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Sodium Site Exchange and Migration in Polar Stuffed Cristobalite Framework Structure

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Table of Contents. Two-dimensional exchange ²³Na NMR spectroscopy evidences sodium site exchange phenomenon in polar stuffed Cristobalite *o*-Na₂MgSiO₄ which triggers the long-range sodium migration.

ABSTRACT:

The study of ionic dynamics in solids is essential to understand and develop modern energy technologies. Here we study the ionic dynamics of orthorhombic Na₂MgSiO₄, an interesting case of a polar stuffed cristobalite-type structure that contains two inequivalent Na sites within the channels of the magnesium-silicate tetrahedral framework. Its preparation by solid state reaction method favours the presence of ~ 2% of Na vacancies, converting it into pure sodium ionic conductor with an optimized ionic conductivity of ~10⁻⁵ S cm⁻¹ at 200 °C. The macroscopic migration has been characterized through impedance spectroscopy and molecular dynamics simulation which proves the pure sodium ionic character of the compound through hopping between Na1 and Na2 sites, forming three-dimensional migration zig-zag shaped paths. High-resolution solid–state ²³Na magic MAS–NMR spectroscopy is employed to characterize the local structure and microscopic dynamics of sodium ion transport in Na₂MgSiO₄. Remarkably, variable temperature ²³Na MAS NMR and two-dimensional exchange spectroscopy evidence for the first time a sodium site exchange phenomenon at room temperature, which further triggers the sodium ionic conduction at elevated temperatures.

1 Introduction

Ionic migration in solid state materials is a key physical phenomenon in numerous technologically important compounds such as solid state electrolytes used in batteries,¹ electrochromic displays,² sensors³ or solid oxide fuel cells (SOFCs).⁴ It is important to understand the interplay between ionic migration and structural chemistry, which helps improving the ionic conduction and guiding the materials design. Electrochemical impedance measurements (EIS) are frequently used to characterize the macroscopic long-range diffusional migration coupled to molecular dynamic (MD) simulations.^{5, 6} Nevertheless, detailed knowledge of the microscopic dynamic processes (set by the distance between neighboring atoms sites, i.e. at atomic and/or molecular scale) underlying ionic conductance is still required in order to fully understand and predict the behavior of ionic conducting materials.

Among the different techniques characterizing solid state inorganic materials at local scale, solid state magic angle spinning nuclear magnetic resonance (MAS NMR) is a powerful tool. It has been widely employed to explore the local environment of certain atoms allowing, for instance, the concrete crystallographic description of unit cells, local structure around defects and so on. The application of this technique has been further enhanced by the possibility to acquire MAS NMR spectra at variable temperatures (VT MAS NMR),⁷ which expands the possibility to study the dynamic processes of mobile species present in solid state ionic conductors as a function of temperature.⁸ Moreover, for materials possessing multiple sites for mobile ions, it enables distinguishing ions moving on inequivalent sites that can lead to multiple possible pathways for diffusion. For this kind of materials, the clearest sign of diffusive dynamics is the merging of spectral peaks, initially assigned to atoms in different structural environments, with increasing temperature.⁹ The information on the ionic dynamics by NMR spectroscopy can be

further extended using two-dimensional exchange NMR spectroscopy (2D EXSY)¹⁰ that allows detecting exchange processes between ions on multiple sites during the initial migration stages. These techniques to study the ionic dynamics in solids have been frequently employed for characterizing the dynamics in lithium ion conductors.^{11, 12} Nevertheless, studies about the dynamics in Na ion conductors still remain scarcely explored even when they are potential materials as solid state electrolytes for batteries.

In this study, we investigate the sodium dynamics in stuffed cristobalite Na₂MgSiO₄ using EIS and MD simulations coupled to NMR spectroscopy. This compound exhibits a tetrahedral framework based on corner-linked $[SiO_4]^{4-}$ and $[MgO_4]^{6-}$ units with two inequivalent sodium sites stuffing the channels formed in the tetrahedral network. It was initially identified as potential sodium ion conductor due to its framework structure, where three-dimensional (3D) conductivity might be expected.^{13, 14} Two different polymorphs have been reported at room temperature: firstly Baur synthesized it hydrothermally and described its structure in the nonpolar *Pn* space group (*m*-Na₂MgSiO₄),¹³ then Withers et al.¹⁵ prepared it following a sol-gel route and described its structure by using the polar *Pna*2₁ space group (*o*-Na₂MgSiO₄). Both orthorhombic and monoclinic structures contain an ordered Si/Mg distribution but differ in the pattern of tetrahedral tilting (Figure 1). Herein we obtain the polar *o*-Na₂MgSiO₄ phase and demonstrate remarkable sodium site exchange at room temperature between the two inequivalent Na sites, the subsequently triggered sodium migration and its mechanism at elevated temperatures in this polar stuffed cristobalite.



Figure 1. Tetrahedral framework structure comparison between m-Na₂MgSiO₄ (a) and o-Na₂MgSiO₄ (b) as described by Baur (ICSD 15618) and Withers et al. (ICSD 84562), respectively. Shaded blue and grey tetrahedra contain Si and Mg, respectively. The O atoms are illustrated by red balls atoms.

2 Methods

2.1 Synthesis

Na₂MgSiO₄ has been synthesized by solid state chemistry route. Starting reagents were high purity Na₂CO₃ (99.99 %, Aladdin), MgO (99.8 %, Aladdin) and SiO₂ (99.99 %, Aladdin). Both Na₂CO₃ and MgO were initially dried at 350 °C and 1000 °C overnight, respectively. The starting materials were weighed according to the stoichiometry and mixed in ethanol with an agate mortar and pestle. The dried mixtures were pressed under 4 Tons in a 12 mm diameter cylinder and calcined at 975 °C for 12 h for three times with intermediate grindings and pressings. The same batch of Na₂MgSiO₄ was used for the subsequent characterization techniques. Additionally, in order to maximize the ionic conductivity while preserving the polar orthorhombic phase, Na_{2-2x}Mg_{1-x}Si_{1+x}O₄ (x = 0.015, 0.025) compounds were prepared following the same method through substitution of Mg for Si to induce sodium vacancies. In order to prepare pellets for EIS measurements, the powder was mixed with polyvinyl acetate (PVA) and then sintered at 975 °C.

2.2 Characterization

Second-harmonic generation (SHG) measurements were carried out on the powders using the Kurtz–Perry method.¹⁶ The measurements were performed with a Q-switched Nd:YAG laser at a wavelength of 1064 nm. The intensity of the frequency-doubled output emitted from the samples was collected by a photomultiplier tube and compared to crystalline KH₂PO₄ (KDP, serving as reference).

Differential scanning calorimetry (DSC) data was acquired on a PerkinElmer STA 8000 simultaneous analyzer. About 20 mg of o-Na₂MgSiO₄ was placed in an Al₂O₃ crucible, heated at a rate of 10 °C min⁻¹ from room temperature to 800 °C, and then cooled to room temperature at the same rate under flowing air.

The identification of the crystalline phases was initially determined by laboratory X-ray powder diffraction (XRD), which were recorded on a Bragg–Brentano D8 Advance Bruker diffractometer (Cu K α radiation) equipped with a LynxEye XE detector over an angular range of $10^{\circ} < 2\theta < 130^{\circ}$. Variable temperature X-ray diffraction (VT–XRD) data were collected on Panalytical X'Pert PRO powder diffratometer (Cu $K\alpha$ radiation) equipped with HTK1200N furnace over the 15–65°2 θ range every 50 ° Cfrom RT up to 800 °C using 5°C/min heating and cooling rates, and allowing 3 min for temperature equilibration before collecting the data for 30 min at each set point.

High-intensity and high-resolution synchrotron X-ray powder diffraction (SPD) data was recorded on the 11-BM diffractometer at the Advanced Photon Source, Argonne National

Laboratory (USA). For this experiment, a sample of nominal composition Na_2MgSiO_4 was used. SPD data was acquired over the 0.5–50°20 range with a 0.001 step size at room temperature.

Constant-wavelength (λ =1.2289 Å) neutron powder diffraction (NPD) data were collected at ambient temperature over the 10–120° 2 θ range at 2 θ intervals of 0.05° on the 3T2 diffractometer at LLB-Orphée source (France).

The diffraction patterns were analysed using the Rietveld method implemented in the Topas-Academic software¹⁷ to identify the corresponding phase and estimate the amount and distribution of Na content. The bond valence sum calculation for the Na–O, Si–O and Mg–O bonds obtained from Rietveld analysis were calculated according to the following expression^{18, 19}

$$\sum v_i = \exp[(R_{ij} - d_{ij})/b] \qquad (1)$$

where v_i is the bond valence bond between the cation i and the anion *j*, b is a universal constant (0.37), d_{ij} is the observed bond length and the R_{ij} values have been taken as 1.803, 1.693 and 1.64 Å for Na–O, Mg–O and Si–O bonds, respectively as defined by Brown and Altermatt.¹⁸

The compositional homogeneity of the samples was analyzed by SEM and energy-dispersive X-ray spectroscopy (EDS) using a ZEISS Gemini SEM 300 scanning electron microscope (SEM, Germany), equipped with OXFORD EDS analyzer (ULTIM MAX 170, U.K.).

Solid state ²³Na and ²⁹Si NMR spectra were obtained using Bruker AVANCE I 400 (9.4 T), AVANCE III HD 750 (17.6 T), and AVANCE NEO 850 (20 T) NMR spectrometers. Chemical shifts were referenced to a 0.1 M NaCl solution for ²³Na and tetramethylsilane (TMS) for ²⁹Si. ²³Na magic-angle spinning (MAS) NMR spectra were acquired at 17.6 T and 20 T using MAS probes from Bruker, with 4 mm and 1.3 mm diameter rotors at the spinning rates of 12.5 kHz and 60 kHz, respectively. ²⁹Si MAS NMR data were collected using a Bruker AVANCE I 400 (9.4 T) spectrometer with a single pulse experiment while samples were spun at a frequency of 7

kHz in 7 mm Bruker probe. For ²⁹Si $\pi/2$ pulse width was 5 μ s with a recycle delay of 7200 s. The NMR parameters (chemical shifts, chemical shift anisotropies, asymmetry parameters, line widths, and quadrupolar parameters) were fitted using the DMfit program.²⁰ Spin–lattice relaxation (SLR) times (T_1) were obtained using a saturation recovery pulse sequence and the data were fitted to a stretched exponential of the form

$$1 - \exp[-(\tau/T_1)^{\alpha}]$$
 (2)

where τ are variable delays and α is the stretch exponential coefficient (between 0.85 and 1).

Variable temperature ²³Na MAS NMR data (up to 730 °C) were acquired on a Bruker AVANCE III 750 MHz NMR spectrometer using the diode laser-heated NMR probe head described previously.²¹ The sample was sandwiched between two layers of ground KBr which allowed for monitoring of the effective sample temperature through the shift of ⁷⁹Br resonance signal of KBr.²² In the high temperature laser heating setting, the sample is contained in an aluminum nitride container placed in a zirconia rotor and immobilized with two O-rings made from the sodium-containing ceramic that give some secondary signals in the high temperature experiments.

²³Na 2D exchange spectroscopy (EXSY) experiments were conducted at 17.6 T in a 4 mm rotor MAS probe while spinning at 12.5 kHz using the echo compensated EXSY pulse sequence developed by Messinger et al.²³ The acquisition conditions were as follows: $\pi/2$ pulse 7.75 µs, mixing time 100 ms (1250 and 5000 rotor periods), recycle delay (d_1) 2 s, with quadrature detection in indirect dimension.

Alternating–current (AC) impedance spectroscopy measurements were performed with a Solartron 1260 frequency response analyzer over a frequency range of $10^{-1} - 10^7$ Hz within the 100–800 ° Ctemperature range. Platinum paste was coated on the opposite faces of the pellets

(80% relative density) and heat treated at 600 °C for 30 minutes to burn out the organic components in the paste to form platinum electrodes. The impedance data analysis was carried out through the equivalent circuit fitting using the ZView software.²⁴ The activation energies (E_a) for ionic migration were determined from the slope of the conductivity curves by linear fitting of the Arrhenius equation:

$$\log() = \log A - (E_a/2.303 RT)$$
 (3)

where *A* is the pre–exponential factor, *R* is the gas constant, and *T* is the temperature in Kelvin.

2.3 Theoretical calculations

Density functional Theory (DFT) calculations for NMR parameters were performed using the CASTEP DFT code,²⁵ employing the gauge including projector augmented wave (GIPAW) NMR calculations²⁶ following the methodology previously reported.²⁷

The atomistic static–lattice and molecular dynamics (MD) atomistic simulations have been carried out on the basis of the interatomic potential approach employing the General Utility Lattice Program (GULP) program.^{28, 29} The Buckingham potential function was employed to describe the interactions between ions with the shell model³⁰ to describe the polarizability for the structure modeling of Na₂MgSiO₄. A three-body potential term was employed to describe the O-Si-O interaction.³¹ The potentials parameters used for the atomistic simulations are summarized in Table S1,^{31, 32} and tested well to reproduce the crystal structure of Na₂MgSiO₄ (Table S2). The MD simulation for sodium migration in Na_{2-2x}Mg_{1-x}Si_{1+x}O₄ were performed with the DL_POLY³³ code in the NVT ensemble with a Berendsen³⁴ thermostat on a 3×6×5 supercell model containing 2862 atoms (with the inclusion of the shell model) and having compositions of Na₂MgSiO₄ and Na_{1.95}Mg_{0.975}Si_{1.025}O₄. The system was equilibrated under a constant pressure of 1 atm for 100 ps, adding up to 1,250,000 time steps with a time step of 0.08 fs before initiating

the main MD simulations for 100 ps at specific temperatures within the 600 °C to 1000 °C temperature range. The substitutions of Mg with Si were randomly distributed in the supercell, resulting in Na vacancies uniformly distributed. The Visual Molecular Dynamic package³⁵ was adopted to view the migration pathways of Na ions, and the data of mean square displacements (MSDs) were exported by using nMoldyn code.³⁶ Sodium diffusion coefficients were obtained from the MSDs plots as a function of simulation time through linear fitting.

3 Results and Discussion

3.1 Long range structure.

The as-prepared Na₂MgSiO₄ compound exhibits weak second harmonic generation response (~0.125 × KDP at room temperature), confirming a non-centrosymmetric structure. The laboratory XRD pattern of Na₂MgSiO₄ can be correctly indexed in the *Pna*2₁ space group with $a \sim 10.89$ Å, $b \sim 5.29$ Å, $c \sim 7.06$ Å, similar to the unit cell reported by Withers et al (ICSD 84562).¹⁵ This indexation was confirmed by high-resolution SPD pattern recorded at the 11BM beam line, with no evidence for symmetry reduction to the maximal non-isomorphic subgroups P1a1, Pn11 or $P112_1$. These data revealed a strong and systematic asymmetry in the peak profiles, which is present throughout the pattern and is strongly *hkl*-dependent. In particular, the h00 peaks were found to exhibit substantial asymmetry with long tails towards high angles, whilst 00l exhibited similar tails towards low-angles, but the 0k0 peak were almost completely symmetrical (Figure S1). This is indicative of a complex microstrain with *a*- and *c*-parameters (which lie orthogonal to the channels in the framework) strained by compression and elongation respectively, with little apparent strain along the *b*-axis (which runs parallel to the channels, Figure 2a). The peak asymmetry was not apparent in the corresponding neutron diffraction pattern, due to the lower resolution of the instrument. The homogeneity of the sample was then

checked by EDS-SEM analysis which further evidenced a homogeneous distribution of elements at m-scale (see Figure S2) and no segregation of elements could be detected. We performed chemical analysis in different grains giving the average at% 46 ± 1 , 27 ± 1 , 27 ± 2 for Na, Mg and Si, respectively. Thus, the EDS-SEM data further confirmed that our sample is well reacted as well as the presence of some Na defect probably arising from Na volatilization which is commonly reported in the preparation of Na-based ceramics at high temperatures.^{14, 37, 38}



Figure 2. (a) [010] projections of the average *o*-Na₂MgSiO₄ structure emphasizing the tetrahedral framework. Grey and blue colors represent the Mg and Si atoms, respectively. Navy blue and yellow green colors indicate Na1 and Na2, respectively. (b) Representative portion of the Rietveld plots from SPD (R_{wp} = 9.16%, R_p = 7.11%, χ^2 = 1.01) and (c) NPD (R_{wp} = 2.61%, R_p

= 2.01%, χ^2 = 3.00) data using the multiple cells (seven) based on the *Pna*2₁ structure model. Experimental (circles), fitted (dark green line) and difference curve (purple). The sets of vertical lines correspond to the reflection positions for each one of the seven unit cells of o-Na₂MgSiO₄ employed. Overall reliability factors: R_{wp} = 5.72%, R_p = 5.26%, χ^2 = 1.44. d) Normalized refined lattice parameter distribution and unit cell weighting data obtained from SPD-NPD combined refinement.

We then carried out a combined Rietveld refinement using the SPD and NPD datasets. To account for the peak asymmetry in the SPD pattern, we constructed a multiple-unit-cell Rietveld model starting from the published Pna21 structure of Na2MgSiO415 as explained in supplementary Note 1. Figure 2b depicts a representative view of the SPD final refinement (a general view can be found in Figure S3). Figure 2c presents the final refinement of the NPD pattern. The resulting refined structure represents a weighted average of the seven unit cells, whose refined lattice parameter distribution and weighting is shown in Figure 2d. The final refined composition was Na_{1.968(3)}Mg_{0.968(4)}Si_{1.032(4)}O₄, leading to a mechanism in which sodium volatilization induces sodium vacancies that are compensated by Si substitution for Mg according to Na_{2-2x}Mg_{1-x}Si_{1+x}O₄. The multi-cell refinement results indicate that the sample contains a small sub-micro compositional distribution arising from uncontrollable Na volatilization that is strongly coupled to an anisotropic local distortion of the crystal structure that shortens/elongates the a/c lattice parameters respectively (see Figure 2d), while the b parameter is only weakly affected because the channels are distributed along this [010] direction (Figure 2a). This is possible because of the high flexibility of the framework structure through the tetrahedral connections even when the Si/Mg tetrahedra are quite rigid.³⁹ Overall this causes the unit cell volume to be strained towards smaller values, which is consistent with the presence

of vacancies on the Na sites as well as the Si⁴⁺ substitution for the larger Mg²⁺. It is worth noting that careful examination of the final refinement of SPD data evidenced a set minor unindexed reflections (Figure S4). Although we could not assign them to any known compound within the Na₂O-MgO-SiO₂ phase diagram, it is consistent with the proposed mechanism in which sodium vacancies from sodium volatilization are compensated by the substitution of Si for Mg, which drove the final composition of the sample into a multiple-phase coexistence. The validity of our multiple-unit-cell model was further corroborated with the preparation of two additional non-stoichiometric compounds belonging to the Na_{2-2x}Mg_{1-x}Si_{1+x}O₄ system, *i.e.* x = 0.015, 0.025 which exhibit the same unit cell parameters evolution (smaller *a* axis, longer *c* axis but nearly unchanged *b* axis and overall unit cell volume reduction, Figure S5a-b).

Table S1-2 gather the refined structural parameters and main bond distances of the main phase. The two inequivalent Na sites hosted within the channels in the -cristobalite structure exhibits similar number of vacancies (~1% and 2% in Na1 and Na2, respectively). Both are coordinated by twelve oxygens, although a careful observation of the Na–O bond lengths indicates the presence of five and four nearest oxygen to Na1 and Na2 sites, respectively, within 2.31–2.75 Å as first coordination spheres, while the rest are placed further than 3 Å. Therefore, considering Na1 as five coordinated and Na2 as four coordinated seems reasonable given their respective bond valence sums (BVS) 0.88 and 1.08 close to unit.

3.2 Short range structure

²⁹Si MAS NMR. The room temperature ²⁹Si MAS NMR spectrum (Figure S6) mainly contains one Si site in very symmetrical tetrahedral Si(4Mg) environment, confirming an ordered Si/Mg distribution in the structure. The value of the ²⁹Si chemical shift, –69.3 ppm, and the full width at half-maximum (FWHM), 76 Hz (1.0 ppm), are close to those obtained by Withers et al., *i.e.*, –

68.9 ppm and FWHM = 1.2 ppm.¹⁵ In addition, there is another weak resonance around -75.3ppm. This signal may be attributed to the secondary phase observed by SPD. Nevertheless, we have dismissed it because the relative amount of Si in different environments is 94/6 according to the ²⁹Si MAS NMR spectrum fit, while the amount of the impurity phase has been estimated as 0.3 wt% from the SPD pattern (calculated from the A_{020}/A_i ratio, being A_{020} the integrated area of the (020) reflection and A_i the integrated area of the unindexed peak at 2 ~10.7°- Figure S4). More likely, the presence of this signal can be assigned to the existence of a Si(3Mg, 1Si) site as described by Withers et al. for the non-stoichiometric compounds in Na_{2-2x}Mg_{1-x}Si_{1+x}O₄ system (i.e., -74.9 ppm).¹⁵ Moreover, the relative amount of Si in different environment (Si(4Mg) and Si(3Mg, 1Si)) determined as 94/6 is in reasonably good agreement to the refined composition, 97.8(3)/3.2(3). This further supports that some Si moves inside the Mg tetrahedra to compensate the charge imbalance due to Na volatilization. Theoretically, Si substitution for Mg in the fully ordered structure of Na₂MgSiO₄ would have led to formation of Si(0Mg, 4Si) and Si(3Mg, 1Si) centers. Nevertheless, the presence of additional environments (e.g. Si(0Mg, 4Si), Si(1Mg, 3Si) or Si(2Mg, 2Si)) were not detected, which indicates that small amount of Si substitution for Mg in the ordered Na₂MgSiO₄ avoids condensation of SiO₄ tetrahedra. This may interrupt locally ordered Mg and Si distribution in Na₂MgSiO₄, which accounts for the swelling feature on the background of NPD data (Figure 2c).

²³Na MAS NMR. The room temperature ²³Na MAS NMR spectrum revealed the presence of two Na sites with relative multiplicities of 1:1 with well–defined quadrupolar line shapes (Figure 3), as expected. From the fitted spectrum, the chemical shifts (δ_{iso}) and quadrupolar parameters (quadrupolar constant, C_Q and asymmetry parameter, η_Q) were obtained and the assignment of ²³Na resonances to sodium crystallographic sites based on the DFT computations considering the

Mg/Si fully ordered structural model is given in Table 1. Na1 site exhibits a larger quadrupolar coupling C_Q (3.3 MHz) than that observed for Na2 (2.1 MHz), confirming the more distorted environment observed from the SPD data refinement (Figure 3).



Figure 3. Experimental (blue) and simulated (red) ²³Na MAS NMR spectra at 20 T and 60 kHz recorded at room temperature. The fitting of the individual ²³Na resonances at Na1 and Na2 sites are shown in purple and green, respectively. Inset: Sodium coordination polyhedra in Na1 and Na2 sites. The bond lengths have been obtained from Rietveld combined refinement on NPD-SPD data.

Table 1. ²³Na isotropic chemical shifts (δ_{iso}), quadrupolar constants (C_Q), integrated intensity (I) and asymmetry parameters (η_Q) obtained from the simulation of the ²³Na MAS at 17.6 T and 20 T and calculated ²³Na NMR parameters (isotropic magnetic shielding constant σ_{iso} , chemical shifts $\delta_{isocalc}$, quadrupolar constant $|C_{Qcalc}|$, and asymmetry parameter η_{Qcalc}) for *o*-Na₂MgSiO₄.

Site	$\delta_{ m iso} \pm 0.2 m ppm$	Co ±0.2MHz	$\eta_{\rm Q}$ ±0.1	I(%) ±2%	σ_{iso} ppm	$\delta_{ ext{isocalc}} \ ext{ppm}$	C _{Ocalc} MHz	$\eta_{ ext{Qcalc}}$
Na1	16.0	3.3	0.74	50	537.9	16.1	3.86	0.72
Na2	28.7	2.1	0.42	50	521.6	29.8	2.33	0.32

3.3 Macroscopic ionic migration

Electrochemical impedance spectroscopy (EIS). The electrical behavior characterization of the sample was undertaken by EIS. At temperatures below ~150 C, the spectra exhibit non-well defined shapes in the low frequency range (see Figure S7). Between ~150 - 400 °C, the spectra consist of one depressed semicircle arc in the high frequency region and one Warburg-type inclined line in the low frequency region with capacitances of $>10^{-8}$ F·cm⁻¹ at 10⁻¹ Hz,⁴⁰ indicative of ionic conduction. Figure 4a depicts the complex impedance plot of Na₂MgSiO₄ recorded at 154 ° C, showing one asymmetric and depressed semicircular arc in high frequency region. In the C' vs. f plot two overlapping plateaus can be apparently discerned within $10^7 - 10^2$ Hz region (Figure 4a, inset): the one in $10^2 - 10^3$ Hz has capacitances ~2 pF·cm⁻¹ and the other one within $10^5 - 10^6$ Hz has smaller capacitances ~0.5 pF·cm⁻¹. Both capacitance plateaus lie with the bulk responses,⁴⁰ *i.e.* that the semicircular arc actually comprises two bulk components and may be indicative of the electrical inhomogeneous nature of the as-prepared Na₂MgSiO₄, probably due to partial Na volatilization. Apart from the overlapping bulk response arc and electrode response line, no apparent grain boundary response was observed in the complex impedance plot. Therefore deconvolution of complex impedance plot was performed through equivalent circuit fitting with two bulk and one electrode components, which is shown in Figure S8. The high frequency part of the bulk impedance data is a series combination of one bulk component consisting of R_{b1} , C_{b1} , CPE₁ in parallel and the other bulk component consisting of R_{b2} , C_{b2} in parallel. The fit gives $R_{b1} = 1.0(1) \times 10^7 \,\Omega \cdot \text{cm}, C_1 = 4.3(3) \times 10^{-13} \text{ Fem}^{-1}, R_{b2}$

~ $2.9(2) \times 10^{6} \Omega \cdot cm$ and $C_2 \sim 1.9(1) \times 10^{-12} \text{ F} \cdot cm^{-1}$. It must be indicated that a CPE element is not included for the second element since its values are not meaningful and, indeed, a satisfactory fit was obtained. While increasing the temperature, the bulk response arc disappeared gradually and above ~400 °C only electrode response was identified. The electrode response recorded at

650 ° Cdoes not show any evident collapse after changing the atmosphere from pure nitrogen to pure oxygen, in contrast with the typical behavior to partial oxygen pressure exhibited by oxide ion conductors (Figure 4b).^{5, 41-43} Moreover, its ionic conductivity is maintained at different pO_2 values (Figure 4b-inset). Therefore, the ionic conduction in *o*-Na₂MgSiO₄ arises from sodium cations instead of oxide ions.

Figure 4c depicts the Arrhenius plots for the bulk conductivities of o-Na₂MgSiO₄. It exhibits $\sigma_{\rm b}$ ~ 10^{-5} S·cm⁻¹ at 300 °C, in good agreement with the values reported by Baur¹³ and Shannon,¹⁴ and at 800 °C its conductivity reaches $\sim 5 \times 10^{-3}$ S·cm⁻¹. Slight curvature was observed on the Arrhenius plot of bulk conductivity: the E_a is ~0.73(1) eV within 100 – 500 °C and then decreases at higher temperatures (~0.51(2) eV between 500 - 800 °C). This evolution cannot be attributed to any structure change, as either VTXRD or DSC data do not evidence any structure rearrangement apart from the enlargement of the unit cell parameters and Na–O bonds (Figure S9-S10, respectively). The change of E_a may be related to a local order-disorder transition of sodium vacancies at high temperature, where the sodium ions become progressively more mobile. Such local order-disorder of small amount of sodium vacancies is not enough to change significantly the environments of sodium ion in the structure. Therefore, it is hard to deduce the information on the order-disorder of sodium vacancies from the variable-temperature ²³Na MAS NMR data described later. Moreover, the conductivity can be further improved by inducing more sodium vacancies through varying the stoichiometry allowing for the Mg substitution for Si in the tetrahedral network according to the Na_{2-2x}Mg_{1-x}Si_{1+x}O₄ formula proposed above and achieves ~ 10^{-5} S·cm⁻¹ at 200 °C for the *o*-Na_{1.95}Mg_{0.975}Si_{1.025}O₄ nominal composition (Figure 4c) while conserving the polar $Pna2_1$ structure.



Figure 4. (a) Experimental (open circles) and calculated (red line) complex impedance plot of *o*-Na₂MgSiO₄ at 154 °C in air and its deconvolution to two bulk components (in blue and green lines). The numbers denote the logarithms of the selected frequencies marked by filled circles. The inset shows the capacitance as a function of frequency of *o*-Na₂MgSiO₄ at 154 °C. (b) Complex impedance plots for *o*-Na₂MgSiO₄ at 650 °C under N₂, air and O₂. The inset depicts the pO_2 dependency of the conductivity of Na₂MgSiO₄ at 650 °C. (c) Arrhenius plots of total conductivity for *o*-Na_{2-2x}Mg_xSi_{1+x}O₄ (*x* = 0, 0.015, 0.025) compositions.

MD simulation. Atomistic static lattice and MD simulations were carried out on the basis of the interatomic potential method for both *o*-Na_{1.95}Mg_{0.975}Si_{1.025}O₄ and *o*-Na₂MgSiO₄ nominal compositions. Figure 5a shows the calculated MSD values of Na, Mg, Si and O atoms in *o*-Na_{1.95}Mg_{0.975}Si_{1.025}O₄ obtained at 1000 °C (generally higher temperatures than the experimental ones are required for the simulations to observe the long-range migration paths within the limited simulated time 100 ps). Only Na ions migrate through the structure; Mg, Si and O ions vibrate slightly around their lattice positions without long–range migration, in good agreement with the sodium ionic conduction character revealed by impedance spectroscopy. The sodium diffusion coefficients, calculated from the MSD values of Na ions, fall in the range of ~ 10^{-8} – 10^{-6} cm²·s⁻¹ within the 700-1300 °C temperature range (Figure S11) and the calculated *E*_a obtained from the

Arrhenius plot of the sodium diffusion coefficients is 0.69(2) eV, which is between the experimental values in the high (0.73(1) eV) and low (0.51(2) eV) temperature ranges. The streaming and overlapping of positions of sodium ions originally at Na1 and Na2 sites in their scatter plots (Figure 5b), together with their MSD values suggest that both Na sites are involved in, and equally contribute to the sodium ion migration mainly through hopping between Na1 and Na2 sites, forming three-dimensional migration zig-zag shaped paths. For comparison, the MSD values for Na ions in Na₂MgSiO₄ (red dashed line in Figure 5a), *i.e.* free of vacancies, are remarkably lower than those calculated for Na_{1.95}Mg_{0.975}Si_{1.025}O₄ at the same temperature. This further supports that the presence of Na vacancies is strictly necessary to observe ionic conductivity in Na₂MgSiO₄.



Figure 5. (a) MSD values of Na, Mg, Si and O atoms in Na_{1.95}Mg_{0.975}Si_{1.025}O₄ as a function of time from the MD simulations at 1000 °C. The inset depicts the MSD values of Na1 and Na2 atoms. Green dashed line indicates the MSD for Na in vacancy-free Na₂MgSiO₄ compound at 1000 °C. (b) Scatter plots of sodium ion positions showing the diffusion paths consisting of both Na1 and Na2 sites at 1000 °C (marked by dashes lines) from MD simulations. Blue and greenish-olive dots denote sodium ions originally placed at Na1 and Na2 sites respectively.

3.4 Microscopic ionic migration.

VT ²³Na MAS NMR spectroscopy. Since the pure sodium ionic conduction nature is demonstrated in o-Na₂MgSiO₄, the sodium migration dynamics at local scale was monitored by VT ²³Na MAS NMR spectroscopy. As aforementioned, the spectrum of o-Na₂MgSiO₄ recorded under a magnetic field of 17.6 T and spinning at 5 kHz exhibits two well-separated quadrupolar lines expressing second order line shapes. It is worth mentioning that when MAS NMR measurements were performed at room temperature, and without temperature regulation, the actual sample temperatures were 49±2 °C at 5 kHz (7 mm rotor) MAS and 52±2 °C at 12.5 kHz (4 mm rotor) due to frictional heating. Considering the 1D experiments (Figure 6a), we observed that the temperature increase from 49 °C up to 730 °C first led to the collapse of the quadrupolar line shapes merging to a single broad spectrum at 175-200 °C that then sharpened into an averaged line up to 730 °C. This behavior is closely related to what was observed for nepheline (a trydimite-type framework) at 500 °C and it is indicative of an ion exchange process.^{9, 44} As indicated by the VT-XRD data, this tendency cannot be attributed to any structure change within the analyzed temperature range. We have estimated the activation energy for Na migration from ²³Na SLR time measurements (T_1) (Figure S12). The linear slope indicates an $E_a = 0.321$ eV for fluctuations causing relaxation, which is much smaller than that obtained from conductivity measurements (0.51(2)-0.73(1) eV), probably because that NMR relaxation is occurring not only through cationic motion required for diffusion and conductivity, but also via local fluctuation of Na environment (via local ion hopping back and forth between sites and/or by motion of nearby defects). This may lead to lower activation energies than those needed for through-going (longrange) ionic diffusion alone.^{11, 47-49} It is worth noting that both techniques tends to reflect different averaging of the underlying distribution of dynamic processes and it is not surprising to

observe such a mismatch. For instance, similar phenomenon of lower activation energy observed by NMR than those needed for the longer-range sodium ionic diffusion was also noticed in the glassy phase from the sodium-doped $SrSiO_3$ compositions⁴⁵ or other crystalline sodium-based materials such as $Na_2B_{10}H_{10}^{46}$ or Na_3InCl_6 .⁴⁷ Therefore the lower activation energy observed by NMR may indicate that local hopping mechanism different from the long-range migration takes place in the sample.

²³Na 2D EXSY experiments. In order to better understand the dynamics of sodium mobility in the material at moderate temperature and before the merging of quadrupolar line-shapes, we performed ²³Na 2D EXSY experiments at 52 °C and 100 °C. The EXSY experiments (Figure 6b) allow evidencing slow ionic jumps between the ²³Na crystallographic sites at the time scale of hundred of ms exchange or mixing time. Non-exchanged Na atoms do not change frequency during the mixing time but they sign on the diagonal which is homogeneously broadened by spin-spin relaxation (T_2) . Na atoms exchanging with a neighboring site will sign as off-diagonal (direct exchange between different structural positions) or on-diagonal cross-peaks (relayed exchange). Thanks to the specific line shapes of the cross-peaks correlating two second order broadened line-shapes, we implemented a simulation of the different components in DMfit (available from version #20200728 available on request) and to retrieve their respective contributions (integrals of diagonal and cross peaks).²⁰ At 52 °C and 100 ms exchange we observe 83 % of non-exchanged Na atoms, 17 % of exchange between Na1 and Na2 sites and negligible self exchange; at 100 °C and 100 ms exchange, we observe at 23% of non-exchanged Na atoms, 48% of exchange between Na1 and Na2 sites and 28 % self exchange (Table S5).



Figure 6. (a) ²³Na NMR spectra recorded at 17.6 T and MAS 5 kHz as a function of temperature. The weak ²³Na signals between -5 and -2 ppm (red circle) arise from the ceramic O–rings used

to fix the AlN sample holder in place within the zirconia rotor. (b-c) 23 Na 2D EXSY NMR spectra (left) obtained with 100 ms mixing time, (b) at 52 °C; (c) at 100 °C and their fitting results (right).

3.5 Global sodium migration mechanism in o-Na2MgSiO4

According to our data covering the sodium dynamics from macroscopic to microscopic range, an atomistic point of view of the Na migration can be understood as follows. Four Na1 atoms located in the second coordination sphere of the Na2 atom and also four Na2 atoms are placed in the second coordination sphere of the Na1 atom. Since the distances between two different sodium sites (3.12; 3.23; 3.30; 3.36 Å) are shorter than the distances between the same sites (from 4.77 up to 5.61 Å), we only observe exchange between Na1 and Na2 sites in the ²³Na EXSY NMR spectrum at 52 °C (this step is illustrated in Figure 7a). Upon slight heating (100 ° G, the sodium atoms start to migrate-diffuse within and between the channels of the magnesium–silicate framework. Then, with a further increase in temperature, the movement of atoms becomes very fast and we cannot distinguish these two states anymore on the NMR timescale as inferred from ²³Na EXSY NMR data and the long range diffusion of sodium atoms takes place though the 3D framework of the structure mainly along the paths composed of alternate Na1 and Na2 sites, as exemplified in Fig 7b.



Figure 7. (a) <010> view of the sodium site exchange at low temperatures. Si, Mg, Na1 and Na2 are represented as blue, grey, navy blue and greenish spheres, respectively. (b) <010> view of Na₂MgSiO₄ refined structure emphasizing the voids along (010) direction. The macroscopic Na migration at high temperatures along the [100] (blue), [010] (red) and [001] (purple) directions are indicated. The interconnected Na atoms are placed at distances ranging ~ 3.12-3.36 Å.

4 Conclusions

The sodium dynamics have been studied and described for the first time in polar stuffedcristobalite *o*-Na₂MgSiO₄ through a complementary set of techniques (EIS, MD simulations, (VT) ²³Na MAS NMR and ²³Na EXSY NMR). *o*-Na₂MgSiO₄ exhibits two inequivalent Na sites and it should not exhibit ionic conduction on the strictly stoichiometric composition, according to molecular dynamics simulations. Yet, the synthesis route employed results partial Na volatilization (~2 at%) that converts the material into a pure ionic conductor exhibiting moderated ionic conductivity (~ 10^{-5} S cm⁻¹ at 300 °C) that can be further improved by inducing additional Na vacancies while keeping the polar structure according to Na_{2-2x}Mg_{1-x}Si_{1+x}O₄ formula. 1D ²³Na VT MAS NMR and ²³Na EXSY NMR clearly evidence that a sodium site exchange at ~ R*T* takes place triggering the sodium ionic conduction at higher temperatures. This work demonstrates the sodium site exchange phenomenon for first time by ²³Na EXSY NMR and paves the way to possible further identification of sodium ionic conductors from complex structures as 3D stuffed-cristobalite ones exhibiting multiple sites for mobile ions.

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ASSOCIATED CONTENT

Supporting Information. SPD diffraction refinements; SEM-EDS elemental mapping for Na₂MgSiO₄; Laboratory XRD patterns of *o*-Na_{2-2x}Mg_{1-x}Si_{1+x}O₄ samples and normalized refined lattice parameters; ²⁹Si MAS NMR spectrum; Complex impedance plot at 60 °C; Equivalent circuit to fit the EIS spectrum recorded at 154 °C; VTXRD, DSC and unit cell evolution within

30-750 ° C Arrhenius plot of the sodium diffusion coefficient; ²³Na spin–lattice relaxation times plotted against 1000/T; Structural parameters and selected interatomic distances at room temperature; Interatomic potential parameters used for the MD simulations and experimental and calculated structural parameters; Multiple-unit-cell Rietveld model description.

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