

Enhanced Ion Conduction in $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$ via Anion Doping

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Cite This: <https://dx.doi.org/10.1021/acs.chemmater.0c00025>

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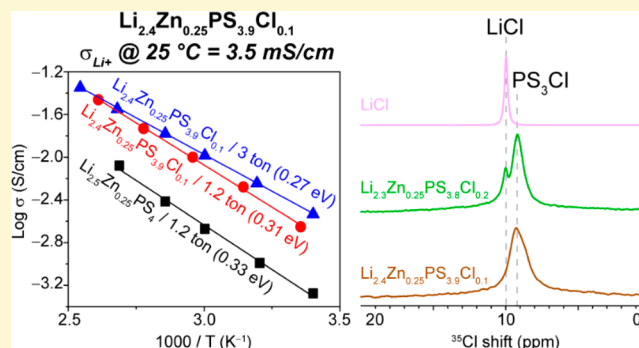
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ABSTRACT: $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$ was computationally predicted to have superionic conductivity of over 50 mS/cm. However, experimental efforts so far have only yielded ionic conductivities on the order of 10^{-4} S/cm, due to difficulty in producing phase-pure crystalline products. Here, to improve phase purity and enhance ionic conductivity, Cl⁻ doping is employed to synthesize $\text{Li}_{1+2x-y}\text{Zn}_{1-x}\text{PS}_{4-y}\text{Cl}_y$, which is shown to stabilize $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$ from decomposing to Li_3PS_4 . $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$ which shows the highest ionic conductivity among the $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$ samples is chosen as an example to study. After incorporating only a small amount of Cl, the ionic conductivity increases from 0.6 mS/cm for $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$ to 3.5 mS/cm for $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ at 25 °C. In addition, the activation energy is reduced from 0.33 to 0.27 eV. The phase purity and fine structure of $\text{Li}_{1+2x-y}\text{Zn}_{1-x}\text{PS}_{4-y}\text{Cl}_y$ are determined with XRD, and ^7Li , ^{31}P , ^{35}Cl solid-state NMR characterizations. The experimental data confirm the success of Cl⁻ doping. In addition, $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ exhibits a low electronic conductivity of 1.21×10^{-9} S/cm and a wide electrochemical stability window.



INTRODUCTION

All-solid-state batteries are promising to offer enhanced safety and energy density.^{1–3} Developing highly conductive solid electrolytes is essential to high performance all-solid-state batteries, especially for high-power applications. Researchers continue to search for new types of solid electrolytes with high ionic conductivities and stability.^{4,5} Wang et al. discovered the body-centered cubic feature in several superionic sulfide conductors and used it as a screening descriptor for potential new ionic conductors.³ Following this work, Richards et al. investigated a sulfide material LiZnPS_4 with a body-centered cubic sublattice with density functional theory (DFT) and ab initio molecular dynamics (AIMD).⁵ Taking both thermal stability and ionic conductivity into account, the optimized composition is $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$, in which excess Li^+ partially replaces Zn^{2+} . It is predicted to have a high ionic conductivity of more than 50 mS/cm and a low activation energy of 0.14 eV.

Recently, experimental efforts have been made to prepare $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$, aiming to achieve the predicted ionic conductivity. To prepare $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$ via solid state reaction, nanosized ZnS is needed to improve the reactivity at relatively low temperature.⁶ Impurities were found within all the $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$ samples, such as $\text{Li}_4\text{P}_2\text{S}_6$ with small x values and Li_3PS_4 with large x values. The impurities led to much lower conductivity than the computed value. The highest ionic conductivity obtained from reported experiments is only 0.84 mS/cm with $x = 0.75$ ($\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$). Naoki et al. also prepared $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$ from LiZnPS_4 and Li_3PS_4 , in order to

suppress the evaporation of S and thus formation of $\text{Li}_4\text{P}_2\text{S}_6$ impurity.⁷ But crystallinity decreases with increasing x , and the highest ionic conductivity achieved is 0.57 mS/cm at $x = 0.625$ ($\text{Li}_{2.25}\text{Zn}_{0.375}\text{PS}_4$).

In order to prepare highly conductive $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$, phase pure crystalline products with large x values are needed. In addition, Cl⁻ doping in the S^{2-} sites is shown as a good strategy to enhance ionic conductivity of sulfide-based electrolytes. Kanno et al. reported a Cl⁻ doped $\text{Li}_{9.54}\text{Si}_{1.74}\text{P}_{1.44}\text{S}_{11.7}\text{Cl}_{0.3}$, which delivers a high ionic conductivity of 25 mS/cm at room temperature.⁸ Cl⁻ was also introduced to tetragonal Na_3PS_4 , which has significantly enhanced the ionic conductivity.^{9,10} Here, we introduce a small amount of LiCl to partially replace Li_2S as one precursor, which is shown to suppress the formation of Li_3PS_4 and enhance crystallinity. The optimized conductivity reaches 3.5 mS/cm for $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$, which is about 4 times the values reported for $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$.⁶ $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ also exhibits significantly enhanced stability with a low electronic conductivity and a large electrochemical stability window.

Received: January 4, 2020

Revised: March 14, 2020

Published: March 16, 2020

EXPERIMENTAL SECTION

ZnS nanoparticles were prepared by a precipitation method using $\text{Zn}(\text{CH}_3\text{COO})_2$ and Na_2S as the source of Zn and S.¹¹ Li_2S , P_2S_5 , and LiCl were purchased from Sigma-Aldrich and LiCl was dried under dynamic vacuum at 200 °C for 12 h before use. Li_2S , nanosized ZnS , P_2S_5 , and LiCl were mixed with a $\text{Li}:\text{P}:\text{S}:\text{Cl}$ molar ratio of $(1 + 2x - y):(1 - x):(4 - y):y$ and manually ground for 10 min. The premixed powders were then placed in a ZrO_2 jar (two ZrO_2 balls; 10 mm) and ball-milled (Spex 8000M) for 20 h under vacuum. After ball-milling, the mixed powders were pelletized into a disk (6 mm in diameter and 1 mm in thickness) and sintered at 260 °C for 12 h under Ar. All operations were performed under the protection of Ar gas in a glovebox (Mbraun , $\text{H}_2\text{O} < 0.5$ ppm, $\text{O}_2 < 0.5$ ppm).

Powder XRD experiments were performed using a Philips X'Pert powder diffractometer at 45 kV and 40 mA with $\text{Cu K}\alpha$ radiation ($\lambda = 1.5406$ Å). The as-synthesized pellets were ground gently into fine powders for a short time and then placed on a zero-background holder, which was sealed with a Kapton film to avoid air exposure. The patterns were collected at room temperature with a 2θ range from 10° to 80° and a scan rate of $2^\circ/\text{min}$.

^6Li , ^7Li , and ^{31}P magic-angle-spinning (MAS) NMR experiments were performed on a Bruker Avance-III 500 spectrometer at the Larmor frequency of 73.6, 194.4, and 202.4 MHz, respectively. A spinning rate of 25 kHz was used for all the experiments. A single pulse sequence was employed to acquire all ^6Li and ^7Li NMR spectra with a solid 90° pulse length and a recycle delay of 4.75 μs and 500 s and 3.35 μs and 5 s for ^6Li and ^7Li NMR, respectively. ^{31}P NMR was recorded with a rotor-synchronized spin-echo sequence with a 90° pulse length of 4.2 μs and a recycle delay of 200 s. ^6Li and ^{31}P NMR spectra were calibrated to $\text{LiCl}(\text{s})$ at -1.1 ppm¹² and 85% $\text{H}_3\text{PO}_4(\text{l})$ at 0 ppm, respectively. ^{35}Cl MAS NMR experiments were performed on a Bruker 830 spectrometer using a home-built 3.2 mm probe¹³ at the Larmor frequency of 81.3 MHz. The rotor-synchronized spin-echo sequence with a solid 90° pulse length of 2.9 μs and a recycle delay of 20 s was employed. ^{35}Cl NMR spectra were calibrated to $\text{LiCl}(\text{s})$ at 9.93 ppm.

Variable-temperature (VT) solid-state ^7Li MAS NMR experiments were carried out on a Bruker Avance III spectrometer (11.75 T) with a ^7Li Larmor frequency at 194.319 MHz and a spinning rate of 5 kHz. VT ^7Li MAS NMR spectra of $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ were acquired using the Bruker Laser probe¹⁴ from 25 to 300 °C. The temperature was calibrated with the ^{79}Br NMR shift of KBr .¹⁵ After adjusting the temperature to the desired value before each acquisition, 5 min (during heating) or 15 min (during cooling) was allowed to reach thermal equilibration. One-dimensional ^7Li MAS NMR spectra were collected by a 90° single-pulse excitation of 3.15 μs and with a recycle delay of 5 s. Sixteen scans were accumulated for each spectrum. After each acquisition of a 1D ^7Li MAS NMR spectrum, ^7Li longitudinal NMR relaxation time (T_1) was measured with an inversion–recovery pulse sequence. ^7Li chemical shift was referenced to 1 M $\text{LiCl}(\text{l})$ at 0 ppm. All the analyses of NMR spectra and ^7Li T_1 times were performed on Bruker Topspin v.4.0.3.

The ionic conductivity of $\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$ was measured at 25 °C using Ac electrochemical impedance spectroscopy (EIS) in the frequency range from 5 MHz to 1 Hz with a potential of 50 mV. To extract the activation energy of ion conduction in $\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$, variable-temperature impedance measurements were then performed from room temperature to 120 °C in a CSZ microclimate chamber. The as-synthesized $\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$ disks were sandwiched by indium foils (4.7 mm in diameter) and sealed in a home-built cylindrical cell¹⁶ for all measurements. The electronic conductivity of $\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$ was measured by applying a voltage of 3 V on $\text{In}|\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y|\text{In}$ and measuring current until it becomes stable. Cyclic voltammetry measurements were carried out on $\text{Li}|\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}|\text{Stainless steel}$ between -0.3 and 5 V at a scanning rate of 10 mV/s.

RESULTS AND DISCUSSION

The space group of pristine LiZnPS_4 is I_4^- , with Li^+ fully occupying 2a sites and Zn^{2+} fully occupying 2b sites. After partly replacing Zn^{2+} with Li^+ ($\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$), the 2b sites are shared by Li^+ and Zn^{2+} , and extra Li^+ occupies 2d sites (Figure 1a). Previous work⁶ has shown that Li_3PS_4 and Li_2PS_3 are two

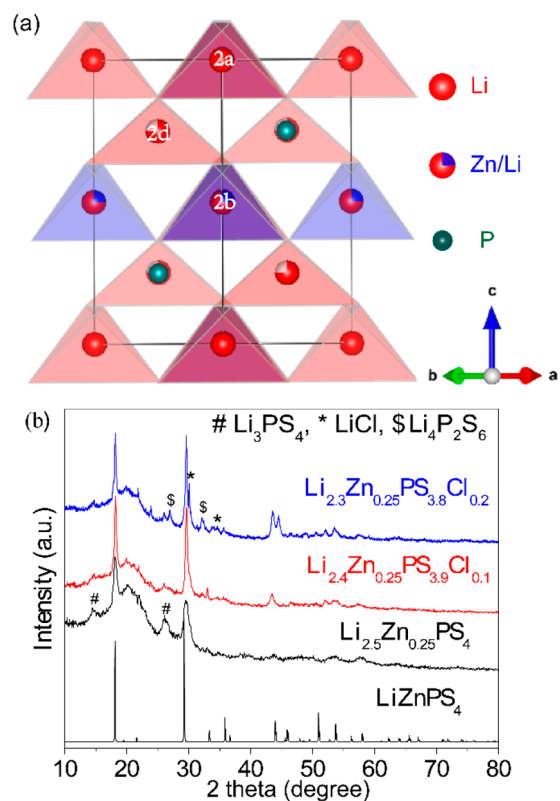


Figure 1. (a) Crystal structure of $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$, (b) XRD patterns of $\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$ ($y = 0, 0.1, 0.2$); the standard XRD pattern of LiZnPS_4 is shown as a reference.

observed impurities accompanying the synthesis of $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$. This work suggests when $x \leq 0.5$, the main impurity is $\text{Li}_4\text{P}_2\text{S}_6$ (Figure S1a), with P^{5+} partially reduced to P^{4+} . When the x value increases to 0.75 ($\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$), $\text{Li}_4\text{P}_2\text{S}_6$ almost disappears and Li_3PS_4 emerges as the main impurity phase. This is consistent with computational studies reporting that when the amount of Zn reduces, more Li occupies the 2b and 2d sites.⁵ The structure becomes unstable and is likely to decompose to Li_3PS_4 and $\text{Li}_{1+2x-2\delta}\text{Zn}_{1-x+\delta}\text{PS}_4$. Another change with higher x values is the peak intensity decreases in the XRD pattern, which infers lower crystallinity. The conductivity of $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$ is maximized at $x = 0.75$ ($\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$) (Figure 2), which is about 0.6 mS/cm at 25 °C (Figure S1b), which agrees very well with previous findings.^{6,7}

To further improve the ionic conductivity of $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$, anion-doping with Cl^- is employed. Cl^- replacement of S^{2-} will lead to slight Li deficiency, structural distortion, and changes in Li^+ –host interactions. Therefore, $\text{Cl}^- \rightarrow \text{S}^{2-}$ exchange will affect the stability of the resulting structure and impact Li^+ transport. As shown in Figure 1b, the amount of Li_3PS_4 impurity is greatly reduced and the crystallinity is significantly enhanced after Cl^- doping in $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$. However, the amount of Cl dopant is limited due to the stable PS_4^{3-} . When y increases to 0.2 in $\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$, not all

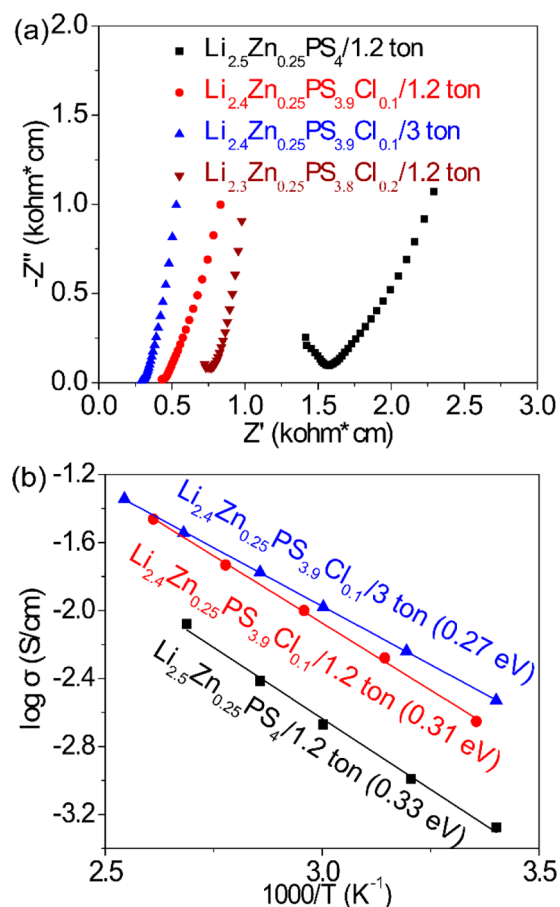


Figure 2. AC impedance spectra (a) and Arrhenius plots (b) of $\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$ ($y = 0, 0.1, 0.2$) samples with varied Cl^- content and prepared under a pressure of 1.2 tons or 3 tons.

Cl^- can be incorporated into $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$, and the remaining LiCl exists as an impurity ($\text{Li}_{2.3}\text{Zn}_{0.25}\text{PS}_{3.8}\text{Cl}_{0.2}$). In addition, with $y > 0.1$, the impurity of Li_2PS_3 forms after sintering, which will further lower the ionic conductivity.

With enhanced phase purity and crystallinity after Cl^- doping ($\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$), the conductivity also exhibited significant improvement (Figure 2). When $y = 0.1$ in $\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$, the conductivity increases to about 2.2 mS/cm at 25 $^\circ\text{C}$ ($\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}/1.2$ ton) and further improves to be 3.5 mS/cm with higher pressure ($\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}/3$ ton). Beyond $y = 0.1$, the ionic conductivity decreases due to the increased formation of impurities with low ionic conductivities including $\text{Li}_4\text{P}_2\text{S}_6$. Besides improvement in the ionic conductivity, activation energy is also reduced with Cl doping. The activation energy of $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$ is about 0.33 eV, which decreases to 0.31 eV in $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ and further decreases to 0.27 eV with higher pressure ($\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}/3$ ton). The enhanced total ionic conductivity and decreased activation energy with relatively high pressure are due to reduced grain boundary resistance for ion transport.^{17,18}

In addition to high ionic conductivity, low electronic conductivity is also important for solid electrolytes, in order to minimize self-discharge, enhance stability, and reduce dendritic lithium formation.¹⁹ The DC polarization curves of $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$ and $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ under an applied voltage of 3 V are shown in Figure 3a, based on which the electronic conductivities of $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$ and $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ are

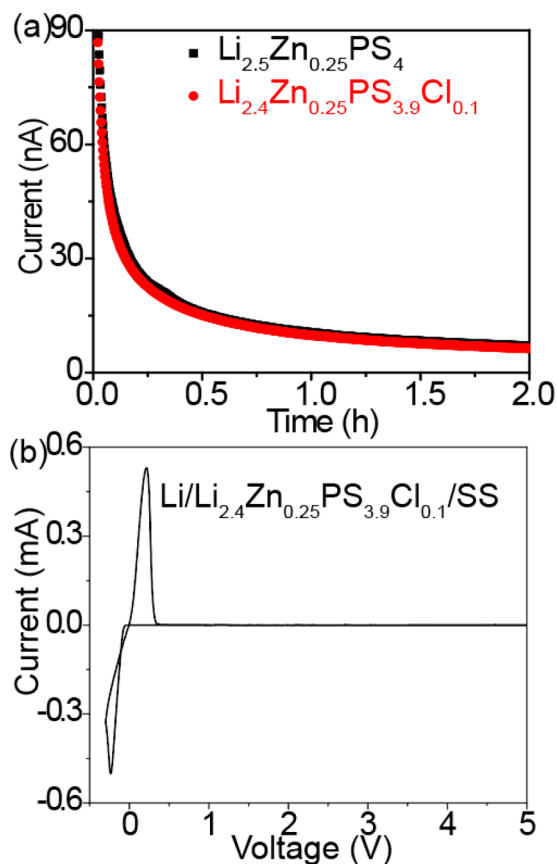


Figure 3. (a) DC polarization curves of $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$ and $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ under 3 V (diameter of In current collector is 4.7 mm, thickness is 1 mm), and (b) cyclic voltammogram for the $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ (SS represents stainless steel).

calculated to be 1.35×10^{-9} S/cm and 1.21×10^{-9} S/cm, respectively, much lower than the electronic conductivity of widely studied solid-state electrolyte material $\text{Li}_7\text{La}_3\text{Zr}_2\text{O}_{12}$.¹⁹

To evaluate the electrochemical stability, the cyclic voltammogram of the $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ is measured and shown in Figure 3b. The open circuit voltage of $\text{Li}/\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}/\text{Stainless steel}$ is ~ 2 V, and the cell is first discharged to -0.3 V vs Li/Li^+ . During reduction, only one process happens at about -0.2 V (vs Li/Li^+), which is the plating of Li on stainless steel. According to previous reports, decomposition of $\text{Li}_{2.25}\text{Zn}_{0.375}\text{PS}_4$ to form Li_2S , Zn_3P_2 , and ZnP_2 happens when the potential is less than 1 V vs Li/Li^+ .⁷ Further reactions leading to the formation of electronically conductive Zn and/or Li–Zn alloy, which allows non-self-limiting SEI formation at the Li–LZPS interface or other parasitic reactions in the bulk electrolyte, compromising the electrochemical stability of LZPS.^{7,16} Lower Zn^{2+} content in $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$ is expected to eventually help reduce the total electronic conductivity of the SEI layer between $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$. Here, no obvious decomposition peak is detected during the discharge process, which suggests improved stability of $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ against Li metal. The improved stability against Li metal is due to the lower electronic conductivity after Cl doping and reduced Zn amount compared to $\text{Li}_{2.25}\text{Zn}_{0.375}\text{PS}_4$. During the charge process, Li stripping occurs at about 0.2 V and no apparent oxidations are observed up to 5 V (vs Li/Li^+), indicating good

electrochemical stability of $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ at high voltages, which makes it compatible with most commercial cathodes.

To probe the local structure and Li^+ mobility, solid-state NMR is employed (Figure 4–7). The ^6Li NMR spectrum of

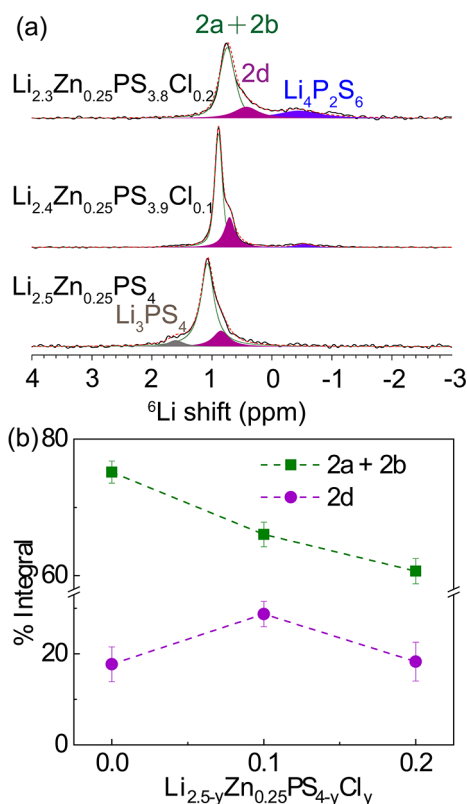


Figure 4. ^6Li NMR spectra of $\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$ ($y = 0, 0.1, 0.2$) samples (a), and the quantification results of different Li sites (b).

$\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$ exhibits two main peaks at around 0.9 and 1.2 ppm, respectively. Based on the structure of $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$, the difference in 2a, 2b, and 2d Li sites is very small and only manifests in the second coordination shell: 2a+2b is surrounded by Li^+ and P^{5+} while 2d is surrounded by Li^+ and Zn^{2+} . The 2a+2b ^6Li NMR resonance is expected to appear at lower field (higher ppm values) based on electron charge distribution compared with the 2d ^6Li resonance. Therefore, the 0.9-ppm ^6Li resonance is assigned to Li at 2d sites while the 1.2-ppm resonance is assigned to Li at 2a and 2b sites. The area integrals of the Li NMR resonances are proportional to the actual Li occupancies at different structural sites, which are used to perform quantitative site occupancy analysis. Detailed site fractions based on spectra analysis are shown in Figure 4b. For an ideal $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$, the fraction of Li^+ at 2d site should be within the range of $[x/(1+2x), 1/(1+2x)]$, i.e., [30%, 40%] for $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$, calculated based on two extreme cases, 2a+2b fully occupied or 2d fully occupied. But according to ^6Li NMR, the Li_{2d} fraction is only around 20%, which means the real composition is not $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$ but $\text{Li}_{2.5-2\delta}\text{Zn}_{0.25+\delta}\text{PS}_4$ with a lower predicted ionic conductivity.⁵ A similar phenomenon has been observed with neutron diffraction.⁶ The Li_{2d} fraction increases to about 30% in $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$, which is close to the theoretical ratio and suggests higher purity. In addition, Li_3PS_4 almost disappears and thus further confirms the minimization of Li_3PS_4 formation with Cl doping. However, a small amount of

$\text{Li}_4\text{P}_2\text{S}_6$ appears, especially in $\text{Li}_{2.3}\text{Zn}_{0.25}\text{PS}_{3.8}\text{Cl}_{0.2}$, which agrees with the XRD results.

Furthermore, variable-temperature ^7Li solid state NMR is used to study the Li^+ mobility and thermal stability of $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ (Figure 5 and 6). Figure 5 summarizes one-

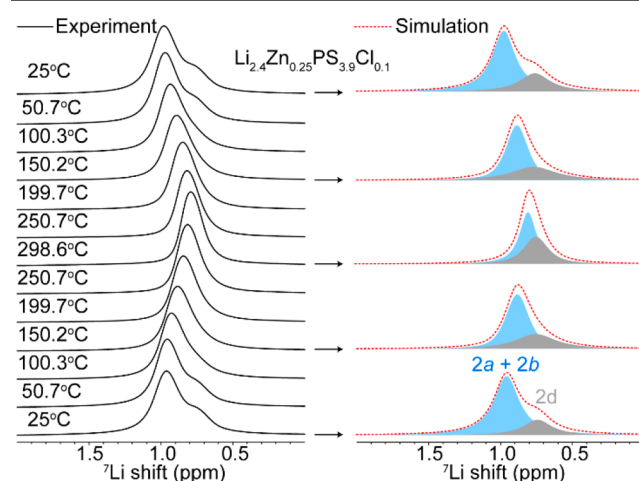


Figure 5. Variable-temperature ^7Li NMR spectra of $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ acquired at a MAS rate of 5 kHz. Selected spectral simulations are shown on the right.

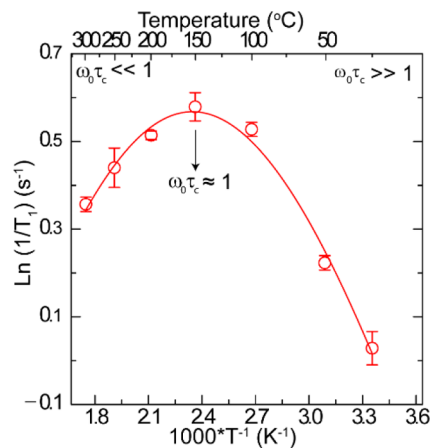


Figure 6. Variable-temperature ^7Li T_1 NMR relaxation time measurements of $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$. The solid line represents the fit using the BPP expression.¹⁸

dimensional ^7Li MAS NMR spectra of $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ acquired within the temperature range from 25 °C to 298.6 °C. Selective spectral analyses are presented in the right panel of Figure 5. At room temperature, the Li fraction at 2d sites (Figure 5, right panel) adds up to 30%, which leaves 70% of lithium to collectively distribute among 2a and 2b sites; this result is in line with ^6Li MAS (25 kHz) NMR (Figure 4). Upon heating $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$, a characteristic feature that signifies chemical exchange between Li^+ at different sites (2a + 2b vs 2d) is identified. The ^7Li signals (2a + 2b and 2d) migrate toward each other with increasing temperature and nearly coalesce as a single peak resonating at the center-of-mass with respect to the two signals (2a + 2b and 2d) at 298.6 °C.

Upon cooling, Li^+ kinetics slows down and the merged ^7Li peaks split into two partially resolved resonances at room

temperature as was identified before heating (Figure 5), suggesting good thermal stability under 300 °C.

Li ion dynamics are further investigated with variable-temperature ^7Li T_1 NMR relaxation time measurements. As shown in Figure 6, the dependence of NMR relaxation rates (i.e., $R_1 = 1/T_1$ in Hz) on temperatures presents a so-called “NMR rate peak”, which is induced by a Li^+ diffusional process.¹⁸ The rate maximum appears at ~ 150 °C, at which the Li^+ jump rate can be estimated with $\omega_0\tau_c \approx 1$. Since ^7Li T_1 measurements are performed with a ^7Li Larmor frequency $\omega_0/2\pi = 194.319$ MHz, τ_c is calculated as 8.2×10^{-10} s at 150 °C. If we assume the jump distance, a , between $2a/2b$ and $2d$ (3.69 Å) remains unvaried at 150 °C, we obtain a diffusion coefficient $= 2.8 \times 10^{-7}$ cm² s⁻¹ through the Einstein–Smoluchowski equation, $D = a^2/(6\tau_c)$ (3D diffusion⁵). This diffusivity extracted from ^7Li T_1 measurements is lower than the computed value on this material ($\sim 6 \times 10^{-6}$ cm² s⁻¹).⁵ In addition, fitting the diffusion-induced ^7Li rate peak (Figure 6) using the BPP model²⁰ yields an activation energy (E_{HT}) of only 0.13 eV (Table S1), which is close to the computational results (0.14 eV) but significantly lower than that of 0.27 eV probed by EIS.⁵ NMR T_1 relaxation rate is typically sensitive to ion dynamics on a time scale that is much shorter than what impedance accesses;²¹ thus, higher activation energy revealed by EIS most likely relates to macroscopic diffusion while NMR T_1 relaxation sheds light on microscopic diffusion and/or localized ion dynamics. As a result, the low activation energy obtained from the high- T flank ($\omega_0\tau_c \ll 1$) of the ^7Li rate peak (Figure 6) is associated with local ion movement. It is worth noting that in order to enable a 3D diffusion in $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$, successful hopping from $2a$ sites to $2b$ sites is necessary and this diffusion process should be treated as a rate-determining step of the overall ion conduction.

Lastly, our BPP fitting (see eqs 1 and 2) results in a β value of 1.76 for the “asymmetric” ^7Li rate peak.

$$R_1(\omega) = C \left[\frac{\tau_c}{1 + (\omega_0\tau_c)^\beta} + \frac{4\tau_c}{1 + (2\omega_0\tau_c)^\beta} \right] \quad (1)$$

$$\tau_c^{-1} = \tau_{c,0}^{-1} \exp(-Ea/k_bT) \quad (2)$$

$R_1 = 1/T_1$ = relaxation rate (^7Li)

C = constant

τ_c = correlation time

$\tau_{c,0}$ = pre-factor

ω_0 = Larmor frequency (^7Li)

β ($1 < \beta \leq 2$) = an exponent to specify the frequency dependence of the rate peak on both the low- T and high- T side.^{20,22}

A β value of 2 is in general found with uncorrelated 3D motion as was originally demonstrated in the case of liquid.²³ However, solid electrolytes with structural disorder often show a β value < 2 ,^{24,25} which suggests a strong correlation effect²⁶ induced by Coulomb interactions.²⁷ Therefore, a β value of 1.76 for $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ indicates a correlated 3D Li^+ motion, in which the Li^+ dynamics are affected by correlation effects originating from structural disorder and the $\text{Li}^+ - \text{Li}^+$, $\text{Li}^+ - \text{S}^{2-}$, and $\text{Li}^+ - \text{Cl}^-$ interactions. Incorporation of halogens into PS_4^{3-} anionic sublattice has been shown to effectively perturb their interactions with positively charged mobile carriers such as Li^+/Na^+ .^{10,28} In $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$, we expect to observe such correlation effects due to the formation of $\text{PS}_3\text{Cl}^{2-}$, and the broken symmetry in PS_4^{3-} anionic sublattice

may indirectly facilitate Li-ion conduction by softening the $\text{Li}-\text{S}$ interactions.²⁹

In addition to $^6, ^7\text{Li}$ NMR, ^{31}P and ^{35}Cl NMR spectra are also acquired and shown in Figure 7. $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$ only has

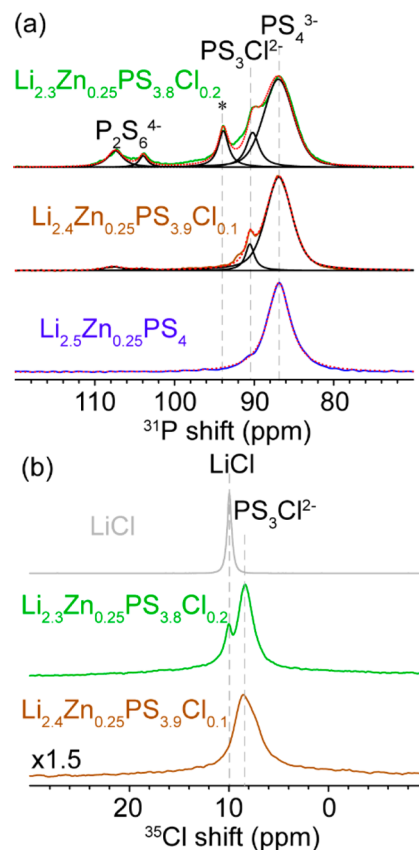


Figure 7. ^{31}P (a) and ^{35}Cl (b) NMR spectra of different $\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$ samples ($y = 0, 0.1, 0.2$); * denotes an unknown impurity. Dotted red lines in (a) are simulated spectra.

one broad peak at around 87 ppm, which is from the PS_4 framework. After doping with Cl, S is partially replaced by Cl and then a fraction of the PS_4 framework changes to PS_3Cl , which shifts the ^{31}P resonance to the left. Another two peaks at 103 ppm and 106 ppm appear with Cl doping, which are from the impurity of Li_4PS_6 .^{10,30} In $\text{Li}_{2.3}\text{Zn}_{0.25}\text{PS}_{3.8}\text{Cl}_{0.2}$, more impurities form, suggested by the extra peak at 93 ppm. For $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$, the amount of Li_4PS_6 is small. The ratio of PS_3Cl is about 10%, confirming 0.1 Cl doped into the $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ successfully. ^{35}Cl NMR spectra of $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ only have one peak, which is from Cl in $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$. The amount of Cl in the nominal $\text{Li}_{2.3}\text{Zn}_{0.25}\text{PS}_{3.8}\text{Cl}_{0.2}$ is calculated to be about 0.14 based on the ^{35}Cl NMR, and the remaining Cl exists as LiCl impurity (Figure 7b). The ^{31}P and ^{35}Cl NMR proves the success of Cl doping and the maximum y value in $\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$ is determined to be 0.14.

CONCLUSION

In summary, we have successfully replaced a small amount of S^{2-} in $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$ with Cl^- , which enhances the ionic conductivity by 4-fold to about 3.5 mS/cm at room temperature. Our investigation reveals three reasons why Cl^- doping enhances the ionic conductivity: (1) improved phase

purity, (2) enhanced crystallinity of $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$, and (3) lowered electrostatic attraction between Li^+ and the anion framework. For Li-rich $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$ with high x ($x > 0.75$), it tends to decompose to Li_3PS_4 at high temperature; therefore, low temperature or Li deficiency helps stabilize $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$. With a small amount of Cl doping in $\text{Li}_{2.5}\text{Zn}_{0.25}\text{PS}_4$, the Li^+ amount decreases, the impurity of Li_3PS_4 is observed to decrease, and the crystallinity is enhanced. ^{31}P NMR and ^{35}Cl NMR prove the successful doping of Cl into the structure of $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$, and the maximum y value in $\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$ is determined to be between 0.1 and 0.2. The electrochemical stability of $\text{Li}_{2.4}\text{Zn}_{0.25}\text{PS}_{3.9}\text{Cl}_{0.1}$ is evaluated with cyclic voltammetry, which shows improved stability compared to $\text{Li}_{2.25}\text{Zn}_{0.375}\text{PS}_4$, due to the reduced content of Zn and decrease in electronic conductivity.

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.chemmater.0c00025>.

XRD patterns and AC impedance of $\text{Li}_{1+2x}\text{Zn}_{1-x}\text{PS}_4$ ($x = 0.5, 0.75, 0.85$), variable-temperature AC impedance of $\text{Li}_{2.5-y}\text{Zn}_{0.25}\text{PS}_{4-y}\text{Cl}_y$ ($y = 0, 0.1$).

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<https://pubs.acs.org/doi/10.1021/acs.chemmater.0c00025>

Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

The authors acknowledge the support from National Science Foundation (DMR-1847038). All solid-state NMR experiments were performed at the National High Magnetic Field Laboratory, which is supported by National Science Foundation Cooperative Agreement No. DMR-1644779 and the State of Florida.

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