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# **Fe-P-S electrodes for all-solid-state lithium secondary batteries using sulfide-based solid electrolytes**

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## Abstract

Although lithium-sulfur batteries are expected to be the next-generation high-capacity battery of choices, the sulfur electrodes studied so far contain large amounts (typically 30–60 wt%) of carbon additives, resulting in low energy density. In this study, a Fe-P-S-based material is prepared and characterized for the use as a sulfur electrode with a low content of carbon additives. The electrode material based on Fe-P-S is synthesized by the mechanical milling of FePS<sub>3</sub> and elemental sulfur. The ball-milled 70FePS<sub>3</sub>·30S (wt%) electrode comprises FeS<sub>2</sub> particles of ~30 nm in size and amorphous P-S. The all-solid-state battery operated at 100 °C exhibits a reversible capacity of more than 625 mAh g<sup>-1</sup> for 50 cycles at 0.51 mA cm<sup>-2</sup> (~0.1 C) in spite of the low carbon content (2 wt%) in the electrode. During the discharge-charge cycle, the sulfur component is confirmed to be both reduced and oxidized, cyclically.

KEYWORDS: Lithium-sulfur battery; All-solid-state battery; Fe-based composite cathode material; High-capacity electrode

## Introduction

Lithium-ion secondary batteries have been used as power sources for portable electric devices because of their high operating voltage, high energy density, and good cycleability [1–3]. Further increase in the energy density of the batteries can be achieved by high-voltage and/or high-capacity electrodes. Elemental sulfur has attracted much attention as a high-capacity cathode material for lithium secondary batteries [4–6]. Although sulfur as a cathode has a low discharge potential ( $\sim 2.1$  V vs. Li), the theoretical capacity is  $1672 \text{ mAh g}^{-1}$ , which is approximately ten times higher than the capacity of  $\text{LiCoO}_2$ . Thus, the energy density of the lithium batteries can be increased significantly by using a high-capacity electrode composed of sulfur. One of the disadvantages of the sulfur active material is its insulating nature [7]. To form sufficient electron conducting paths, large amount of carbon additives, typically 30-60 wt% [8,9], must be mixed with the sulfur of the electrode. The another disadvantage of the sulfur electrode active material is the rapid degradation in the battery when using liquid electrolytes due to dissolution of lithium polysulfide formed in the battery operation into the liquid electrolytes.

All-solid-state lithium secondary batteries using sulfur electrodes and non-flammable inorganic solid electrolytes have the advantage of high safety [10]. Needless to say,

lithium polysulfides do not dissolve into solid electrolytes, leading to long cycle life of the battery with the sulfur electrode. However, in all-solid-state batteries, the formation of sufficient lithium-ion paths as well as a large contact area between electrodes and electrolytes is difficult to achieve. To form the sufficient paths and a large contact area, solid electrolytes must be mixed with the sulfur electrode to form composite electrodes. In the same way, electron conduction paths are also formed by mixing carbon additives with the sulfur electrode. Since all the components in the all-solid-state batteries are solids, it is not easy to create enough conduction paths so that the all-solid-state battery can only be operated at or below a moderate discharge-charge rate (for  $\sim 0.1$  C). For example, all-solid-state batteries using sulfur composite electrodes such as Cu-S [10] and C-FeS<sub>2</sub>-S [11] were operated at only 0.02 C even though they showed a high capacity. Although it is known that the carbon content in the sulfur electrode should be as low as possible in order to increase the volumetric and gravimetric energy density of batteries, the amount of conductive carbon used in the sulfur battery is hardly below 10 wt.% due to insulating nature of sulfur. Based on the best of our knowledge, there is no literature reporting a discharge-charge rate  $> 0.1$  C of all-solid-state batteries using sulfur composite electrodes with a low carbon content less than 10 wt.%. Properties of the all-solid-state lithium-sulfur batteries using sulfur or sulfur composite cathode materials reported in previous

literatures are tabulated in Table S1. The low C-rate may be related to low ionic and electronic conductivities in the sulfur electrode. Interestingly, these conductivities can be enhanced by increasing the temperature. High-temperature operation, which is the advantage of a solid electrolyte over a liquid electrolyte because of no evaporation nature of the solid electrolyte, would allow an all-solid-state battery to exhibit a discharge-charge profile with a reasonable discharge and charge rate.

Metal sulfides (e.g.  $\text{TiS}_2$  [12],  $\text{MnS}$  [13],  $\text{FeS}$  [14],  $\text{FePS}_3$  [15,16],  $\text{NiS}$  [17],  $\text{NiPS}_3$  [15,18],  $\text{CuS}$  [19], and  $\text{MoS}_2$  [20]) have been investigated as electrode active materials for lithium secondary batteries. Among them, metal sulfides such as  $\text{TiS}_2$  and  $\text{FePS}_3$ , which have layered structures, exhibit the lithium-ion insertion/extraction reaction [21]. The electronic conductivities of  $\text{TiS}_2$  and  $\text{FePS}_3$  are  $1.2 \times 10^2$  and  $\sim 10^{-5}$   $\text{S cm}^{-1}$ , respectively [22,23]. The theoretical capacities of  $\text{TiS}_2$  and  $\text{FePS}_3$  are 239 and 220  $\text{mAh g}^{-1}$ , respectively, which are lower than those of other metal sulfides showing conversion reactions. To increase the capacity, we focused on mixing the metal sulfide with sulfur electrode active material. The sulfur composite electrode which contains the metal sulfide has a moderate electronic conductivity and is expected to possess electron paths without the addition of large amounts of carbon additives. The large volume change of the sulfur electrode, which is responsible for the degradation for these types of batteries, can be

reduced by the inclusion of metal sulfides because of the smaller volume change of the metal sulfides. Moreover, stable electrode-electrolyte interfaces are expected to be formed even at high temperatures because of the similarity of the composition of sulfur in the sulfur composite electrodes and sulfide-based solid electrolytes. For the S-based batteries using liquid electrolytes, some composites such as  $\text{Li}_2\text{S}-M$  ( $M = \text{Co}, \text{Fe}$ ) [24],  $\text{Li}_2\text{S}-\text{MS}_2$  ( $M = \text{Ti}, \text{V}, \text{Zr}, \text{Mo}$ ) [25,26],  $\text{Li}_2\text{S}-\text{FeS}_x$  [27], and  $\text{Li}_2\text{S}-\text{FePS}_3$  [28] have been developed as cathode materials. Transition-metal polysulfides (*e.g.*  $\text{MS}_3$  ( $M = \text{Ti}$  [29,30],  $\text{Nb}$  [31],  $\text{Mo}$  [32,33]),  $\text{Li}_2\text{TiS}_3$  [34],  $\text{MoS}_{3.4}$  [35],  $\text{MS}_4$  ( $M = \text{Ti}$  [36],  $\text{V}$  [37],  $\text{Nb}$  [31]),  $\text{Li}_2\text{WS}_4$  [38],  $\text{Li}_3\text{NbS}_4$  [34],  $M_3\text{S}_4$  ( $M = \text{Fe}$  [39],  $\text{Ni}$  [40]),  $\text{NbS}_5$  [31], and  $\text{MoS}_{5.7}$  [41]) have attracted much attention as another family of cathode materials possessing high theoretical capacity based on the redox reaction of sulfur and/or transition metals.

In addition to studies on these composite cathode materials, the composite cathode materials that are suitable for all-solid-state batteries such as  $\text{Li}_2\text{S}-\text{Cu}$  [42],  $\text{Li}_2\text{S}-\text{P}_2\text{S}_5-\text{Cu}$  [43], and  $\text{C}-\text{FeS}_2-\text{S}$  [11] have also been studied. Considering the more abundant element in the earth's crust, an iron-based metal sulfide would be economical to be used as the additive for a sulfur electrode. However, in the all-solid-state batteries, few composite cathode materials consisting of sulfur and an iron-based metal sulfide have been investigated so far [11]. Moreover, there are not many reports on the operation of the all-

solid-state batteries using sulfide-based solid electrolytes at a high temperature (100 °C) [44–47], even though the evaluation of the battery performance at 100 °C is useful for the practical all-solid-state lithium-sulfur batteries in terms of the faster lithium dissolution-deposition process at the high temperature than a room temperature.

In the present study, we developed new electrode systems for all-solid-state batteries by simple mixing an iron-based metal sulfide with sulfur, and investigated the performance of the new electrode materials in batteries at 100 °C. Among iron-based metal sulfides, we focused on FePS<sub>3</sub> containing a P-S unit, which has been reported as an electrode active material [48,49]. Since the addition of P<sub>2</sub>S<sub>5</sub> to a sulfur electrode has been reported to improve the performance of batteries [50], the mixture of FePS<sub>3</sub> with the sulfur electrode in our batteries is expected to obtain superior performance.

In this study, new Fe-P-S electrodes were synthesized by mechanical milling of FePS<sub>3</sub> and S, and the electrochemical performance of the Fe-P-S electrodes with a liquid or solid electrolyte was investigated.

## **Experimental**

Fe-P-S electrodes were synthesized by mechanical milling of FePS<sub>3</sub> and sulfur (Kanto Chemical, 99.5%). FePS<sub>3</sub> was synthesized by heating a mixture of iron powder (Wako

Chemical, 99.9%), red phosphorus (Kanto Chemical, 98.0%), and sulfur (Kanto Chemical, 99.5%) [16,48]. The synthesized FePS<sub>3</sub> and elemental sulfur were mixed using an agate mortar with weight ratio of 100:0, 70:30, and 50:50, respectively. Then, the mixtures were mechanically milled using a planetary ball mill (Fritsch, Pulverisette 7) with a zirconia pot (45 mL volume) and 500 zirconia balls ( $\phi = 4$  mm) in a dry Ar atmosphere at 510 rpm for 24 h. To identify the crystalline structures of the resultant powder of 70FePS<sub>3</sub>·30S electrode, an X-ray diffraction (XRD) pattern was recorded using an X-ray diffractometer (Miniflex 600, Rigaku) under Cu-K $\alpha$  radiation. For this measurement, a sample stage was covered with Kapton film to avoid undesired reaction of the samples with air. A Raman spectrum of the milled powder of 70FePS<sub>3</sub>·30S electrode was measured using a Raman spectrometer (XploRA, Horiba) with a green laser (Wavelength: 532 nm) to identify the structural units. For the Raman measurement, the sample was sealed in an evacuated capillary glass tube. The morphology and elemental mapping were studied using focused ion beam–scanning electron microscopy (FIB-SEM; JIB-4600, JEOL) and scanning transmission electron microscopy (STEM; HD-2000, Hitachi) in conjunction with an energy dispersive spectroscopy (EDS) system. The stability of the 70FePS<sub>3</sub>·30S electrode at high temperature was evaluated by thermogravimetric and differential thermal analysis (TG-DTA; Thermo Plus TG 8120, Rigaku). The electronic conductive of the 70FePS<sub>3</sub>·30S

electrode was measured using a symmetric stainless steel/70FePS<sub>3</sub>·30S/stainless steel cell.

All-solid-state batteries (Li-In/75Li<sub>2</sub>S·25P<sub>2</sub>S<sub>5</sub> (mol%) glass/Fe-P-S) were assembled as described in a previous paper [51]. 75Li<sub>2</sub>S·25P<sub>2</sub>S<sub>5</sub> (mol%) glass solid electrolytes were prepared by mechanical milling of Li<sub>2</sub>S (Mitsuwa Chemical, 99.9%) and P<sub>2</sub>S<sub>5</sub> (Aldrich, 99%) [52]. The Li<sub>2</sub>S and P<sub>2</sub>S<sub>5</sub> powders were mixed together using an agate mortar, and the mixture was placed into ZrO<sub>2</sub> pots (45 mL) with 500 ZrO<sub>2</sub> balls ( $\phi = 4$  mm). The pots were set in a planetary ball mill apparatus (Fritsch, Pulverisette 7), and mechanical milling was performed at 510 rpm for 24 h. Cathode composite electrodes were prepared by mixing the Fe-P-S electrode samples, 75Li<sub>2</sub>S·25P<sub>2</sub>S<sub>5</sub> (mol%) glass, and vapor-grown carbon fibers (VGCF, Showa Denko) in a weight ratio of 69:29:2 (wt%). To measure the electronic conductivity of the composite electrode, a symmetric stainless steel/composite electrode/stainless steel cell was constructed. The composite electrodes (10 mg) and the 75Li<sub>2</sub>S·25P<sub>2</sub>S<sub>5</sub> (mol%) glass solid electrolytes (120 mg) were placed into a polycarbonate tube ( $\phi = 10$  mm), and pressed under 360 MPa. A Li-In alloy foil was attached to the bilayer pellet consisting of the cathode composite electrode and the solid electrolyte layers by uniaxially pressing them together at 120 MPa. These pellets were sandwiched by two stainless-steel disks as current collectors. The assembled all-solid-state batteries were discharged and charged under a constant current density of 0.51 mA cm<sup>-2</sup> at different

temperatures (25, 60, 80, and 100 °C) using a discharge-charge measuring device (Scribner Associates, 580 battery-type system). The discharge-charge measurement was initiated with discharge. To evaluate the rate performance of the battery, the constant current densities from 0.51 to 8.15 mA cm<sup>-2</sup> were used for the discharge cycles.

To compare the performance of the all-solid-state battery with that of the battery using liquid electrolytes, the battery (Li / Fe-P-S) using liquid electrolytes was also constructed.

In a typical processing, the Fe-P-S electrode sample of 70FePS<sub>3</sub>-30S, polyvinylidene fluoride (PVDF), and Super P were mixed using an agate mortar in a weight ratio of 75:15:10 (wt%). Subsequently, 1-methyl-2-pyrrolidone (NMP) was added to the mixture to prepare a slurry. This slurry was loaded onto a copper foil and dried at 100 °C for 24 h in a vacuum. A CR2016 coin-cell battery was constructed using the dried composite cathode foil as a working electrode, porous propylene separator (Celgard 2500), 1 M LiPF<sub>6</sub> in ethylene carbonate (EC)/dimethyl carbonate (DMC)/diethyl carbonate (DEC) (1:1:1, wt/wt/wt%) as a liquid electrolyte, and Li foil as reference and counter electrodes. The weight of the 70FePS<sub>3</sub>-30S sample was 0.575 mg cm<sup>-2</sup>. The cell was discharged and charged at a constant current density of 65.6 mA g<sup>-1</sup> (~0.1 C) at room temperature (typically 25–30 °C), using a discharge-charge machine (LANHE CT2001A). The discharge-charge measurement was initiated with discharge. To examine the reaction

mechanism of the 70FePS<sub>3</sub>·30S electrode, the products of the 70FePS<sub>3</sub>·30S electrode composite electrode after the charge cycles were investigated by the XRD measurement. To reveal the redox reaction of the 70FePS<sub>3</sub>·30S electrode during discharge-charge cycles, the S *K*-edge X-ray absorption near edge structure (XANES) spectra of the cathode composite electrode in the all-solid-state batteries were measured before and after discharge/charge. XANES spectra were measured at the BL6N1 (proposal No. 201705058) of the Aichi Synchrotron Center, Aichi Science & Technology Foundation. K<sub>2</sub>SO<sub>4</sub> was used as a standard sample for XANES measurement to calibrate the spectra data. Their intensity was normalized as zero and one for the intensity at 2466.4 eV and 2489.5 eV, respectively. These processes were all performed under a dry Ar atmosphere.

## **Results and Discussion**

Figure 1 shows XRD patterns of the Fe-P-S electrode of 70FePS<sub>3</sub>·30S. For comparison, the XRD spectra of pyrite FeS<sub>2</sub>, reagent-grade sulfur, and the prepared FePS<sub>3</sub> are also shown in Figure 1. The diffraction peaks of the XRD pattern of 70FePS<sub>3</sub>·30S were attributed to pyrite FeS<sub>2</sub> (ICSD #316) without peak shift. Considering the composition of the 70FePS<sub>3</sub>·30S electrode, this result indicates that the 70FePS<sub>3</sub>·30S electrode is composed of the pyrite FeS<sub>2</sub> phase and amorphous phase(s) containing phosphorus and

sulfur.

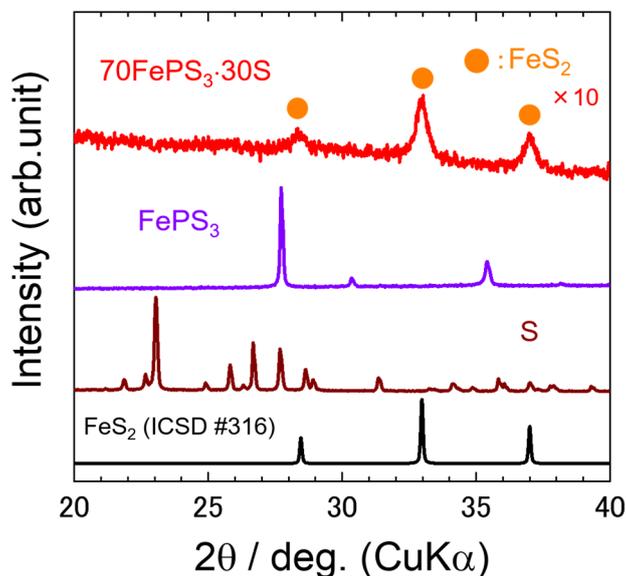


Figure 1. XRD patterns of the Fe-P-S electrode of 70FePS<sub>3</sub>·30S, FePS<sub>3</sub> before mechanical milling, reagent-grade pure sulfur and PDF file of FeS<sub>2</sub>.

Figure 2 shows the Raman spectrum of the 70FePS<sub>3</sub>·30S electrode. Raman bands present at 337 and 373 cm<sup>-1</sup> are similar to those attributed to S<sup>2-</sup> in FeS<sub>2</sub> [53]. These results are in agreement with the XRD result, in which only the FeS<sub>2</sub> pyrite phase is detected. In the Raman spectrum, a band at 412 cm<sup>-1</sup> is also noted. It has been reported that P-S bonds in P<sub>2</sub>S<sub>6</sub><sup>4-</sup>, P<sub>2</sub>S<sub>7</sub><sup>4-</sup>, and PS<sub>4</sub><sup>3-</sup> units show Raman bands at 390, 410, and 425 cm<sup>-1</sup>, respectively [54,55]. Therefore, the observed band at 412 cm<sup>-1</sup> could be attributed to the P-S bond of the P<sub>2</sub>S<sub>7</sub><sup>4-</sup> unit in amorphous phase(s) containing phosphorus and sulfur.

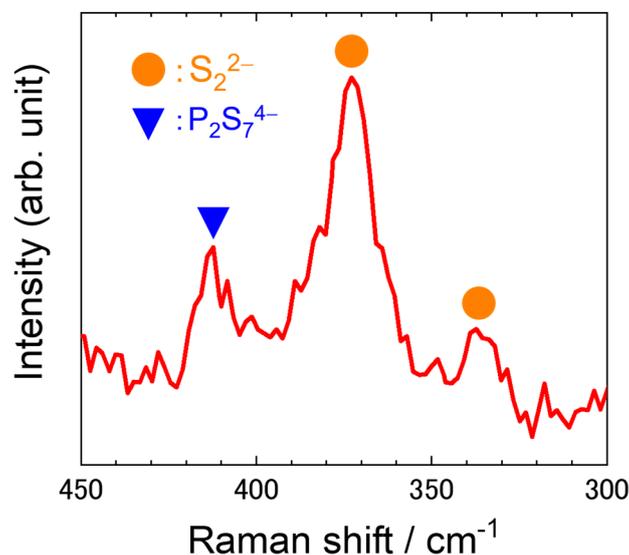


Figure 2. Raman spectrum of the 70FePS<sub>3</sub>·30S electrode.

Figure 3 (a) through (e) present an SEM and a HAADF-STEM image, and EDS mapping of Fe, P, and S of the 70FePS<sub>3</sub>·30S electrode, respectively. The particle size of the 70FePS<sub>3</sub>·30S electrode is in the range of 1–5 μm (Figure 3 (a)). The STEM image of the 70FePS<sub>3</sub>·30S electrode particle as shown in Figure 3(b) reveals bright and dark features, suggesting that the 70FePS<sub>3</sub>·30S particle is composed of two or more component phases. Indeed, iron and phosphorus in the 70FePS<sub>3</sub>·30S particle are non-uniformly distributed (Figure 3 (c) and (d)) while sulfur is uniformly distributed (Figure 3 (e)). In the mapping image of iron (Figure 3 (c)), the distribution of iron corresponds mainly to the bright regions of the HAADF-STEM image (Figure 3 (b)), with an average size of

approximately 30 nm. Considering SEM and STEM images, and the XRD results, 70FePS<sub>3</sub>·30S contains FeS<sub>2</sub> with a particle size of approximately 30 nm. However, we cannot deny the possibility of P doping into FeS<sub>2</sub>, because a very similar case of P-doped CoS<sub>2</sub>, isostructural to FeS<sub>2</sub>, has been reported [56,57]. Other components in the 70FePS<sub>3</sub>·30S particle, which correspond to darker regions in Figure 3 (b), are amorphous phase(s) composed of phosphorus and sulfur. The electronic conductivity of the 70FePS<sub>3</sub>·30S electrode was  $3 \times 10^{-7}$  S cm<sup>-1</sup> at room temperature and  $6 \times 10^{-6}$  S cm<sup>-1</sup> at 100 °C, which were much higher than that of sulfur ( $5 \times 10^{-18}$  S cm<sup>-1</sup> at 20 °C) [7]. In the TG-DTA curves (Figure S1), no significant change in weight and heat were seen below 120 °C, suggesting that the 70FePS<sub>3</sub>·30S electrode neither showed evaporation and melt.

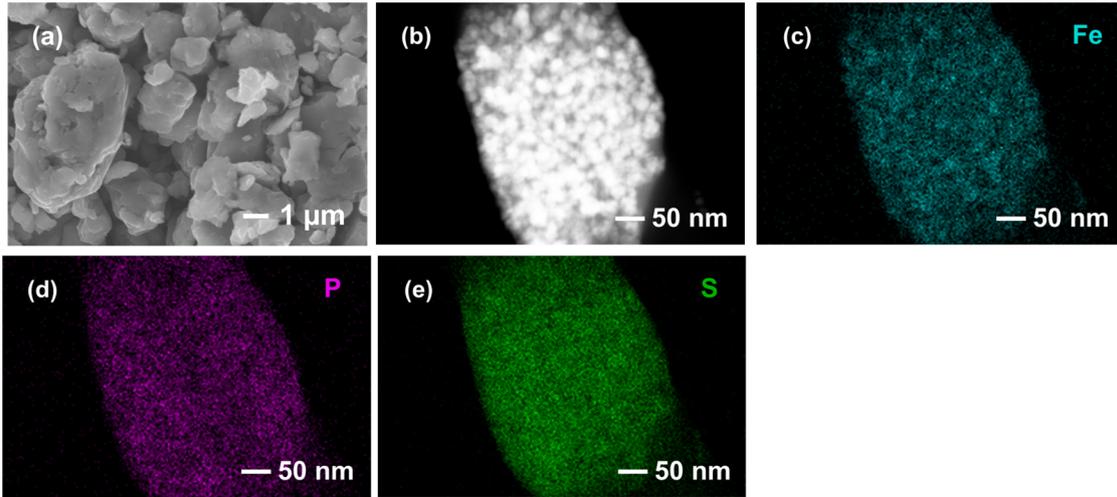


Figure 3. (a) SEM image, (b) HAADF-STEM image, and EDS mapping of (c) Fe, (d) P, and (e) S of the 70FePS<sub>3</sub>·30S electrode.

Figure 4 shows discharge-charge curves of the all-solid-state battery using the 70FePS<sub>3</sub>·30S electrode at 25 °C (Figure 4 (a)) and the battery with 70FePS<sub>3</sub>·30S and the liquid electrolytes at room temperature (typically 25–30 °C) (Figure 4 (b)). The current densities of 0.51 mA cm<sup>-2</sup> (~0.1 C) for the all-solid-state battery and of 65.6 mA g<sup>-1</sup> (~0.1 C) for the battery with liquid electrolytes were used. The all-solid-state battery exhibits discharge-charge capacity of less than 1 mAh g<sup>-1</sup> at 25 °C. The low capacity could be attributed to the low electronic conductivity of 5×10<sup>-7</sup> S cm<sup>-1</sup> of the composite electrode with Li-P-S electrolyte and 2 wt% carbon additive. In contrast, the first charge capacity of the battery using the liquid electrolyte is 414 mAh g<sup>-1</sup>. Note that the capacity of the battery with liquid electrolytes cannot be directly compared with that of the all-solid-state

battery because of the different amounts of carbon additives. More importantly, its capacity, however, degraded rapidly with repetition of the discharge-charge cycles. From the 10th cycle onwards, the discharge capacity is less than  $15 \text{ mAh g}^{-1}$  because of the dissolution of the polysulfide [58].

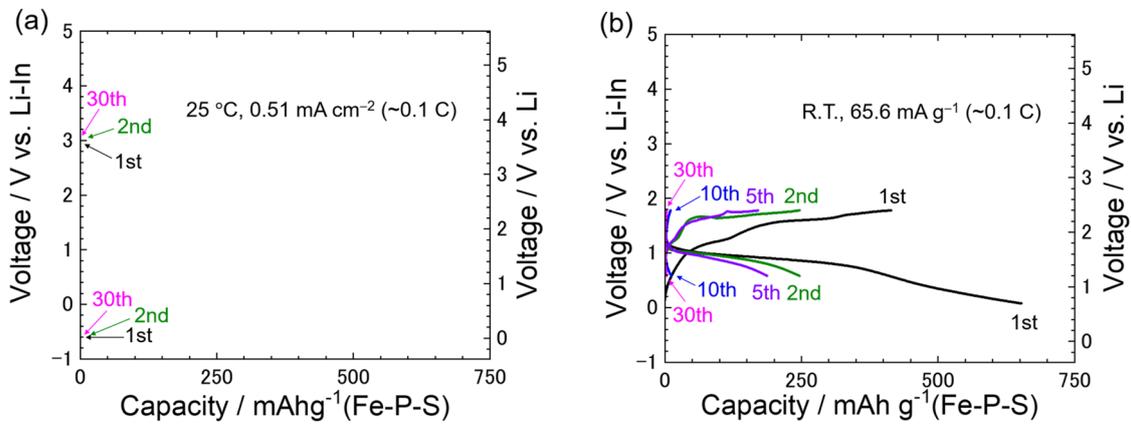


Figure 4. Discharge-charge curves of (a) all-solid-state battery using the 70FePS<sub>3</sub>-30S electrode at 25 °C and (b) battery with the 70FePS<sub>3</sub>-30S and liquid electrolytes at room temperature (typically 25–30 °C). In the all-solid-state battery, the cut-off voltage was set to  $-0.62 \text{ V vs. Li-In}$  for discharging and  $3.0 \text{ V vs. Li-In}$  for charging. On the other hand, the battery with liquid electrolytes was firstly discharge to a capacity of  $656 \text{ mAh g}^{-1}$ . From the first charge cycle onwards, the cut-off voltage was set to  $1.78 \text{ V vs. Li-In}$  for charging and  $0.58 \text{ V vs. Li-In}$  for discharging.

In order to achieve large discharge-charge capacity at a reasonable rate, we examined

the discharge-charge performance of the all-solid-state battery at 100 °C. In the preliminary experiment of the discharge-charge measurements at different temperatures (25, 60, 80, and 100 °C), shown in Figure S2, the all-solid-state battery heated at 100 °C exhibits the high reversible capacity of 625 mAh g<sup>-1</sup> for the first cycle. Figure 5 (a) shows discharge-charge curves of the all-solid-state battery with the 70FePS<sub>3</sub>·30S electrode at 100 °C, under a current density of 0.51 mA cm<sup>-2</sup> (~0.1 C). 1 C-rate was defined as 5.8 mA cm<sup>-2</sup> required to (dis)charge the battery in 1 hour. In the discharge-charge measurement, the battery is firstly discharged with a capacity of 656 mAh g<sup>-1</sup><sub>(Fe-P-S)</sub>. Subsequently, the battery is charged to 3.0 V vs. Li-In. The capacity of 656 mAh g<sup>-1</sup><sub>(Fe-P-S)</sub> is calculated by using the theoretical capacities of FePS<sub>3</sub> (220 mAh g<sup>-1</sup>) and sulfur (1672 mAh g<sup>-1</sup>) with the weight ratio of 70:30, but the theoretical capacity of the 70FePS<sub>3</sub>·30S electrode is different to this capacity (656 mAh g<sup>-1</sup>). The theoretical capacity can be estimated based on final products of FeS<sub>2</sub> and amorphous P-S. If 70 wt% FePS<sub>3</sub> and 30 wt% sulfur (1 mol FePS<sub>3</sub> and 2.4 mol sulfur) in the original composite react completely by mechanical milling process, the weight ratio of FeS<sub>2</sub> and amorphous P-S can be estimated to be 46:54 (wt%) based on this reaction. The theoretical capacity of FeS<sub>2</sub> is 894 mAh g<sup>-1</sup>. In regarding to the amorphous P-S, the composition of P and S can be estimated to be 1:3.4 (mol) based on the reaction of 70 wt% FePS<sub>3</sub> with 30 wt% sulfur.

If the amorphous  $\text{PS}_{3.4}$  reacts with Li completely and  $\text{Li}_3\text{P}$  and  $\text{Li}_2\text{S}$  are formed, the theoretical capacity can be calculated to be  $1876 \text{ mAh g}^{-1}$ . By using the estimated weight ratio (46:54 (wt%)) and these theoretical capacities ( $\text{FeS}_2$ :  $894 \text{ mAh g}^{-1}$  and  $\text{PS}_{3.4}$ :  $1876 \text{ mAh g}^{-1}$ ), the theoretical capacity of the  $70\text{FePS}_3\cdot 30\text{S}$  electrode would be  $1423 \text{ mAh g}^{-1}_{(\text{Fe-P-S})}$ . The battery shows stable reversible discharge-charge behavior for 50 cycles in spite of the low content of the carbon additive (2 wt%) in the  $70\text{FePS}_3\cdot 30\text{S}$  composite electrode.

During the first discharge curve, three plateaus around at 0.9, 1.7, and 1.8 V vs. Li-In are observed. Sulfur has been reported to show two plateaus [59,60]. The plateau at  $\sim 2.3$  V vs. Li is attributed to the reduction of sulfur to  $\text{Li}_2\text{S}_n$  ( $n > 4$ ). The another plateau at  $\sim 2.1$  V vs. Li corresponds to the further reduction of  $\text{Li}_2\text{S}_n$  to  $\text{Li}_2\text{S}_2$  or  $\text{Li}_2\text{S}$ . The observed plateaus at around 1.7 and 1.8 V vs. Li-In in a range of capacities from 0 to  $\sim 280 \text{ mAh g}^{-1}$  can be attributed to the redox reaction of sulfur in amorphous phase(s) containing phosphorus and sulfur. On the other hand, the plateaus at  $\sim 1.0$  V vs. Li-In in a range of capacities from  $\sim 280$  to  $\sim 660 \text{ mAh g}^{-1}$  can correspond to the redox reactions of  $\text{S}_2^{2-}$  (Eq. 1) in  $\text{FeS}_2$  because  $\text{FeS}_2$  exhibits a discharge plateau at around 1.5 V vs. Li, corresponding to around 0.9 V vs. Li-In, in the redox reactions of  $\text{S}_2^{2-}$ , as shown in the following equation [61,62].



After the first cycle, the discharge capacity corresponding to the redox reaction of sulfur in the amorphous phase(s) is around 280 mAh g<sup>-1</sup>. The sulfur utilization based on the additional amount of sulfur is 56%. A possible reason for the moderate sulfur utilization is that the electron path in the composite electrode is still insufficient to increase the sulfur utilization drastically; the electronic conductivity of the composite electrode was 1×10<sup>-5</sup> S cm<sup>-1</sup> at 100 °C.

On the other hand, the discharge capacity attributable to the redox reactions of S<sub>2</sub><sup>2-</sup> in FeS<sub>2</sub> is around 380 mAh g<sup>-1</sup>. The utilization based on the amount of FeS<sub>2</sub> is 92%. From the 1st to 10th cycle, the discharge profile changed gradually, and the discharge voltage in a range of capacities from ~220 to ~450 mAh g<sup>-1</sup> slightly increased. For the batteries using FeS<sub>2</sub>, the discharge voltage has been reported to increase from the second cycle onwards because of the complex reaction of FeS<sub>2</sub> forming FeS<sub>x</sub> and S partially [61,62]. In the complex reaction, FeS<sub>x</sub> and S are formed partially. In the XRD pattern of the 70FePS<sub>3</sub>·30S electrode composite electrode after the charge cycles (Figure S3), the peaks can be assigned as S, Fe<sub>7</sub>S<sub>8</sub>, Li<sub>2</sub>FeS<sub>2</sub>, Li<sub>4</sub>P<sub>2</sub>S<sub>6</sub>, and the unknown phase(s) are observed. This result for the products of S, Fe<sub>7</sub>S<sub>8</sub>, and Li<sub>2</sub>FeS<sub>2</sub> is consistent with the previously reported complex reaction of FeS<sub>2</sub> forming FeS<sub>x</sub> and S [61,62]. Li<sub>4</sub>P<sub>2</sub>S<sub>6</sub> and unknown

phase(s) could be formed by the reaction of the amorphous P-S with  $\text{Li}^+$ . From these results, the observed profile change can be attributed to not the redox reaction of sulfur in amorphous P-S but mainly the redox reaction in  $\text{FeS}_2$ .

Figure 5(b) shows the cycle performance of the all-solid-state battery using the  $70\text{FePS}_3\cdot 30\text{S}$  electrode. The reversible capacity is more than  $625 \text{ mAh g}^{-1}$ , and the charge capacity at the 50th cycle is  $653 \text{ mAh g}^{-1}$ . At  $100 \text{ }^\circ\text{C}$ , the battery shows better cycle performance than at  $25 \text{ }^\circ\text{C}$ . The battery exhibited the discharge capacity of larger than  $581 \text{ mAh g}^{-1}$  at the current density of  $2.03 \text{ mA cm}^{-2}$  while the battery hardly showed discharge-charge behavior at the current density of equal or higher than  $4.07 \text{ mA cm}^{-2}$  (Figure S4). We also investigated the effect of composition ratios of  $\text{FePS}_3$  and S on the capacities of all-solid-state batteries operated at  $100 \text{ }^\circ\text{C}$ , and find that  $70\text{FePS}_3\cdot 30\text{S}$  shows higher capacity than those of  $100\text{FePS}_3\cdot 0\text{S}$  and  $50\text{FePS}_3\cdot 50\text{S}$  (Figure S5).

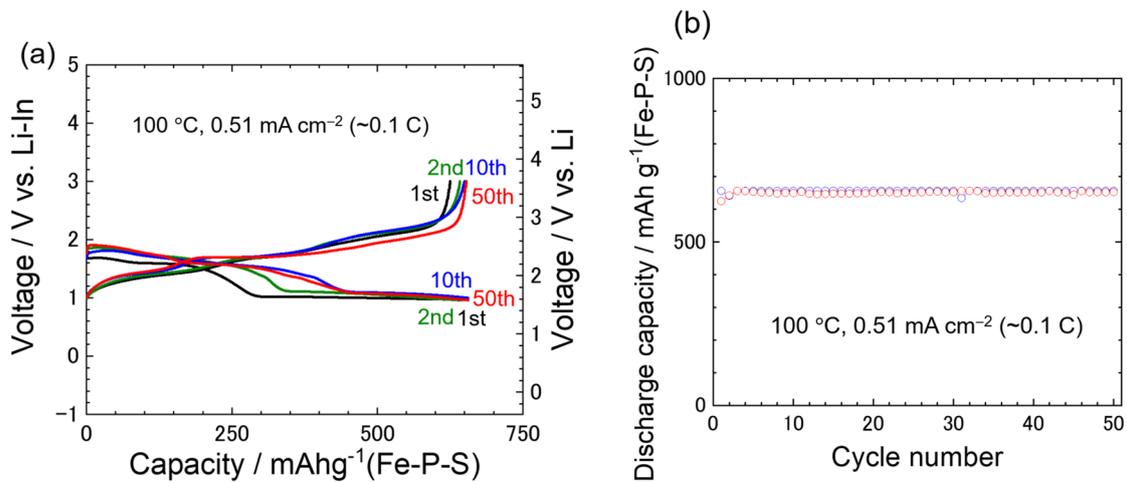


Figure 5. (a) Discharge-charge curves and (b) cycle performance of the all-solid-state

battery using the 70FePS<sub>3</sub>·30S electrode at 100 °C under a current density of 0.51 mA cm<sup>-2</sup> (~0.1 C). Discharge and charge capacities are represented by blue and red dots, respectively. For discharging, the cut-off voltage was set to 0.9 V vs. Li-In and the cut-off capacity was set to 656 mAh g<sup>-1</sup>. For charging, the cut-off voltage was set to 3.0 V vs. Li-In for charging.

To clarify the discharge-charge behavior, the redox reaction of sulfur in the 70FePS<sub>3</sub>·30S electrode was further investigated. Figure 6 shows the sulfur *K*-edge XANES spectra of the 70FePS<sub>3</sub>·30S composite electrode at different discharge-charge conditions marked as colored circles in a first discharge-charge curve of the all-solid-state battery using the 70FePS<sub>3</sub>·30S electrode. For comparison, the spectra of the 75Li<sub>2</sub>S·25P<sub>2</sub>S<sub>5</sub> glass solid electrolyte (Figure 6 (A)) and sulfur (Figure 6 (B)) are also shown. Moreover, the intensity of the sulfur component at 2471.8 eV, at which shows a strong absorption of elemental sulfur, is shown on the lower right side in Figure 6 to discuss the chemical state of the sulfur component during the discharge-charge cycle. The samples contain some amounts of 75Li<sub>2</sub>S·25P<sub>2</sub>S<sub>5</sub> glass of a solid electrolyte layer. The absorption at 2471.8 eV is attributed to sulfur, whereas the absorption at 2470.8 eV is probably attributable to the 75Li<sub>2</sub>S·25P<sub>2</sub>S<sub>5</sub> glass. The absorption of the 75Li<sub>2</sub>S·25P<sub>2</sub>S<sub>5</sub> glass cannot be used for

discussion in detail, because some amount of  $75\text{Li}_2\text{S}\cdot 25\text{P}_2\text{S}_5$  glass of a solid electrolyte layer was unintentionally mixed in the composite electrode samples when the measurement samples were collected. Before discharge and charge, the sulfur component is confirmed (left side in Figure 6 (a)). At 1.1 V vs. Li-In during the first discharge, the peak intensity of the sulfur component decreases, as shown in the intensity at 2471.8 eV (right lower side in Figure 6). This suggests that the sulfur component be chemically reduced during the discharge cycle up to 1.1 V vs. Li-In. After the first discharge, the peak intensity of the sulfur component is not significantly changed. This means the chemical state of sulfur was not significantly altered during the discharge from 1.1 V vs. Li-In to after the first discharge. After the first charge, the peak intensity of the sulfur component becomes stronger, as shown in the intensity at 2471.8 eV (right lower side in Figure 6). The change of peak intensity indicates that the sulfur component is oxidized during the charge cycle. Note that the profile before discharge and charge is not completely consistent with those after the first charge. This suggests that the initial state of sulfur is not totally recovered after the charge cycle. The absorption edge energy, which is estimated at the energy of the normalized absorption intensity of 0.5, is between 2469.4 and 2469.6 eV in the spectra of the Fe-P-S composite electrodes at the all discharge-charge conditions. The absorption edge energy is almost the same as that of

75Li<sub>2</sub>S·25P<sub>2</sub>S<sub>5</sub> glass (2469.6 eV). This is because the samples contain some amounts of 75Li<sub>2</sub>S·25P<sub>2</sub>S<sub>5</sub> glass. Considering the effect of the solid electrolyte inclusion on the absorption edge energy, the redox reaction of the sulfur cannot be discussed based on the absorption edge energy. In the post-edge region of the Fe-P-S composite electrodes, the peak is observed at 2478.6 eV before discharge and charge, which is similar to that of (B) sulfur [28]. On the other hand, at 1.1 V vs. Li-In during the first discharge, the peak is observed at 2476.0 eV, which is consistent with that of (A) 75Li<sub>2</sub>S·25P<sub>2</sub>S<sub>5</sub> glass. These results indicate the sulfur component is chemically reduced during the discharge cycle up to 1.1 V vs. Li-In. After the first discharge and the first charge cycles, the same peak at 2476.0 eV is observed. This suggests that the sulfur component does not show the redox reaction from the discharge cycle below 1.1 V vs. Li-In to the end of the first charge cycle. Note that this result may be affected by the solid electrolyte inclusion in the samples. From the change of the absorption of sulfur *K*-edge XANES spectra, it is reasonable to conclude that the sulfur component is reduced and oxidized during the discharge-charge cycles.

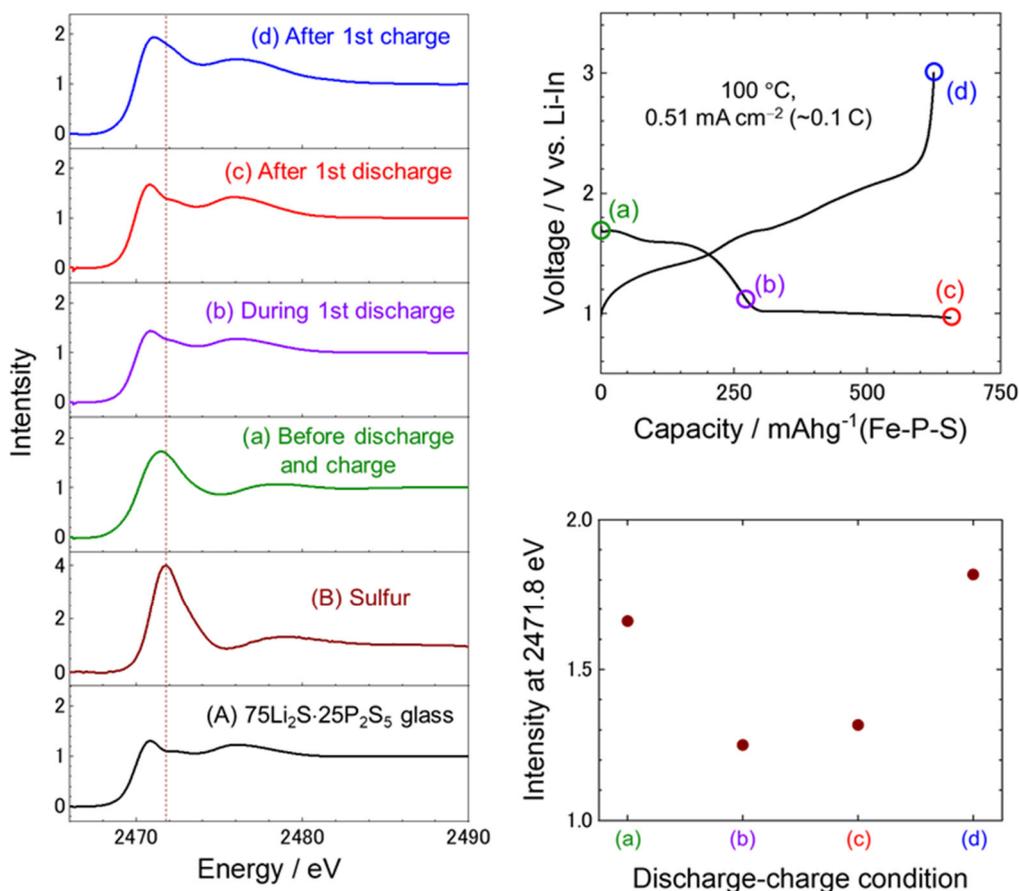


Figure 6. Sulfur *K*-edge XANES spectra of the 70FePS<sub>3</sub>·30S composite electrode (a) before discharge and charge, (b) at 1.1 V vs. Li-In during the first discharge, (c) after the first discharge, and (d) after the first charge cycle. For comparison, the spectra of (A) the 75Li<sub>2</sub>S·25P<sub>2</sub>S<sub>5</sub> glass solid electrolytes and (B) sulfur are shown. The brown dotted line indicates the energy at 2471.8 eV corresponding to the sulfur component. First discharge-charge curve of the all-solid-state battery using the 70FePS<sub>3</sub>·30S electrode is also shown in the right upper figure. Colored circles are from the S *K*-edge XANES spectra of the 70FePS<sub>3</sub>·30S composite electrode, as measured (a) before discharge and charge, (b) at 1.1 V vs. Li-In during the first discharge, (c) after the first discharge, and (d) first charge

cycles. Intensities at 2471.8 eV at each discharge-charge condition are shown in the right lower panel.

In this study, the all-solid-state battery using the Fe-P-S electrode synthesized from 70 wt % FePS<sub>3</sub> and 30 wt% sulfur exhibits a reversible capacity of more than 625 mAh g<sup>-1</sup>(Fe-P-S) based on the redox reaction of sulfur in spite of low content of carbon additive (only 2 wt%) in the electrode. This is because the addition of FePS<sub>3</sub> into sulfur helped to ensure the electronic path in the electrode. The electronic conductivity of the 70FePS<sub>3</sub>·30S electrode is 3×10<sup>-7</sup> S cm<sup>-1</sup> at 27 °C and 6×10<sup>-6</sup> S cm<sup>-1</sup> at 100 °C. To compare with other all-solid-state lithium-sulfur batteries previously reported, Table S1 shows details of our battery and these batteries. The discharge capacity of 625 mAh g<sup>-1</sup>(Fe-P-S) in our battery using the Fe-P-S electrode is lower than those of some other all-solid-state batteries shown in Table S1 since the Fe-P-S electrode can contain partially inactive iron and phosphorus. Note that the difference of the construction conditions of the batteries prevents the quantitative comparison of these electrochemical properties. On the other hand, we believe all-solid-state battery shows reasonably good moderate rate performance (~0.1 C) and the moderate discharge capacity (625 mAh g<sup>-1</sup>) at 100 °C in spite of not only low carbon content but also high active material content. In addition,

because of extremely low carbon content and introduction of phosphorus, which presumably help lithium-ion conductivity, our battery shows high energy density based on volume of the composite electrode even after repeated discharge-charge cycles.

## **Conclusion**

Fe-P-S electrode active materials were synthesized by mechanical milling of FePS<sub>3</sub> and elemental sulfur. The resultant 70FePS<sub>3</sub>-30S powder with particles of about 1–5 μm diameter was composed of FeS<sub>2</sub> with the size of 30 nm, and the amorphous P-S inclusions. In the discharge-charge measurement at 100 °C, at which liquid electrolytes are unstable, the all-solid-state battery using the 70FePS<sub>3</sub>-30S composite electrode, containing only 2 wt% carbon additives, showed excellent reversible discharge-charge behavior. The reversible capacity was more than 625 mAh g<sup>-1</sup>(Fe-P-S) for 50 cycles at 0.51 mA cm<sup>-2</sup> (~0.1 C) in spite of the low content of carbon additive in the composite electrode. This capacity was based on a redox reaction of the sulfur component. Thus, the addition of an

iron-based metal sulfide to elemental sulfur can provide moderate electronic conductivity to sulfur, and the carbon additive content in the sulfur composite electrode can be thereby decreased. The Fe-P-S electrode with low content of carbon would be useful for a high-capacity electrode for all-solid-state lithium-sulfur batteries.

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