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Removal of phosphorus and fluorine from wastewater containing PF_6^- via accelerated decomposition by Al^{3+} and chemical precipitation for hydrometallurgical recycling of lithium-ion batteries†

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During hydrometallurgical recycling of lithium-ion batteries (LIBs), one important challenge is the efficient treatment of wastewater containing LiPF_6 used as a lithium salt in the LIBs. The difficulty of the treatment is attributed to the persistence of PF_6^- in aqueous solutions. In this study, the accelerated decomposition of PF_6^- by Al^{3+} at an elevated temperature and the removal of phosphorus and fluorine by chemical precipitation were attempted. These reactions were analyzed using a pH electrode and fluoride-ion selective electrode, and by a distillation method for total fluorine analysis, ICP-AES, ion chromatography, XRD, and WDS. The results showed that when 10 mM LiPF_6 aqueous solution containing 100 mM $\text{Al}_2(\text{SO}_4)_3$ was kept at 90 °C for 24 h, more than 90% of the PF_6^- was decomposed into PO_4^{3-} and F^- . The produced PO_4^{3-} and F^- were coprecipitated with $\text{Ca}_6\text{Al}_2(\text{SO}_4)_3(\text{OH})_{12}$ (ettringite) by adding sufficient $\text{Ca}(\text{OH})_2$. The concentrations of the total phosphorus and total fluorine in the supernatant after precipitation were 0.028 mM and 0.77 mM, respectively. Here, the pH after the decomposition of 10 mM PF_6^- decreases to around 1 due to the formation of H^+ during the decomposition, which may be too low for some practical cases. For this problem, the decomposition of PF_6^- in various pre-mixed solutions of $\text{Al}_2(\text{SO}_4)_3$ and $\text{Ca}(\text{OH})_2$ was also examined. As a result, when the prepared molar ratio was $\text{Al}/\text{Ca} > 2/3$, the decomposition of PF_6^- proceeded, and the pH decrease accompanying the decomposition was alleviated due to the buffer effect of the $\text{Al}(\text{OH})_3$ precipitate.

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Water impact

The method for faster decomposition of PF_6^- , which is persistent in aqueous water, is required for wastewater treatment. The efficient decomposition of PF_6^- and the removal of phosphorus and fluorine were achieved using $\text{Al}_2(\text{SO}_4)_3$ and $\text{Ca}(\text{OH})_2$. This method is applicable for treating wastewater exhausted in hydrometallurgical recycling of lithium-ion batteries.

1. Introduction

The production of lithium-ion batteries (LIBs) is exponentially increasing year by year towards a decarbonized and sustainable society, which simultaneously leads to the generation of a large number of spent LIBs. In order to recover various valuable elements such as lithium (Li), cobalt (Co), and nickel (Ni) contained in LIBs, efficient recycling methods for spent LIBs are required.^{1–3} The recycling processes of spent LIBs currently in commercial use are

mainly classified as pyrometallurgical and hydrometallurgical processes.^{4–6} In the pyrometallurgical process, spent LIBs are roasted or melted at various temperatures depending on the companies.^{7,8} One of the advantages of the pyrometallurgical process is the safe treatment of the flammable organic solvents and active Li remaining in the negative electrode by combustion. On the other hand, the necessity of the treatment of the fluorine-containing gases remains a problem. In addition, Li transfers to the slag under some melting conditions, and the recovery cost of Li could be high. The recycling processes of spent LIBs at room temperature (R.T.) including the hydrometallurgical process are considered as a solution to the problems of the pyrometallurgical process. When recycling of spent LIBs is conducted without roasting, treatment of the fluorine-containing gases is not necessary, and Li remains in the

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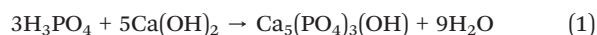
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positive electrode or electrolyte. The most important challenge is the safe deactivation of spent LIBs containing flammable organic solvents and active Li that may cause ignition. One of the deactivation methods proposed in the past is comminution in water or an inert atmosphere. The first reported method of comminution of spent LIBs in water at R.T., to the author's knowledge, was a patent by Asaka Riken.⁹ Later, other technologies were reported for comminution in water, for example, by Retrie¹⁰ and LiCycle,¹¹ and for comminution without water under a CO₂ or Ar atmosphere by Recupyl.¹² Our group proposed the submerged comminution in water in an inert atmosphere, especially in lime water (saturated calcium hydroxide (Ca(OH)₂) solution), to deactivate the spent LIBs even at a charged state.¹³ The submerged comminution in water is superior in terms of low ignition risk to comminution in the gas phase, but treatment of wastewater containing electrolytes of LIBs, *i.e.*, organic solvents such as carbonate ester and Li salts such as LiPF₆, is required.^{13–15}

Phosphorus (P) and fluorine (F) in aquatic ecosystems cause eutrophication and health hazards, respectively, and effluent standards for wastewater are established in many countries.^{16–18} For example, the concentrations of P and F in industrial wastewater are regulated at 16 ppm and 8 ppm, respectively, in the representative Japanese standards. Wastewater containing P is mainly treated by chemical precipitation, crystallization, and biological treatment.^{16,17} In the chemical precipitation, P is removed by being coprecipitated with aluminum hydroxide (Al(OH)₃) or iron hydroxide (Fe(OH)₃) using chlorides or sulfates of aluminum(III) or iron(III). In the crystallization, P is removed as precipitates such as hydroxyapatite (Ca₅(PO₄)₃(OH)), reaction (1) by introducing calcium compounds such as Ca(OH)₂.



Wastewater containing F is typically treated by chemical precipitation, crystallization for example by forming calcium fluoride (CaF₂, reaction (2)), membrane treatment, and adsorption methods.^{18–20}



It is difficult to treat wastewater containing hexafluorophosphate ion (PF₆[−]) produced by the ionization of LiPF₆ by chemical precipitation or crystallization because PF₆[−] is persistent in aqueous solution^{21,22} and does not usually form insoluble precipitates with common metal cations as far as we know. The precipitation with non-metallic compounds and the decomposition into PO₄^{3−} and F[−] are candidate treatment methods for wastewater containing PF₆[−]. As a method of the former, pyridine (C₅H₅N) is reported to form an insoluble precipitate with PF₆[−] in reaction (3); to the best of our knowledge, this is the only method that can precipitate with PF₆[−] in aqueous solution without decomposition.²³



However, the use of pyridine has several problems such as toxicity to humans, flammability, and odorousness. Furthermore, the treatment of pyridine-containing wastewater is costly because combustion or adsorption treatment is required.²⁴

On the other hand, some methods to decompose PF₆[−] into PO₄^{3−} and F[−] were reported according to reaction (4).^{22,25}



One decomposition method patented by some Japanese companies is adding 2–25 wt% hydrochloric acid (HCl) or 35 wt% sulfuric acid (H₂SO₄) into wastewater containing PF₆[−] and heating at 50–100 °C for 0.5–5 h.^{26–28} In this method, the acid concentration is so strong and the operation temperature is so high that highly corrosion-resistant components are required for the treatment tanks. In addition, the vapor pressure of HF increases under heated acidic conditions,²⁹ and the treatment of HF-containing gas could be also required. Another decomposition method for PF₆[−] is the addition of compounds containing cations working as hard acids such as Zr⁴⁺, Th⁴⁺, Al³⁺, and Be²⁺.³⁰ The reaction temperatures in the report were only around R.T., and the decomposition rate was not sufficient; for example, the half-lives of PF₆[−] in 1.0 mol L^{−1} (M) HCl dissolving 1.0 M Zr⁴⁺, 1.5 M Th⁴⁺, 2.0 M Al³⁺, and 1.5 M Be²⁺ were 78 h, 213 h, 767 h, and 1050 h, respectively. The above decomposing methods need improvement in terms of reagent cost and treatment time.

In this study, the accelerated decomposition of PF₆[−] by Al³⁺ was attempted at elevated temperature. Chemical precipitation of PO₄^{3−} and F[−] produced by the decomposition of PF₆[−] was carried out by adding Ca(OH)₂. As shown in Fig. 1, two types of procedures were investigated. In procedure A (Fig. 1(a)), LiPF₆ solutions containing aluminum sulfate (Al₂(SO₄)₃) are heated at 90 °C for 24 h to decompose PF₆[−], followed by adding Ca(OH)₂ to remove the produced PO₄^{3−} and F[−]. In procedure B (Fig. 1(b)), LiPF₆ solutions containing both Al₂(SO₄)₃ and Ca(OH)₂ at various prepared concentrations are heated at 90 °C for 24 h to determine the best conditions for treating PF₆[−]. The supernatants and precipitates of the respective samples are analyzed by various methods. The advantages and disadvantages of procedures A and B are discussed.

2. Methods

2.1. Experimental procedures

Decomposition of PF₆[−] in procedure A. In procedure A, a sample solution containing 10 mM LiPF₆ and 100 mM Al₂(SO₄)₃ was prepared by dissolving LiPF₆ (Fujifilm Wako Pure Chemical Corporation, 98.0+%) and Al₂(SO₄)₃·14–18H₂O (Kanto Chemical Co., Inc., GR) in deionized water (DI, Organo Corporation, Pure Light, <0.1 μS cm^{−1}). As LiPF₆ powder reacts readily with moisture in air, it was weighed in a dry Ar glove box and then quickly dissolved in DI water in air. Sample solutions containing 10 mM LiPF₆ with various pH levels were prepared by adding a suitable amount of HCl





Fig. 1 Procedures (a) A and (b) B for the decomposition of PF₆⁻ and chemical precipitation of PO₄³⁻ and F⁻.

solution (Nacalai Tesque, Inc., GR, 35%) to examine the effect of pH. Also, a sample solution containing 10 mM LiPF₆ and 100 mM Al(NO₃)₃ was prepared by using Al(NO₃)₃·9H₂O (Fujifilm Wako Pure Chemical Corporation, GR) to examine the effect of anions. The prepared sample solutions were placed in polypropylene (PP) screw containers with lids and were kept at R.T. or 90 °C for 24 h in a water bath (As One Corporation, HWA-50A). The concentrations of F⁻ and the pH were then measured after cooling to R.T. The decomposition percentage of PF₆⁻ was evaluated by quantifying the concentration of F⁻, because F⁻ was produced as the decomposition of PF₆⁻ proceeds according to reaction (4).

Chemical precipitation after decomposition of PF₆⁻ by Al³⁺ in procedure A. The saturated Ca(OH)₂ slurry was prepared by adding Ca(OH)₂ powder (Nacalai Tesque, Inc., GR) more than its solubility (0.16 g/100 g H₂O, 21.6 mM at 298 K) into DI water. After the sample solution containing 10 mM LiPF₆ and 100 mM Al₂(SO₄)₃ was kept at 90 °C for 24 h, the saturated Ca(OH)₂ slurry was mixed with the sample solutions at a volume ratio of 1:1. The mixed solution was placed at R.T. for 1 h. Then, the supernatant and precipitate were separated by centrifugation, and the precipitate was dried under vacuum at R.T. The removal percentages of P and F were calculated according to eqn (5).

$$[\text{removal percentages}] (\%) = \left(1 - \frac{[\text{amount of substance of P or F in supernatant}] (\text{mol})}{[\text{amount of substance of P or F in prepared solution}] (\text{mol})} \right) \times 100 \quad (5)$$

The pH and concentrations of the total P and total F in the supernatant were measured. The precipitate was subjected to phase identification by X-ray diffraction (XRD) and elemental analysis by wavelength dispersive spectroscopy (WDS).

Decomposition of PF₆⁻ and chemical precipitation in the aqueous solutions containing both Al₂(SO₄)₃ and Ca(OH)₂ at various prepared concentrations in procedure B. In procedure B, sample solutions containing 10 mM LiPF₆, 0–200 mM Al₂(SO₄)₃, and 108 mM Ca(OH)₂ were prepared by dissolving each reagent in DI water. These sample solutions were then kept at 90 °C for 24 h. The formation of precipitates was observed in each sample. In addition to the same analysis of the supernatant and precipitate as procedure A, quantification of the concentration of each anion was conducted.

2.2. Analysis

Measurement of pH using a pH electrode. The pH was measured using a pH electrode (Horiba, Ltd., 9632-10D), which was calibrated with standard solutions (pH 4.01, 6.86, and 9.18) according to the National Institute of Standard and Technology (NIST) standard methods. The pH measurements of the heated samples were performed after cooling to R.T.

Measurement of the F⁻ concentration using a fluoride-ion selective electrode. The F⁻ concentration in each sample solution was measured using a fluoride-ion selective electrode (Horiba, Ltd., 6561S-10C). Calibration was performed with 1 and 10 ppm or 10 and 100 ppm F⁻ standard solutions prepared using a 1000 ppm F⁻ standard solution (Fujifilm Wako Pure Chemical Corporation, Japan Calibration Service System (JCSS)). Before the measurements with the fluoride-ion selective electrode, the sample solutions were mixed with a total ionic strength adjustment buffer (TISAB) solution in a 1:1 volume ratio. Two types of TISAB solutions were prepared following American Society for Testing and Materials (ASTM) D1179 method B. TISAB 1 was prepared by dissolving 57 mL acetic acid (Nacalai Tesque, Inc., GR), 58 g NaCl (Nacalai Tesque, Inc., GR), and 0.3 g trisodium citrate dihydrate (Kanto Chemical Co., Inc., GR) in DI water to a volume of 1 L, and used for the sample solutions without Al³⁺. TISAB 2 was prepared by dissolving 84 mL HCl solution, 242 g tris(hydroxymethyl)aminomethane (Nacalai Tesque, Inc., GR), and 230 g sodium (+)-tartrate dihydrate (Fujifilm Wako Pure Chemical Corporation, GR) in DI water to a volume of 1 L, and used for the sample solutions containing Al³⁺ to mitigate the interfering effect of Al³⁺. As a preliminary test, sample solutions dissolving Al₂(SO₄)₃ and NaF were mixed with TISAB 1 or 2 solution and measured with a fluoride-ion selective electrode (Table S1†).



The results showed that the interfering effect of Al^{3+} was almost negligible in TISAB 2 when the Al^{3+} concentration was below 10 mM. Therefore, the sample solutions were diluted with DI water before the measurements so that the concentration of Al^{3+} was less than 10 mM.

Measurement of the total F concentration by the distillation method. For the determination of the total F concentration in the sample solutions, distillation was performed according to Japanese Industrial Standard (JIS) K 0102 for the quantification of the total F in various fluorine compounds. 1.0 mL of the sample solution, 0.2 g SiO_2 powder (Fujifilm Wako Pure Chemical Corporation, GR), 0.2 mL phosphoric acid (Nacalai Tesque, Inc., GR, 85%), 8 mL perchloric acid (Nacalai Tesque, Inc., GR, 60%), about 8 mL DI water, and some boiling stones were placed in a distillation flask. The temperature of the distillation flask was controlled at 145 ± 5 °C with steam blowing. The distillate was collected up to approximately 50 mL through a Liebig cooling pipe with cooling water. After adding DI water to the collected distillate to a volume of 100 mL, the F^- concentration was analyzed using the fluoride-ion selective electrode. The measured value may be lower than the true concentration because it depends on the recovery percentage of distillation.

Measurement of the total P and total Al concentrations by ICP-AES. Concentrations of the total P and total Al in sample solutions were determined by inductively coupled plasma atomic emission spectroscopy (ICP-AES, Hitachi High-Tech Science Corporation, SPS3520UV). The wavelengths used for the P and Al measurements were 177.496 nm and 237.312 nm, respectively. Standard solutions were prepared by diluting a standard solution of 1000 ppm P (Fujifilm Wako Pure Chemical Corporation, for water analysis) and 1000 ppm Al (Nacalai Tesque, Inc., SP) with 1 M HCl solution. The calibration curves were measured at four points respectively with the concentrations of 0 (blank), 10, 30, and 50 ppm P, and 0 (blank), 5, 10, and 20 ppm Al. Sample solutions were diluted with 1 M HCl solution to fit within the calibration range of the aimed elements. The average of two times measured values was used for the results.

Measurement of concentration of anions (PF_6^- , F^- , PO_4^{3-} , PO_2F_2^-) by ion chromatography. The quantification of anions in solution was performed by suppressed ion chromatography (Shimadzu Corporation, HIC-ESP) with a column (Shimadzu Corporation, Shim-pack IC-SA2), a suppressor (Shimadzu Corporation, ICDS-40A), and an electrical conductivity detector (Shimadzu Corporation, CDD-10AVP). A guard column (Shimadzu Corporation, Shim-pack IC-SA2(G)) and line filter (Shimadzu Corporation, A-356) were attached upstream of the analytical column. Column temperature was kept at 50 °C. The eluent was prepared by dissolving 3.6 mM Na_2CO_3 (Fujifilm Wako Pure Chemical Corporation, GR) and 3.4 mM NaHCO_3 (Fujifilm Wako Pure Chemical Corporation, GR) in DI water and flowed at a flow rate of 1.0 mL min^{-1} . The sample solutions to be measured were diluted one hundredfold with the eluent, and 20 μL of diluted solution was injected using a manual injector. The anions that could be formed during the

decomposition of PF_6^- are F^- , PO_4^{3-} , PO_2F_2^- , and PO_3F^{2-} , and their concentrations were measured. The calibration curves were measured at three or more concentrations for each anion using standard sample solutions prepared using LiPF_6 , LiPO_2F_2 (Tokyo Chemical Industry Co., Ltd., >98.0%), and $\text{Na}_2\text{PO}_3\text{F}$ (Thermo Fisher Scientific K.K.), 1000 ppm PO_4^{3-} standard solution (Fujifilm Wako Pure Chemical Corporation, JCSS), 1000 ppm F^- standard solution, and 1000 ppm SO_4^{2-} standard solution (Fujifilm Wako Pure Chemical Corporation, JCSS). The ion chromatograms of the respective standard solutions used in this study are shown in Fig. S1.† In this study, PO_3F^{2-} could not be quantified for sample solutions containing SO_4^{2-} because their peaks overlap each other. The noise level was calculated by the average of the difference between the maximum and minimum values, accounting for drift, in each 0.5 min section without any peaks following the ASTM standards. The signal/noise (S/N) ratios of 3.3 and 10 were set as the detection limit and the limit of quantification, respectively.

Phase identification of precipitates by XRD and elemental analysis by WDS. An X-ray diffractometer (XRD, PANalytical, X'pert Pro, Cu-K α line, 45 kV, 40 mA, θ -2 θ method) was used for the phase identification of the precipitates. For the elemental analysis of the precipitates, wavelength dispersive spectroscopy (WDS, 15 kV, 50 nA, ZAF method) was performed with a field emission-electron probe micro analyzer (FE-EPMA, JEOL Ltd., JXA-8530F). Powdered samples were fixed on carbon tape, and carbon-coated with a carbon coater (Meiwafosis Co., Ltd., CADE-E) to give conductivity. In the mapping analysis, spot measurements were carried out at $0.5 \mu\text{m} \times 0.5 \mu\text{m}$ intervals for a $60 \mu\text{m} \times 43 \mu\text{m}$ area.

3. Results

3.1. Decomposition of PF_6^- in aqueous solution under various conditions

The F^- concentration indicates the decomposition progress of PF_6^- . The measured concentrations of F^- and pH are shown in Fig. 2, where 60 mM F^- corresponds to the complete decomposition reaction of 10 mM LiPF_6 . The F^- concentration in each sample kept at R.T. is only less than 2 mM, but 54–56 mM F^- are detected for sample solutions kept at 90 °C in HCl solution with pH 1, 100 mM $\text{Al}_2(\text{SO}_4)_3$ solution, and 100 mM $\text{Al}(\text{NO}_3)_3$ solution. On the other hand, about 10 mM F^- is detected in HCl solution with pH 3, which is approximately equal to the pH in 100 mM $\text{Al}_2(\text{SO}_4)_3$ or 100 mM $\text{Al}(\text{NO}_3)_3$ solution before keeping. These results suggest that not only H^+ but also Al^{3+} accelerate the decomposition of PF_6^- at elevated temperatures. In $\text{Al}_2(\text{SO}_4)_3$ and $\text{Al}(\text{NO}_3)_3$ solutions, pH changes from around 3 to 1 are observed due to the formation of H^+ according to reaction (4).

3.2. Removal of P and F by adding $\text{Ca}(\text{OH})_2$ after decomposition of PF_6^- by Al^{3+} (procedure A)

According to procedure A, the removal of P and F was examined by adding the saturated $\text{Ca}(\text{OH})_2$ slurry after the decomposition of PF_6^- in $\text{Al}_2(\text{SO}_4)_3$ solution at 90 °C for 24 h.



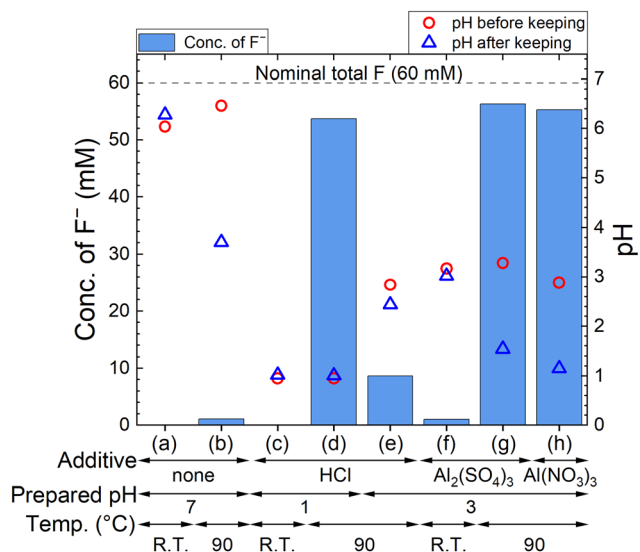


Fig. 2 Measured concentrations of F^- after keeping 10 mM $LiPF_6$ + none, HCl, $Al_2(SO_4)_3$, or $Al(NO_3)_3$ solution at R.T. or 90 °C for 24 h, and pH before and after keeping. The additive, prepared pH, and temperature are (a) none, 7, and R.T., (b) none, 7, and 90 °C, (c) HCl, 1, and R.T., (d) HCl, 1, and 90 °C, (e) HCl, 3, and 90 °C, (f) 100 mM $Al_2(SO_4)_3$, 3, and R.T., (g) 100 mM $Al_2(SO_4)_3$, 3, and 90 °C, and (h) 100 mM $Al(NO_3)_3$, 3, and 90 °C, respectively.

The removal percentages of P and F at 1 h after adding $Ca(OH)_2$ are shown in Fig. 3. In Exp. #01, the pH after adding 150 mM of $Ca(OH)_2$ is 4.3, and the removal percentages are 99% for P and 57% for F. On the other hand, the pH after adding 400 mM of $Ca(OH)_2$ is 10.6 in Exp. #02, and the removal percentages reach higher than 97% both for P and F. The removal percentages of P and F are enough or close to achieve the Japanese industrial effluent standards; the

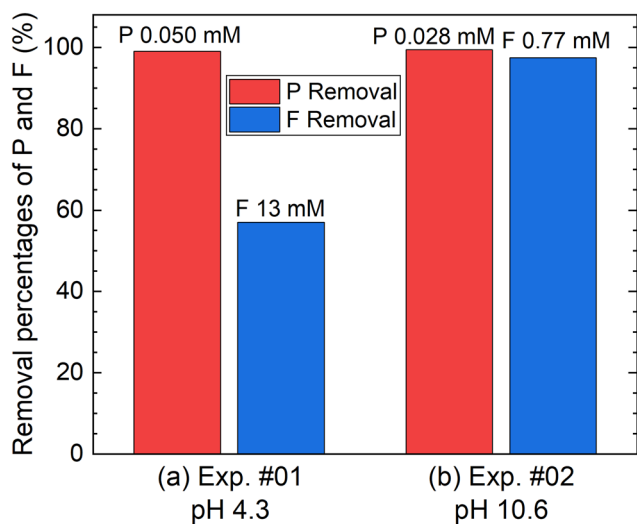


Fig. 3 Removal percentages of P and F after keeping the 10 mM $LiPF_6$ + 100 Al mM $Al_2(SO_4)_3$ solution at 90 °C for 24 h following the addition of (a) 150 mM or (b) 400 mM of saturated $Ca(OH)_2$ slurry at a volume ratio of 1:1 according to procedure A. The numbers above the bar graph indicate the measured concentrations of the supernatant of the total P or total F.

standards of 16 ppm and 8 ppm correspond to 0.516 mM for P and 0.421 mM for F, respectively.

The XRD pattern of the precipitate obtained in Exp. #02 is shown in Fig. 4. Strong peaks of $Ca_6Al_2(SO_4)_3(OH)_{12} \cdot 26H_2O$ (ettringite) and weak peaks of $Ca(OH)_2$ are identified. The formation of ettringite is expected according to reaction (6).



To evaluate the formation of a compound containing P and F, the precipitate was analyzed by FE-EPMA and WDS. The obtained scanning electron microscope (SEM) image and elemental distribution mappings are shown in Fig. 5. Sulfur, Ca, and Al coexist in the same grains, and they are expected to be ettringite. P and F are also observed in the ettringite, and PO_4^{3-} and F^- are suggested to be coprecipitated with ettringite.

3.3. Removal of P and F via decomposition of PF_6^- in $Al_2(SO_4)_3$ + $Ca(OH)_2$ solutions (procedure B)

According to procedure B, the decomposition of PF_6^- and the removal percentages of P and F were examined in $Ca(OH)_2$ + $Al_2(SO_4)_3$ solutions. The removal percentages of P and F in the supernatant kept at 90 °C for 24 h are shown in Fig. 6(a) and the pH levels before and after keeping are shown in Fig. 6(b). The removal percentage of P significantly increases when the $Al_2(SO_4)_3$ concentration is higher than 70 mM and reaches 85–98% at 100 mM. The removal percentage of F also shows a similar trend, reaching about 40% at 100 mM. As shown in Fig. 6(b), the pH decreases from about 4 to 2.5–3.5 before and after keeping for 24 h at 100 mM. For the representative conditions, the concentrations of the

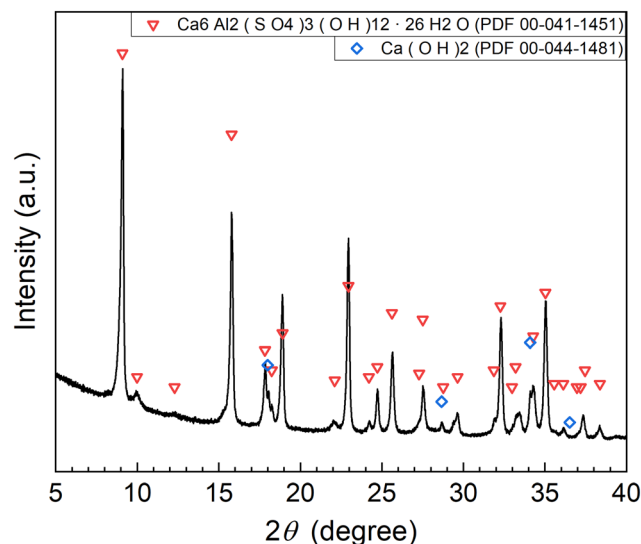


Fig. 4 XRD pattern of the precipitate obtained after keeping the 10 mM $LiPF_6$ + 100 Al mM $Al_2(SO_4)_3$ solution at 90 °C for 24 h following the addition of 400 mM $Ca(OH)_2$ as the saturated slurry (Exp. #02) according to procedure A.



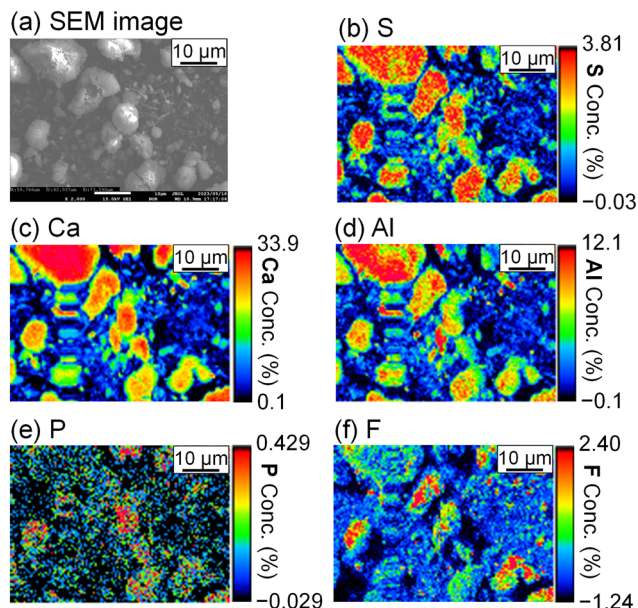


Fig. 5 (a) SEM image and WDS mappings for (b) S, (c) Ca, (d) Al, (e) P, and (f) F of the precipitate obtained after keeping the 10 mM LiPF_6 + 100 Al mM $\text{Al}_2(\text{SO}_4)_3$ solution at 90 °C for 24 h following the addition of 400 mM of saturated $\text{Ca}(\text{OH})_2$ slurry (Exp. #02) according to procedure A.

respective anions are shown in Table 1 and the ion chromatograms are shown in Fig. S2.† The precipitates were formed under all conditions. The difference of the total P concentrations calculated from the sum of species by chromatography with those measured by other methods is not significantly high, and the same can be correct for the total F concentrations. Other chemical species containing P or F which are not listed in Table 1 are expected to be therefore almost absent. When the prepared concentration of $\text{Al}_2(\text{SO}_4)_3$ is 0 or 50 mM (Exp. #03 and #04), the concentration of PF_6^- in the supernatant after keeping is 9.6 or 9.0 mM, and only small amounts of other anions containing P or F are detected. On the other hand, at 100–150 mM (Exp. #05 and #06), PF_6^- is not detected but PO_4^{3-} and F^- are significantly detected with a large decrease in the total P and total F concentrations. In procedure B, Exp. #03 and #04 suggest that PF_6^- is stable at 0–50 mM, and Exp. #05 and #06 demonstrate that PF_6^- is decomposed and P and F are precipitated. It is noted that there is no significant pH decrease as observed in procedure A.

The XRD patterns of the obtained precipitates are shown in Fig. 7. Depending on the added amount of $\text{Al}_2(\text{SO}_4)_3$, $\text{Ca}(\text{OH})_2$, ettringite, and/or $\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$ (gypsum) are identified. The formation of ettringite and gypsum is expected according to reactions (6) and (7), respectively.



The stoichiometric molar ratios of Al/Ca in reactions (6) and (7) are 1/3 and 2/3, respectively, and the compounds in the precipitate are expected to depend on the molar ratio of

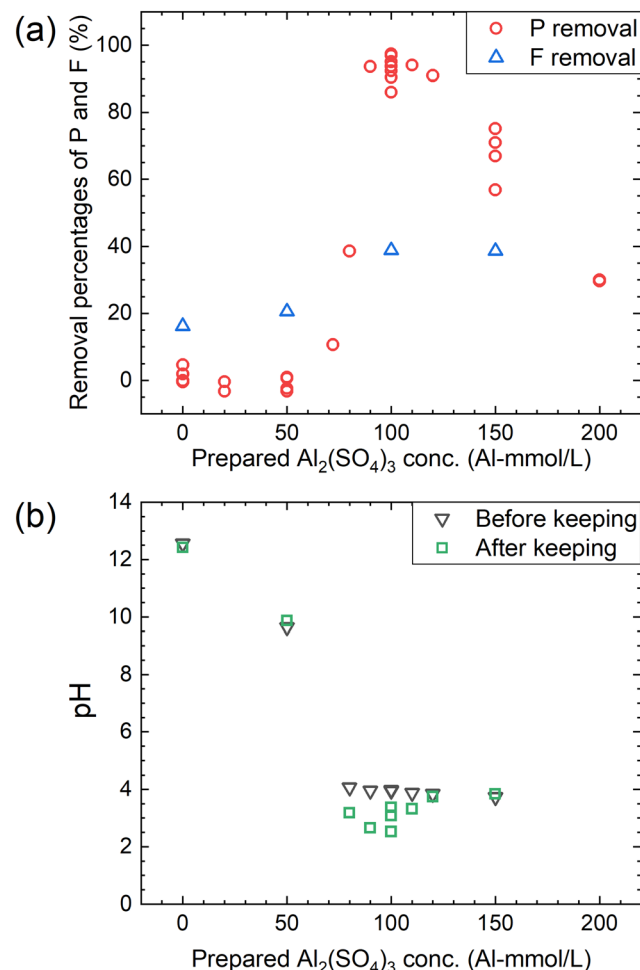


Fig. 6 (a) Removal percentages of P and F after keeping the 10 mM LiPF_6 + 0–200 Al mM $\text{Al}_2(\text{SO}_4)_3$ + 108 mM $\text{Ca}(\text{OH})_2$ solutions at 90 °C for 24 h according to procedure B and (b) the pH before and after keeping.

Al/Ca at the prepared concentrations. In view of this, ettringite and excess $\text{Ca}(\text{OH})_2$ are expected to be in the precipitate for $\text{Al}/\text{Ca} < 1/3$, ettringite, gypsum, and $\text{Al}(\text{OH})_3$ for $1/3 < \text{Al}/\text{Ca} < 2/3$, and gypsum and $\text{Al}(\text{OH})_3$ for $\text{Al}/\text{Ca} > 2/3$. In Fig. 7, ettringite and $\text{Ca}(\text{OH})_2$ are detected by XRD for $\text{Al}/\text{Ca} = 0.19$ (Exp. #07), ettringite and gypsum for $\text{Al}/\text{Ca} = 0.46$ (Exp. #04), and only gypsum for $\text{Al}/\text{Ca} = 0.93$ (Exp. #05) and 1.4 (Exp. #06); the compounds are detected as expected from the stoichiometric ratios of Al/Ca in reactions (6) and (7) except for $\text{Al}(\text{OH})_3$. In procedure B, the removal percentages of P and F are maximum (86% for P and 38% for F) when $\text{Al}/\text{Ca} = 0.93$ (Exp. #05). To evaluate the formation of a compound containing P and F, SEM and WDS were conducted for the precipitate obtained in Exp. #05 and the results are shown in Fig. 8. Only gypsum is detected by XRD, and the grains where S and Ca coexist are expected to be gypsum. The Al grains independently distributed are also observed and may be $\text{Al}(\text{OH})_3$ in amorphous form. Here, P and F are observed in the amorphous $\text{Al}(\text{OH})_3$, suggesting that PO_4^{3-} and F^- are coprecipitated with amorphous $\text{Al}(\text{OH})_3$.



Table 1 Results of measurements of the supernatant obtained in procedure B

Exp. #	Prepared concentration			Measured concentration (mM)						Removal rate (%)		pH			
	LiPF ₆ (mM)	Ca(OH) ₂ (mM)	Al ₂ (SO ₄) ₃ (Al mM)	Ion chromatography ^a				ICP ^b	Distillation ^c	ICP	Distillation	Before	After		
				PF ₆ ⁻	F ⁻	PO ₄ ³⁻	PO ₂ F ₂ ⁻	Total P	Total F					Total P	Total F
03	10	108	0	9.6	0.42	n.d. ^d	<0.057 ^e	9.6	58	10	50	0.0	17	12.6	12.5
04	10	108	50	9.0	{0.19} ^f	<0.13 ^e	n.d. ^d	9.0	54	9.9	48	1.0	20	9.6	10.2
05	10	108	100	n.d. ^d	43	0.58	n.d. ^d	0.58	43	1.4	37	86	38	4.0	2.7
06	10	108	150	n.d. ^d	51	2.4	{0.087} ^f	2.4	51	2.9	37	71	38	3.8	3.8

^a Measured by ion chromatography. ^b Measured by ICP-AES. ^c Measured by the distillation method. ^d Below the detection limit. ^e Below the limit of quantification. ^f {} means "out of the concentration range in the calibration curve".

The measured concentrations of the total Al in the supernatants of each sample kept at 90 °C for 24 h according to procedure B are shown in Fig. 9. The total Al concentration is below the detection limit at Al/Ca < 2/3, but it increases at Al/Ca > 2/3. Excess Al³⁺ for reaction (7) is present in the supernatant when Al/Ca is greater than the stoichiometric molar ratio in reaction (7), 2/3, and the decomposition of PF₆⁻ is expected to proceed due to the excess Al³⁺ (Exp. #05 and #06).

4. Discussion

4.1. Comparison of procedures A and B

The features of the reactions in procedures A and B are schematically illustrated in Fig. 10. The progress of the

decomposition of PF₆⁻ and the coprecipitation with PO₄³⁻ and F⁻ depends on the molar ratio of Al/Ca at the prepared concentration of Ca(OH)₂ and Al₂(SO₄)₃. The decomposition of PF₆⁻ is accelerated by Al³⁺ where Ca = 0 (procedure A) or Al/Ca is greater than 2/3 (procedure B). In procedure A, Exp. #02 showed that the addition of Ca(OH)₂ at the molar ratio of Al/Ca = 0.25 leads to the precipitation removal of PO₄³⁻ and F⁻. The molar ratio of Al/Ca = 0.25 corresponds to the area of Al/Ca < 1/3, where ettringite is formed with excess Ca(OH)₂ at pH 12. In procedure B, some PO₄³⁻ and F⁻ produced by the decomposition are coprecipitated with Al(OH)₃ at Al/Ca > 2/3 without additional Ca(OH)₂.

Here, Exp. #02 showed that the removal percentages of P and F were higher than 97% in procedure A, but in procedure B, Exp. #05 showed that the removal percentages were only 86% for P and 38% for F. It is noted that the molar ratios of Al/Ca in Exp. #02 and #05 are 0.25 and 0.93, respectively. Experimental results show that the removal percentages of P

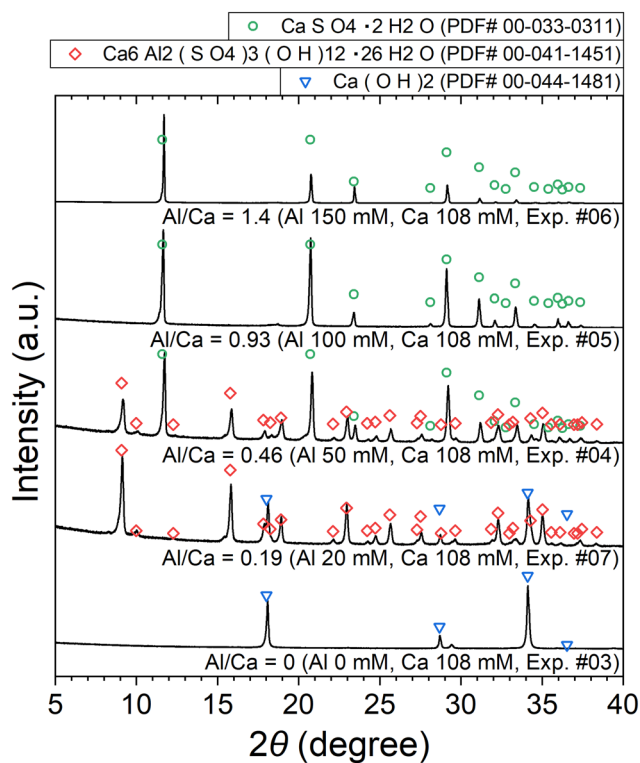


Fig. 7 XRD patterns of the precipitates obtained after keeping the 10 mM LiPF₆ + 0–150 Al mM Al₂(SO₄)₃ + 108 mM Ca(OH)₂ solutions at 90 °C for 24 h (Exp. #03–07) according to procedure B.

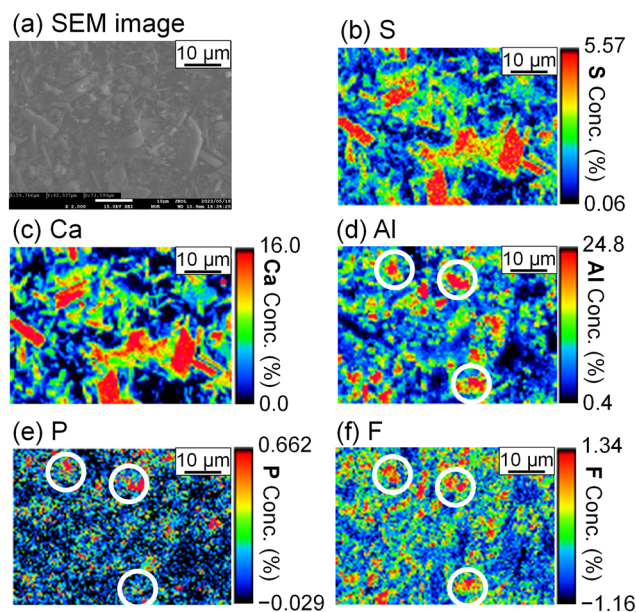


Fig. 8 (a) SEM image and WDS mappings for (b) S, (c) Ca, (d) Al, (e) P, and (f) F of the precipitate obtained after keeping the 10 mM LiPF₆ + 100 Al mM Al₂(SO₄)₃ + 108 mM Ca(OH)₂ solution at 90 °C for 24 h (Exp. #05) according to procedure B. The white circles are representative grains where Al, P, and F coexist.



Table 3 Input value of concentrations of chemical species adopted for the equilibrium calculation

Chemical species	Input value of concentrations	
	Condition A	Condition B
Ca(OH) ₂	From 0 mM to 250 mM in 500 steps	108 mM
Al ₂ (SO ₄) ₃	50 Al mM	From 0 Al mM to 200 Al mM in 500 steps
Li ⁺	5 mM	10 mM
PF ₆ ⁻	0 mM	10 mM for Al/Ca < 2/3 0 mM for Al/Ca > 2/3
PO ₄ ³⁻	5 mM	0 mM for Al/Ca < 2/3 10 mM for Al/Ca > 2/3
F ⁻	30 mM	0 mM for Al/Ca < 2/3 60 mM for Al/Ca > 2/3

Al₂(SO₄)₃ at each concentration. As a simplified condition of the experimental results of this study, condition B used here is that PF₆⁻ is not decomposed at all for Al/Ca < 2/3 and is decomposed completely for Al/Ca > 2/3. The full text of the input data is summarized in the ESI.†

The molar amounts of precipitates equilibrated for each concentration are shown in Fig. 11. The graphs shown in Fig. 11(a) and (b) show the results for conditions A and B, respectively. Under condition A, gypsum, gibbsite, and

hydroxylapatite are formed corresponding to Exp. #01, and ettringite, fluorite, Ca(OH)₂, and hydroxylapatite to Exp. #02, as shown in Fig. 11(a). On the other hand, the experimental results indicated that neither fluorite nor hydroxylapatite was observed by the analysis of XRD (Fig. 4), and WDS (Fig. 5) analysis showed that P and F are coprecipitated with ettringite. The difference between the calculation and experiment results is possibly due to the ion exchange ability of ettringite. The ion exchange behavior of SO₄²⁻ in ettringite with other anions in aqueous solutions such as PO₄³⁻ or F⁻ was reported.^{36,37} The ion exchange reaction may be superior in terms of thermodynamics or kinetics to the formation of fluorite and/or hydroxylapatite.

Under condition B, Ca(OH)₂ is formed corresponding to Exp. #03, ettringite and Ca(OH)₂ to Exp. #07, gypsum, gibbsite, and ettringite to Exp. #04, and gypsum and gibbsite to Exp. #05 and 06, as shown in Fig. 11(b). These results are consistent with the experimental results in section 3.3, except for the coprecipitation of PO₄³⁻ and F⁻ with Al(OH)₃. The coprecipitation of PO₄³⁻ and F⁻ in this study was expected to proceed by surface adsorption on formed Al(OH)₃, as these anions are reported to precipitate by surface adsorption on Al(OH)₃.³⁸⁻⁴⁰

5. Conclusion

In this study, the decomposition of PF₆⁻ in aqueous solution was demonstrated to be accelerated when Al³⁺ was contained in the solution at an elevated temperature. The decomposition of PF₆⁻ is negligibly slow when the solution contained no additive or the solution temperature was room temperature, but it was accelerated when the solution at pH 1 or containing 100 mM Al³⁺ at pH 3 was kept at 90 °C for 24 h. In procedure A, after PF₆⁻ was decomposed by Al³⁺, the produced PO₄³⁻ and F⁻ were removed from the precipitate by adding sufficient Ca(OH)₂ and the removal percentages of P and F were higher than 97%. The precipitate was ettringite, and PO₄³⁻ and F⁻ were coprecipitated with it. The decomposition of PF₆⁻ was accompanied by a pH decrease due to the formation of H⁺. In procedure B, PF₆⁻ was decomposed in pre-mixed solutions of Al₂(SO₄)₃ and Ca(OH)₂ at the prepared molar ratio of Al/Ca > 2/3, and the pH decrease during the decomposition of PF₆⁻ was alleviated. The alleviation of the pH decrease is expected to be due to the buffer effect of the Al(OH)₃ precipitate formed by the reaction of Al₂(SO₄)₃ and Ca(OH)₂. The produced PO₄³⁻ and F⁻ were coprecipitated with Al(OH)₃, but the removal percentages were only 86% for P and 38% for F. For higher removal percentages of P and F, the addition of sufficient Ca(OH)₂ is expected to be required after the decomposition.

Author contributions

All authors conceived the ideas. T. M. contributed to the execution of the experiments, wrote the manuscript, and performed the analysis. All authors reviewed the manuscript.



Fig. 11 Molar amounts of precipitates equilibrated for each concentration of (a) added Ca(OH)₂ under condition A and (b) added Al₂(SO₄)₃ under condition B shown in Table 3. The numbers #0x (x = 1–7) in the graph correspond to the same conditions as Exp. #0x.



Conflicts of interest

There are no conflicts to declare.

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