

Cite this: *RSC Appl. Interfaces*, 2024,  
1, 1419

## Preparation and characterization of new solid electrolytes $\text{Na}_{3-x}\text{Zn}_{1-x}\text{Al}_{1+x}\text{S}_4$ †

Tomoya Otono, Hamdi Ben Yahia, Chie Hotehama, Kota Motohashi, Atsushi Sakuda \* and Akitoshi Hayashi \*

Sulfide solid electrolytes, known for their high ionic conductivity and formability, are key materials for the practical use of all-solid-state sodium batteries. In this study, new sulfide solid electrolyte materials,  $\text{Na}_{3-x}\text{Zn}_{1-x}\text{Al}_{1+x}\text{S}_4$  ( $x \leq 0.2$ ), were prepared via a self flux synthesis route, using reagents such as  $\text{Na}_2\text{S}$ , Zn, Al, and S. The new materials were characterized using X-ray powder diffraction, Raman spectroscopy, and electrochemical impedance spectroscopy.  $\text{Na}_{3-x}\text{Zn}_{1-x}\text{Al}_{1+x}\text{S}_4$  formed a solid solution up to  $x = 0.2$  and crystallized with a  $\beta\text{-Ca}_3\text{Ga}_2\text{N}_4$ -type structure. As the Al content increased, the number of sodium vacancies also increased, resulting in improved ionic conductivity. Among  $\text{Na}_{3-x}\text{Zn}_{1-x}\text{Al}_{1+x}\text{S}_4$  samples,  $\text{Na}_{2.9}\text{Zn}_{0.9}\text{Al}_{1.1}\text{S}_4$  exhibited the highest ionic conductivity of  $4.5 \times 10^{-6} \text{ S cm}^{-1}$  at 25 °C and lowest activation energy of 32 kJ mol<sup>-1</sup>. Furthermore, the  $\text{Na}_{2.9}\text{Zn}_{0.9}\text{Al}_{1.1}\text{S}_4$  phase was relatively stable when exposed to humid air, which facilitated its practical use in all-solid-state sodium batteries.

Received 31st July 2024,  
Accepted 10th September 2024

DOI: 10.1039/d4lf00275j

rsc.li/RSCApplInter

## Introduction

Sodium secondary batteries are promising for large-scale energy storage devices owing to the abundance of sodium resources and low cost.<sup>1–3</sup> All-solid-state sodium secondary batteries are expected to meet both safety requirements and low cost.<sup>4</sup> Solid electrolytes are key materials for the practical application of all-solid-state sodium batteries.<sup>5,6</sup> Among these, sulfide solid electrolytes exhibit the highest ionic conductivities and formabilities.<sup>7</sup> We previously reported that  $\text{Na}_3\text{PS}_4$  glass-ceramics exhibited an ionic conductivity of  $2.0 \times 10^{-4} \text{ S cm}^{-1}$  and that all-solid-state sodium cells with  $\text{Na}_3\text{PS}_4$  glass-ceramics operated reversibly at 25 °C for the first time.<sup>8</sup> Various sulfide electrolytes, such as  $\text{Na}_{11}\text{Sn}_2\text{PS}_{12}$  and  $\text{Na}_3\text{SbS}_4$ -based electrolytes, have been investigated.<sup>9–17</sup> Particularly,  $\text{Na}_{2.88}\text{Sb}_{0.88}\text{W}_{0.12}\text{S}_4$  solid electrolytes, where Sb in  $\text{Na}_3\text{SbS}_4$  was partially replaced by W, exhibited the highest ionic conductivity of  $1.25 \times 10^{-1} \text{ S cm}^{-1}$  at 25 °C.<sup>17</sup> Sulfide electrolytes generate  $\text{H}_2\text{S}$  gas when exposed to humid air. To increase the stability in humid environments, substitutions of sulfur with oxygen and nitrogen in sulfide electrolytes have been explored.<sup>18,19</sup> Furthermore, electrolytes using a softer acid as a central element were also investigated.<sup>20–22</sup> Among them,  $\text{Na}_3\text{SbS}_4$ -based electrolytes exhibited the highest ionic

conductivity and stability under humid environments.<sup>12–16</sup> However,  $\text{Na}_3\text{SbS}_4$ -based electrolytes had low reduction stabilities and narrow electrochemical windows.<sup>23–25</sup> To construct safe all-solid-state sodium batteries, sulfide electrolytes with high ionic conductivity, high tolerance to humidity, and high reduction stability are required.

Recently,  $\text{Na}_3\text{ZnGaS}_4$ -based sulfide electrolytes have been reported.<sup>26–29</sup>  $\text{Na}_3\text{ZnGaS}_4$  with a framework of corner-shared  $(\text{Zn}/\text{Ga})_4\text{S}_{10}$  super-tetrahedra structure exhibits an ionic conductivity of  $3.0 \times 10^{-7} \text{ S cm}^{-1}$  at room temperature, and its ionic conductivity can be increased to over  $10^{-4} \text{ S cm}^{-1}$  by tuning the relative ratio of Zn/Ga and introducing sodium vacancies.<sup>27</sup> When the  $\text{Na}_3\text{ZnGaS}_4$ -based electrolytes were exposed to humid air,  $\text{H}_2\text{S}$  gas was not generated, and X-ray powder diffraction (XRPD) patterns did not change, suggesting that  $\text{Na}_3\text{ZnGaS}_4$ -based electrolytes have high humidity stability.<sup>27–29</sup> Moreover, the  $\text{Na}_2\text{Sn}$  symmetric cell of  $\text{Na}_3\text{ZnGaS}_4$ -based electrolytes operated stably, suggesting that among sulfide solid electrolytes,  $\text{Na}_3\text{ZnGaS}_4$ -based electrolytes have relatively high stability for negative electrodes.<sup>27–29</sup> Among the samples of  $\text{Na}_3\text{ZnMS}_4$  ( $M = 13\text{th}$  group elements), the structures of  $\text{Na}_3\text{ZnGaS}_4$  and  $\text{Na}_3\text{ZnInS}_4$  have been previously reported;<sup>30</sup> however, the structure of  $\text{Na}_3\text{ZnAlS}_4$  has not been reported to date.  $\text{Na}_3\text{ZnAlS}_4$ -based electrolytes are expected to exhibit high ionic conductivities, similar to those of  $\text{Na}_3\text{ZnGaS}_4$ -based electrolytes. Moreover, aluminum is the second most abundant resource in the Earth's crust,<sup>3</sup> and  $\text{Na}_3\text{ZnAlS}_4$ -based electrolytes can be synthesized at low costs.

$\text{Na}_3\text{ZnAlS}_4$ -based electrolytes can be synthesized using sodium polysulfides. Sulfide electrolytes can be synthesized

Department of Applied Chemistry, Graduate School of Engineering, Osaka Metropolitan University, 1-1 Gakuen-cho, Naka-ku, Sakai, Osaka 599-8531, Japan.  
E-mail: saku@omu.ac.jp, akitoshihayashi@omu.ac.jp; Fax: +81 72 2549334;  
Tel: +81 72 2549331

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4lf00275j>



using sodium polysulfides *via* an ambient-pressure heat treatment without a sealing process. Moreover, sodium polysulfides are used as self-fluxes, and aluminum elements can be used as aluminum sources instead of  $\text{Al}_2\text{S}_3$ , which is challenging to obtain as a high-purity starting material.<sup>17</sup>

In this study, we synthesized the new material  $\text{Na}_3\text{ZnAlS}_4$  with the  $\beta\text{-Ca}_3\text{Ga}_2\text{N}_4$ -type structure using sodium polysulfides.  $\text{Na}_{3-x}\text{Zn}_{1-x}\text{Al}_{1+x}\text{S}_4$  samples, in which sodium vacancies were introduced to increase the ionic conductivity, were also synthesized and their ionic conductivities and moisture stabilities were evaluated.

## Experimental section

### Synthesis

The  $\text{Na}_{3-x}\text{Zn}_{1-x}\text{Al}_{1+x}\text{S}_4$  ( $x = 0, 0.1, 0.15, \text{ and } 0.2$ ) samples were prepared *via* a simple heat treatment synthesis route using stoichiometric mixtures of  $\text{Na}_2\text{S}$  (99.1%; Nagao), Zn (99.99%; Fujifilm Wako Chem.), Al (99.9%; Kojundo Chem.), and S (99.99%; Kojundo Chem.). The reagents were placed in carbon crucibles and heated in an electric furnace at 750 °C for 17 h under ambient pressure, followed by quenching at room temperature. During synthesis, sodium polysulfide melts were produced and reacted with Zn and Al.  $\text{Na}_{3-x}\text{Zn}_{1-x}\text{Al}_{1+x}\text{S}_4$  ( $x = 0, 0.1, 0.15, 0.2$ ) samples did not melt at 750 °C. All processes were performed inside a glove box under a dry Ar atmosphere.

### Characterization

The X-ray powder diffraction (XRPD) patterns of the prepared materials were measured with a Rigaku SmartLab diffractometer with a  $\text{Cu-K}\alpha$  radiation source, using Bragg-Brentano geometry. X-ray data were collected in a  $2\theta$  range of 5.0–120.0° in steps of 0.01°. The data were refined by Le Bail profile analysis using JANA2006 software,<sup>31</sup> and crystal structures were confirmed by Rietveld refinement. In the first step, we performed a Le Bail profile fitting analysis. In this step, the cell parameters, the origin shift, the background, and the profile function (pseudo-Voigt) were refined. Once good  $R$  values were obtained, we fixed all these parameters as the initial cell parameter. In the second step, we analyzed the crystal structure. We included the atomic positions from the isostructural compound  $\text{Na}_3\text{ZnGaS}_4$ , and we refined their coordinates, their thermal parameters and the scale factor. Once the reliability factors converged to low values, we refined all parameters including those of the profile.

The Raman spectra of the samples were measured with a Raman spectrophotometer (LabRAM HR-800; Horiba) using a 532 nm solid-state laser. The samples were placed in a vessel, sealed airtight with a cover glass, and placed in a glovebox under a dry Ar atmosphere.

The morphology and the elements distribution were observed using scanning electron microscope (SEM) (SU8220; Hitachi High Technologies) equipped with an energy dispersive X-ray spectroscopy system (EDS) (EMAX Evolution; Horiba Ltd.).

The ionic conductivities were determined using an impedance analyzer (Solartron; 1260) *via* an alternating

current (AC) impedance measurement in the frequency range of 0.1 Hz to 10 MHz, under an applied voltage of 50 mV. Measurements were performed using compressed powder pellets (diameter: 10 mm; thickness: 1 mm). Gold thin films (diameter: 10 mm) were deposited as ion-blocking electrodes on both faces of the pellets using a quick coater (Quick Coater SC-701; Sanyu Electron). Each pellet was sealed in a laminate-type pouch cell to prevent exposure to air.

The humidity stability of the  $\text{Na}_{2.9}\text{Zn}_{0.9}\text{Al}_{1.1}\text{S}_4$  electrolyte was evaluated by exposing the powder to an air flow at 70% relative humidity for 30 min at room temperature. Subsequently, the powder was analyzed using XRPD and Raman spectroscopy.

## Results and discussion

The XRPD pattern of  $\text{Na}_3\text{ZnAlS}_4$ , shown in Fig. 1, is significantly similar to those of  $\text{Na}_3\text{ZnGaS}_4$  and  $\text{Na}_3\text{ZnInS}_4$ . This suggests that the three phases were isostructural. Consequently, by adjusting the cell parameters of  $\text{Na}_3\text{ZnGaS}_4$ , a full pattern matching was performed successfully. This yielded the refined cell parameters of  $a = 12.9206(1)$  Å,  $c = 18.5822(2)$  Å, and  $V = 3102.12(4)$  Å<sup>3</sup> for  $\text{Na}_3\text{ZnAlS}_4$ . This analysis revealed the presence of impurity peaks at 18.6°, 24.7°, 26.5°, 38.7°, 38.8°, 83.2°, and 83.4°. These peaks were excluded, and Rietveld refinement was performed by using the atomic positions of  $\text{Na}_3\text{ZnGaS}_4$  as the starting model. Only the Ga atom at the 32g Wyckoff position was replaced by Al. With isotropic atomic displacement parameters (ADPs), the reliability factors converged to satisfactory values. However, the Na2 atom displayed a much larger ADP than Na1 (0.087 Å<sup>2</sup> vs. 0.035 Å<sup>2</sup>). Therefore, anisotropic ADPs were applied to Na2 atoms. This resulted in the final reliability factors, refined atomic positions, and anisotropic ADPs listed in Tables 1–3, respectively. Fig. 2 shows the excellent agreement between the experimental and calculated patterns.

The crystal structure of  $\text{Na}_3\text{ZnAlS}_4$  is shown in Fig. 3.  $\text{Na}_3\text{ZnAlS}_4$  crystallizes in a  $\beta\text{-Ca}_3\text{Ga}_2\text{N}_4$ -type structure.<sup>32</sup> The structure was composed of (Zn/Al) $\text{S}_4$  tetrahedra that shared corners and formed (Zn/Al) $\text{S}_{10}$  clusters (Fig. 3b). These

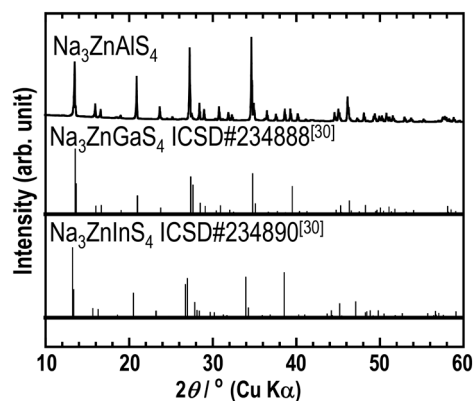


Fig. 1 Experimental XRPD pattern of  $\text{Na}_3\text{ZnAlS}_4$  and calculated patterns of  $\text{Na}_3\text{ZnInS}_4$  and  $\text{Na}_3\text{ZnGaS}_4$ .



**Table 1** Crystallographic data and structure refinement for Na<sub>3</sub>ZnAlS<sub>4</sub>

Crystal data	
Chemical formula	Na <sub>3</sub> ZnAlS <sub>4</sub>
Mr	289.6
Crystal system space group	Tetragonal <i>I</i> 4 <sub>1</sub> / <i>acd</i>
Temperature (K)	293
<i>a</i> (Å)	12.9206(1)
<i>c</i> (Å)	18.5822(2)
<i>V</i> (Å <sup>3</sup> )	3102.12(4)
<i>Z</i>	16
Data collection	
Radiation type	Cu-Kα
Diffractometer	Rigaku SmartLab
Geometry	Bragg-Brentano
2θ values (°)	2θ <sub>min</sub> = 9.94 2θ <sub>max</sub> = 120 2θ <sub>step</sub> = 0.02
Refinement	
<i>R</i> factors and goodness of fit	<i>R</i> <sub>p</sub> = 0.037 <i>R</i> <sub>wp</sub> = 0.053 <i>R</i> <sub>exp</sub> = 0.014 <i>R</i> ( <i>F</i> ) = 0.057 <i>R</i> <sub>B</sub> = 0.097 χ <sup>2</sup> = 14.516
No. of parameters	46

clusters also shared corners and formed a 3D-(Zn/Al)<sub>4</sub>S<sub>8</sub> framework (Fig. 3c), within which sodium atoms were located. The interatomic distances and bond valence sums (BVS)<sup>33</sup> of Na<sub>3</sub>ZnAlS<sub>4</sub> are listed in Table 4.

In the Zn/AlS<sub>4</sub> tetrahedra, the Zn/Al-S distances varied from 2.281 Å to 2.341 Å with an average <*d*<sub>Zn/Al-S</sub>> value of 2.308 Å. This value is lower than the average <*d*<sub>Zn/In-S</sub>> value of 2.408 Å observed for Na<sub>3</sub>ZnInS<sub>4</sub>.<sup>30</sup> This is in good agreement with the increase in the ionic radii of four coordinated aluminum and indium atoms from 0.39 Å to 0.62 Å.<sup>34</sup>

The Na1 and Na2 atoms occupied 32g Wyckoff positions, and both atoms were octahedrally coordinated to the sulfur atoms. Furthermore, the Na1S<sub>6</sub> octahedra were more regular in shape than the Na2S<sub>6</sub> octahedra, even though the average <*d*<sub>Na1-S</sub>> and <*d*<sub>Na2-S</sub>> distances of 3.003 and 3.035 Å, respectively, were similar (Table 4 and Fig. 3d). These values are similar to the average <*d*<sub>Na-S</sub>> of 3.029 and 3.074 Å observed in the Na<sub>3</sub>ZnInS<sub>4</sub> compound.<sup>30</sup> In the three Na<sub>3</sub>ZnMS<sub>4</sub> compounds (M = Al, Ga, In), the first sodium atom exhibited larger displacement parameters than the second

**Table 3** Anisotropic displacement parameters (Å<sup>2</sup>) of Na<sub>3</sub>ZnAlS<sub>4</sub>

Atom	<i>U</i> <sub>11</sub>	<i>U</i> <sub>22</sub>	<i>U</i> <sub>33</sub>	<i>U</i> <sub>12</sub>	<i>U</i> <sub>13</sub>	<i>U</i> <sub>23</sub>
Na2	0.081(5)	0.075(5)	0.105(5)	-0.006(3)	-0.057(4)	0.012(4)

sodium atom (*i.e.*, *U*<sub>eq</sub> = 0.0366 Å<sup>2</sup> and *U*<sub>eq</sub> = 0.088 Å<sup>2</sup> in Na<sub>3</sub>ZnGaS<sub>4</sub> and *U*<sub>eq</sub> = 0.0453 Å<sup>2</sup> and *U*<sub>eq</sub> = 0.1213 Å<sup>2</sup> in Na<sub>3</sub>ZnInS<sub>4</sub>). This indicates that the first sodium atom is more mobile than the second one. This was also suggested by Hwang *et al.* in their study on Ca-substituted Na<sub>3</sub>ZnGaS<sub>4</sub> compounds.<sup>28</sup> BVS calculations yielded values of 0.90, 1.11, 2.23, and 3.03 for Na1, Na2, Zn1, and Al1, respectively. Only the Zn atom was slightly over bonded. The crystal density of Na<sub>3</sub>ZnAlS<sub>4</sub> was calculated to be 2.49 g cm<sup>-3</sup>.

Fig. S1† shows the SEM-EDS images of Na<sub>3</sub>ZnAlS<sub>4</sub> electrolyte. On the EDS mappings, the constituent elements Na, S, Zn and Al were almost uniformly distributed.

To increase the ionic conductivity of Na<sub>3</sub>ZnAlS<sub>4</sub>, Na vacancies were created by partially replacing Zn with Al. This led to a new composition, Na<sub>3-x</sub>Zn<sub>1-x</sub>Al<sub>1+x</sub>S<sub>4</sub> (*x* = 0.1, 0.15, and 0.2). The XRPD patterns of the compounds are shown in Fig. S2.† All the patterns appear similar, confirming the formation of new phases. Furthermore, when *x* increased, the diffraction peak 440 at 2θ = 39.5° shifted to higher angles, whereas the peak 004 at 2θ = 19.1° shifted to lower angles. This indicates that the tetragonal cell parameters *a* and *b* decrease, while *c* increases with *x*. Furthermore, full pattern-matching analyses confirmed these observations. The evolution of the cell parameters as a function of *x* is shown in Fig. 4 and Table S1.†

For the Na<sub>3-x</sub>Zn<sub>1-x</sub>Al<sub>1+x</sub>S<sub>4</sub> (*x* = 0, 0.1, 0.15, and 0.2) compounds, the lowest and highest ionic conductivities at 25 °C were measured for the compositions *x* = 0 and 0.1, respectively. Therefore, for clarity, only the temperature dependences of the ionic conductivities of *x* = 0 and 0.1 are shown in Fig. 5. The Nyquist plot of Na<sub>2.9</sub>Zn<sub>0.9</sub>Al<sub>1.1</sub>S<sub>4</sub> (*x* = 0.1) at 29.8 °C is shown in Fig. S3.† The semicircle in the high-frequency region and spike in the low-frequency region indicate that Na<sub>2.9</sub>Zn<sub>0.9</sub>Al<sub>1.1</sub>S<sub>4</sub> is a typical ionic conducting material. Bulk and grain boundary contributions were not separated; thus, the ionic conductivities were calculated from the total resistance *R*, and this indicates that the Na<sub>3-x</sub>Zn<sub>1-x</sub>Al<sub>1+x</sub>S<sub>4</sub> electrolytes possess excellent formability, which enables them to exhibit high ionic conductivity even when pressed at room temperature. The temperature dependence

**Table 2** Atomic coordinates and isotropic displacement parameters (Å<sup>2</sup>) of Na<sub>3</sub>ZnAlS<sub>4</sub>

Atom	Wyck.	Occ.	<i>x</i>	<i>y</i>	<i>z</i>	<i>U</i> <sub>iso/eq</sub>
Na1	32g	1	0.1143(3)	0.6357(3)	0.87500	0.042(2)
Na2	32g	1	0.1333(3)	0.9131(3)	0.9845(2)	0.087(3)
Zn1	32g	0.5	0.12142(13)	0.67045(14)	0.06233(12)	0.0168(7)
Al1	32g	0.5	0.12142(13)	0.67045(14)	0.06233(12)	0.0168(7)
S1	32g	1	0.25000	0.5819(3)	0.00000	0.0240(13)
S2	32g	1	0.04505(19)	0.5403(2)	0.12609(18)	0.0244(11)
S3	32g	1	0.00000	0.75000	0.98698(18)	0.0139(12)



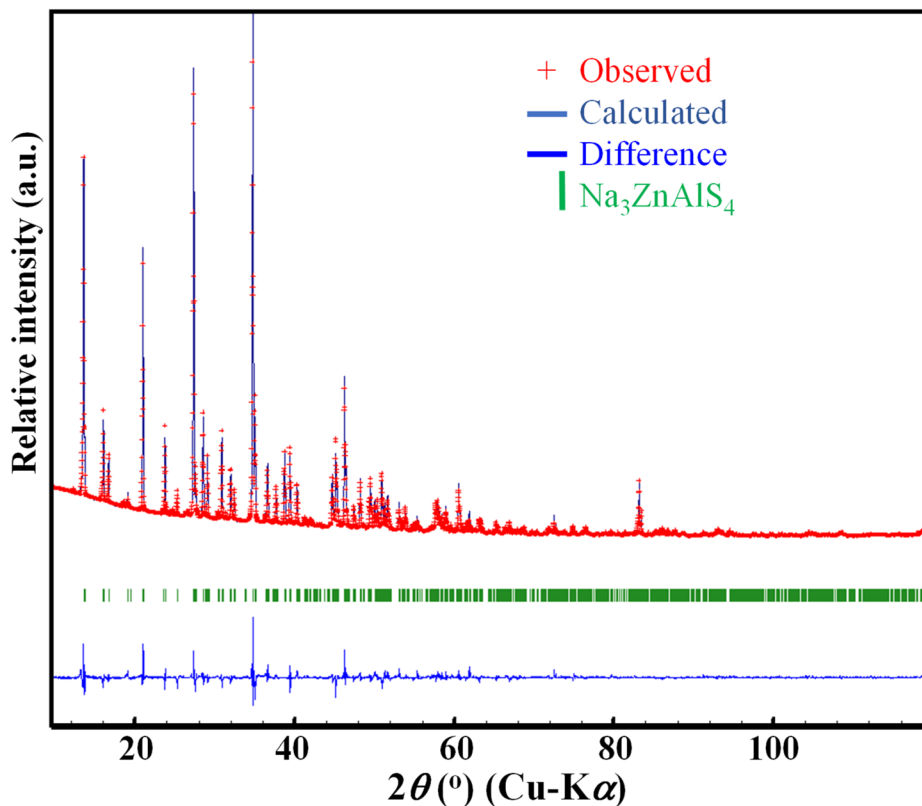


Fig. 2 Observed (cross), calculated (solid line), and difference (bottom) XRPD patterns (Cu-K $\alpha$  radiation) resulting from Rietveld refinement for Na<sub>3</sub>ZnAlS<sub>4</sub>. The impurity peaks were excluded.

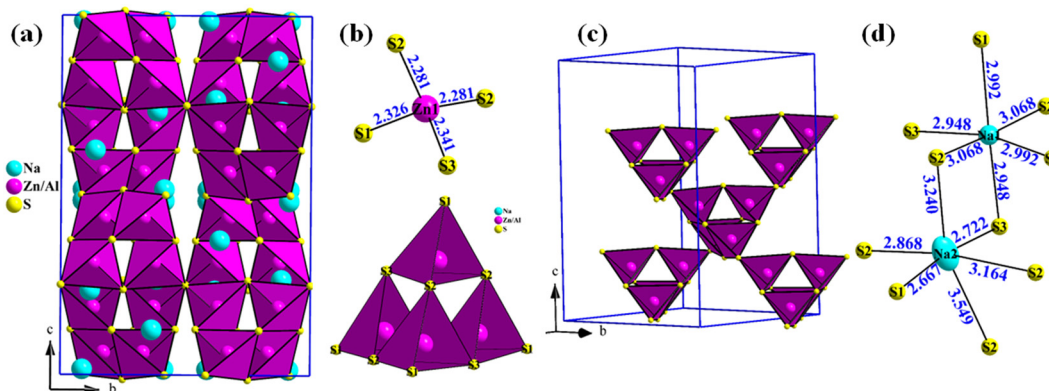


Fig. 3 (a) Projection view of the crystal structure of Na<sub>3</sub>ZnAlS<sub>4</sub> along [100], (b) (Zn/Al)<sub>4</sub>S<sub>10</sub> cluster, (c) (Zn/Al)<sub>4</sub>S<sub>8</sub> framework, and (d) coordination spheres of Na1 and Na2 atoms.

of the ionic conductivities obeyed Arrhenius' law; therefore, the activation energies were obtained from the linear fit of  $\log(\sigma)$  vs.  $1000/T$ . The ionic conductivities at 25 °C and activation energies are summarized in Table 5.

At 25 °C, the Na<sub>3</sub>ZnAlS<sub>4</sub> sample exhibited an ionic conductivity of  $3.8 \times 10^{-7}$  S cm<sup>-1</sup> ( $E_a = 37$  kJ mol<sup>-1</sup>), whereas Na<sub>2.9</sub>Zn<sub>0.9</sub>Al<sub>1.1</sub>S<sub>4</sub> exhibited an ionic conductivity of  $4.5 \times 10^{-6}$  S cm<sup>-1</sup> ( $E_a = 32$  kJ mol<sup>-1</sup>), which is one order of magnitude higher than that of Na<sub>3</sub>ZnAlS<sub>4</sub>. This was due to the

introduction of Na vacancies, which enhanced the ionic conductivity. A similar phenomenon was observed in Na<sub>3-x</sub>Zn<sub>1-x</sub>Ga<sub>1+x</sub>S<sub>4</sub>-based electrolytes. In a previous study, it was shown that vacancies were introduced at the Na2 sites, which had large ADPs, and the conductivity increased through Na2–Na2 pathways.<sup>27,28</sup>

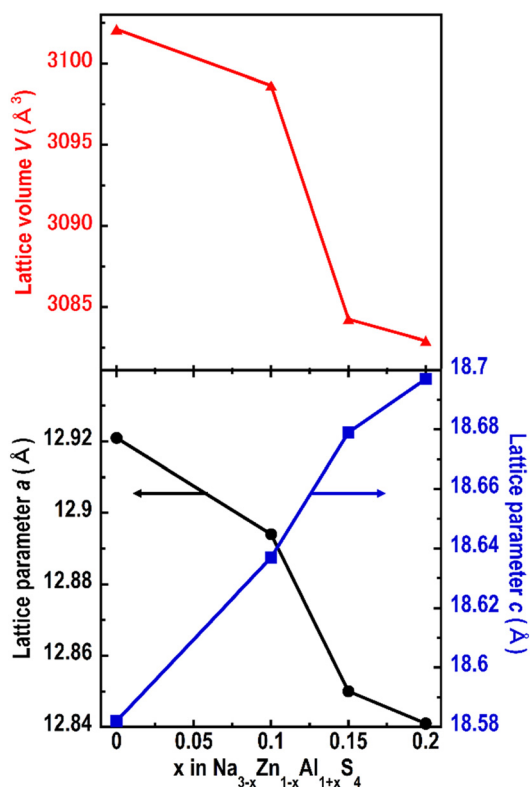
Fig. 6 and 7 show the XRPD patterns and Raman spectra of Na<sub>2.9</sub>Zn<sub>0.9</sub>Al<sub>1.1</sub>S<sub>4</sub> before and after exposure to humid air (relative humidity: 70%) for 30 min. The



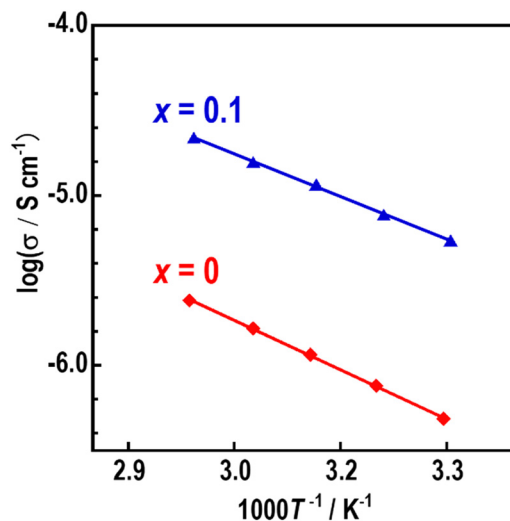
**Table 4** Interatomic distances (Å) and BVS for Na<sub>3</sub>ZnAlS<sub>4</sub>

	Distance
Na1-S3 (x2)	2.948(3)
Na1-S1 (x2)	2.992(2)
Na1-S2 (x2)	3.068(5)
	<3.003>*0.90
Na2-S1	2.667(5)
Na2-S3	2.722(4)
Na2-S2	2.868(5)
Na2-S2	3.164(5)
Na2-S2	3.240(5)
Na2-S2	3.550(5)
	<3.035>*1.11
Zn/Al1-S2	2.281(3)
Zn/Al1-S2	2.282(4)
Zn/Al1-S1	2.326(3)
Zn/Al1-S3	2.341(3)
	<2.308>
	*2.23 for Zn
	*3.08 for Al

\*Bond valence sum, B. V. =  $e^{(r_0-r)/b}$  with the following parameters:  $b = 0.37$ ,  $r_0(\text{Na}^{\text{I}}-\text{S}) = 2.3$  Å,  $r_0(\text{Zn}^{\text{II}}-\text{S}) = 2.18$  Å, and  $r_0(\text{Al}^{\text{III}}-\text{S}) = 2.13$  Å. The average distances are expressed as <>.

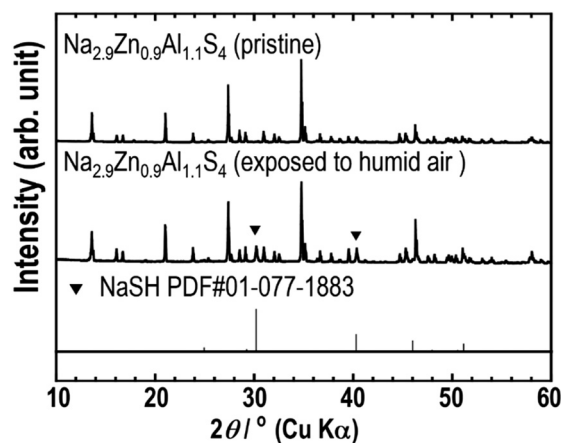
**Fig. 4** Evolution of the cell parameters as a function of  $x$  in Na<sub>3-x</sub>Zn<sub>1-x</sub>Al<sub>1+x</sub>S<sub>4</sub> ( $x = 0, 0.1, 0.15$ , and  $0.2$ ).

XRPD pattern of Na<sub>2.9</sub>Zn<sub>0.9</sub>Al<sub>1.1</sub>S<sub>4</sub> exposed to humid air was almost identical to that of pristine Na<sub>2.9</sub>Zn<sub>0.9</sub>Al<sub>1.1</sub>S<sub>4</sub>. However, new impurity peaks, attributable to NaSH, appeared. This was confirmed *via* Raman spectroscopy. On the Raman spectra, the peaks between 100 and 550

**Fig. 5** Temperature dependence of the ionic conductivities ( $\sigma$ ) of Na<sub>3</sub>ZnAlS<sub>4</sub> ( $x = 0$ ) and Na<sub>2.9</sub>Zn<sub>0.9</sub>Al<sub>1.1</sub>S<sub>4</sub> ( $x = 0.1$ ).**Table 5** Ionic conductivities at 25 °C ( $\sigma_{25}$ ) and activation energies ( $E_a$ ) of Na<sub>3-x</sub>Zn<sub>1-x</sub>Al<sub>1+x</sub>S<sub>4</sub>

$x$	$\sigma_{25}$ (S cm <sup>-1</sup> )	$E_a$ (kJ mol <sup>-1</sup> )
0	$3.8 \times 10^{-7}$	37
0.1	$4.5 \times 10^{-6}$	32
0.15	$3.5 \times 10^{-6}$	32
0.2	$3.8 \times 10^{-6}$	36

cm<sup>-1</sup>, which were attributed to pristine Na<sub>2.9</sub>Zn<sub>0.9</sub>Al<sub>1.1</sub>S<sub>4</sub>, did not change after exposure to humid air, even though the peak attributed to S-H (ref. 35) appeared at around 2550 cm<sup>-1</sup>. The previous study<sup>27,28</sup> indicated that the Na<sub>3</sub>ZnGaS<sub>4</sub> compound has a high humidity stability due to its open-channel structure with a network of corner-shared super-tetrahedra. Similarly, the relatively

**Fig. 6** XRPD patterns of pristine Na<sub>2.9</sub>Zn<sub>0.9</sub>Al<sub>1.1</sub>S<sub>4</sub> and Na<sub>2.9</sub>Zn<sub>0.9</sub>Al<sub>1.1</sub>S<sub>4</sub> exposed to air.

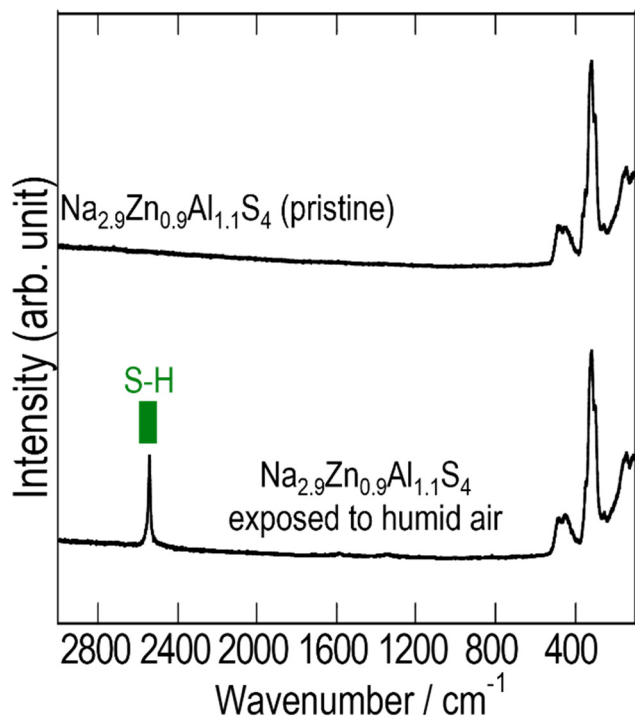


Fig. 7 Raman spectra of pristine  $\text{Na}_{2.9}\text{Zn}_{0.9}\text{Al}_{1.1}\text{S}_4$  and  $\text{Na}_{2.9}\text{Zn}_{0.9}\text{Al}_{1.1}\text{S}_4$  exposed to air.

high humidity stability of  $\text{Na}_{2.9}\text{Zn}_{0.9}\text{Al}_{1.1}\text{S}_4$  when exposed to humid atmosphere for a short time can be correlated with its crystal structure.

## Conclusions

The new solid electrolytes ( $\text{Na}_{3-x}\text{Zn}_{1-x}\text{Al}_{1+x}\text{S}_4$  ( $x \leq 0.2$ )) were prepared for the first time *via* a simple heat treatment using sodium polysulfides. These materials are isostructural with  $\text{Na}_3\text{ZnGaS}_4$ . As the aluminum content increased from 1 to 1.2, the cell volume decreased almost linearly, suggesting the formation of a solid solution. The sodium vacancies introduced by tuning the relative Zn/Al ratio in  $\text{Na}_{3-x}\text{Zn}_{1-x}\text{Al}_{1+x}\text{S}_4$  enhanced the ionic conductivities, and  $\text{Na}_{2.9}\text{Zn}_{0.9}\text{Al}_{1.1}\text{S}_4$  exhibited the highest conductivity of  $4.5 \times 10^{-6} \text{ S cm}^{-1}$  at 25 °C. Moisture stability tests showed that  $\text{Na}_{2.9}\text{Zn}_{0.9}\text{Al}_{1.1}\text{S}_4$  was relatively stable when exposed to a humid atmosphere.

## Data availability

The data supporting this article have been included as part of the ESI.†

## Conflicts of interest

There are no conflicts of interest to declare.

## Acknowledgements

This work was supported by JSPS KAKENHI Grant Number JP23H02071 and the 2023 Osaka Metropolitan University (OMU) Strategic Research Promotion Project (Priority Research).

## References

- B. L. Ellis and L. F. Nazar, *Curr. Opin. Solid State Mater. Sci.*, 2012, **16**, 168–177.
- H. Pan, Y.-S. Hu and L. Chen, *Energy Environ. Sci.*, 2013, **6**, 2338.
- N. Yabuuchi, K. Kubota, M. Dahbi and S. Komaba, *Chem. Rev.*, 2014, **114**, 11636–11682.
- H.-L. Yang, B.-W. Zhang, K. Konstantinov, Y.-X. Wang, H.-K. Liu and S.-X. Dou, *Adv. Energy Sustainability Res.*, 2021, **2**, 2000057, DOI: [10.1002/aesr.202000057](https://doi.org/10.1002/aesr.202000057).
- A. Hayashi, *Electrochemistry*, 2023, **91**(10), 101002, DOI: [10.5796/electrochemistry.23-00096](https://doi.org/10.5796/electrochemistry.23-00096).
- H. Ahmad, K. T. Kubra, A. Butt, U. Nisar, F. J. Iftikhar and G. Ali, *J. Power Sources*, 2023, **581**, 233518.
- A. Sakuda, A. Hayashi and M. Tatsumisago, *Sci. Rep.*, 2013, **3**, 2261.
- A. Hayashi, K. Noi, A. Sakuda and M. Tatsumisago, *Nat. Commun.*, 2012, **3**, 856.
- M. Duchardt, U. Ruschewitz, S. Adams, S. Dehnen and B. Roling, *Angew. Chem.*, 2018, **130**, 1365–1369.
- Z. Zhang, E. Ramos, F. Lalère, A. Assoud, K. Kaup, P. Hartman and L. F. Nazar, *Energy Environ. Sci.*, 2018, **11**, 87–93.
- F. Tsuji, K. L. Hoh, K. H. Kim, A. Sakuda, M. Tatsumisago, S. W. Martin and A. Hayashi, *J. Ceram. Soc. Jpn.*, 2021, **129**, 323–328.
- A. Banerjee, K. H. Park, J. W. Heo, Y. J. Nam, C. K. Moon, S. M. Oh, S. T. Hong and Y. S. Jung, *Angew. Chem.*, 2016, **128**, 9786–9790.
- H. Wang, Y. Chen, Z. D. Hood, G. Sahu, A. S. Pandian, J. K. Keum, K. An and C. Liang, *Angew. Chem.*, 2016, **128**, 8693–8697.
- L. Zhang, D. Zhang, K. Yang, X. Yan, L. Wang, J. Mi, B. Xu and Y. Li, *Adv. Sci.*, 2016, **3**, 1600089, DOI: [10.1002/advs.201600089](https://doi.org/10.1002/advs.201600089).
- F. Tsuji, S. Yubuchi, A. Sakuda, M. Tatsumisago and A. Hayashi, *J. Ceram. Soc. Jpn.*, 2020, **128**, 641–647.
- F. Tsuji, N. Masuzawa, A. Sakuda, M. Tatsumisago and A. Hayashi, *ACS Appl. Energy Mater.*, 2020, **3**, 11706–11712.
- A. Nasu, T. Otono, T. Takayanagi, M. Deguchi, A. Sakuda, M. Tatsumisago and A. Hayashi, *Energy Storage Mater.*, 2024, **67**, 103307.
- T. A. Yersak, Y. Zhang, F. Hao and M. Cai, *Front. Energy Res.*, 2022, **10**, 882508, DOI: [10.3389/fenrg.2022.882508](https://doi.org/10.3389/fenrg.2022.882508).
- A. Fukushima, A. Hayashi, H. Yamamura and M. Tatsumisago, *Solid State Ionics*, 2017, **304**, 85–89.
- G. Sahu, Z. Lin, J. Li, Z. Liu, N. Dudney and C. Liang, *Energy Environ. Sci.*, 2014, **7**, 1053–1058.



- 21 T. Kimura, A. Kato, C. Hotehama, A. Sakuda, A. Hayashi and M. Tatsumisago, *Solid State Ionics*, 2019, **333**, 45–49.
- 22 K. Kanazawa, S. Yubuchi, C. Hotehama, M. Otoyama, S. Shimono, H. Ishibashi, Y. Kubota, A. Sakuda, A. Hayashi and M. Tatsumisago, *Inorg. Chem.*, 2018, **57**, 9925–9930.
- 23 H. Tang, Z. Deng, Z. Lin, Z. Wang, I.-H. Chu, C. Chen, Z. Zhu, C. Zheng and S. P. Ong, *Chem. Mater.*, 2018, **30**, 163–173.
- 24 T. Takayanagi, A. Nasu, F. Tsuji, K. Motohashi, A. Sakuda, M. Tatsumisago and A. Hayashi, *J. Ceram. Soc. Jpn.*, 2022, **130**, 22017.
- 25 H. Gamo, N. H. H. Phuc, H. Muto and A. Matsuda, *ACS Appl. Energy Mater.*, 2021, **4**, 6125–6134.
- 26 S. Balijapelly, Q. Zhang, P. Sandineni, A. Adhikary, S. Mohapatra, S. Sundaramoorthy, N. Gerasimchuck, A. V. Chernatynskiy and A. Choudhury, *ACS Appl. Energy Mater.*, 2021, **4**, 7942–7951.
- 27 S. Han, J. Y. Seo, W. B. Park, A. B. Ikhe, S. Y. Choi, S. C. Han, K.-S. Sohn and M. Pyo, *J. Mater. Chem. A*, 2022, **10**, 25039–25046.
- 28 E. H. Hwang, J. Y. Seo, W. B. Park, S. Y. Kang, K.-S. Sohn and M. Pyo, *J. Power Sources*, 2023, **581**, 233511.
- 29 J. Y. Seo, W.-B. Park, S. Y. Kang, Y.-K. Lee, K.-S. Sohn and M. Pyo, *Energy Storage Mater.*, 2024, **65**, 103123.
- 30 R. Chen, X. Wu and Z. Su, *Dalton Trans.*, 2018, **47**, 15538–15544.
- 31 V. Petříček, M. Dušek and L. Palatinus, *Z. Kristallogr. - Cryst. Mater.*, 2014, **229**, 345–352.
- 32 S. J. Clarke and F. J. Disalvo, *J. Alloys Compd.*, 1998, **274**, 118–121.
- 33 I. D. Brown and D. Altermatt, *Acta Crystallogr., Sect. B: Struct. Sci.*, 1985, **41**, 244–247.
- 34 R. D. Shannon, *Acta Crystallogr., Sect. A: Cryst. Phys., Diffraction, Theor. Gen. Crystallogr.*, 1976, **32**, 751–767.
- 35 P. Bazylewski, R. Divigalpitiya and G. Fanchini, *RSC Adv.*, 2017, **7**, 2964–2970.

