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Tuning mobility and stability of lithium ion conductors based on lattice dynamics†

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Lithium ion conductivity in many structural families can be tuned by many orders of magnitude, with some rivaling that of liquid electrolytes at room temperature. Unfortunately, fast lithium conductors exhibit poor stability against lithium battery electrodes. In this article, we report a fundamentally new approach to alter ion mobility and stability against oxidation of lithium ion conductors using lattice dynamics. By combining inelastic neutron scattering measurements with density functional theory, fast lithium conductors were shown to have low lithium vibration frequency or low center of lithium phonon density of states. On the other hand, lowering anion phonon densities of states reduces the stability against electrochemical oxidation. Olivines with low lithium band centers but high anion band centers are promising lithium ion conductors with high ion conductivity and stability. Such findings highlight new strategies in controlling lattice dynamics to discover new lithium ion conductors with enhanced conductivity and stability.

Introduction
Replacing organic liquid electrolytes1,2 with solid lithium ion conductors in lithium ion batteries enables the use of the metallic lithium to markedly boost the energy density3–5 and also increases battery safety.5 Current research and development of solid-state lithium ion batteries has been catalyzed by recent breakthroughs in solid lithium ion conductors that have ion conductivities6–9 rivaling that of conventional organic liquid electrolytes.1 Although a number of structural families such as lithium superionic conductors (LISICON),10 garnets,11 NASICON-like (sodium superionic conductor),12 and perovskites14 have ion conductivities approaching that of liquid electrolytes, an increasing number of experimental15 and computational16,17...
studies show that fast solid lithium ion conductors are not stable against lithium ion battery electrodes. Of significance, no known fast solid lithium conductors is stable against lithium ion positive electrode materials,16 highlighting critical needs to search for new solid lithium conductors with high lithium ion conductivity and stability.

Increasing ion mobility and stability of lithium solid conductors is not straightforward and progress in the past decades has been achieved primarily by trial and error. Structural and chemical tuning via isovalent or aliovalent substitution of cation10,11 and/or anion18–20 in given structural families has led to steady increase in the lithium ion conductivity, and recent discovery of superionic lithium ion conductors.6–9 Recently a number of structure-based ion conductivity descriptors have been proposed to aid and accelerate the design of new superionic conductors including the volume of the unit cell10,13,21 the volume of lithium diffusion pathway,22 the anion in the structure,21 and the structure of the anion sublattices.23 For example, increasing lattice volume in the LISICON,10 NASICON-like,13 or perovskite14,21 structure can enhance ion conductivity and reduce activation energy.21 In addition, changing the anions by moving down in the periodic table (e.g. Li₁₀Ge₃P₂X₁₂, (X = O, S and Se)27 or Li₃PX₄ (X = O²⁻ and S²⁻)) and/or arranging anions in a body centered cubic lattice25 can increase lithium ion conductivity by reducing activation energy. Unfortunately, most of these descriptors have limited predictive power when applied across different structure families and chemistry of lithium ion conductors.21 Direct computation of Li-ion conductivity using ab initio molecular dynamics (AIMD) has also been pursued as a way to discover/design new materials with higher ion conductivity.25,26 However, this method is computationally very demanding not to mention various practical limitations such as the need to extrapolate ionic conductivity from high temperature to room temperature. Moreover, no design principle is established for the stability of lithium solid conductors. Limited predictive power of reported conductivity descriptors and the lack of stability descriptor hampers the search for new lithium-ion conductors with enhanced conductivity and stability in order to meet all the requirements of solid-state lithium-ion batteries. In this article, we report a new approach to design lithium ion mobility and stability of lithium ion conductors using their lattice dynamical properties. We relate lithium ion mobility to lattice dynamics based on the following hypothesis: small energetic barrier for lithium ion migration (enthalpy of migration) is associated with large displacement amplitude from their equilibrium sites (Fig. 1a), rendering greater probability for lithium ions to explore multiple energy minima. Large excursions of mobile ions away from the equilibrium position are enabled in the soft atomic potential and also are associated with low phonon frequencies considering the Einstein model, where the amplitude of thermal displacement $u$ of the ion is inversely proportional to the square of the frequency $\omega_e$ using:27 $\langle |u|^2 \rangle = 3k_BT/m\omega_e^2$, having $k_B$, $T$ and $m$ denote the Boltzmann constant, temperature and the mass of the mobile species, respectively. In contrast, with a high migration barrier, the mobile species oscillate close to their equilibrium position with small thermal displacement amplitude and high frequency, and have low probability of hopping to adjacent sites. The hypothesis is supported by the following observations. The self-diffusion coefficient and the activation energy of metal atoms in body center cubic metal strongly correlate with the frequency of the longitudinal acoustic mode along the (111) direction (at reduced wavevector $q = 2/3$), being higher with lower frequency.28 Second, the computed migration enthalpy of oxygen diffusion via interstitial in rare-earth Ruddlesden–Popper phases shows strong positive correlation with reduced (more negative) force constant of the soft mode associated with the rotation of AO₆ octahedra (A = rare-earth metal ions).29 Third, Wakamura has shown that the activation...
energy in Ag⁺, Na⁺Cu⁺ and F⁻ superionic conductors decreases with decreasing frequency of low-energy optical mode⁴⁰ supporting the idea that low phonon frequency can be associated with high probability of ions hopping to the neighboring sites.¹¹

We elected to study a series of lithium ion conductors in the LISICON family derived from Li₃PO₄, to test the hypothesis that their lattice dynamics can greatly influence lithium ion mobility and stability. Lithium ion conductivity can be increased by up to 15 orders of magnitude¹⁰,¹¹,¹₈,²₀,²¹ via cation and anion substitutions in this structural family relative to Li₃PO₄ (Fig. 1d). We also included Olivine structure which has the same (distorted) hexagonal close packed of anion sublattice as the LISICON compounds but differ in occupancy of Lithium ions which are located in octahedral sites as opposed to tetrahedral sites in LISICON structure. Although substitution (doping) and defects may alter migration pathway and mechanism and the activation energy, we have selected lithium ion conduction in LISICON and Olivine, having similar one-dimensional channels of Li ions within hexagonal anion sublattice. Previous work has shown that in Olivine, these one-dimensional channels serve as the dominant pathway for lithium migration and govern ion conduction in this structure.⁵⁰ In Lisicon, although the diffusion pathway is three-dimensional,¹¹ previous study has shown that the enthalpies of vacancies migration along each crystallographic direction are very similar and on the order of 0.7 eV in both γ-LiₓPO₄ and β-LiₓPO₄.²² It should be mentioned that depending on the concentration of mobile specie, the dominant diffusion mechanism might be interstitial instead of vacancy diffusion considered here. Nevertheless, we believe that the trends that we propose here remain valid as long as one considers the same mechanism with similar diffusion pathway regardless of the compound chemistries and structures. Isowalent substitution of oxygen by sulfur anion can enhance the ionic conductivity by six orders of magnitude, having reduced activation energy from 1.3²² to 0.52 eV,²⁰ and additional four orders of magnitude by aliovalent substitution of phosphorus by germanium, with decreasing activation energy from 0.52 to 0.21 eV.¹⁸ In this work, we systematically study the lattice dynamics of LISICONs derived from LiₓPO₄ by measuring the phonon density of states (DOS) of 17 compounds and one olivine compound as well as computing the phonon DOS of more than 20 compounds which are isostructural to LISICON and 6 olivines. The reader is referred to the Table S1 in (ESI†) for a complete list of compounds included in this study. From these data, combined with the measured activation energy, computed enthalpy of lithium ion migration of LISICONs and computed potential for electrochemical oxidation, the descriptors for lithium ion mobility and stability for oxidation were proposed.

### Experimental and computation methods

All phonon DOS measurements were made at the Wide-angular range chopper spectrometer (ARCS) at the Spallation Neutron Source at Oak Ridge National Labs. Approximately 5 g of the samples were packed into aluminum canisters, which were sealed with a vanadium ring, unless spectra at elevated temperatures were measured where a qualitative (aluminum on aluminum) seal was used. All samples were sealed under argon. Temperatures were varied from 10 K to 600 K for select samples. Unless otherwise noted, the samples were measured at 100 K. The incident neutron energies were varied between 60, 100, and 200 meV. The resulting data were analyzed using the Mantid DGS Reduction package. For oxides, the background fraction was set as 1, the fraction of the total scattering to subtract as a constant background was 0.0–0.25 unless otherwise noted, the cutoff for the elastic peak removal was set as 10 meV, 3 bins after the cutoff were averaged to get the value of the density of states near the cutoff, the estimate of the Debye cutoff and the limit on the energy range were set depending on the calculated phonon density of states and where the last feature in the experimental density of states was noted (these parameters were used to remove any high energy background), and the range in wave vector that was used was from 7–12 Å⁻¹. For sulfides, the parameters were the same except the range of wave vectors used was from 4–9 Å⁻¹.

For computations, we used density functional theory based on the Perdew–Burke–Ernzerhof (PBE) generalized gradient approximation to the Projector Augmented Wave (PAW) method.³⁵ Migration barriers for a lithium ion hopping were calculated using the climbing-image nudged elastic band method in a 2 × 2 × 1 supercell for LISICON (space group Pnma) while for the LISICON (space group Pmna), a 2 × 2 × 2 supercell was employed. A 2 × 2 × 2 k-point grid was used and the cutoff of the kinetic energy was set to the default values as set in the pseudopotential files. For phonon calculations, the same supercells were used with finer k-point grid (3 × 3 × 3) and a higher energy cutoff (520 eV) in order to obtain more accurate values of the force. We have also used a higher cut-off of 700 eV for phonon DOS calculations but haven’t found any significant change (Fig. S1, ESI†) indicating that phonon DOS calculations are already well converged at 520 eV of cut-off energy. We have also tested LDA functional for phonon DOS calculations and found that the main effect of is to shift all the modes especially the high-energy feature in the DOS to higher frequencies resulting in an upward shift of band center (Fig. S1, ESI†) in agreement with previous study.³⁷ However, the magnitude of the shift is small and fairly constant across different chemistries and crystal structures suggesting that LDA functional will result in a rigid shift to slightly higher energy but will not affect the trend. Finite displacement method was used for phonon calculations and the total as well as the atom-projected DOS were extracted using the phonopy package.³⁸

To quantify the average vibrational frequency of a given material, we defined the ‘phonon band center’ which is the phonon frequency weighted by the DOS. Mathematically, it is written as:

$$
\omega_{\text{av}} = \frac{\int \omega \times \text{DOS} (\omega) d\omega}{\int \text{DOS} (\omega) d\omega}
$$

If we replace DOS(\omega) in this expression by the total phonon DOS, we obtain what we called ‘total phonon band center’. Similarly, if we replace DOS(\omega) by one of the atom-projected
DOS for instance Li-projected DOS, we obtain the ‘Lithium phonon band center’ which can also be viewed as the centroid of the Li-projected phonon DOS. The stability windows were computed following the method proposed by Richard et al.16 using the data from Materials project database39 and Pymatgen software package.40

The stability window was computed by constructing the grand potential phase diagram and varying the chemical potential of Li until the grand potential of the electrolytes were above the convex hull.41 Measured phonon DOS of 17 LISICON and LISICON-like compounds can be found in Fig. 2–4 and Fig. S2 (ESI†). The measured phonon DOS at 0 K (the lower spectrum) is in quantitative agreement with the DOS computed from DFT at 0 K (the upper spectrum of each panel in Fig. 2) is in quantitative agreement with the DOS computed from DFT at 0 K (the upper spectrum of each panel in Fig. 2). The measured phonon DOS data were obtained after background correction, obtaining lower and higher energy cut-off and details can be found in Fig. S4 (ESI†). The computed DOS were weighted by neutron scattering cross section of the different elements that make up the structures. The details of this procedure can be found in previous work.67

**Results and discussions**

**Influence of chemical substitution and temperature on phonon DOS of lithium conductors**

Cation and anion substitutions in the Li3PO4 structural family were shown to induce systematic changes by examining measured phonon DOS of 17 LISICON-like compounds and computed phonon DOS of 20 LISICONs. The measured and computed phonon DOS of Li3PO4 (Pnma), Li3PS4 (Pnma2), Li3Ge0.4P0.6O4 (Pnma) and Li1.25Ge0.25P0.75S4 (Pnma) are shown as example in Fig. 2a–d, and those of the other LISICON-like compounds can be found in Fig. 4 and Fig. S2, S3 of the (ESI†). The phonon DOS measured at 100 K (the upper spectrum of each panel in Fig. 2) is in quantitative agreement with the DOS computed from DFT at 0 K (the lower spectrum). Substitution of oxygen by sulfur led to significant softening of phonon modes. Li3PO4 (Pnma) in Fig. 2a and Li1.4Ge0.4P0.6O4 (Pnma) in Fig. 2c were found to have higher energy phonon DOS than Li3PS4 (Pnma2) in Fig. 2b and Li1.25Ge0.25P0.75S4 (Pnma) in Fig. 2d. Similar lattice softening were found for other sulfides relative to oxides in Fig. S2 and S3 (ESI†). Contribution from lithium ion vibrations (lithium-projected phonon DOS) in the measured total phonon DOS was identified by comparison with the computed atom-projected DOS, as shown in Fig. 2 and Fig. S3 (ESI†). Generally speaking, the computed phonon DOS of LISICONs show that low-energy, mid-energy and high-energy features originate mainly from the vibrations of anion sublattice, lithium sublattice and anion units involving non-mobile, structural cations (Fig. S3, ESi†) such as PO4, respectively. Of significance, Li3PO4 (Pnma) was found to have phonon DOS peaks in the range from 40–70 meV, which primarily came from lithium ion vibrations, as shown in Fig. 2a and Fig. S3 (ESI†). Substituting oxygen in Li3PO4 (Pnma) by sulfur to form Li3PS4 (Pnma and Pnma2) led to a down-shifting (softening) of lithium ion vibrations from 40–70 meV to 40–50 meV (Fig. 2b and Fig. S2, S3, ESI†).
observed for Li3PS4 upon increasing temperature from 100 K to supported by marked broadening of lithium phonon modes involving non-mobile, structural cations, to the of lithium band center but large downshifting of high-energy phonons with increasing temperature. Therefore, the broadening of phonon DOS for LISICONs was found to correlate with decreasing measured activation energy for lithium ion conductivity. Increasing temperature has negligible influence on this phonon band center, which is supported by temperature-dependent phonon DOS in Fig. 4b and c, showing no softening of phonon frequencies with increasing temperature. Downshifting the total phonon band center was shown to markedly lower measured activation energy of stoichiometric LISICONs, as shown in Fig. 5a. Replacing oxygen with sulfur from Li10PO4 (Pnm21) to Li10PS4 (Pnm21) led to the largest downshift of the total phonon band center from ~70 to ~40 mV, which was correlated with reduction of activation energy from 1.4 eV24 to 0.5 eV20 in Fig. 5a. Measured activation energy of Li10PO4 (Pnm1), Li10VO4 (Pnm21), Li10Ge0.4P0.6O4 (Pnm21) and Li10SnS0.5O4 (x = 0, 0.2, 0.4, 0.6 and 1) obtained in this study are in agreement with previous EIS measurements.10,24,44,45 As the measured activation energy of stoichiometric LISICONs consists of both enthalpy of defect formation and enthalpy of migration, such a correlation for stoichiometric LISICONs does not explicitly imply lowered lithium migration barrier with softening of the total phonon band center. We further show that the measured activation energy for 8 substituted LISICONs with partial lithium occupancy, which corresponds largely to lithium ion migration enthalpy, was decreased with softening of the total phonon band center. Li10Ge0.4P0.6O4 (P62/nmc) and Li10SnS0.5O4 (chemically similar to Li10GeP2S12, Li10SnP2S12 (P62/nmc)) were found to have the lowest activation energy of ~0.25 eV comparable to best superionic conductors5–9 with the lowest measured total band centers of ~40 mV. Another interesting observation that we can draw from Fig. 5a is the clear separation between stoichiometric and non-stoichiometric compounds which follows two distinct trend lines whose difference is essentially related to the enthalpy of defect formation. It is well-known in perovskites that the enthalpy of migration of oxygen ions decreases with decreasing enthalpy of vacancy formation.48 Although, in this study, we do not consider the energetic of lithium defect formation energy, in light of the data shown in Fig. 5a, one might hypothesize that the enthalpy of Lithium defect formation might also correlate with phonon band center. As the total phonon band center can be weighted considerably for the vibrations of non-mobile species such as structural cations and anions, we further sought correlations between the band center of lithium-projected phonon DOS and activation energy.

Correlation between Li-band center and enthalpy of migration

Aliovalent cation substitution induces broadening of features in the phonon DOS, without significant softening. Phonon peaks, including the peak that primarily came from lithium ion vibrations in the range of 40–70 meV, was broadened systematically with increasing Ge substitution in Li1-xGe0.65V0.35O4, where x = 0, 0.2, 0.4, 0.8 and 1 (Li10V0.4O4, LGV020, LGV040 and LGV080 for x = 0.2, 0.4 and 0.8 respectively) measured at 100 K. (b) The phonon DOS of Li10PS4 (Pnma) measured at 100, 200 and 300 K, which shows marked broadening at ~50 meV. (c) The phonon DOS of Li3PO4 (Pnma) measured at 100, 200 and 300 K, which shows little variation with temperature. (d) The phonon DOS of Li32SnP4S12 that are featureless measured at 100, 200 and 300 K. The measured phonon DOS data were obtained after background correction, optimizing lower and higher energy cut-off and details can be found in Fig. S4 (ESI†).
In addition, we computed the enthalpy of migration, which would allow us to systematically examine both stoichiometric LISICONs without partial occupancy and substituted LISICONs with partial occupancy. The downshifting of lithium phonon band center or average lithium ion vibration frequency was found to correlate with reduced migration barrier and thus greater lithium mobility, supporting the hypothesis described in Fig. 1a. Lithium ion migration enthalpy, which was defined as the difference between transition state energy and that of the initial/final configuration, was computed using nudge elastic band (NEB) calculations. Identical jump sequence along the diffusion pathway shown in Fig. 5b inset was used to compute the enthalpy of lithium ion migration even though this jump sequence might not be associated with the lowest migration enthalpy. In addition to 7 stoichiometric LISICON and one substituted LISICON in Fig. 3, we included 8 others stoichiometric LISICONs in the inorganic crystal structure database (ICSD) in Fig. 5b, which represent all stoichiometric LISICONs (excluding those containing transition metals) and two computed structures (Li$_3$CdSiS$_4$ and Li$_3$CdGeSe$_4$) to complete the series of Li$_x$CdX$_4$ ($X = Si$, Ge and Sn) and Li$_x$CdGeY$_4$ ($Y = O$, S and Se). Decreasing computed lithium band center of 17 stoichiometric structures (Li$_2$CdSiS$_4$ and Li$_2$CdGeSe$_4$) to complete these series of LISICONs and one substituted LISICON Li$_{1.25}$Ge$_0.25$P$_{0.75}$S$_4$ (chemically similar to Li$_{10}$GeP$_2$S$_{12}$) was found to have the lowest measured (Fig. 5a) and computed enthalpy (Fig. 5b) of lithium ion migration of ~0.2 eV, which is comparable to that of conventional liquid electrolytes used in lithium ion batteries$^1$ and fast lithium ion conductors such as Li$_{10}$P$_3$S$_{11}$,$^9$ Li$_{1.25}$Ge$_{0.25}$P$_{0.75}$S$_4$ (chemically similar to Li$_{10}$GeP$_2$S$_{12}$) were found to correlate well with the computed lithium band center at 0 K.

**Correlation between anion band center and oxidation potential**

Lowering the anion phonon band center was found to greatly reduce the stability of these LISICONs against electrochemical oxidation, as shown in Fig. 6. The oxidation potential was defined as the potential above which LISICONs will be oxidized electrochemically by lithium ion removal from the lattice, which was computed using a thermodynamic approach as reported by Richards et al.$^{16}$ Replacing oxygen with sulfur in the LISICON structure had the largest downshift of the anion phonon band center and largest reduction in the oxidation potential. While oxidative stability can be understood in term of electronic structures of the electrolytes which correlate with the thermodynamic driving force to oxidize the materials,$^{44}$ the correlation between the oxidation potential and the anion band center
shown in Fig. 6 might be kinetics in origin as lowering anion band center will make anion sublattice more mobile, thus lowering the kinetic barrier for the (decomposition) reactions. Moreover, the stability of a compound can be governed by kinetic properties (migration barriers) and thermodynamics properties. Well-known principles of understanding controlling kinetics based on thermodynamics include the Evans–Polanyi principle or linear free energy relation, which relates the activation energy of chemical reactions (kinetics property) to the enthalpy of reactions (thermodynamics property). We believe that the correlation between the oxidation potential and anion band center is another instance of this type of ‘kinetic–thermo-dynamic’ relationships, which is thought in the following ways: as Li-band centers correlate with the enthalpy of lithium migration we argue that lowering the anion-band centers correlate with smaller enthalpy of anion migration (kinetic property), which promotes the kinetics of reactions. This kinetics property, by the Evans–Polanyi principle, correlates with oxidative stability (thermodynamic property) as measured by the oxidation potential. We recognize that more studies are needed to substantiate this idea. Nevertheless, we believe that this work will spur interests from the community to further explore this concept and its potential implications. Softening of the anion band center from Li$_3$PO$_4$ ($Pmm2_1$) to Li$_3$PS$_4$ ($Pmna$) by ~30 meV is accompanied with reduction of the oxidation potential from ~4 to ~2.5 V vs. lithium. The computed oxidation potential of Li$_3$PO$_4$ ($Pmm2_1$) and Li$_3$PS$_4$ ($Pmna$) are in agreement with previous computation.$^{16,17}$ The computed oxidation potential of Li$_{3.25}$Ge$_{0.25}$P$_{0.75}$S$_4$ is 0.21 V vs. lithium, which is close to the measured oxidation potential of chemically similar Li$_{10}$Ge$_{3/2}$P$_{11/2}$.$^{48}$ In addition, the oxidation potential of LISICONs was found to correlate strongly with computed anion band center (goodness of fits $R^2 = 0.97$) and to a lesser extent with lithium band center (goodness of fits $R^2 = 0.92$, Fig. S8a, ESIf) and total band center (goodness of fits $R^2 = 0.93$, Fig. S8b, ESIf). There was no correlation between the reduction potential and the phonon band centers (Fig. S8c–e, ESIf), whose physical origin requires further studies. We also seek correlation between oxidation potential and oxygen/sulfur p-band center which have been used to understand the stability of perovskites under oxygen evolution reaction condition.$^{49}$ However, as can be seen from Fig. S9a and b (ESIf), there is no clear correlation between p-band center and the oxidation potential of the anion band center although we can note that the oxidation potential is largely determined by the anion chemistry (oxides vs. sulfides) as reported previously in the literature$^{44}$ and that oxides tend to have lower anion p-band center than sulfides as expected. It should be noted that the absence of correlation might also be due, at least partly, to the uncertainty in the computed p-band center arising from the underestimation of DFT band gap in these insulating compounds.$^{33}$ Therefore, the anion band center is proposed as one descriptor for electro-chemical oxidation stability of lithium ion conductors.

Of significance, fast lithium ion conductors$^{18}$ based on LISICONs such as Li$_{3.25}$Ge$_{0.25}$P$_{0.75}$S$_4$ have low enthalpy of lithium ion migration and softened lithium phonon DOS, which is accompanied with downshifted anion band center (Fig. S8f, ESIf). The correlation between lowered migration barrier with softened average lithium vibration frequency in Fig. 5b, and between lowered oxidative stability with softened average anion vibration frequency in Fig. 6 highlight a trade-off between lithium ion mobility and oxidation stability for the design of lithium ion conductors. Moreover, extending the concept in Fig. 1a to anion mobility, lowered anion band centers can be accompanied by increased anion mobility, which can promote any solid-state reaction kinetics with electrode materials. Therefore, the interplay between lattice dynamics and ion mobility and stability highlights the need and opportunities to search for fast lithium ion conductors having low lithium band center but high anion band center which exhibit high ion conductivity and high oxidative stability in lithium ion batteries.

**Overcoming the trade-off between mobility and stability**

In search of lithium ion conductors with such lattice dynamics characteristics, we further examine Olivines that have the same hexagonal anion sublattice as the LISICON structure (Fig. 7c). The computed enthalpy of lithium ion migration and lithium band center for all the stoichiometric Olivines without transition metal ions listed in the ICSD (except LiCdPO$_4$, LiScGeO$_4$ and LiTmSiO$_4$) are included in Fig. 7a, which demonstrates lithium band center as descriptor for lithium ion mobility in the two structural families (LISICONs and Olivines). Olivines, exhibit low lithium band centers and migration barriers comparable to that of LISICON thiophosphates (Fig. 7a) but high anion band centers comparable to those of LISICON phosphates (Fig. 7b).
**Conclusion**

In this study, we account for the first time, the trend in ionic mobility and electrochemical oxidation stability of lithium ion conductors from one common physical origin, lattice dynamics, or more precisely, atomic vibrational frequencies of ion conductor constituents, thus highlighting the critical role played by the lattice dynamics in governing the lithium ion conductivity and stability of lithium-ion conductors. Unlike previous studies on lattice dynamics in ion conductors (especially Ag⁺ conductors) which focus on the total phonon DOS (such low-energy peak in x-AgI phonon DOS) or zone-center phonons as measured from infrared and Raman spectroscopy, our study clarifies the influence of lattice dynamics on ionic conductivity by separating the effect of different sublattice and in particular the special role of the mobile species sublattice dynamics in the observed ionic mobility. The phonon DOS have been collected from a series of lithium conductors in the LISICON family using INS and computed by first-principles simulations to identify specific spectral features corresponding to the vibrations of sublattices involving lithium, anion and nonmobile structural cations. Replacing oxygen with sulfur greatly downshifts the lithium vibrational features in the phonon DOS. The phonon DOS of Ag⁺ conductors in the LISICON family were found to be lower than in LISICON. All the computed enthalpies of migration were calculated using the standard climbing image nudge elastic band method and the phonon band center is defined as the average phonon frequency weighted by phonon DOS. The blue and orange colours refer respectively to the HT phase (space group Pnma) and the LT phase (space group Pmmn), respectively while the red colour refers to the Olivine compounds. The filled circles are compounds that are known in the ICSD and/or computed in this work, where the lithium ion conductivity has not been measured experimentally. For more details, please refer to Table S1 (ESI†).

Computed migration barriers of Olivines studied here such as LiMgPO₄ are comparable to those of other Olivine materials such as LiMPO₄ (M = Mn, Fe, Co and Ni, 0.1–0.4 eV) [50] and LiMXO₄ (main group M²⁺–X⁵⁺, M²⁺–X⁵⁺; M = Mg, Ca, Sr, Ba, Sc, Y, Al, In, Ga and rare-earth elements; X = Si, Ge, Sn, P, As and Sb). In addition, the computed migration barrier (~0.3 eV) of LiMgPO₄ is in good agreement with NMR measurements for the site-to-site hopping (0.3–0.5 eV). Generally speaking, while compounds having high Li mobility tend to have low stability, Olivine family was found to be an exception to this trend, which was rationalized in terms of the low Li-band center but high anion band center in these compounds (Fig. 7b). For example, new compositions such as LiMgAsO₄ and LiInGeO₄ could potentially have high ion conductivity (Fig. 7a) and stability (Fig. S10, ESI†). Unfortunately, the one-dimensional nature of diffusion pathway in the olivine structure makes it prone to anti-site defects which can result in an apparent activation energy much higher than the intrinsic migration barrier. For example, the long-range diffusion barrier measured by EIS for LiMgPO₄ is 0.7–1.0 eV in previous work and also in this study, much higher than the intrinsic migration barrier (~0.3 eV). Further studies are needed to increase the long-range ion conductivity of these Olivines by reducing anti-site defects using smaller particle sizes and/or having Li-excess in the lattice, which would potentially lead to the development of lithium superionic conductors with high stability against electrochemical oxidation.
determined in this study do not explicitly take into account the specific vibration pattern of each mode. In addition, future experimental and computational studies should exploit the interplay between lattice dynamic and ion conductivity and stability for lithium ion conductors of different structural families\(^{21}\) beyond LISICON and Olivine. Moreover, further studies are needed to extend and examine the universality of such descriptors based on lattice dynamics to understand and potentially control ion mobility and stability of other ionic conductors such as Na\(^+\), Cu\(^+\) or O\(^2-\) conductors. For example, copper sulfide (Cu\(_2\)Se) with superionic Cu\(^+\) conductivity at room temperature is known to have low-energy phonon DOS and lattice thermal conductivity.\(^{59}\) Lattice dynamics was also found to play important role in the Na\(^+\) conductors Na\(_3\)PS\(_4\)\(_x\)Se\(_x\)\(^{60}\) as well as O\(^2-\) conductors Nd\(_2\)NiO\(_4\)\(_x\)\(^{61}\) and brownmillerites Sr(Fe,Co)O\(_2\).\(^{62}\)

Lastly, although extensive database for material phonon DOS is not yet available due to high computational cost to compute phonon DOS, recent study has shown that it is possible to use a machine learning model to predict phonon-related properties such as vibrational free energy and entropy with high accuracy at a greatly reduced computational cost.\(^{63}\) Future advance in machine-learning computational material design represents an exciting opportunity to use this approach to explore more compositional space and to discover new ion conductors.\(^{64-66}\)

**Author contributions**

S. M., Y. S.-H. and J. C. B. proposed the concept. J. C. B. and H.-H. C. prepared the oxide materials. R. K. provided the sulphides samples. S. M., J. C. B., D. B. and O. D. performed the phonon DOS measurements and analysis. S. M. performed all the DFT calculations with L. G.’s help for the NEB calculations. S. M. and Y. S.-H. wrote the manuscript. All authors contributed to the discussion and revision of the manuscript.

**Conflicts of interest**

There are no conflicts to declare.

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**References**
