centre for Electrochemical Energy Storage (GISEL) - INSTM, Via Giusti 9, Firenze 50121, Italy
- **Ana B. Muñoz-García** - National Reference Centre for Electrochemical Energy Storage (GISEL) - INSTM, Via Giusti 9, Firenze 50121, Italy; Department of Physics “E. Pancini”, University of Naples Federico II, Compl. Univ. Monte Sant’Angelo, Via Cintia 21, Naples 80126, Italy; ORCID <https://orcid.org/0000-0002-9940-7358>
- **Ben Breitung** - Institute of Nanotechnology, Karlsruhe Institute of Technology (KIT), Hermann-von- Helmholtz Platz 1, 76344 Eggenstein-Leopoldshafen, Germany
- **Michele Pavone** - Department of Chemical Science, University of Naples Federico II, Compl. Univ. Monte Sant’Angelo, Via Cintia 21, Naples 80126, Italy; National Reference Centre for Electrochemical Energy Storage (GISEL) - INSTM, Via Giusti 9, Firenze 50121, Italy; ORCID <https://orcid.org/0000-0001-7549-631X>
- **Claudio Gerbaldi** - GAME Lab, Department of Applied Science and Technology (DISAT), Politecnico di Torino, Corso Duca degli Abruzzi 24, Torino 10129, Italy; National Reference Centre for Electrochemical Energy Storage (GISEL) - INSTM, Via Giusti 9, Firenze 50121, Italy; ORCID <https://orcid.org/0000-0002-8084-0143>

Notes

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Data Availability

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All data supporting the findings of this study are available within the article and its Supplementary Information.

