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Enhanced Cycling Performance of Ni-rich Positive Electrodes (NMC) in Li-ion Batteries by Reducing Electrolyte Free-solvent Activity

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Abstract

The interfacial (electro-)chemical reactions between electrode and electrolyte dictate the cycling stability of Li-ion batteries. Previous experimental and computational results have shown that replacing Mn and Co with Ni in layered LiNi\textsubscript{x}Mn\textsubscript{y}Co\textsubscript{1-x-y}O\textsubscript{2} (NMC) positive electrodes promotes the dehydrogenation of carbonate-based electrolytes on the oxide surface, which generates protic species to decompose LiPF\textsubscript{6} in the electrolyte. In this study, we utilized this understanding to stabilize LiNi\textsubscript{0.8}Mn\textsubscript{0.1}Co\textsubscript{0.1}O\textsubscript{2} (NMC811) by decreasing free-solvent activity in the electrolyte through controlling salt concentration and salt dissociativity. Infrared spectroscopy revealed that highly concentrated electrolytes with low free-solvent activity had no dehydrogenation of ethylene carbonate, which could be attributed to slow kinetics of dissociative adsorption of Li\textsuperscript{+}-coordinated solvents on oxide surfaces. The increased stability of the concentrated electrolyte against solvent dehydrogenation gave rise to high capacity retention of NMC811 with capacities greater than 150 mAhg\textsuperscript{-1} (77 % retention) after 500 cycles without oxide-coating, Ni-concentration gradients, or electrolyte additives.

1. Introduction

LiNi\textsubscript{x}Mn\textsubscript{y}Co\textsubscript{2}O\textsubscript{2} (NMC) are promising positive electrode materials in high-energy Li-ion batteries for electric vehicle applications.\textsuperscript{1-5} Understanding and controlling the (electro-)chemical reactions at the interface between NMC and the electrolyte is the key to enable stable cycling of these high-energy Li-ion batteries.\textsuperscript{6-8} Increasing Ni content in the NMC series from LiNi\textsubscript{1/3}Mn\textsubscript{1/3}Co\textsubscript{1/3}O\textsubscript{2} (NMC111) to Ni-rich NMC such as LiNi\textsubscript{0.6}Mn\textsubscript{0.2}Co\textsubscript{0.2}O\textsubscript{2} (NMC 622) and LiNi\textsubscript{0.8}Mn\textsubscript{0.1}Co\textsubscript{0.1}O\textsubscript{2} (NMC811) can greatly increase discharge capacity\textsuperscript{1,2,9} but at the expense of capacity loss during cycling,\textsuperscript{1,9-11} accompanied by lower voltage onset of O\textsubscript{2} and CO\textsubscript{2} gas evolution.\textsuperscript{9,12} Even though
upper voltage limits used to cycle Ni-rich NMC are much lower than the electrochemical oxidation voltage limit (~5 V Li) of conventional electrolytes, which typically consist of 1.0 M LiPF$_6$ dissolved in cyclic carbonates such as ethylene carbonate (EC) and linear carbonates (e.g. ethyl methyl carbonate - EMC), greater decomposition of solvent and salt decomposition has been reported for NMC cycling with increased Ni content. The enhanced electrolyte degradation found for Ni-rich oxides has been attributed to greater reactivity between the electrolyte and charged oxide surfaces and/or singlet-oxygen liberated from Ni-rich oxides during charging. Recent density functional theory (DFT) calculations have shown that the thermodynamic driving force for dissociative adsorption of carbonate solvents increases with greater M-O covalency by having more later transition metals such as Ni or less lithium in the layered structure, where carbonate solvents including ethylene carbonate (EC) and ethyl methyl carbonate (EMC) dehydrogenate on surface oxygen of Li$_x$MO$_2$ to form a surface hydroxyl group (e.g. C$_3$O$_3$H$_3^+$–O$_{\text{surface}}$ and H$^+$–O$_{\text{surface}}$). This proposed carbonate dehydrogenation is supported by ex-situ Fourier transform infrared (FT-IR) and Raman spectroscopy experiments while the protic species generated from carbonate dehydrogenation can further react with LiPF$_6$ to produce less-fluorine coordinated species including PF$_3$O and PF$_2$O$_2^-$ as revealed from X-ray photoelectron spectroscopy (XPS) and nuclear magnetic resonance (NMR) results. This proposed mechanism – oxide-mediated electrolyte oxidation – is in agreement with previous reports that tuning Ni-rich NMC surface through a less-M-O-covalent coating or creating a Ni-poor protective layer can greatly increase capacity retention of Ni-rich NMC electrodes. For example, recent FT-IR measurements have shown that an Al$_2$O$_3$-coating on NMC622 can reduce carbonate dehydrogenation and LiPF$_6$ decomposition, which is associated with greater cycling stability than uncoated NMC622. Moreover, altering the electrolyte compositions by including additives or increasing salt concentrations can enhance capacity retention of Ni-
rich NMC. Inaba et al.44–49 have shown that increasing LiBF$_4$ concentration in propylene carbonate (PC) to ~3.5 M (7.25 mol/kg), NMC532 can retain a capacity of 185 mAh/g after 50 cycles to 4.6 $V_{\text{Li}}$ while that in ~1 M decays to 140 mAh g$^{-1}$.45 In addition, by increasing LiBF$_4$ concentration in DMC to ~ 4 M (8.67 mol/kg), NMC811 exhibited a capacity of 190 mAh/g after 50 cycles to 4.3 $V_{\text{Li}}$ while that in ~1 M decays to 140 mAh g$^{-1}$.44 The change in solvation structure and the proportion of free solvent can affect the electrochemical stability of the solvent. In the presence of Li$^+$ solvation, DFT calculations$^{50–53}$ and molecular dynamics (MD)$^{50}$ studies show that the solvent molecule is stabilized by bond formation with Li$^+$, which stabilizes the solvent’s electron lone-pairs, decreasing the electron energy level at the highest occupied molecular orbital (HOMO) states and increasing the oxidative stability of the solvents.$^{50}$ The stabilization of the solvent through Li$^+$ coordination is further supported experimentally by several research groups.$^{44–47,54–57}$ However, as mentioned previously, upper voltage limits used to cycle Ni-rich NMC (4.4–4.6 $V_{\text{Li}}$)$^{9,37}$ are much lower than the simple electrochemical oxidation potential (~5 $V_{\text{Li}}$) of even conventional electrolytes solutions (~1 M).13 Thus, the physical origin of enhanced stability due to increased salt concentration is still not well understood, which is the focus of this study. In the present manuscript, we propose a new concept, where Li-coordination with carbonates – greatly decreases the kinetics of dehydrogenation supported by DFT than free carbonates, which is responsible for increased capacity retention during cycling of NMC811.

Here we examine this oxide-mediated electrolyte oxidation mechanism$^{14,16,17,37,58}$ by employing highly concentrated carbonate-based electrolytes and electrolyte salts of different dissociation constants, and test if having fewer free carbonate molecules can potentially reduce the electrolyte reactivity with Ni-rich NMC. Highly concentrated electrolytes that typically have a salt molar concentration greater than ~3 M and a molar ratio of Li:solvent less than 1:4, have shown radically different properties such as reduced volatility/flammability$^{55,59–64}$,
greater electrochemical stability window,\textsuperscript{65–69} enhanced rate performance,\textsuperscript{56,68,70–72} and altered interfacial reactivity\textsuperscript{61,66–68,70,73–79} from conventional electrolytes (\textsim{1} M).\textsuperscript{13,80} These changes have been attributed to the reduction of free solvent molecules that do not coordinate with Li ions.\textsuperscript{62,81–83} For example, Inaba \textit{et al.} first reported that lithium can be intercalated reversibly into graphite in a concentrated electrolyte of propylene carbonate (PC) in 2003\textsuperscript{84} while conventional PC-based electrolytes failed to form a stable SEI with graphite.\textsuperscript{13} In addition, Winter and Placke \textit{et al.}, showed that the oxidative stability expanded to \textsim{6} \textsubscript{V}_{\text{Li}} with 4 M LiPF\textsubscript{6} in DMC.\textsuperscript{85} Moreover, Watanabe \textit{et al.} have reported that the oxidation potential of ether-based electrolytes, having lithium bis(trifluoromethanesulfonyl)imide: LiTFSI with triglyme and tetraglyme, can be increased from \textsim{4.0} \textsubscript{V}_{\text{Li}} in \textsim{0.2} M to \textsim{4.5} \textsubscript{V}_{\text{Li}} in \textsim{3} M by removing free ether solvent molecules (1:1 molar mixture of LiTFSI with glyme), where enhanced stability against oxidation is associated with lower HOMO energy level of Li\textsuperscript{+}-coordinated solvents.\textsuperscript{55}

In this work, to understand the mechanism of improved stability through changing electrolyte solvation structures, we quantify the amounts of free carbonate solvents when increasing LiPF\textsubscript{6} salt concentration from 0.2 to 3.1 M and by changing anions from BF\textsubscript{4}\textsuperscript{−}, PF\textsubscript{6}\textsuperscript{−} to TFSI\textsuperscript{−} with increasing lithium salt dissociation constant\textsuperscript{86–88} in 1.0 M electrolytes using Raman spectroscopy. The amounts of free carbonate solvents in 1.0 and 3.1 M were discussed and correlated with capacity loss and impedance growth of NMC811 during cycling with these electrolytes. The dissociativity of the salt in solvent also largely dictates the amount of free Li\textsuperscript{+} in the electrolyte as well as the Li\textsuperscript{+} coordination of the solvent species. The association strength of different common anions used in lithium-ion battery salts can be ranked as TFSI\textsuperscript{−}~PF\textsubscript{6}\textsuperscript{−}~AsF\textsubscript{6}\textsuperscript{−}~ClO\textsubscript{4}\textsuperscript{−}~BF\textsubscript{4}\textsuperscript{−},\textsuperscript{53,89,90} where TFSI\textsuperscript{−} and PF\textsubscript{6}\textsuperscript{−} are the most dissociative with more free Li\textsuperscript{+} ions when dissolved in a carbonate electrolyte, and BF\textsubscript{4}\textsuperscript{−} is the most associative with more cation-anion pairs.
instead of free Li\(^+\). In addition, carbon-free and binder-free oxide-only electrodes charged in these two electrolytes (1 and 3.1 M) to different voltages in the first cycle were analyzed by FT-IR, Raman spectroscopy and XPS to identify oxidized or decomposed products of carbonate and salt, from which the role of free carbonate solvents on the electrolyte reactivity with NMC surface, and implications on the capacity loss and impedance of NMC during cycling are discussed.

2. Methods

2.1 Experimental methods

**Materials** 1.0 M (mol/L) LiPF\(_6\) in a 3:7 wt:wt ethylene carbonate (EC): ethyl methyl carbonate (EMC) (LP57, battery-grade, BASF), LiPF\(_6\) (>99.99%, battery-grade, Aldrich), LiBF\(_4\) (>99.99%, ultra-dry-grade, Aldrich), EC (battery-grade, BASF), EMC (battery-grade, BASF) were used as received. The electrolytes (except for LP57) were prepared by simple mixing of LiPF\(_6\) or LiBF\(_4\) and EC or EMC in an Argon-filled glove box (MBraun, < 0.5 ppm of H\(_2\)O and O\(_2\)). The residual water contents in the electrolytes were measured by Karl Fischer titration and were less than 20 ppm.

**Viscosity, density and ionic conductivity** The viscosities and densities of the electrolytes were measured by using a Stabinger viscometer (SVM3001, Anton Paar). Ionic conductivity was measured by using the complex impedance method with the conductivity cell (CONPT-BTA, Vernier). The cell constant of the conductivity probe was calculated by measuring the conductivity of a standard solution (0.01 M KCl aqueous solution, 1.413 mS/cm at 25 °C, VWR). The density, viscosity and ionic conductivity for the concentrations of LiPF\(_6\) (0, 0.22, 0.43, 1.0, 1.3, 2.0, 2.4 and 3.1 M) in EC/EMC, and LiClO\(_4\) and LiBF\(_4\) (1.0 M) in EC/EMC solutions at 25 °C were measured, where details are shown in **Table S1** and **Table**
S2, respectively. In addition, the viscosity, density and ionic conductivity of 1.5 and 1.0 M LiPF₆ solution in pure EC or EMC at 25 °C are shown for comparison in Table S3, where the viscosity and density of EC was measured at a supercooled liquid state at 25 °C as the melting point of EC is ~35 °C. Moreover, activation energy was calculated from temperature dependent data of viscosity and ionic conductivity measurements, where the plots were fitted only for the high-temperature region (Figure S1).

Oxide synthesis and electrode preparation NMC111 (LiNi₀.₃₃Mn₀.₃₃Co₀.₃₃O₂) was prepared as reported previously.¹⁴ NMC622 and NMC811 were obtained from ECOPRO, South Korea. Both NMC622 and NMC811 had no-coating and no-Ni concentration gradient within individual oxide particles, which was used to examine the intrinsic stability/reactivity trend of the electrode-electrolyte interface.

The carbon-free and binder-free pellet electrodes were prepared as reported previously.¹⁴ The pellet electrodes were used for DRIFT and Raman measurements. The carbon-free binder-free oxide powder electrodes were also prepared as reported previously.¹⁴ The carbon-free, binder-free powder electrodes were used for XPS measurements. The active material loading was ~2.4 mg/cm².

The composite positive electrodes for battery cycling and EIS measurements were composed of 85 wt% of NMC811 as an active material, carbon black (5 wt % KS6 and 2 wt% Super P, both from Timcal) as an electrically conductive additive, and poly(vinylidene fluoride) (8 wt%, PVDF, Kynar) as a polymer binder. The mesh reference electrode was also composed of 80 wt% of Li₄Ti₅O₁₂ (LTO, Itasco, >99.5 %), 10 wt% of acetylene black (C-55, Chevron) and 10 wt% of PVDF. Detail electrode preparation method can be found in previous report.⁵⁸ Each electrode was punched with a 1/2 inch diameter (1.27 cm diameter) for NMC811
(on Al foil) and 18 mm diameter for Li$_4$Ti$_5$O$_{12}$ mesh reference. The active material loading was $\sim$2.4 mg/cm$^2$ for composite electrodes and $\sim$1 mg/cm$^2$ for Li$_4$Ti$_5$O$_{12}$ mesh reference electrodes, respectively. After compression at 6.3 T/cm$^2$, the thickness of NMC811 composite electrodes was ca. 10 $\mu$m, packing density was ca. 3.2 g$_{NMC-C-PVDF}$/cm$^3$ and porosity was $\sim$18 %. Particle size and specific surface area of NMC811 used in this work is $d = 5$-$10$ $\mu$m and 0.334 m$^2$/g, respectively.

**Electrochemical measurements** Electrochemical behavior of the carbon-free, binder-free electrodes and pellet electrodes were confirmed by galvanostatic measurements in two-electrode cells (Tomcell type TJ-AC, disk type two-electrode cell). Cells were assembled in an argon-filled glovebox (MBraun, <0.5 ppm of H$_2$O and O$_2$) and comprised a lithium metal foil negative electrode (15 mm diameter) and the positive electrode (1/2 inch (12.7 mm) diameter), separated by two pieces of polyolefin separator (Celgard 2325, 21 mm diameter) or a piece of glass fiber separator (Whatman GF/A, Aldrich, 21 mm diameter, dried at 150 $^\circ$C in vacuum for overnight prior to use, glass fiber separator was used for concentrated electrolyte, 3.1 M LiPF$_6$ in EC/EMC solution, due to issues with wettability for polyolefin separators), with 100 $\mu$L of different electrolytes. The cells were charged with different end-of-charge potentials (4.1, 4.2, 4.4, 4.6 and 4.8 V$_{Li}$) at a C/100 rate and held at the potential for 5 hours at room temperature with VMP3 potentiostat (Biologic), based on the theoretical capacity calculated assuming full delithiation ($\text{LiNi}_x\text{Mn}_y\text{Co}_{1-x-y}\text{O}_2 \rightarrow \text{Li}^+ + e^- + \text{Ni}_x\text{Mn}_y\text{Co}_{1-x-y}\text{O}_2$, $\sim$275 mAh g$^{-1}$, 1C = 275 mA g$^{-1}$).

For electrochemical impedance spectroscopy (EIS) and battery cycling experiments with composite electrodes, three-electrode cells were assembled in an argon-filled glove box
with Li metal foil (15 mm diameter), Li$_4$Ti$_5$O$_{12}$ mesh reference electrode (18 mm diameter), and NMC811 composite electrode (1/2 inch (12.7 mm) diameter) with Whatman GF/A separators (19 mm diameter). A mesh reference electrode was used to avoid an artefactual EIS response.$^{58,91,92}$ A detailed cell configuration can be found in the previous work.$^{58}$ NMC811 working electrodes were then charged at 27.5 mA/g (C/10 rate, 27.5 mA/g corresponds to $\sim$65 $\mu$A/cm$^2$ for average loading density of $\sim$2.4 mg/cm$^2$) and held at potentials of 3.9, 4.0, 4.1, 4.2, 4.3, 4.4, 4.5 and 4.6 V$_{Li}$ for 1 hour, and relaxed for 1 hour. After each relaxation, EIS measurements were carried out at the open circuit potential with 10 mV amplitude and frequency range from $\sim$10$^{-2}$ to 10$^{6}$ Hz. In this work, due to high R$_{HF}$ on NMC811 electrodes (at some conditions it cannot even form a complete semicircle within this frequency range, down to 100 mHz) and overlapped and depressed semicircles likely due to low ionic conductivity and high viscosity in concentrated solution, we did not fit the obtained Nyquist plots to obtain R$_{HF}$ and R$_{LF}$ values and rather chose to discuss through the Nyquist plots to keep a rigorous discussion. After the 1st cycle charge-discharge (4.6 – 2.0 V$_{Li}$, charge sequence coupled with EIS) at C/10 was performed in a three-electrode configuration, faster cycling tests (2nd-99th cycle) were then conducted with a two-electrode configuration (NMC811 as working electrode and Li metal as counter electrode) with CCCV - constant current (1C) / constant voltage (cut-off current at 0.05C) – charging condition and 1C discharge condition with cut-off potential between 4.6 V$_{Li}$ and 2.0 V$_{Li}$. At the 100th cycle, three-electrode 0.1C charge-discharge with EIS measurement was conducted again to evaluate resistance growth compared with the 1st cycle. After the 101st cycle, two-electrode fast cycling with CCCV condition (1C-0.05C for charge, 1C for discharge) was conducted again. The 1st and 100th three-electrode cycling with EIS measurement was conducted at 25 °C in an environmental
chamber (SU-241, Espec) and fast two-electrode cycling (2-99th cycle and after 101th cycle) was conducted at room temperature. Coin-cell cycling was also conducted as previously reported.\textsuperscript{37}

**Raman spectroscopy** Raman spectra of the electrolyte solutions and charged pellet electrodes were measured using a microscope Raman spectrometer system (LabRAM HR, Horiba) with 532 and 632 nm laser excitations, which was calibrated using a silicon standard. All the pellet electrodes and most of the electrolyte solution samples were measured with 532 nm laser excitation unless otherwise noted. The spectra were accumulated 5 times (pellet electrode) or 10 times (electrolyte solution sample) to achieve a sufficiently high signal-to-noise ratio. Detail Raman sample preparation method can be found in the previous report.\textsuperscript{14}

For solution samples, after baseline correction of the entire spectral range, the integrated intensity (area) in the wave number range of 850 – 920 cm\(^{-1}\) (including both free- and bound-EC, and EMC band) was normalized. The Raman spectra of the solutions were deconvoluted using a Gaussian-Lorentzian (pseudo-Voigt) function as shown in Figure S2. The molar fraction of free/bound species was computed from integral intensities of corresponding Raman peaks. Assuming that Raman scattering coefficients for free- and bound-carbonate are identical and constant in the whole concentration range\textsuperscript{93}, \( \frac{I_{\text{bound}} - \text{EC}}{I_{\text{free}} - \text{EC} + I_{\text{bound}} - \text{EC}} = \frac{c_{\text{bound}} - \text{EC}}{c_{\text{EC}}} = \frac{n_{\text{EC}} \cdot c_{\text{LiPF}}}{c_{\text{EC}}} \)

\[
\frac{I_{\text{bound}} - \text{EMC}}{I_{\text{free}} - \text{EMC} + I_{\text{bound}} - \text{EMC}} = \frac{c_{\text{bound}} - \text{EMC}}{c_{\text{EMC}}} = \frac{n_{\text{EMC}} \cdot c_{\text{LiPF}}}{c_{\text{EMC}}}
\]
\[ n_{EC} + n_{EMC} = n_{total} \]
Where \( c_{EC} \) and \( c_{EMC} \) [M] are the molar concentration of EC and EMC, \( c_{\text{bound-EC}} \) and \( c_{\text{bound-EMC}} \) [M] are the concentration of bound-EC and bound-EMC, \( c_{\text{LiPF}_6} \) [M] is the concentration of LiPF\(_6\), \( n_{EC} \) and \( n_{EMC} \) are the number of solvated EC and EMC per one Li\(^+\) (\( c_{\text{bound-EC}/c_{\text{Li}^+}} \) or \( \text{Li}^+:\text{bound-EC} = 1:n_{EC} \)) and \( c_{\text{bound-EMC}/c_{\text{LiPF}_6}} \) (\( \text{Li}^+:\text{bound-EMC} = 1:n_{EMC} \)), respectively, and \( n_{total} \) is the total solvation number of Li\(^+\) from both EC and EMC. Herein, \( I_{\text{bound}} / I_{\text{total}} \) is the same as \( c_{\text{bound}} / c_{\text{total}} \). It should be noted that EC:EMC of 30:70 wt/wt corresponds to 33.6:66.4 mol/mol.

**FT-IR spectroscopy** The Fourier transform infrared (FT-IR) spectra of the materials were obtained on an FT-IR Tensor II (Bruker) equipped with a deuterated triglycine sulfate (DTGS) detector inside an argon-filled glovebox. The species formed on the active material during (electro)chemical processes were analyzed with the diffuse reflectance infrared Fourier transform (DRIFT) accessory (Praying Mantis, Harrick scientific products). Detailed experimental condition can be found elsewhere.\(^\text{14}\) All FTIR spectra of the liquid samples were recorded using a single reflection attenuated total reflection (ATR) accessory (Pike Vee-Max II, Pike Technologies) with a Ge prism (Pier optics) at an incident angle of 45 degrees.

**X-ray photoelectron spectroscopy (XPS) measurements** All the XPS spectra were collected using a PHI 5000 VersaProbe II (ULVAC-PHI, INC.) using a monochromatized Al K\(_\alpha\) source and a charge neutralizer. Detail procedures can be found in the previous publication.\(^\text{14}\) Adventitious carbon at 285 eV (C1s spectra) was used for calibration of all XPS spectra. The chemical compositions, binding energies and full width at half maximum of all spectra can be found in Tables S4–S7.
2.2 Computational methods

**Raman – FTIR simulations** The Raman and FTIR spectra were simulated by computing the vibrational frequencies of solvent molecules, salt ions and solvent-ions complexes in an implicit solvation model (PCM\textsuperscript{9}), and dimethyl sulfoxide (DMSO, dielectric constant $\varepsilon = 47$) was used as solvent. We used the B3LYP functional and 6-311++G** basis set, as implemented in the Gaussian (g09) suite\textsuperscript{94}.

**Interaction/adsorption energy calculations** The EC C-H bond dissociation energy and the interaction energy of Li\textsuperscript{+} with EC were estimated by considering complexes of one Li ion with 1-4 EC molecules in vacuum and in implicit DMSO solvent. We used the B3LYP functional for EC bond dissociation energy and Perdew Burke-Enzerhof (PBE) functional for Li\textsuperscript{+}-EC interaction\textsuperscript{95} and 6-31+G* basis set.

The adsorption energy of EC on the surface of LiMO\textsubscript{2} (M = Mn, Co and Ni) and Mn- and Co-doped LiNiO\textsubscript{2}, and the thermodynamic reaction profiles for EC and Li (EC)\textsubscript{3} on LiNiO\textsubscript{2} were computed by periodic plane wave calculations, with projector augmented wave potentials, and PBE functional, as implemented in the Vienna ab initio simulation package (VASP).\textsuperscript{96,97} We employed a DFT+U approach\textsuperscript{98,99} with effective Hubbard $U$ values $U_{\text{eff}} = 4.0$ eV for Mn, 3.3 eV for Co and 6.4 eV for Ni.\textsuperscript{100,101} EC was adsorbed on the (10\overline{1}4) surface, with a (4 × 2) surface unit cell and a five-layer slab, where the three bottom layers were kept fixed at the bulk positions, while the coordinates of the two uppermost layers and the adsorbates were fully relaxed. The slabs were separated by a vacuum layer of at least 13 Å. Additional details can be found in the previous work\textsuperscript{16}.
3. Results and Discussion

Salt-concentration-dependent Free-solvent activity

Concentration-dependent Raman spectra in the range from 850 to 965 cm\(^{-1}\) revealed the EC ring breathing modes with and without Li\(^+\) coordination\(^\text{14}\) and the O-CH\(_3\)/O-C\(_2\)H\(_5\) stretching modes of EMC with and without Li\(^+\) coordination,\(^\text{67}\) as shown in Figure 1 (a). Pure EC/EMC (30/70 wt\%) solution showed one sharp band at \(~892\) cm\(^{-1}\), corresponding to the EC ring breathing mode,\(^\text{14}\) and one broad band centered at \(~935\) cm\(^{-1}\), corresponding to the O-CH\(_3\)/O-C\(_2\)H\(_5\) stretching mode of EMC, where the nature of the broad band\(^\text{93}\) most likely come from two peaks (\(~930\) and \(~938\) cm\(^{-1}\)) due to the asymmetric structure of EMC. With increasing concentration of LiPF\(_6\) in the solution, the band intensity of EC ring breathing (\(~892\) cm\(^{-1}\)) was found to decrease monotonically, which was replaced gradually by another sharp band at a higher wavenumber (\(~904\) cm\(^{-1}\)), with a clear isosbestic point at \(~899\) cm\(^{-1}\). Similarly, the band intensity of O-CH\(_3\)/O-C\(_2\)H\(_5\) stretching in EMC (\(~935\) cm\(^{-1}\)) was reduced gradually with increasing LiPF\(_6\) concentration, which was coupled with the appearance and growth of another broad band at \(~946\) cm\(^{-1}\) with a clear isosbestic point at \(~940\) cm\(^{-1}\). These two new bands centered at \(~904\) and \(~946\) cm\(^{-1}\) can be assigned to Li\(^+\)-coordinated-EC and EMC. No visible change was found for the peak centered at \(~880\) cm\(^{-1}\) that is associated with EMC molecules (can be seen in pure solvent spectrum, Figure S3), which will not be discussed further. Moreover, the appearance and growth of Li\(^+\)-coordinated-EC and -EMC bands with increasing LiPF\(_6\) concentration were accompanied with increasing intensity of a new band centered at \(~742\) cm\(^{-1}\), which can be attributed to the P-F symmetric stretching vibration of PF\(_6\)\(^\text{-102}\), as shown in Figure 1(b). At LiPF\(_6\) concentrations greater than Li:Carbonate = 1:5 (2.0 M), a peak shoulder at the high wavenumber side appeared, indicating the formation of complexes between Li ions and PF\(_6\) anions or contact ion pair (CIP). With increasing LiPF\(_6\) concentration,
the distance between the anions and cations decreases in the solutions and ion-pair formation/aggregation occurs. Therefore, the band at ~742 cm\(^{-1}\) was deconvoluted into one component of free PF\(_6^-\) at ~742 cm\(^{-1}\) at concentrations below Li:Carbonate = 1:5 (2.0 M) while above, two different components, free PF\(_6^-\) at ~742 cm\(^{-1}\) and Li\(^+\)-bound PF\(_6\) at ~744 cm\(^{-1}\) were obtained (see details in Figure S2), which later increased with increasing LiPF\(_6\) concentration.
**Figure 1** Normalized Raman spectra of LiPF$_6$ in EC:EMC (3:7 wt/wt) solution in the range (a) carbonate bands (850 - 965 cm$^{-1}$) and (b) PF$_6^-$ band (700-760 cm$^{-1}$) measured at 25 °C. (c) Ionic conductivity, viscosity and estimated molar fractions of free/bound-EC, free/bound-EMC ($I_{\text{free-EC}}/I_{\text{total-EC}}$, $I_{\text{free-EMC}}/I_{\text{total-EMC}}$), and Li$^+$-bound PF$_6^-$ ($I_{\text{bound-PF6}}/I_{\text{total-PF6}}$) in the LiPF$_6$ in EC/EMC solution based on the ratio of integral intensities of Raman peaks. All the $R^2$ values for the deconvolution are higher than 0.987 and linear fits for calculating solvation number are higher than 0.992. Detail deconvoluted spectra were shown in **Figure S2**. (d) Normalized Raman spectra of 1.0 M LiPF$_6$, ClO$_4$ and BF$_4$ in EC:EMC (30:70 wt%) solution in the range 850 - 965 cm$^{-1}$ (carbonate bands) measured at 25 °C. Note that the region higher than 910 cm$^{-1}$ for LiClO$_4$ sample was removed due to severe overlap of ClO$_4$ peak and EMC peak. (e) Ionic conductivity and free EC fraction in 1.0 M LiPF$_6$, ClO$_4$ and BF$_4$ in EC:EMC (3:7wt) solution.

The fractions of carbonates and PF$_6^-$ anions with and without Li$^+$ coordination were quantified from integrated intensities ($I$) of corresponding Raman bands, where Raman scattering coefficients for free- and bound Carbonates were assumed to be identical and constant.$^{77,93}$ **Figure 1(c)** shows that the fractions of free-EC and free-EMC decrease while the fraction of Li$^+$-coordinated EC and EMC increases monotonically with increasing molar ratio of LiPF$_6$/Carbonate ($c_{\text{Li}}/c_{\text{Carbonate}}$), which were obtained from the ratios of $I_{\text{free}}/I_{\text{total}}$ and $I_{\text{bound}}/I_{\text{total}}$, respectively. The fraction of Li$^+$-bound EC is higher than that of Li$^+$-bound EMC, which is consistent with having much higher polarity of EC (dielectric constant of 90$^{103}$) compared to EMC (dielectric constant of 3$^{103}$). Of significance to note is that the molar fractions of free EC and EMC were very small (combined fractions equal to ~0.07 or ~7 mol%) for the most concentrated solution of LiPF$_6$/Carbonate of 1:3 (3.1 M), indicating negligible free solvent activity, in comparison to that of the conventional electrolyte concentration of 1.0 M (0.5 or 50 mol%). The fraction dependence of Li$^+$-bound EC and EC on the molar ratio of LiPF$_6$/Carbonate (the slope in **Figure 1c**) was used to estimate the solvation number of Li$^+$, which fitting was limited to concentrations below Li:Carbonate = 1:5 (2.0 M) where LiPF$_6$ is fully dissociative (**Figure 1b**). Li$^+$ was found to be coordinated by ~1.6 for EC and ~2.4 for EMC, giving rise to a total solvation number of ~4, which is in agreement with previous reports of Li$^+$ solvation numbers around 4.$^{75,81,104}$ The result is further
supported by having negligible fractions of Li\(^+\)-bound PF\(_6^-\) at concentrations below Li:Carbonate = 1:5 (2.0 M) estimated from the integrated intensities of free (~742 cm\(^{-1}\)) and bound PF\(_6^-\) (~744 cm\(^{-1}\)) components (Figure 1b), which is consistent with the highly dissociative nature of PF\(_6^-\) anions. At higher concentrations, 0.13 and 0.57 of PF\(_6^-\) interacted directly with Li\(^+\) for Li:Carbonate ratios of 1:4 (2.4 M) and 1:3 (3.1 M), indicating stronger CIP formation/aggregation. Therefore, Raman spectroscopy reveals that Li\(^+\) is coordinated as \([\text{Li(EC)}_{1.64}(\text{EMC})_{2.37}]^+\) in the EC:EMC (3:7 wt/wt or 1:2 mol/mol) solutions with LiPF\(_6\) concentrations equal to or less than 2.0 M. In contrast, at the highest concentration having Li:Carbonate of 1:3 (3.1 M), nearly all the EC and EMC (>93 mol%) are coordinating to Li\(^+\) and more than half of PF\(_6^-\) is also participating Li\(^+\) solvation, giving rise to complexes resembling \([\text{Li(EC)}_{1}(\text{EMC})_{2}(\text{PF}_6)]\) and \([\text{Li(EC)}_{1}(\text{EMC})_{2}]^+\).

Replacing more dissociative LiPF\(_6\) with less dissociative LiBF\(_4\), the Li-coordinated EC ring breathing mode feature (~904 cm\(^{-1}\)) was found to decrease in intensity while that of the free EC ring breathing band (~892 cm\(^{-1}\)) increased, indicating an increased fraction of free EC molecules. As solvent and anion (both Lewis bases) are competing with each other to solvate (interact) to Li\(^+\) (strong Lewis acid) in the electrolyte solution,\(^{81,105}\) fully dissociative PF\(_6^-\) at 1.0 M (Figure 1b) does not disturb the solvation of Li\(^+\) by carbonate molecules whereas less dissociative BF\(_4^-\) can bind with lithium ions and results in more free carbonate solvents. The B-F stretching mode of 1.0 M LiBF\(_4\) in EC/EMC solution (Figure S4) shows free-BF\(_4^-\) at 776 cm\(^{-1}\) and Li\(^+\)-bound -BF\(_4^-\) at 774 cm\(^{-1}\),\(^{47}\) which intensity was used to quantify Li\(^+\)-bound-BF\(_4^-\) as 0.26 or 26 mol% in the 1.0 M EC/EMC (30/70 w/w%) electrolyte. Further support of more free solvent molecules in 1.0 M BF\(_4^-\) electrolyte came from ATR measurements (Figure S5), which shows the Li-coordinated EC and EMC C=O features (1774 cm\(^{-1}\) and 1720 cm\(^{-1}\)) both increase in intensity relative to free EC (1807 cm\(^{-1}\)) and EMC C=O (1744 cm\(^{-1}\)) features. By integrating the free EC
(892 cm\(^{-1}\)) and Li-coordinated EC (904 cm\(^{-1}\)) band intensities in the Raman spectra of 1.0 M LiBF\(_4\) electrolyte, we further quantified the fraction of free EC compared to Li\(^+\)-coordinated EC in Figure 1e, which revealed a higher fraction of free EC by replacing LiPF\(_6\) (0.51) with LiBF\(_4\) (0.73).

The ionic conductivity was found to first increase with greater LiPF\(_6\) concentration and then decrease at concentrations greater than (up to 1.3 M), as shown in Figure 1c. This nonlinear concentration-dependent conductivity trend is similar to those found previously for non-aqueous electrolyte solutions with ether\(^{55,60,61,106,107}\), carbonate\(^{67}\) and other solvents.\(^{70,75}\) The maximum conductivity found for concentrations at \(\sim\)1 M can be explained by increasing charge carrier concentrations including \([\text{Li(EC)}_{1.64}(\text{EMC})_{2.37}]^+\) and PF\(_6^-\) but reduced ion mobility at concentrations greater than \(1.3 \text{ M (Figure 1c)}\). The proposed ion mobility reduction is supported by markedly increased electrolyte viscosity (Figure 1c), which results from electrostatic drag associated with solvated lithium ions, \([\text{Li(EC)}_1(\text{EMC})_2]^+\) and PF\(_6^-\) and the formation of ion-paired complexes, \([\text{Li(EMC)}_1(\text{EMC})_2(\text{PF}_6)]\). These two competing effects result in a maximum ionic conductivity at around 1 M.\(^{61,75,107,108}\) Moreover, the change in the solvation structure directly affects the change in the ionic conductivity due to a decrease in charge carrier (Li\(^+\)) concentration through the formation of ion-pair (neutral) species, where LiPF\(_6\) has the highest conductivity and lowest free-carbonate activity (\(i.e.\) highest dissociation) compared to LiClO\(_4\) and LiBF\(_4\) salt with the same concentration, due to stronger aggregation of those more Lewis basic anions (BF\(_4^-\) and ClO\(_4^-\)) to Li\(^+\).

The influence of Free-solvent activity on the cycling stability of Ni-rich NMC811

Using the most concentrated electrolyte (3.1 M LiPF\(_6\) in EC/EMC) with negligible free solvent molecules (Figure 1c) led to enhanced capacity retention of NMC811 upon cycling.
compared to electrolytes with lower LiPF$_6$ concentrations (1.0 and 2.0 M LiPF$_6$ in EC/EMC), as shown in Figure 2a. NMC811 was found to have comparable initial discharge capacities of ~220 mAh g$^{-1}$ (C/10 and charged to 4.6 V$_{Li}$) and ~200 mAh g$^{-1}$ (1 C and 4.6 V$_{Li}$) among all the electrolytes tested in this study. Much greater capacity retention was found for the most concentrated electrolyte upon subsequent cycling at 1 C than the other electrolytes with lower LiPF$_6$ concentrations (1.0 and 2.0 M LiPF$_6$ in EC/EMC in Figure 2a). The greater discharge capacities of NMC811 cycled in the most concentrated electrolyte (215 mAh g$^{-1}$ in Figure 2a) were confirmed at C/10 after 100 cycles than those tested in 2.0 M (175 mAhg$^{-1}$) and 1.0 M (163 mAhg$^{-1}$). After 500 cycles (1 C and 4.6 V$_{Li}$), the most concentrated electrolyte (3.1 M) shows reversible capacity ~150 mAh g$^{-1}$ or ~75% capacity retention while the other electrolytes with lower LiPF$_6$ concentrations (1.0 and 2.0 M LiPF$_6$ in EC/EMC) had electrode capacity fading below 100 mAh g$^{-1}$. Similar capacity loss was found for cycling NMC811 in EC-only or EMC-only electrolytes such as 1.5 M LiPF$_6$ in EC (1.5 M LiPF$_6$ in EC is liquid and 1.0 M LiPF$_6$ in EC is solid) and EMC-only (1.0 M LiPF$_6$ in EMC) in Figure S7 to that of 1.0 M LiPF$_6$ in EC/EMC. The greater capacity retention with the most concentrated electrolyte was further shown for cycling NMC811 at 1 C to 4.4 V$_{Li}$ than the electrolyte with 1.0 M LiPF$_6$ (Figure S8). Moreover, the Coulombic efficiency of NMC811 (quantified as the ratio of discharge capacity over charge capacity, 99.7% averaged from the 3rd to 100th cycle) in the most concentrated electrolyte (3.1 M) was greater than the other electrolytes with lower concentrations (98.9% in 1.0 M and 98.6% in 2.0 M), as shown in Figure S9. These observations suggest that reducing free carbonate activity by replacing free EC or EMC by Li$^+$-coordinated EC or EMC can lead to enhanced capacity retention and Columbic efficiency upon cycling of NMC811 in the 3.1 M (0.07 fraction of both free EC and EMC) than that of the 1.0 M LiPF$_6$ (0.51 and 0.65 fraction of free EC and EMC,
respectively) electrolyte. The greater capacity retention of NMC811 cycled in the 3.1 M LiPF$_6$ electrolyte is accompanied with smaller impedance growth as reflected by lower gaps between charge and discharge voltages during cycling (Figure 2b) in comparison to that cycled in the 1.0 M LiPF$_6$ electrolyte (Figure 2c). The charge and discharge voltage profiles and discharge capacity at the 50th cycle (red in Figure 2b) were comparable to those of the 3rd cycle in the 3.1 M electrolyte. In contrast, the charge and discharge voltage profiles were altered considerably with the 1.0 M electrolyte in comparison to those of the 3rd cycle, showing nearly four times greater voltage polarization, and much lower discharge capacity than that of the 3rd cycle.

Figure 2 Cycling performance of NMC811 composite electrodes with different electrolyte solutions. (a) The discharge capacity and charge-discharge profile with (b) 3.1 M LiPF$_6$ in EC/EMC and (c) 1.0 M LiPF$_6$ in EC/EMC upon charging to 4.6 V$_{Li}$. At the 1st and 100th cycle, the cells were measured in a three-electrode configuration, which composed of NMC811|Li$_4$Ti$_5$O$_{12}$-mesh|Li (see experimental section in detail), galvanostatically charged at 27.5 mA/g (C/10) and subject to EIS measurements (Nyquist plots are available in Figure 3). From the 2nd to the 99th cycle and after 101st cycle, the cells were tested with a two-electrode configuration that consisted NMC811 as the working electrode and Li metal as the counter electrode upon cycling between 4.6
and 2.0 \textit{V}_{\text{Li}}, which included first charging to 4.6 \textit{V}_{\text{Li}} under 1C, subsequently voltage holding at 4.6 \textit{V}_{\text{Li}} until the current reached C/20, then discharging at 1C to 2.0 \textit{V}_{\text{Li}}.

EIS measurements of NMC811 composite electrodes in the NMC811|Li\textsubscript{4}Ti\textsubscript{5}O\textsubscript{12}-mesh|Li three-electrode cells obtained in the 1\textsuperscript{st} and 100\textsuperscript{th} cycles at C/10 further supported smaller impedance growth during cycling in the most concentrated electrolyte (3.1 M) than that of 1.0 M. As shown in Figure 3, two semicircles were observed in the Nyquist plots for Li intercalation into composite electrodes like previous work.\textsuperscript{58,109} The high-frequency semicircle (left, $10^3 - 10^4$ Hz) has been assigned to impedance at the electrified interface (predominantly lithium ion migration and adsorption/desorption in the pore structure of composite electrode\textsuperscript{58}) while the low-frequency semicircle (right, $10^0 - 10^1$ Hz) corresponds to the combination of charge transfer resistance at the electrode-electrolyte interface (equivalent circuit can be found in the inset of Figure 3c).\textsuperscript{58} In the first cycle, although a higher resistance was observed for the low-frequency semicircle ($10^0 - 10^1$ Hz) in the most concentrated solution compared to 1.0 M solution (Figure 3ab) at 4.0 \textit{V}_{\text{Li}}, suggesting greater charge-transfer resistance\textsuperscript{58} in 3.1 M solution, a smaller increase in the low-frequency impedance was found with increasing potential from 4.0 to 4.5 \textit{V}_{\text{Li}} for the 3.1 M than the 1.0 M electrolyte. It is interesting to note in Figure 3a, high-frequency resistance ($10^3 - 10^4$ Hz) and low-frequency resistance ($10^0 - 10^1$ Hz) were comparable below 4.2 \textit{V}_{\text{Li}} while \textit{R}_{LF} suddenly became ~10 times larger when the potential increased from 4.2 to 4.3 \textit{V}_{\text{Li}}, indicating severe resistance growth on the NMC811 surface in the 1.0 M electrolyte, which is consistent with greater capacity loss upon cycling to 4.3 \textit{V}_{\text{Li}} and greater.\textsuperscript{110} At the 100th cycle (Figure 3cd), the low-frequency impedance with 3.1 M solution grew slightly (~100 \textit{\Omega} at 4.0 \textit{V}_{\text{Li}}) and exhibited comparable potential dependence to that in the first cycle. In contrast, the low-frequency (charge transfer) impedance with 1.0 M grew significantly after 100 cycles, where the potential
dependence of the impedance observed in the first cycle disappeared. These observations are in agreement with higher capacity retention and lower overpotential observed during cycling in the most concentrated electrolyte (3.1 M) compared to electrolytes of lower salt concentrations shown in Figure 2. Although the cycling experiments were carried out with two-electrode configuration, this three-electrode EIS results clearly shows that the capacity decay is mainly caused by NMC positive electrode side. In addition, two-electrode cell cycling of Li/NMC111 in the conventional electrolyte: 1 M LiPF$_6$ in EC:EMC has much reduced capacity loss than that of Li/NMC811 in this work and in previous work, indicating that the lithium metal electrode and its interfacial changes during cycling is not responsible for large capacity loss noted for Li/NMC811. Further support came from EIS measurements of potential holds at 4.4 V$_{Li}$ for 100 hours in the 3.1 M and 1.0 M LiPF$_6$ electrolytes, shown in Figure S10. Both high-frequency and low-frequency impedances were found not to change with time for 100 hours with the 3.1 M LiPF$_6$ EC/EMC electrolyte shown in Figure S10b, which is consistent with the higher capacity retention in cycling results (Figure 2a). On the other hand, the low-frequency impedance was found to continuously increase with time in the 1.0 M electrolyte while there was no significant changes for the high-frequency impedance upon a potential hold at 4.4 V$_{Li}$ (Figure S10a), which is in agreement with the assignment of high-frequency impedance to the electrified interface resistance associated with ion adsorption and desorption in the composite pore structure. Not only did NMC811 show a much greater low-frequency impedance of ~400 Ω under comparable conditions to that of other positive electrode materials such as composite Li$_x$CoO$_2$ (~20 Ω) for R$_{LF}$ at 4.4 V$_{Li}$, but the low-frequency impedance also increased sharply with time in Figure S10a, which is consistent with capacity loss and overpotential growth during cycling shown in Figure 2b and Figure 2c, respectively. The bulk resistances in concentrated electrolyte (3.1 M solution) and 1.0 M solution are ~10 ohm and
~3 ohm, respectively, while the charge-transfer resistances in both electrolytes are ~200 ohm at 4.3 V\textsubscript{Li} at the first cycle and increase over cycling (Figure 3). Thus, the rate limiting step of this system would be the charge transfer reaction at the NMC811-electrolyte interface.

Figure 3 Nyquist plots for EIS measurement on NMC811 composite electrode in NMC811|Li\textsubscript{4}Ti\textsubscript{5}O\textsubscript{12}-mesh|Li three electrode cell with 1.0 M LiPF\textsubscript{6} / EC-EMC (3:7 wt/wt) at (a) 1st cycle and (b) 100th cycle, and with 3.1 M LiPF\textsubscript{6} / EC-EMC (3:7 wt/wt) at (c) 1st cycle and (d) 100th cycle at different potential at 25 °C. At 1st and 100th cycle, the cell was operated with three-electrode configuration and galvanostatically charged at 27.5 mA/g (0.1C) and hold potential for 1 hour at each potential (3.9, 4.0, 4.1, 4.2, 4.3, 4.4, 4.5 and 4.6 V\textsubscript{Li}), then relax 1 hour before EIS sequence. From 2nd to 99th cycle, the cell was operated with two-electrode configuration (NMC811 as working electrode and Li metal as counter electrode) and cycle between 4.6 and 2.0 V\textsubscript{Li} with CCCV condition (1C charge to 4.6 V\textsubscript{Li} and hold at 4.6 V\textsubscript{Li} until the current reach to C/20, then discharge at 1C to 2.0 V\textsubscript{Li}, cycle performance can be found in Figure 2).
The influence of Free-solvent activity on the surface chemistry of cycled NMC811

**Figure 4** DRIFT spectra of (a) C=O stretching region for Li$_{x}$Ni$_{0.8}$Mn$_{0.1}$Co$_{0.1}$O$_{2}$ (NMC811) for pristine pellets (no electrolyte exposure) and after charged to 4.1, 4.2, 4.4, and 4.6 $V_{\text{Li}}$ in 1.0 M (=LP57) and 3.1 M LiPF$_{6}$ in EC/EMC (3:7 w:w). (b) Simulated FT-IR spectra of C=O vibration shown in EC, Li$^{+}$-coordinated EC, dehydrogenated EC (deH-EC), Li$^{+}$-coordinated dehydrogenated EC, EMC, vinylene carbonate (VC), and oligomers possibly formed through EC dehydrogenation (including, from top to bottom, C$_{3}$H$_{8}$O$_{3}$, C$_{5}$H$_{8}$O$_{4}$, C$_{6}$H$_{10}$O$_{4}$, C$_{6}$H$_{10}$O$_{3}$). (c) C=O stretching region for Li$_{x}$Ni$_{0.6}$Mn$_{0.2}$Co$_{0.2}$O$_{2}$ (NMC622), and Li$_{x}$Ni$_{0.8}$Mn$_{0.1}$Co$_{0.1}$O$_{2}$ (NMC811) for pristine pellets (no electrolyte exposure) and after charged to 4.1, 4.2, 4.4, and 4.6 $V_{\text{Li}}$ in 1.0 M LiPF$_{6}$ in EC/EMC (3:7 w:w) and 1.0 M LiBF$_{4}$ in EC/EMC (3:7 w:w). The example electrochemistry profiles of NMC622, and 811 pellet electrodes in different electrolytes are shown in **Figure S11**.

To examine the reactivity of the electrolyte towards the NMC oxide surface, surface analysis for charged NMC was conducted via FT-IR DRIFT and XPS measurements. DRIFT measurements showed that there was a decrease in the oxide-electrolyte reactivity induced by a decrease of free
solvent by increasing LiPF$_6$ concentration. Figure 4a shows the C=O stretching region of pristine NMC 811 and charged electrodes to 4.1 V$_{Li}$, 4.2 V$_{Li}$, 4.4 V$_{Li}$, 4.6 V$_{Li}$ from DRIFT measurements, and that of the pristine electrolyte (1.0 M LiPF$_6$ in EC/EMC 30/70 w/w%) from the ATR measurements, which comprises of three major peaks at ~1807, ~1773, and ~1744 cm$^{-1}$, corresponding to the C=O stretching mode of EC, Li$^+$-coordinated EC, and EMC, respectively.$^{14}$ Upon charging to 4.6 V$_{Li}$ in 3.1 M, no visible blueshift was observed for the C=O stretching modes of NMC811, revealing no evidence of EC dehydrogenation on NMC 811 in the most concentrated region with low free-carbonate activity. In addition, Raman spectroscopy experiments for charged NMC811 pellet electrodes charged to 4.6 V$_{Li}$ in 3.1 M LiPF$_6$ solution failed to detect characteristic peaks at ~732 cm$^{-1}$ and 907 cm$^{-1}$ attributed to dehydrogenated EC (ring distortion and ring breathing mode of dehydrogenated Li$^+$-EC, respectively)$^{14}$ (Figure S13), further supporting that the most concentrated electrolyte suppresses carbonate oxidative dissociation. In contrast, upon charging NMC811 to 4.2 V$_{Li}$ in 1.0 M LiPF$_6$ in EC/EMC, a blueshift was observed for the peaks at 1807 and 1773 cm$^{-1}$ which can be attributed to EC-derived species formed on charged oxide surfaces through the dehydrogenation of CH$_2$-hydrogen in EC.$^{14}$ This assignment is supported by the calculated spectra for the C=O stretching mode of dehydrogenated EC coordinated with Li$^+$ (1811 cm$^{-1}$) and/or oligomers derived from dehydrogenated EC such as C$_5$H$_8$O$_4$ (1806 cm$^{-1}$) and C$_6$H$_{10}$O$_3$ (1791 cm$^{-1}$) in Figure 4b (chemical structures of C$_5$H$_8$O$_4$ and C$_6$H$_{10}$O$_3$ are also shown). This blueshift cannot be attributed to EMC or EMC-derived features as C=O stretching of EMC (Figure 4b middle panel) is located at much lower wavenumbers (around 1740 cm$^{-1}$). In addition, as DRIFT is an ex-situ analysis after grinding charged pellet electrode with KBr, the bound carbonate and free carbonate can be exchanged during sample preparation. This would explain why bound EC exhibits a blue shift even if free carbonate is dehydrogenated. As further evidence,
a simple mixture of LiPF$_6$ and carbonates prepared with the same procedure showed a decrease in the free EC band and an increase in the Li$^+$-EC band, which indicates that there is an exchange between free/bound species during the grinding process. In summary, DRIFT spectra on charged electrodes revealed that with decreasing LiPF$_6$ concentration (increasing free-carbonate activity), we lowered the onset potential for solvent dissociative adsorption on oxide surfaces, and increased the solvent and oxide reactivity.

Switching from LiPF$_6$ to less dissociative LiBF$_4$ to have more free solvent molecules than Li$^+$-coordinated molecules, DRIFT measurements showed increased oxide-electrolyte reactivity with charged NMC622 and 811. Upon charging to 4.4 V$_{Li}$ for NMC622 in 1.0 M LiBF$_4$ electrolyte, a blueshift is observed for the peaks at 1807 and 1773 cm$^{-1}$ whereas no significant changes are observed for NMC622 even when charged to 4.6 V$_{Li}$ in 1.0 M LiPF$_6$ electrolyte (Figure 4c). A similar blueshift is observed for NMC811 charged to 4.2 V$_{Li}$ and above in LiPF$_6$ and charged to 4.1 V$_{Li}$ and above in LiBF$_4$. Such blueshifts can be attributed to EC-derived species formed on charged oxide surfaces through the dehydrogenation of CH$_2$-hydrogen in EC.$^{14}$ This observation indicates that 1.0 M LiBF$_4$ electrolyte shows lower onset voltages for solvent dehydrogenation compared with more dissociative LiPF$_6$ on NMC622, as well as NMC811, supporting the argument that electrolytes with lower free-carbonate activity show increased stability against dehydrogenation on the oxide surface. This argument is in agreement with NMC811 cycled with 1.0 M LiPF$_6$ having greater capacity retention than 1.0 M LiBF$_4$ (Figure S14).
Figure 5 Concentration dependent XPS spectra of the (a) F1s and (b) O1s photoemission lines collected from carbon-free, binder-free NMC811 electrodes. C1s and P2p spectra are available in Figure S15. Atomic percentages from XPS spectra for the pristine and charged carbon-free, binder-free NMC811 electrodes in (c) 3.1 M LiPF$_6$ and (d) 1.0 M LiPF$_6$ in EC: EMC 3:7 (wt:wt). The electrodes were charged to 4.1, 4.2, 4.4 and 4.6 V$_{Li}$ at 2.75 mA/g (C/100) and held at the given potential for 5 hours using 1.0 M LiPF$_6$ in EC: EMC (3:7 wt:wt) and 3.1 M LiPF$_6$ in EC: EMC (3:7 wt:wt) solution and compared with pristine electrodes. All the spectra were normalized by fixing the C1s photoemission peak of adventitious carbon (285 eV). The example electrochemistry profiles of NMC811 carbon-free binder-free powder electrodes are shown in Figure S16.

The increased electrolyte stability with decreasing free solvent activity is further supported by XPS results. XPS spectra of F 1s and O 1s (Figure 5ab) along with C1s, P2p, Ni2p, Co2p, Mn2p, Li1s/Co3p spectra (Figure S15 and Figure S17) were collected from carbon-free, binder-free NMC811 electrodes charged to different potentials of the first charge in the 3.1 and 1.0 M LiPF$_6$ electrolyte. The F1s spectra can be deconvoluted into three species: metal fluorides/oxyfluorides (LiM$_x$F$_y$O$_z$) around 685.1 eV, Li$_x$PF$_y$O$_z$ species around 686.6 eV and LiPF$_6$ salt at 688.3 eV. O1s spectra were also fitted to components at 529.3 eV, 531 eV, 532 eV, 533.4 eV, which can be attributed to O lattice$^{17,112,113}$, ROLi (semicarbonate and polyether species) $^{114,115}$,
semicarbonates (ROCO$_2$Li) with the CO$_3$ group$^{116,117}$, O-C=O bond as in esters or single bond between a carbon and oxygen (C-O) like in polyethers$^{112,114,118}$. NMC811 electrodes charged to 4.1 and 4.2 V$_{Li}$ in the 3.1 M LiPF$_6$ electrolyte were found to have less coverage of LiM$_{x}$F$_y$O$_z$ (685.1 eV)$^{14}$ and more coverage of Li$_x$PF$_3$O$_2$ (686.6 eV) and/or LiPF$_6$ (688.3 eV) in the F1s spectra (Figure 5a) relative to those charged in the 1.0 M LiPF$_6$ electrolyte (Figure 5a). This observation is in agreement with the lower oxygen lattice component of LiM$_{x}$F$_y$O$_z$ (529.3 eV,$^{14}$ Figure 5b) than oxygenated species such as Li$_x$PF$_3$O$_2$ (534.8 eV, Figure 5b) in the O1s spectra relative to those found for the 1.0 M LiPF$_6$ electrolyte,$^{14,17,37}$ where the O lattice peak of LiM$_{x}$F$_y$O$_z$ (529.3 eV,$^{14}$ Figure 5b) for charged NMC811 in both electrolytes was shifted to higher binding energy together with greater binding energy of the Ni2p peak (Figure S17) with increasing charging voltage. As the formation of lithium nickel oxyfluoride (LiM$_{x}$F$_y$O$_z$) has been attributed to oxide-mediated electrolyte oxidation,$^{14,17}$ specifically reactions among charged NMC811, LiPF$_6$ and protic species resulted from EC dehydrogenation on charged NMC811$^{14,17}$, these XPS results suggest that charged NMC811 in the 3.1 M electrolyte is more stable against oxide-mediated electrolyte oxidation$^{14,17}$ up to 4.2 V. This hypothesis is supported by much less coverage of oxidized carbon species such as C-O (Figure 5b-d) found on charged NMC811 in the 3.1 M electrolyte than the 1.0 M electrolyte, indicative of the oxidation of carbonate solvents. Unfortunately, with an increase in voltage to 4.6 V$_{Li}$, the intensity of LiM$_{x}$F$_y$O$_z$ (685.1 eV) in the F1s (Figure 5a and Figure 5cd) edge became greater for charged NMC811 in the 3.1 M LiPF$_6$ electrolyte, indicating greater reactivity with the electrolyte than for voltages 4.2 V and lower. Nevertheless, XPS results of charged NMC811 in the 3.1 M electrolyte (7% free solvents) to 4.2 V and lower, showing less coverage of lithium metal oxyfluorides and oxidized carbon species than that in the 1.0 M electrolyte (~50% free solvents), support that lithium-coordinated carbonate
solvents are more stable against oxide-mediated dehydrogenation and electrolyte oxidation than free carbonate solvents.

**Figure 6** XPS spectra of the O1s and F1s photoemission lines for NMC111 (a), NMC622 (b) and NMC811 (c) for the pristine carbon-free, binder-free electrodes and after charging to 4.1, 4.2, 4.4, and 4.6 V\textsubscript{Li} with 1.0 M LiBF\textsubscript{4} in EC: EMC (3:7 wt:wt) electrolyte. The C1s, B1s, Mn2p, Ni2p and Li1s/Co3p XPS spectra is shown in Figure S18, Figure S19 and Figure S20. Atomic percentages from XPS spectra for F in metal fluorides/oxyfluorides, LiBF\textsubscript{x} and LiPF\textsubscript{y}O\textsubscript{z} (from F1s region) in the pristine and charged carbon-free, binder-free (d) NMC111, (e) NMC622 and (f) NMC811 electrodes at 4.1, 4.2, 4.4 and 4.6 V\textsubscript{Li} using 1.0 M LiPF\textsubscript{6} or LiBF\textsubscript{4} in EC: EMC 3:7 (wt:wt). Other atomic percentages are available in Figure S21. The deconvolution parameters are shown in Table S5, Table S6 and Table S7. The representative electrochemical profile is shown in Figure S22. The original spectra in 1.0 M LiPF\textsubscript{6} electrolyte are available in previous publication.¹⁴
Further support for the stability of lithium-coordinated carbonate solvent molecules against oxide-mediated dehydrogenation and electrolyte oxidation compared to free carbonate molecules came from XPS results of charged NMC111, 622 and 811 in a 1.0 M LiBF₄ electrolyte, which has more free solvent molecules than 1.0 M LiPF₆ (Figure 1e). XPS analysis of O1s and F1s spectra (Figure 6a-c along with C1s, B1s, Mn2p, Ni2p, Co2p and Li1s/Co3p) of carbon-free, binder-free NMC 111, 622 and 811 electrodes charged to 4.1, 4.2, 4.4 and 4.6 V_{Li} in 1.0 M LiBF₄ EC/EMC electrolyte (Figure S18, Figure S19 and Figure S20) were compared to those charged in 1.0 M LiPF₆ EC/EMC electrolytes as reported previously. The F1s spectra can be deconvoluted to three species: metal fluorides/oxyfluorides (LiMₓFᵧOₓ) around 685 eV, LiBFₓ species around 686.5 eV and LiBF₄ salt at 688.5 eV. The F1s intensity of NMC 111 charged with LiBF₄ electrolyte is higher than NMC 111 charged with LiPF₆ electrolyte (Figure 6) suggesting greater reactivity with BF₄⁻ than PF₆⁻ associated with EC dehydrogenation with the 1.0 M LiBF₄ electrolyte (with more free solvents, 73%). This argument is supported by the Ni2p and O1s spectra of NMC 111 charged with LiBF₄ electrolyte, where the Ni2p spectra shifted to higher binding energy from 854.5 eV to 856.0 eV with increasing potential, indicating the formation of lithium nickel oxyfluoride species, and the O lattice peak shift to higher binding energy with increasing potential. Similarly, NMC 622 charged in 1.0 M LiBF₄ electrolyte revealed higher intensity of LiMₓFᵧOₓ-like species than NMC 622 charged with the LiPF₆ electrolyte whereas the LiMₓFᵧOₓ-like species have comparable atomic percentages for NMC 811 charged with LiBF₄ and LiPF₆ electrolyte. This observation is consistent with the differences noted in the C=O region of the DRIFT spectra of NMC622 and 811 charged with LiBF₄ and LiPF₆ electrolytes. While DRIFT spectra of NMC622 charged with LiBF₄ electrolyte revealed earlier voltage onset of EC dehydrogenation than NMC622 charged with LiPF₆ electrolyte (4.1 V_{Li} with LiBF₄, 4.6 V_{Li} with
LiPF₆), NMC 811 results indicate similar onset potentials for EC dehydrogenation. The contribution of Li₃MₓFᵧOz-like species decreased with increasing potential for NMC 111, 622 and 811 (Figure 6d-f), suggesting surface removal of these F-containing species. The reduction of Li₃MₓFᵧOz-like species on charged NMC electrodes might expose the reactive oxide surface to the electrolyte for subsequent cycles and further degrade/oxidize the electrolyte. Moreover, the intensity of F1s collected from charged NMC622 and 811 are larger than NMC111, indicating greater reactivity towards LiBF₄ as the nickel content increases, which is in agreement with previous work with 1.0 M LiPF₆ electrolyte by Yu et al.¹⁴ Furthermore, unlike LiPF₆ results from our previous work¹⁴, there is almost no sign of formation of B-F-O species. This is due to the differences between hydrolysis of LiPF₆ and LiBF₄ salt. While the hydrolysis of LiPF₆ forms POF₃ species (LiPF₆ + H₂O -> LiF+POF₃+2HF), the hydrolysis of LiBF₄ does not form B-F-O species, but H₃BO₃ species instead (LiBF₄ + H₂O -> H₃BO₃+HF+LiF).¹¹⁹

**Discussion**

Carbonate dehydrogenation reactions can be suppressed (Figure 4) in concentrated electrolytes with lower free-carbonate activity (Figure 1), providing lower resistance growth (Figure 3) and much higher capacity retention for cycling of Ni-rich NMC electrodes (Figure 2). The enhanced capacity retention is not due to the formation of coatings on NMC811 in the concentrated electrolyte (Figure 6). The concentrated electrolyte solutions, which have lower free-carbonate activity, have higher stability against carbonate dehydrogenation. Cycling NMC811 electrodes in the 1st formation cycle in the concentrated electrolyte and replacing this with 1.0 M solution shows almost the same capacity decay as the cycling results with 1.0 M solution (Figure S23), and the changes in F1s spectra together with C1s and O1s spectra indicate the formation of a non-
protective EEI layer on NMC811 surfaces. Thus, we conclude that the higher capacity retention for the cycling experiment in 3.1 M LiPF$_6$ EC/EMC solution is not a result of the formation of a protective layer to passivate further decomposition, and the electrolyte used must always be concentrated to avoid parasitic decomposition reactions of the electrolyte at the electrode surface.

Intuitively, the higher stability could arise from a difference in dehydrogenation energy between free-carbonate and Li$^+$-bound carbonate. However, calculated bond dissociation energies for EC dehydrogenation are 2.10 eV and 2.13 eV in free-EC and Li$^+$-bound-EC, respectively, as computed at the DFT/B3LYP level of theory in implicit solvent, which is almost identical and cannot explain the above observations. We thus hypothesized that the higher stability of Li$^+$-bound carbonates is due to slower reaction kinetics at the oxide surface. The reaction profile for EC surface-mediated oxidative dehydrogenation has been computed by Østergaard et al. on Li$_{0.5}$CoO$_2$ with and without the involvement of a EC chemisorbed reaction intermediate, where the carbonyl carbon forms a bond with a surface oxygen (Figure 7). They reported that the EC chemisorption step lowers the activation barrier compared to a direct proton transfer path, which is kinetically inhibited (0.78 eV and 1.15 eV activation energy with and without the chemisorption step). Based on the requirement for the chemisorption step, which is formed after the carbonyl oxygen of EC coordinates to the metal site of oxide surface (electrophilic attack, Figure 7), we calculated the interaction energy between EC and the metal site of LiMO$_2$ (M=Mn, Co, Ni), Mn-doped LiNiO$_2$ and Co-doped LiNiO$_2$, as well as EC and Li$^+$ ion in the electrolyte solution (ion-dipole interaction) as shown in Table S8. The interaction energy between EC and the oxide surface was found to be -0.5 eV or weaker, while the interaction energy of EC and Li$^+$ (computed in vacuum) ranges from -2.37 eV for Li$^+$EC to -0.78 eV for Li$^+$(EC)$_3$, which indicates that EC preferentially interacts with Li$^+$ in the solution. This result can be understood intuitively, as the
EC-Li⁺ interaction is an ion-dipole interaction while the EC-oxide interaction is a dipole-dipole interaction, which is generally weaker. Therefore, if there are free-carbonate species in the electrolyte solution, this free-carbonate can interact with the oxide surface and form chemisorbed EC with a lower activation barrier for the dehydrogenation reaction. On the other hand, if the activity of free-carbonate is quite low such as the case where a dissociative salt was used in the highly concentrated region, Li⁺-bound-carbonate preferentially interacts with Li⁺ than with the oxide surface, making carbonate dehydrogenation more difficult. The thermodynamic reaction profile for Li(EC)₃ on LiNiO₂ with and without EC chemisorbed reaction intermediate, is reported in Figure 7a. We can see that the pathway involving the chemisorbed intermediate requires the desolvation of EC, which has an energy penalty of more than 1 eV. The proton transfer in the direct dehydrogenation pathway is thermodynamically uphill, and can be kinetically hindered, making the oxidative dehydrogenation of EC in Li(EC)₃ more difficult compared to free EC, where the reaction profile for both pathways is thermodynamically flat or downhill (Figure 7b). Notably, the dehydrogenated states, where dehydrogenated EC is adsorbed on the surface with and without the ring opening, is less thermodynamically stable for Li(EC)₃ compared to free EC, due to the stability of EC coordinated to Li in solution, indicating a much lower driving force for dehydrogenation of EC in Li(EC)₃. This proposed mechanism shows another electrolyte design principle to suppress parasitic reactions at the electrode-electrolyte interface via tuning the solvation structure and activity of free-solvent in the electrolyte solution.
Figure 7 Computed thermodynamic potential energy diagram for surface oxidative dehydrogenation of (a) Li (EC)$_3$ complex and (b) free EC on the (10\text{\overline{1}}4) surface of LiNiO$_2$, with (red line) and without (blue line) the formation of a chemisorbed reaction intermediate. The energies are referenced to the configuration where EC is bound to Li in solution for Li (EC)$_3$ and to EC with the carbonyl oxygen coordinated to the oxide metal site (electrophilic attack) for free EC. The configurations of reactants, products and reaction intermediates are shown in the insets. Li atoms are shown in light green, Ni in light gray, O in red, C in dark gray, and H in light blue. Li$^+$(solvent)$_3$ was identified as one of the most common solvation environments in highly concentrated solution such as 3.1 M LiPF$_6$ in EC/EMC (LiPF$_6$:EC/EMC=1:3) as supported by Raman spectroscopy shown in Figure 1.

Conclusions

We study the effect of electrolyte solvation structure by varying LiPF$_6$ concentration and replacing LiPF$_6$ with less dissociative LiBF$_4$ salt, on NMC111, 622 and 811 electrodes, combining battery cycling, EIS, Raman, FT-IR and XPS measurements. Solution Raman spectra show that the activity of free-carbonate species decreases either through increasing salt concentration or introducing a more dissociative salt such as LiPF$_6$ rather than LiClO$_4$ and LiBF$_4$. In particular, increasing LiPF$_6$ concentration reduced the activity of free-carbonate to the lowest value of 7 mol%
in 3.1 M LiPF$_6$ in EC:EMC (3:7wt) solution (LiPF$_6$:Carbonate = 1:3 mol/mol mixture). NMC811 electrodes cycled in highly concentrated LiPF$_6$ electrolyte exhibited higher capacity retention than electrolytes with a lower salt concentration, which is supported by less charge transfer impedance growth in the concentrated electrolyte as detected by EIS measurements. By combining Raman, DRIFT and XPS of NMC111, 622 and 811 charged in electrolytes with different LiPF$_6$ concentrations and replacing the salt with less dissociative LiBF$_4$ salt, we show evidence for earlier onset of EC dehydrogenation and the formation of protic species in the lower concentration and less dissociative electrolytes. However, by further increasing LiPF$_6$ concentration, carbonate dehydrogenation is suppressed even on the NMC811 surface. An oxide-mediated mechanism of electrolyte chemical oxidation can also be affected by solvation structure, where LiBF$_4$ has higher ionic association strength than LiPF$_6$ salt, having more free EC molecules/less Li$^+$ coordinated EC solvent in the electrolyte to dehydrogenate at earlier potentials. In addition, simply increasing LiPF$_6$ concentration can further suppress dehydrogenation due to much lower free-carbonate activity in the highly concentrated solution. Li$^+$-bound-carbonate molecules are less prone to interact with the transition-metal oxide surface, as the carbonate-Li$^+$ interaction is stronger than the carbonate-transition metal interaction. Therefore, it is unfavorable for carbonate species to oxidatively dehydrogenate on the oxide surface in concentrated solution which would lead to decomposition products on the NMC surface. In contrast, in less concentrated solutions where there is a considerable amount of free-carbonate species, free-carbonates can interact directly with the oxide surface, where they easily dehydrogenate. The resulting decomposition products behave as a resistor for the charge-transfer reaction, while resistance does not increase in concentrated solution, which has low free-carbonate activity. Through these findings, we propose the activity of free-carbonate as a descriptor for the chemical oxidation of EC, which provides insights into
the effect of solvation structure on electrolyte degradation in lithium ion batteries and rational design for novel salt and electrolyte. The concentrated electrolyte with lowest free-solvent activity studied in this work exhibited the highest capacity retention on NMC811 electrodes of >150 mAh$^{-1}$ after 500 cycles upon charging to 4.6 V$_{Li}$ (77% capacity retention and Coulombic efficiency of 99.7% 3-99th ave).

Supporting Information

Supporting Information includes 23 figures and 8 tables and can be found with this article online at xxx.

Arrhenius plots and activation energy for ionic conductivity and viscosity of electrolyte solutions (Figure S1), typical deconvolution example for Raman analysis (Figure S2), Raman spectra for pure solvent (Figure S3), Raman spectra for B-F region in LiBF$_4$ electrolyte (Figure S4), ATR-IR spectra for electrolytes (Figure S5), charge-discharge profile for 1$^{st}$ and 100$^{th}$ cycle where EIS was conducted (Figure S6), Cycling performance in electrolytes with different solvent (Figure S7), Cycling performance with lower cut-off voltage (Figure S8), Coulombic efficiency for charge-discharge experiments (Figure S9), EIS with potential holding (Figure S10), Electrochemical data for pellet electrodes for DRIFT (Figure S11), DRIFT spectra for simple mixture of salt and solvent (Figure S12), Raman spectra for charged pellet electrode (Figure S13), Capacity retention in electrolytes with different salt (Figure S14), NMC811 XPS for C1s and P2p region in concentrated electrolyte (Figure S15), Electrochemical data for carbon-free, binder-free powder electrodes for XPS with different concentration (Figure S16), NMC811 XPS for Li1s, Ni2p, Mn2p and Co2p region in concentrated electrolytes (Figure S17), NMC811, 622 and 111 XPS for C1s, B1s, Li1s, Ni2p, Mn2p and Co2p region in LiBF$_4$ electrolytes (Figure S18, Figure S19, Figure S20), Quantification results for XPS (Figure S21), Electrochemical data for carbon-free, binder-free powder electrodes for XPS with LiBF$_4$ (Figure S22), Cycling performance with replacing electrolyte after formation cycle (Figure S23), Concentration, viscosity, density and ionic conductivity of electrolyte solutions (Table S1, Table S2, Table S3), XPS deconvolution parameters (Table S4, Table S5, Table S6, Table S7), calculated interaction energies (Table S8).
Conflicts of Interest

The authors declare no competing interests.

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Supporting Information for

Enhanced Cycling Performance of Ni-rich Positive Electrodes (NMC) in Li-ion Batteries by Reducing Electrolyte Free-solvent Activity

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Table S 1 Molar ratio of LiPF$_6$:EC/EMC, viscosity ($\eta$), density ($d$), concentration of LiPF$_6$ ($c$) and ionic conductivity ($\sigma$) of LiPF$_6$ in EC:EMC (3:7wt) solutions at 25 °C.

| LiPF$_6$:EC/EMC | [Carbonate]/[Li] | $\eta$ (mPa s) | $d$ (g cm$^{-3}$) | $c$ (mol dm$^{-3}$) | $\sigma$ (mS cm$^{-1}$) |
|-----------------|-------------------|----------------|-------------------|-------------------|------------------|
| 1:3             | 3                 | 57.0           | 1.37              | 3.06              | 1.98             |
| 1:4             | 4                 | 20.0           | 1.32              | 2.41              | 4.17             |
| 1:5             | 5                 | 10.7           | 1.27              | 1.97              | 6.45             |
| 1:8             | 8                 | 4.32           | 1.22              | 1.30              | 9.15             |
| 1:10.5          | 10.5              | 3.00           | 1.19              | 1.00              | 8.88             |
| 1:25            | 25                | 1.47           | 1.14              | 0.43              | 6.63             |
| 1:50            | 50                | 1.22           | 1.12              | 0.22              | 4.15             |

pure EC/EMC (3:7wt) - 0.93 1.10 0

| 1 M LiX in EC/EMC | $\eta$ (mPa s) | $d$ (g cm$^{-3}$) | $\sigma$ (mS cm$^{-1}$) |
|-------------------|----------------|-------------------|------------------|
| LiPF$_6$          | 3.00           | 1.19              | 8.88             |
| LiClO$_4$         | 2.60           | 1.16              | 5.34             |
| LiBF$_4$          | 2.11           | 1.15              | 2.98             |

pure EC/EMC (3:7wt) 0.93 1.10 -

Table S 3 Viscosity ($\eta$), density ($d$) and ionic conductivity ($\sigma$) of LiPF$_6$ solution in different solvent at 25 °C. *Viscosity and density of EC was measured at supercooling liquid state at 25 °C as melting point of EC is ~35 °C. **Donor number of 16 is value for diethyl carbonate and 17.2 is value for dimethyl carbonate.

| $\xi$ | Donor number | $\eta$ (mPa s) | $d$ (g cm$^{-3}$) | $\sigma$ (mS cm$^{-1}$) |
|-------|--------------|----------------|-------------------|------------------|
| 1 M LiPF$_6$ in EC/EMC | 3.00           | 1.19              | 8.88             |
| 1.5 M LiPF$_6$ in EC | 11.00          | 1.44              | 7.19             |
| 1 M LiPF$_6$ in EMC | 1.70           | 1.11              | 4.47             |
| pure EC | 90            | 16.4              | ~2.7*             | 1.33*            |
| pure EMC | 3.0           | 16-17.2**        | 0.62              | 1.01             |
| pure EC/EMC (3:7wt) | 0.93           | 1.10              | -                 |
Figure S 1 Arrhenius plots for (a) ionic conductivity and (b) viscosity for various concentration range of LiPF₆ in EC:EMC (3:7wt) solution. To avoid the effect of glass transition, the plot was fitted only high temperature region since ionic conduction in solution electrolyte is generally governed by Vogel-Tammann-Fulcher (VTF) equation: \( R^\alpha = A \exp(-E_a/(T-T_0)) \), which is glass transition temperature corrected (\( T_0 \): Vogel temperature, glass transition in ideal glasses) Arrhenius relationship, and gives straight line only at high temperature region.[1] Calculated activation energy was shown in (c), which shows linear relationship between ionic conduction and viscosity.

Figure S 2 Typical results on the deconvolutions of Raman spectra by the Gaussian-Lorentzian function. (a) 850 -965 cm\(^{-1}\) (carbonate bands), (b) 700-760 cm\(^{-1}\) (PF₆\(^{-}\) bands) regions for 1.0 M LiPF₆ in EC:EMC (3:7wt) solution and (c) 850 -965 cm\(^{-1}\) (carbonate bands), (d) 700-760 cm\(^{-1}\) (PF₆\(^{-}\) bands) regions for 3.1 M LiPF₆ in EC:EMC (3:7wt) solution. The spectra for 780-900 cm\(^{-1}\) were deconvoluted into 6 bands (~875, 892, ~904, ~929.6, 938 and ~946 cm\(^{-1}\)) and that for 710-780 cm\(^{-1}\) were deconvoluted into 4 bands (~716, ~729, ~741.5, ~744.4 cm\(^{-1}\)).
**Figure S 3** Raman spectra of pure EMC and EC:EMC (3:7 wt:wt) mixtures measured at 25 °C.

**Figure S 4** Raman spectra of 1.0 M LiBF₄ in EC:EMC (3:7wt) solution in the B-F stretching range from 750 to 790 cm⁻¹ measured at 25 °C, with 632 nm laser to suppress fluorescence effect to avoid overlapping relatively weak B-F stretching signal. The band at 765.8 cm⁻¹ can be assigned to free-BF₄ and at 773.9 cm⁻¹ to bound-BF₄ to Li⁺ in the solution. Estimated molar fractions of Li⁺-bound BF₄⁻ ($I_{\text{bound-BF}_4} / I_{\text{total-BF}_4}$) in the solution based on the ratio of integral intensities is 26 %. The $R^2$ values for deconvolution are higher than 0.971.
Figure S 5 Attenuated total reflection (ATR) spectra of 1.0 M LiPF$_6$ (red), LiClO$_4$ (brown) and LiBF$_4$ (black) in EC/EMC (3:7 wt:wt). The bands at 1774 and 1807 cm$^{-1}$ represent the C=O stretching mode of EC molecule with and without Li coordination, and the bands located at 1720 cm$^{-1}$ and 1744 cm$^{-1}$ represent the C=O stretching of EMC molecule.

Figure S 6 Charge-discharge profile with (a) 1.0 M LiPF$_6$ and (b) 3.1 M LiPF$_6$ in EC/EMC in EC/EMC at 1st and 100th cycle conducted with 0.1C condition, while 2nd – 99th cycles were conducted at CC-CV 1C-0.05C condition. The cells were charged at 0.1C and potential was held at 3.9, 4.0, 4.1, 4.2, 4.3, 4.4, 4.5 and 4.6 V$_{Li}$ for 1 hour, and rest 1hour followed by EIS measurement (the dots in the figure are the points EIS measurement were conducted). Slightly larger discharge capacity at 100th cycle compared with the charge capacity is most likely come from residual charge from faster cycling before 99th cycle.
Figure S 7 Cycling performance of NMC811 composite electrodes with different electrolyte solutions upon charging to 4.6 V\textsubscript{Li} (2nd-99th cycle at CC-CV 1C-0.05C condition). At the 1st and 100th cycle, the cells were measured in a three-electrode configuration, which composed of NMC811|Li\textsubscript{4}Ti\textsubscript{5}O\textsubscript{12}-mesh|Li (see experimental section in detail), galvanostatically charged at 27.5 mA/g (0.1C). From the 2nd to the 99th cycle and after 101st cycle, the cells were tested with a two-electrode configuration that consisted NMC811 as the working electrode and Li metal as the counter electrode upon cycling between 4.6 and 2.0 V\textsubscript{Li}. Slightly lower capacity on the EC-only and EMC-only electrolyte than EC/EMC electrolyte can be attributed to the lower ionic conductivity of EC-only and EMC-only electrolyte compared with an EC/EMC mixture electrolyte (Table S 3).

Figure S 8 Charge-discharge cycle performance of NMC811 composite electrodes with different electrolyte solutions at cut-off voltage of 4.4 V\textsubscript{Li}. (a) Concentration dependent capacity retention (2nd-99th cycle at CC-CV 1C-0.05C condition) and charge-discharge profile with (b) 3.1 M LiPF\textsubscript{6} in EC/EMC and (c) 1.0 M LiPF\textsubscript{6} in EC/EMC. NMC811|Li\textsubscript{4}Ti\textsubscript{5}O\textsubscript{12}-mesh|Li three electrode cell was used. At 1st and 100th cycle, the cells were operated with three-electrode configuration and galvanostatically charged at 27.5 mA/g (0.1C) with measuring EIS. From 2nd to 99th cycle and after 101st cycle, the cells were operated with two-electrode configuration (NMC811 as working electrode and Li metal as reference/counter electrode) and cycle between 4.4 and 2.0 V\textsubscript{Li} with constant current – constant voltage (CCCV) condition (1C charge to 4.4 V\textsubscript{Li} and hold at 4.4 V\textsubscript{Li} until the current reach to 0.05C, then discharge at 1C to 2.0 V\textsubscript{Li}). Only fast cycle at 1C/0.05C (CC/CV) from 2nd to 99th cycles were shown.
**Figure S 9** Coulombic efficiency of battery cycling experiment on NMC811 composite electrode with different electrolyte concentration at room temperature. Original charge-discharge curves and capacity retention are shown in **Figure 2**. Coulombic efficiency of EIS cycle (1st and 100th cycle) are not plotted.

**Figure S 10** Nyquist plots for EIS measurement on NMC811 composite electrode in NMC811|Li₄Ti₅O₁₂-mesh|Li three electrode cell with potential held at 4.4 V_Li in (a) 1.0 M and (b) 3.1 M LiPF₆ / EC-EMC (3:7 wt/wt) solution at 25 °C. The cells were galvanostatically charged at 27.5 mA/g (0.1C) and held potential for 100 hours at 4.4 V_Li. Two semicircles were observed in the Nyquist plots for Li intercalation into composite electrodes like previous work.² The high-frequency semicircle (left) has been assigned to impedance at the electrified interface (predominantly lithium ion migration and adsorption/desorption in the pore structure of composite electrode) while the low-frequency semicircle (right) corresponds to the combination of charge transfer resistance at the electrode-electrolyte interface (equivalent circuit can be found in the inset).
Figure S 11  Electrochemical profiles of LiNi_{0.6}Mn_{0.2}Co_{0.2}O_{2} (NMC622) and LiNi_{0.8}Mn_{0.1}Co_{0.1}O_{2} (NMC811) pellet electrodes charged to 4.6 $V_{Li}$ at a 0.01C rate.

Figure S 12  DRIFT spectra of C=O stretching region for LiPF_{6}, EC, EC+LiPF_{6} mixture, Li$_x$Ni$_{0.8}$Mn$_{0.1}$Co$_{0.1}$O$_2$ (NMC811) soaked in EC:EMC (3:7wt) solution, LiPF$_6$ + soaked NMC811 in EC/EMC. Cumulative number of 256 was used at a 4 cm$^{-1}$ resolution. Spectra were subtracted with respect to a reference spectrum obtained with potassium bromide (KBr) powder. Although Li$^+$-EC band and free EC band around 1770 cm$^{-1}$ are overlapping, peak intensity ratio change between free EC and Li$^+$-EC indicates meaningful difference for the amount of free EC and Li$^+$-EC in the DRIFT samples.
**Figure S 13** Raman spectra for charged Li$_{x}$Ni$_{0.8}$Mn$_{0.1}$Co$_{0.1}$O$_2$ (NMC811) to 4.6 V$_{Li}$, where peaks at $\sim$732 and $\sim$907 cm$^{-1}$ is not emerged. The peaks at $\sim$732 and $\sim$907 cm$^{-1}$ can be attributed to the ring distortion and the ring breathing mode of dehydrogenated Li$^+$-EC.

**Figure S 14** Capacity retention as a function of cycles for NMC811 cycled between 4.6 V$_{Li}$ and 2 V$_{Li}$ in 1.0 M LiPF$_6$ in EC/EMC (3:7 w:w) and 1.0 M LiBF$_4$ in EC/EMC (3:7 w:w), with rate of 1 C ($\sim$275 mA/g) by using coin cell.
Figure S 15 Concentration dependent XPS spectra of the (a) C1s and (b) P2p photoemission lines collected from carbon-free, binder-free NMC811 electrodes (O1s and F1s spectra, and quantification results are shown in Figure 5). The electrodes were charged to 4.1, 4.2, 4.4 and 4.6 V_{Li} at 2.75 mA/g (0.01C) and held at the given potential for 5 hours using 1.0 M LiPF_{6} in EC: EMC (3:7 wt:wt) and 3.1 M LiPF_{6} in EC: EMC (3:7 wt:wt) solution and compared with pristine electrodes. All spectra were calibrated with the C1s photoemission peak of adventitious carbon at 285 eV. After subtraction of a Shirley-type background, all the spectra were normalized by fixing the C1s photoemission peak of adventitious carbon (285 eV) to the same value to facilitate comparison between samples. The spectra were assigned with the following contributions. C1s: C-H/C-C (E_{b} = 285 eV), C-O (E_{b}~286.3 eV), C=O/O-C-O (E_{b}~287.6 eV), O=C-O (E_{b}~288.8 eV) and CO_{3} (E_{b}~290.3 eV). P2p: phosphates (E_{b}~134 eV), Li_{x}PF_{y}O_{z} (E_{b}~136 eV) and LiPF_{6} (E_{b}~137.7 eV).
Table S 4 Binding Energies (eV) and Atomic Percentages (%) of the components from C1s, O1s, F1s, P2p, Li1s, Co2p, Mn2p, and Ni2p from the XPS Spectra of NMC 811 carbon-free, binder-free electrodes charged at different potentials using 3.1 M LiPF₆ in EC: EMC (3:7 wt:wt) as an electrolyte. 1.0 M data are available in previous publication.³

| Peak | Species   | Pristine | 4.1V | 4.2V | 4.4V | 4.6V |
|------|-----------|----------|------|------|------|------|
|      | BE (eV)   | FWH M (eV) | At%  | BE (eV) | FWH M (eV) | At%  | BE (eV) | FWH M (eV) | At%  | BE (eV) | FWH M (eV) | At%  |
| C1s  | C-C/C-H   | 285      | 1.4  | 20.6 | 285   | 1.5   | 6.9  | 285   | 1.6   | 21.3  | 285   | 1.6   | 13.8 | 285   | 1.6   | 7.3  |
|      | C-O       | 286.2    | 1.6  | 7.31 | 286.0 | 1.2   | 1.6  | 286.4 | 1.2   | 2.5  | 286.3 | 1.2   | 1.7  | 286.4 | 1.2   | 2.8  |
|      | C=O/O-C-O | 287.6    | 1.8  | 2.17 | 287.1 | 1.8   | 2.9  | 287.5 | 1.6   | 2.6  | 287.0 | 1.5   | 2.6  | 287.6 | 1.5   | 3.2  |
|      | O=C-O     | 288.8    | 1.6  | 5.39 | 288.8 | 2.0   | 1.3  | 289.0 | 1.5   | 1.9  | 288.8 | 1.6   | 2.4  | 288.9 | 1.5   | 1.7  |
|      | CO3       | 290.6    | 1.3  | 2.21 | 290.0 | 1.3   | 1.5  | 290.0 | 2.0   | 1.7  | 290.1 | 2.0   | 1.6  | 290.6 | 2.0   | 1.7  |
| O1s  | O lattice | 529.0    | 1.6  | 6.10 | 529.7 | 1.6   | 1.6  | 529.7 | 1.3   | 0.9  | 529.7 | 1.6   | 2.5  | 529.7 | 1.3   | 4.8  |
|      | ROLi      | 530.9    | 1.8  | 6.85 | 530.9 | 1.5   | 1.2  | 530.9 | 1.7   | 3.9  | 530.9 | 1.5   | 3.2  | 530.7 | 1.8   | 6.4  |
|      | Osurf/CO3/ O-C=O | 531.9 | 1.6  | 17.8 | 532.2 | 1.8   | 7.7  | 532.2 | 1.7   | 11.6 | 532.0 | 1.8   | 10.8 | 532.2 | 1.8   | 5.8  |
|      | C=O/C=O/OP(OR) | 533.6 | 1.5  | 2.22 | 533.4 | 1.7   | 4.3  | 533.4 | 1.4   | 4.1  | 533.4 | 1.6   | 3.7  | 533.4 | 1.4   | 2.7  |
|      | LiPF6Oz   | 534.9    | 1.4  | 0.28 | 534.8 | 1.8   | 1.9  | 534.9 | 1.4   | 0.4  | 534.7 | 1.8   | 0.9  | 534.6 | 1.8   | 2.3  |
| P1s  | LiF       | 685.0    | 1.8  | 1.90 | 685.2 | 1.8   | 11.4 | 685.2 | 1.8   | 10.1 | 685.0 | 1.8   | 11.5 | 685.1 | 1.8   | 16.5 |
|      | LiPF₆     | 687.0    | 1.8  | 0.24 | 687.0 | 1.8   | 15.1 | 687.0 | 2.0   | 5.4  | 686.5 | 2.0   | 10.3 | 686.5 | 1.8   | 7.5  |
|      | LiPF₆     | 688.0    | 2.0  | 1.54 | 688.3 | 2.3   | 12.6 | 688.0 | 2.2   | 7.2  | 688.0 | 2.1   | 5.5  | 688.3 | 2.3   | 4.4  |
|      | phosphate | 133.2    | 2.5  | 0.11 | /     | /     | /    | 134.0 | 2.5   | 0.7  | 134.0 | 2.5   | 0.6  | 134.0 | 1.5   | 0.2  |
| P2p  | LiPF₆     | 135.5    | 2.2  | 0.66 | 135.9 | 2.1   | 2.6  | 135.8 | 2.1   | 2.8  | 135.7 | 2.5   | 1.7  | 135.7 | 2.5   | 1.5  |
|      | LiPF₆     | 138.1    | 0.5  | 0.02 | 137.5 | 2.8   | 0.8  | 138.1 | 2.3   | 0.2  | 137.5 | 2.4   | 0.2  | 138.5 | 2.3   | 0.1  |
| L1s  | NMC/LiF   | 55.3     | 2.2  | 19.2 | 56.2 | 2.2   | 19.0 | 56.2 | 1.8   | 16.2 | 56.1 | 2.1   | 18.6 | 56.3 | 2.1   | 19.4 |
|      | LiPF₆     | /        | /    | /    | /     | /     | /    | /     | /     | /    | /     | /     | /    | /     | /     | /    |
| Co2p | NMC       | 779.3    | 2.2  | 0.31 | /     | /     | /    | 780.9 | 5.0   | 0.6  | 780.5 | 4.5   | 0.8  | 780.1 | 4.2   | 1.0  |
|      | satellite | /        | /    | /    | /     | /     | /    | /     | /     | /    | /     | /     | /    | /     | /     | /    |
| Ni2p | NMC       | 854.3    | 2.1  | 2.3  | 855.9 | 3.3   | 5.0  | 856.3 | 3.9   | 3.7  | 855.9 | 3.0   | 5.2  | 855.3 | 2.7   | 6.6  |
|      | satellite | 861.1    | 4.4  | 7.1  | 862.5 | 4.5   | 4.5  | 859.9 | 4.3   | 5.8  | 858.9 | 4.1   | 6.1  | 858.9 | 4.1   | 6.6  |
| Mn2p | NMC       | 641.7    | 6.9  | 2.7  | 642.0 | 5.0   | 1.4  | 642.5 | 5.0   | 1.8  | 641.6 | 5.0   | 2.4  | 641.1 | 5.0   | 3.9  |
**Figure S 16** Electrochemical profile of NMC811 carbon-free, binder-free electrodes charged at 100/C to 4.6 V\textsubscript{Li} using 3.1 M and 1.0 M LiPF\textsubscript{6} in EC: EMC 3:7 (wt:wt). Overpotential for 3.1 M is most likely due to higher viscosity and lower ionic conductivity. Note that carbon-free, binder-free electrodes were prepared by just drop casting oxide powder on to Al foil and have poor contact.

**Figure S 17** Li1s/Co3p, Ni2p, Mn2p and Co2p XPS spectra of NMC811 carbon-free, binder-free pristine electrodes and after charging at 4.1, 4.2, 4.4, and 4.6 V\textsubscript{Li} using (a) 1.0 M LiPF\textsubscript{6} and (b) 1.0 M in EC:EMC 3:7 (wt:wt).
Figure S 18 C1s and B1s XPS spectra of (a) NMC111, (b) NMC622 and (c) NMC811 carbon-free, binder-free pristine electrodes and after charging at 4.1, 4.2, 4.4, and 4.6 V_{Li} using 1.0 M LiBF_{4} in EC:EMC 3:7 (wt:wt).

Figure S 19 Mn2p and Co2p XPS spectra of (a) NMC111, (b) NMC622 and (c) NMC811 carbon-free, binder-free pristine electrodes and after charging at 4.1, 4.2, 4.4, and 4.6 V_{Li} using 1.0 M LiBF_{4} in EC:EMC 3:7 (wt:wt).
**Figure S 20** Ni2p and Co3p/Li1s XPS spectra of (a) NMC111, (b) NMC622 and (c) NMC811 carbon-free, binder-free pristine electrodes and after charging at 4.1, 4.2, 4.4, and 4.6 V\textsubscript{Li} using 1.0 M LiBF\textsubscript{4} in EC:EMC 3:7 (wt:wt).

**Figure S 21** Atomic percentages of XPS spectra for F in metal fluorides, C-O bonds, C=O/O-C-O, O-C=O and CO\textsubscript{3} bonds (from C1s in Figure S18), and O lattice in the pristine and charged (a) NMC111, (b) NMC622, (c) NMC811 carbon-free, binder-free electrodes at 4.1, 4.2, 4.4, and 4.6 V\textsubscript{Li} using 1.0 M LiBF\textsubscript{4} in EC: EMC 3:7 (wt:wt).
**Figure S 22** Electrochemical profile of (a) NMC111 (black) (b) NMC622 (dark gray) (c) NMC811 (gray) carbon-free, binder-free pristine electrodes charged at 4.6 $V_{Li}$ using 1.0 M LiBF$_4$ in EC: EMC 3:7 (wt:wt).
Table S 5 Binding Energies (eV) and Atomic Percentages (%) of the components from C1s, O1s, F1s, B1s, Li1s, Co2p, Mn2p, and Ni2p from the XPS Spectra of NMC111 carbon-free, binder-free electrodes charged at different potentials using 1.0 M LiBF4 in EC: EMC (3:7 wt:wt) as an electrolyte.

| Peak         | Species                  | BE (eV) | FWHM (eV) | At% | BE (eV) | FWHM (eV) | At% | BE (eV) | FWHM (eV) | At% | BE (eV) | FWHM (eV) | At% |
|--------------|--------------------------|---------|-----------|-----|---------|-----------|-----|---------|-----------|-----|---------|-----------|-----|
| C1s          | C=C/C-H                  | 285.4   | 21.5      | 285 | 1.6     | 10.7      | 285 | 1.6     | 13.0      | 285 | 1.5     | 12.5      | 285 |
|              | C-O                      | 286.4   | 3.7       | 286 | 3       | 1.2       | 286 | 5       | 1.2       | 286 | 5       | 4.3       | 286 | 5       |
|              | C=O/O-C-O                | 287.6   | 3.5       | 287 | 0       | 1.8       | 287 | 5       | 1.8       | 287 | 6       | 1.8       | 287 | 5       |
|              | O=O-C                    | 288.8   | 3.5       | 288 | 8       | 1.6       | 288 | 8       | 2.2       | 288 | 8       | 1.8       | 288 | 9       |
|              | CO3                      | 290.6   | 0.6       | 290 | 6       | 2.0       | 290 | 6       | 0.8       | 290 | 6       | 2.0       | 290 | 6       |
|              | O lattice                | 529.5   | 21.1      | 529 | 7       | 11.2      | 529 | 9       | 16.2      | 529 | 9       | 13.8      | 529 | 9       |
|              | ROLi                     | 531.0   | 6.1       | 530 | 7       | 5.4       | 530 | 9       | 4.8       | 530 | 8       | 4.9       | 530 | 7       |
|              | Osurf/CO3/O-C=O          | 531.8   | 9.3       | 531 | 9       | 8.7       | 532 | 0       | 11.8      | 532 | 1       | 11.0      | 531 | 9       |
|              | C=O/(Li/Getter)          | 533.4   | 20       | 533 | 4       | 3.9       | 533 | 4       | 3.4       | 533 | 7       | 6.8       | 533 | 4       |
|              |                          | /       | /         | /   | /       | /         | /   | /       | /         | /   | /       | /         | /   | /       |
| F1s          | LiF                      | 685.0   | 2.8       | 685 | 4       | 14.5      | 685 | 0       | 9.8       | 685 | 3       | 4.9       | 685 | 0       |
|              | LiBFy                    | 686.6   | 0.9       | 686 | 5       | 5.4       | 686 | 5       | 3.7       | 686 | 6       | 5.4       | 686 | 5       |
|              | LiBF4                    | 688.5   | 1.5       | 688 | 5       | 2.3       | 688 | 5       | 0.9       | 688 | 5       | 1.5       | 688 | 5       |
| B1s          | LiBFy                    | /       | /         | /   | /       | /         | /   | /       | /         | /   | /       | /         | /   | /       |
|              | LiBF4                    | /       | /         | /   | /       | /         | /   | /       | /         | /   | /       | /         | /   | /       |
| Li1s         | NMC                      | 54.4    | 1.3       | 54.4| 1.3     | 11.3      | 55.7| 2.4     | 12.5      | 56.3| 2.3     | 7.7       | 56.3| 2.3     |
|              | Li2CO3/LiF               | /       | /         | /   | /       | /         | /   | /       | /         | /   | /       | /         | /   | /       |
| Co2p         | NMC                      | 779.9   | 2.5       | 779 | 9       | 1.7       | 780 | 4       | 2.7       | 780 | 4       | 2.4       | 780 | 5       |
|              | satellite                | 789.7   | 8.0       | 789 | 5       | 8.0       | /   | /       | /         | /   | /       | /         | /   | /       |
| Ni2p         | NMC                      | 854.1   | 4.3       | 855 | 1       | 2.5       | 855 | 7       | 5.3       | 855 | 7       | 3.0       | 855 | 7       |
|              | satellite                | 861.2   | 6.0       | 862 | 3       | 4.8       | 862 | 3       | 6.0       | 862 | 2       | 6.0       | 862 | 3       |
| Mn2p         | NCM                      | 641.9   | 4.4       | 642 | 3       | 2.5       | 642 | 5       | 5.3       | 642 | 5       | 4.7       | 642 | 5       |

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Table S 6 Binding Energies (eV) and Atomic Percentages (%) of the components from C1s, O1s, F1s, B1s, Li1s, Co2p, Mn2p, and Ni2p from the XPS Spectra of NMC622 carbon-free, binder-free electrodes charged at different potentials using 1.0 M LiBF\textsubscript{4} in EC: EMC (3:7 wt:wt) as an electrolyte.

| NMC622 | Pristine | 4.1V | 4.2V | 4.4V | 4.6V |
|--------|----------|------|------|------|------|
| Peak   | Species  | BE (eV) | FW HM (eV) | At% | BE (eV) | FW HM (eV) | At% | BE (eV) | FW HM (eV) | At% | BE (eV) | FW HM (eV) | At% |
| C1s    | C-C/C-H  | 285  | 1.4 | 24.42 | 285  | 1.6 | 11.5 | 285  | 1.6 | 12.1 | 285  | 1.6 | 11.3 | 285  | 1.6 | 16.9 |
|  | C-O      | 286.2 | 1.5 | 10.33 | 286.2 | 1.6 | 3.2 | 286.3 | 1.2 | 2.3 | 286.5 | 1.2 | 2.7 | 286.5 | 1.2 | 1.1 |
|  | C=O/O=C-O | 287.6 | 1.8 | 4.72 | 287.6 | 1.7 | 1.3 | 287.2 | 1.7 | 2.9 | 287.6 | 1.5 | 1.3 | 287.6 | 1.8 | 6.2 |
|  | O=C-O    | 288.8 | 1.6 | 4.23 | 289.1 | 1.6 | 1.6 | 288.8 | 1.6 | 3.3 | 288.8 | 1.6 | 2.4 | 288.8 | 1.5 | 5.9 |
|  | O3       | 290.0 | 1.3 | 1.26 | 290.6 | 1.7 | 0.5 | 290.5 | 2.0 | 1.1 | 290.3 | 2.0 | 0.8 | 290.6 | 2.0 | 0.9 |
| O1s    | O lattice | 528.9 | 1.5 | 8.33 | 529.6 | 1.3 | 1.6 | 529.4 | 1.3 | 0.9 | 529.3 | 1.3 | 1.5 | 529.4 | 1.3 | 8.5 |
|  | ROLi     | 530.8 | 1.8 | 6.21 | 530.9 | 1.8 | 1.6 | 530.8 | 1.6 | 1.1 | 530.6 | 1.8 | 1.6 | 530.9 | 1.6 | 6.7 |
|  | Osurf/C03/O-C=O | 531.9 | 1.8 | 14.37 | 532.2 | 1.7 | 11.7 | 532.2 | 1.8 | 11.7 | 531.9 | 1.8 | 16.5 | 532.1 | 1.7 | 11.4 |
|  | C-O(O-| 533.5 | 1.6 | 3.12 | 533.4 | 1.8 | 2.9 | 533.6 | 1.8 | 3.9 | 533.4 | 1.8 | 4.0 | 533.9 | 1.8 | 6.7 |
| F1s    | LiF      | / / / | 685.5 | 1.8 | 17.9 | 685.0 | 1.8 | 20.9 | 685.3 | 1.8 | 16.7 | 685.0 | 1.8 | 6.7 |
|  | LiBF\textsubscript{4} | / / / | 686.5 | 1.8 | 6.2 | 687.0 | 1.8 | 2.7 | 686.5 | 1.8 | 3.7 | 687.0 | 2.3 | 6.0 |
|  | LiBF\textsubscript{4} | / / / | 688.5 | 2.3 | 1.9 | 688.5 | 2.3 | 3.9 | 688.4 | 1.8 | 0.5 | 688.5 | 3.4 | 3.7 |
| B1s    | LiBF\textsubscript{4} | / / / | 193.0 | 6.3 | 5.1 | 193.0 | 2.5 | 6.3 | 192.7 | 3.9 | 5.0 | 192.7 | 3.2 | 5.1 |
| Li1s   | NMC      | 54.9 | 3.2 | 15.64 | / / / | / / / | / / / | / / / | / / / | / / / | / / / | / / / | / / / | / / /
|  | Li2CO3/LiF | / / / | 56.2 | 2.1 | 28.5 | 55.6 | 1.8 | 27.6 | 56.0 | 2.2 | 28.2 | 56.1 | 2.2 | 5.8 |
| Co2p   | NMC      | 779.5 | 1.8 | 0.85 | 780.2 | 2.9 | 2.0 | / / / | 780.2 | 2.4 | 1.2 | 780.1 | 1.9 | 2.2 |
|  | satellite | 789.2 | 4.0 | / / / | / / / | / / / | / / / | / / / | / / / | / / / | / / / | / / / | / / /
| Ni2p   | NMC      | 854.3 | 2.0 | 2.98 | 857.0 | 4.7 | 2.2 | 857.3 | 3.0 | 1.8 | 857.0 | 3.0 | 2.2 | 855.6 | 2.9 | 4.1 |
|  | satellite | 861.3 | 4.4 | / / / | / / / | / / / | / / / | / / / | / / / | / / /
| Mn2p   | NCM      | 642.0 | 4.8 | 3.04 | 642.4 | 4.2 | 0.9 | 642.2 | 5.0 | 0.8 | 641.6 | 4.6 | 1.0 | 642.2 | 3.4 | 3.0 |
Table S 7 Binding Energies (eV) and Atomic Percentages (%) of the components from C1s, O1s, F1s, B1s, Li1s, Co2p, Mn2p, and Ni2p from the XPS Spectra of NMC811 carbon-free, binder-free electrodes charged at different potentials using 1.0 M LiBF₄ in EC: EMC (3:7 wt:wt) as an electrolyte.

| Peak     | Species         | NMC 811 | Pristine | 4.1V | 4.2V | 4.4V | 4.6V |
|----------|-----------------|---------|----------|------|------|------|------|
|          |                 | BE (eV) | FW HM (eV) | At% | BE (eV) | FW HM (eV) | At% | BE (eV) | FW HM (eV) | At% | BE (eV) | FW HM (eV) | At% |
| C1s      | C=C/C-H         | 285     | 1.4 | 20.63 | 285 | 1.6 | 10.4 | 285 | 1.6 | 8.4 | 285 | 1.6 | 14.1 | 285 | 1.6 | 9.2 |
|          | C-O             | 286.2   | 1.6 | 7.31 | 286.4 | 1.2 | 1.5 | 286.4 | 1.2 | 1.6 | 286.5 | 1.3 | 2.0 | 286.5 | 1.2 | 3.0 |
|          | C=O/O-C-O       | 287.6   | 1.8 | 2.17 | 287.6 | 1.5 | 1.0 | 287.6 | 1.5 | 1.0 | 287.5 | 1.8 | 1.3 | 287.6 | 1.5 | 2.6 |
|          | O=C-O           | 288.8   | 1.6 | 5.39 | 288.9 | 1.6 | 1.2 | 288.9 | 1.6 | 1.0 | 289.0 | 1.6 | 1.1 | 288.9 | 1.5 | 2.9 |
|          | CO3             | 290.0   | 1.3 | 2.21 | 290.6 | 2.0 | 0.6 | 290.6 | 2.0 | 0.8 | 290.6 | 1.7 | 0.5 | 290.6 | 2.0 | 2.3 |
| O1s      | O lattice       | 529.0   | 1.6 | 6.10 | 529.4 | 1.3 | 1.6 | 529.4 | 1.3 | 0.3 | 529.5 | 1.3 | 4.2 | 529.6 | 1.3 | 7.4 |
|          | ROLi            | 530.9   | 1.8 | 6.85 | 530.5 | 1.5 | 2.4 | 530.5 | 1.6 | 2.6 | 530.8 | 1.5 | 4.0 | 530.9 | 1.5 | 5.8 |
|          | Osurf/C/O-C-O   | 531.9   | 1.6 | 17.81 | 532.2 | 1.8 | 7.2 | 532.2 | 1.8 | 5.8 | 532.2 | 1.8 | 7.2 | 532.2 | 1.8 | 12.0 |
|          | C-O/O-C=O/OP(OR) | 533.6 | 1.5 | 2.22 | 533.4 | 1.8 | 4.7 | 533.4 | 1.7 | 2.8 | 533.4 | 1.8 | 4.4 | 533.9 | 1.8 | 6.4 |
| F1s      | LiF             | 685.0   | 1.8 | 1.90 | 685.4 | 1.7 | 20.4 | 685.4 | 1.7 | 26.8 | 685.4 | 1.7 | 17.5 | 685.4 | 1.8 | 9.9 |
|          | LiₓBxFᵧ         | 687.0   | 1.8 | 0.24 | 686.5 | 1.8 | 2.5 | 686.5 | 2.0 | 4.2 | 686.5 | 2.1 | 3.7 | 686.5 | 2.1 | 4.4 |
|          | LiₓBxFᵧ         | 688.0   | 2.0 | 1.54 | 688.4 | 2.3 | 3.3 | 688.4 | 2.3 | 0.8 | 688.4 | 2.3 | 2.4 | 688.5 | 2.3 | 4.6 |
| B1s      | LiₓBxFᵧ         | /       | /   | /    | 193.3 | 2.7 | 6.9 | 193.3 | 2.6 | 4.1 | 193.3 | 3.3 | 6.6 | 193.3 | 2.9 | 4.9 |
|          | LiₓBxFᵧ         | /       | /   | /    | /     | /   | /   | /     | /   | /   | /     | /   | /   | /     | /   | /   |
| Li1s     | NMC             | 55.3    | 2.2 | 19.27 | /   | /   | /   | /   | /   | /   | /   | /   | /   | /   | /   | /   |
|          | Li2CO3/LiF      | /       | /   | /    | /     | /   | /   | /     | /   | /   | /     | /   | /   | /     | /   | /   |
| Co2p     | NMC             | 779.3   | 2.2 | 0.31 | 780.3 | 2.9 | 0.3 | 780.2 | 3.8 | 0.4 | 780.2 | 2.9 | 0.8 | 780.7 | 3.5 | 1.3 |
|          | satellite       | /       | /   | /    | /     | /   | /   | /     | /   | /   | /     | /   | /   | /     | /   | /   |
| Ni2p     | NMC             | 854.3   | 2.1 | 0.31 | 855.9 | 3.0 | 0.3 | 857.7 | 6.2 | 4.3 | 855.7 | 2.7 | 5.1 | 855.9 | 2.6 | 3.9 |
|          | satellite       | 861.1   | 4.4 | /    | 859.5 | 6.0 | /   | 858.7 | 4.7 | /   | 862.7 | 4.0 | /   | 862.7 | 6.0 | /   |
| Mn2p     | NMC             | 641.7   | 6.9 | 2.7  | 641.6 | 5.0 | 3.3 | 641.7 | 4.6 | 1.1 | 641.8 | 4.6 | 2.4 | 641.7 | 4.4 | 1.0 |
Figure S 23 Cycling performance of NMC811 composite electrodes upon charging to 4.6 V_{Li} (2nd-99th cycle at CC-CV 1C-0.05C condition). The electrode cycled in 3.1 M LiPF_6 in EC/EMC solution for 1st cycle, replaced electrolyte to 1.0 M solution and started 2nd cycle. At the 1st and 100th cycle, the cells were measured in a three-electrode configuration, which composed of NMC811|Li_4Ti_5O_{12}-mesh|Li (see experimental section in detail), galvanostatically charged at 27.5 mA/g (0.1C). From the 2nd to the 99th cycle and after 101st cycle, the cells were tested with a two-electrode configuration that consisted NMC811 as the working electrode and Li metal as the counter electrode. The collected electrode was washed with 1.0 M solution before cell assembly.

Table S 8 Calculated interaction energy between carbonyl oxygen of EC and the metal site of oxide surface (electrophilic attack), for LiMO_2 (M = Mn, Co and Ni) and LiNiO_2 where one surface Ni has been replaced with Co and Mn (Co-doped and Mn-doped LiNiO_2, respectively). The interaction energy of EC with Li^+ ion and Li^+(EC)_n complexes (ion-dipole interaction), computed as the energy of the reaction: Li^+(EC)_{n-1} + EC → Li^+(EC)_n, is also reported.

| Site               | Interaction Energy / eV |
|--------------------|-------------------------|
| LiMnO_2            | Mn                      | -0.19                   |
| LiCoO_2            | Co                      | -0.33                   |
| LiNiO_2            | Ni                      | -0.50                   |
| LiNiO_2            | Li                      | -0.41                   |
| Co-doped LiNiO_2   | Co                      | -0.14                   |
| Mn-doped LiNiO_2   | Mn                      | -0.26                   |
| Li^+(EC)           | -                       | -2.37                   |
| Li^+(EC)_2         | -                       | -1.19                   |
| Li^+(EC)_3         | -                       | -0.78                   |
Reference for Supporting Information

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