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Lifetime limit of tris(trimethylsilyl) phosphite as electrolyte additive for high voltage lithium ion batteries

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The tris(trimethylsilyl) phosphite (TMSPi) is considered as an ideal electrolyte additive for lithium ion batteries. In this work, its positive effect as well as its failure mechanism in a LiPF₆ containing electrolyte was studied by means of selected electrochemical, structural and analytical techniques. The LiNi_{0.5}Co_{0.2}Mn_{0.3}O₂/graphite cells with TMSPi as electrolyte additive were cycled between 2.8 and 4.6 V. Thanks to the compact cathode electrolyte interphase formed by the oxidative decomposition of TMSPi in a freshly prepared TMSPi containing electrolyte, both the discharge capacity and the cycling stability of cells were enhanced. However, our results also show that TMSPi actually reacts with LiPF₆ at room temperature. TMSPi is consumed by this spontaneous reaction after aging for certain time. In addition, a part of the fluorophosphates, generated from the hydrolysis of LiPF₆, is bonded to one or two TMS groups, causing a decrease in the fluorophosphate content in the CEI film. Consequently, the cycling stability of the lithium ion cells with aged TMSPi containing electrolyte deteriorates. The obtained results offer important insights into the practical application of TMSPi, which means that TMSPi can only be used as an effective additive in a freshly prepared LiPF₆ containing electrolyte.

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Introduction

To facilitate the robust development of hybrid electric vehicles (HEVs) and electric vehicles (EVs), lithium ion batteries (LIBs), as one of the most promising energy supplies, are expected to provide “high voltage” (>4.5–4.6 V) to enhance their energy density. A variety of cathode materials with high working potential has been reported so far. However, they are rarely used in commercial cells, mainly due to the poor electrochemical oxidation stability of the traditional LiPF₆ containing carbonate based electrolytes at these high voltages.^{1,2} The oxidative decomposition of the carbonate solvents has been confirmed to produce H₂O and CO₂.³ The generated H₂O further promotes the hydrolysis reactions of LiPF₆ to generate HF,⁴ which induces serious damage of metal oxide cathodes^{5,6} and the solid-electrolyte interphase (SEI) on the graphite anodes.⁷ An economic way to prevent the oxidation of the traditional electrolyte formulations at high voltage is to use specific electrolyte additives to form a cathode-electrolyte interphase (CEI),^{5,8}

stabilize the electrolyte salt,⁹ scavenge the corrosive impurities (HF, PF₅),¹⁰ or to remove the O₂ gas evolved from the cathode material.¹¹

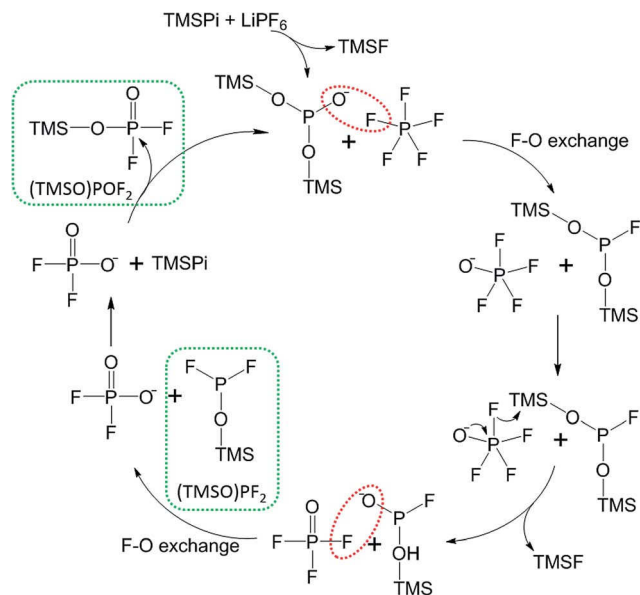
Similar to the SEI formed on the graphite anode, the CEI is considered as an essential protective layer on the cathode surface that helps to prevent further oxidative decomposition of the electrolyte.¹² However, a thick CEI film may deteriorate the cell performance by increasing the interfacial impedance, in particular when it contains more highly resistive components,¹³ e.g. LiF. CEI formation additives are expected to produce a robust and thin CEI layer to enhance both the cycling stability and rate capability.¹⁴

Tris(trimethylsilyl)phosphite (TMSPi, shown in Scheme 1), has been introduced as an effective bifunctional additive to build a CEI upon its oxidative decomposition as well as to scavenge the HF generated during cycling.¹⁵ The cycling stability of LiNi_{0.5}Mn_{1.5}O₄ (ref. 16) and LiNi_{1/3}Co_{1/3}Mn_{1/3}O₂ (ref. 15 and 17) cathodes at high potential (≥4.5 V vs. Li/Li⁺) was significantly improved by 0.5 wt% TMSPi in the electrolyte. Yim *et al.*¹⁸ calculated the oxidation potential (OP) and the reduction potential (RP) based on the Kohn–Sham density functional theory (DFT): TMSPi has an OP as 4.29 V vs. Li/Li⁺, lower than the OP of ethylene carbonate (EC, 6.92 V vs. Li/Li⁺). On the other hand, the RP of TMSPi is much lower than EC.¹⁸ Consequently, the TMSPi would easily undergo an oxidative decomposition on the cathode but less probable reductive decomposition on the

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Scheme 1 A proposed mechanism for the reaction of TMSPi with LiPF₆, without the participation of O₂ and H₂O. The formation of (TMSO)PF₂ and (TMSO)POF₂ is taken as example.

anode. A CEI film can thus be formed to prevent transition metal ion dissolution from the cathode and further oxidative decomposition of the electrolyte.¹⁵ Simultaneously, the trimethylsilyl (TMS) group shows high binding affinity towards HF owing to the formation of a strong Si-F bond. For this reason, TMSPi is also considered as a HF scavenger.

Besides Me₃SiF (TMSF), two more decomposition compounds: Me₃SiOH (TMSOH) and Me₃SiOSiMe₃ (TMSOTMS) have also been observed in the ¹H-NMR spectra of the TMSPi-containing electrolyte,¹⁸ indicating that the highly electrophilic TMS group acts not selectively as a F⁻ receptor, but rather as a receptor for all anions. Hence, it can be suspected that TMSPi may also directly react with the LiPF₆ salt or with the hydrolysis reaction products of LiPF₆. After a certain aging period, the HF scavenging effect of the electrolyte additive may vanish as all TMSPi molecules are continuously consumed by this chemical reaction. In this work, we studied the spontaneous reaction between TMSPi and LiPF₆ and we could identify the products by various analysis methods, including gas chromatography-mass spectrometry (GC-MS) and nuclear magnetic resonance (NMR) spectroscopy. We also investigated the electrochemical performance of the TMSPi containing electrolyte in a LiNi_{0.5}Co_{0.2}Mn_{0.3}O₂/graphite full cell at high cut-off voltage (4.6 V). The oxidation stability analysis revealed that the aged TMSPi containing electrolyte failed to form a protective CEI on the cathode. A detailed interaction between TMSPi and LiPF₆ is shown in this work.

Experimental

Electrolyte and electrode preparation

A 3 : 7 (weight ratio) mixture of ethylene carbonate (EC) and diethyl carbonate (DEC) with 1 M LiPF₆ was used as baseline electrolyte (LP47, UBE, H₂O content ≤10 ppm). 0.5 wt% TMSPi

was added into the baseline electrolyte in an argon filled glovebox (Braun, H₂O and O₂ content ≤0.1 ppm). The “fresh TMSPi containing electrolytes” were prepared shortly prior to each measurement. The “aged TMSPi containing electrolytes” were stored in glovebox at 20 °C for different time periods ranging from 12 hours to 2 months.

LiNi_{0.5}Co_{0.2}Mn_{0.3}O₂ and graphite were used as cathode and anode active materials, respectively. Li metal (Rockwood Lithium®, lithium battery grade) was used as reference electrode (Ø 6 mm). The cathode was composed of 97 wt% LiNi_{0.5}Co_{0.2}Mn_{0.3}O₂ (ATL), 2 wt% carbon black (Super P Li®, Amerys) and 1 wt% polyvinylidene difluoride (PVdF, Solef 5130, Solvay) as binder. The anode was constituted of 95.4 wt% graphite (ATL), 1.5 wt% Super P Li® and 3.1 wt% binder. This binder was a mixture of styrene butadiene rubber (SBR, Lipaton SB 5521, Synthomer) and sodium carboxymethyl cellulose (CMC, Walocel CRT 2000 PPA 12, Dow Wolff Cellulosics®) with a weight ratio of 25 : 6. Both the cathode and anode electrodes were punched into round discs (Ø 12 mm). The mass loading of the cathode was approximately 9.0 mg cm⁻², which meant the total amount of LiNi_{0.5}Co_{0.2}Mn_{0.3}O₂ was ca. 9.9 mg. To balance the cathode/anode capacity ratio, the practical capacity of the LiNi_{0.5}Co_{0.2}Mn_{0.3}O₂ cathode was first determined in the potential range from 2.8 V to 4.6 V vs. Li/Li⁺ with Li metal as counter electrode and a practical capacity of 197.5 mA h g⁻¹ was obtained. The same procedure was applied to determine the practical capacity of the graphite anode between 0.01 and 1.5 V vs. Li/Li⁺, and a specific capacity of 334.5 mA h g⁻¹ at 1C was obtained. The mass loading of the anode was controlled to have 25% extra capacity as the cathode. According to the mass loading of LiNi_{0.5}Co_{0.2}Mn_{0.3}O₂, the total amount of graphite on the anode side was ca. 7.3 mg. In the constant current charge/discharge cycling analysis, the specific current of 197.5 mA g⁻¹ was defined as 1C. 200 µL electrolyte was added into each cell. For each electrolyte formulation, at least three cells were assembled to ensure the reproducibility, which required a capacity deviation below 0.5 mA h g⁻¹ between the parallel cells.

Electrochemical analysis

The cycling stability of the cells was analyzed in a three-electrode Swagelok® T-type cell. The cells were cycled on a Maccor® series 4300 battery cycler at 20 °C. After three initial formation cycles at 0.1C, the cells were charged/discharged at 0.5C in the following cycles. The oxidation stability of the electrolytes was determined by linear sweep voltammetry (LSV) performed by a VMP potentiostat (Biologic® Science Instruments). A platinum disc (Ø 1 mm) was used as working electrode, whereas the counter and reference electrodes were both lithium metal discs (Ø 12 mm and Ø 6 mm, respectively). The scan range was set as 3.0–6.0 V vs. Li/Li⁺ with a scan rate of 0.1 mV s⁻¹.

Qualitative analysis

Qualitative GC-MS experiments were carried out on a GC-MS-QP2010 Ultra System from Shimadzu® equipped with the AOC-5000 Plus autosampler from CTC Analytics. Separation was realized by Rxi®5ms, a nonpolar fused silica column

(30 m × 250 μm × 250 μm) from Restek®. Measurements were performed with helium (purity 6.0) from Westfalen AG as carrier gas. The headspace gas generated from the reaction between LiPF₆ and TMSPi was equilibrated in an agitator at 120 °C for 120 s at 250 rpm. 1 mL gas was injected by a syringe at 120 °C and 500 μL s⁻¹. After injection at 250 °C, the following column oven program was executed and the sample measured in total for 20 min. The temperature was kept at 35 °C for 2 min. and then increased to 60 °C with a rate of 3 °C min⁻¹. Afterwards, the temperature increased to 250 °C with a rate of 31 °C min⁻¹. The mass spectrometer was operated in the electron ionization (EI) mode with an *m/z* scan range of 15–300. The ion source temperature was set to 250 °C, the filament voltage to 70 V and the detector voltage relative to the particular tuning result. As software for compound identification and system control, GC-MS Real Time Analysis and GC-MS Postrun Analysis, which included the National Institute of Standards and Technology (NIST) 08 library, were used.

The water content in the electrolyte was determined by the Karl-Fischer titration method. The measurement was performed by a combination of 801 stirrer and 851 titrator, acquired from Metrohm AG (Herisau, Switzerland).

Surface analysis of the LiNi_{0.5}Co_{0.2}Mn_{0.3}O₂ cathode was performed by X-ray photoelectron spectroscopy (XPS, Axis Ultra HAS, Kratos, GB) with a monochromatic Al K_α source (*hν* = 1486.6 eV) at a 10 mA filament current and a 12 kV filament voltage source energy. Five positions were measured on each sample and each position had an area of 700 μm × 300 μm. The binding energy (BE) of the C 1s peak (C–C, BE = 284.6 eV) was used as an internal reference to calibrate the spectra. All the cells were stopped after 150 cycles in the discharged state. The cathodes were removed out of the cells in a glovebox and transferred into the XPS chamber under argon protection.

To analyze the reaction products of TMSPi and LiPF₆, NMR spectra were recorded on a Bruker Avance III 400 (400.03 MHz, ¹H; 376.37 MHz, ¹⁹F; 161.93 MHz, ³¹P) spectrometer. All measurements were performed at 300 K. Chemical shifts (δ) were reported in parts per million (ppm), and coupling

constants (*J* values) were given in hertz (Hz). All samples were prepared in an argon filled glovebox with pre-dried solvents (H₂O content less than 10 ppm). Fluorinated ethylene propylene (FEP) NMR tubes (Ø 3 mm, Wilmad-labglass) were used to avoid any side reaction between the residual amount of HF and glass. The FEP tubes containing 200 to 300 μL of samples were closed by a polytetrafluorethylene (PTFE) plug and placed in a glass NMR tube (Ø 5 mm) containing 50 μL of acetonitrile-*d*₃ (CD₃CN, 99.8 at% *d*₃, Sigma Aldrich). In this way, any contact between the deuterated solvent and reaction mixtures, which could produce unwanted products, was prohibited. A PTFE NMR cap was used to close the glass NMR tubes which were sealed by Parafilm® afterwards (Fig. 1).

Results and discussion

As reported by Yim *et al.*,¹⁸ the Gibbs free energy of the reaction between TMSPi and LiF was calculated as −34.1 kcal mol⁻¹. It can thus be assumed that TMSPi would influence the equilibrium of the reaction: LiPF₆ ⇌ LiF + PF₅ by removing LiF and continuously promote the decomposition of LiPF₆. As a result, TMSPi would be consumed by this reaction after a certain aging time and would have no longer beneficial effect on the cell performance.

The ¹H-NMR spectra of two aged TMSPi containing electrolytes, which had been stored in glovebox for one week and for 45 days (20 °C), respectively, were measured. Fig. 2a shows the spectrum of pure TMSPi, where CD₃CN was used as solvent. Fig. 2b and c illustrate the ¹H-NMR spectra of LP47 electrolyte with 0.5 wt% TMSPi after one week and 45 days, respectively. In order to avoid any side reaction between CD₃CN and electrolyte components, an experimental set-up, which is shown in Fig. 1, was used. The TMSPi containing electrolyte LP47 was placed in the FEP tube and CD₃CN was filled in the glass tube. The ¹H-NMR spectrum of the TMSPi containing electrolyte after one week of storage shows both signals of TMSPi and TMSF (Fig. 2b). The slight shift of the TMSPi signal from 0.2 ppm (Fig. 2a) to 0.03 ppm (Fig. 2b and c) was caused by using different solvents. As already well known, the multiplet signal in ¹⁹F-NMR spectrum at −157 ppm with ¹*J*_{FH} coupling constant of 7 Hz clearly demonstrates a considerable amount of dissolved TMSF in the solution.¹⁹ The aged TMSPi containing electrolyte, which was stored in a glovebox at 20 °C for 45 days, showed no presence of TMSPi in the electrolyte and only the TMSF could be observed in ¹H-NMR spectrum (Fig. 2c).

The ¹⁹F-NMR spectrum of the TMSPi containing electrolyte aged for 45 days (Fig. 3a) shows different sets of signals between −80 ppm and −86 ppm which are related to different fluorophosphates.²⁰ A doublet at −83.0 ppm in ¹⁹F-NMR spectrum (1 and 1') which correlates with a triplet in ³¹P-NMR spectrum at −20.1 ppm (¹*J*_{PF} = 929 Hz, Fig. 3b) proves the formation of OPF₂(OH). In addition, the ¹⁹F-NMR spectrum (Fig. 3a) shows two sets of doublets at −84.04 ppm and −84.06 ppm with ¹*J*_{PF} coupling constants of 967 (2 and 2') and 1005 (3 and 3') Hz respectively which are assigned to OPF(OMe)₂ and OPF(OEt)₂. However, due to the low signal intensity of the ³¹P-NMR spectrum (Fig. 3b), the related signals could not be determined. In

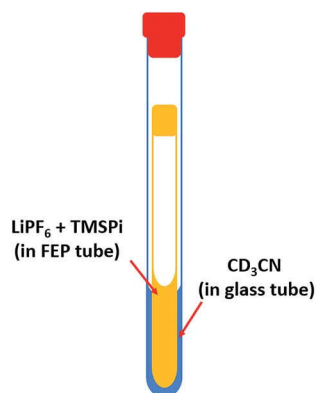


Fig. 1 Schematic plot of the NMR sample set-up to identify the reaction products of LiPF₆ and TMSPi. The LiPF₆ and TMSPi were sealed in the FEP tube; and the CD₃CN was filled in the glass tube as a standard reference.

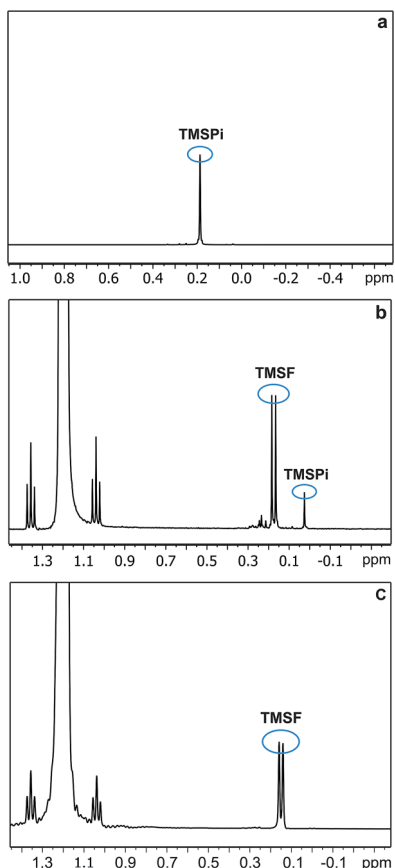


Fig. 2 The ¹H-NMR spectra of pure TMSPi in CD₃CN (a), and 0.5 wt% TMSPi containing LP47 electrolyte stored in a glovebox for one week (b) and for 45 days (c).

Fig. 3a, the singlet at -78.3 ppm cannot be assigned to any product.

The decrease of HF in the TMSPi containing electrolyte has already been confirmed by ¹⁹F-NMR.^{16,21} The TMSPi was thus considered as an effective HF scavenger. However, the high electron affinity of the TMS group may also attract other anions beside F[−]. In this work, a water contaminated LP47 electrolyte was used to evaluate the water scavenging ability of TMSPi. The water content in the contaminated LP47 electrolyte was 50 ppm. 0.5 wt% TMSPi was added and the electrolyte was stored at 20 °C for 12 hours. Afterwards, the water content of both electrolytes with and without TMSPi was measured by Karl-Fischer titration and compared. As summarized in Table 1, the TMSPi-containing electrolyte shows a decreased amount of water, indicating that TMSPi acted as a decontaminating agent to purify the electrolyte by removing the HF and H₂O.

As discussed in the work from Mai *et al.*,¹⁵ the improvement in the cycling stability by using TMSPi was also attributed to a passivating CEI formed by oxidative electrolyte decomposition. The onset potential was at *ca.* 4.1 V *vs.* Li/Li⁺ at a platinum electrode.¹⁵ In this work, as shown in Fig. 2, no TMSPi could be observed after 45 days storage. It is of great importance to find out whether a stable CEI could still be formed after all the TMSPi was consumed by the reaction with HF and H₂O.

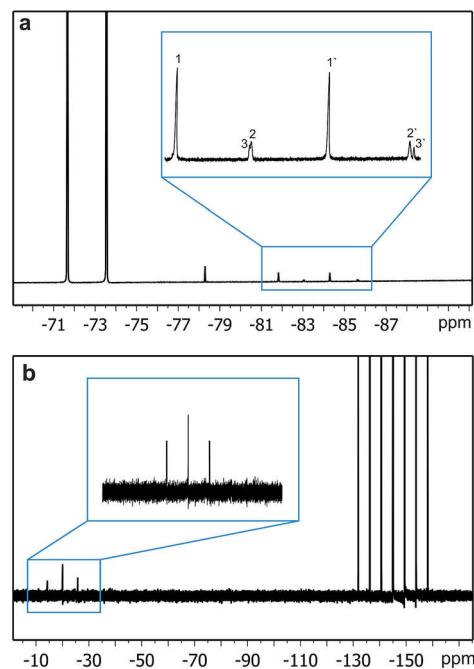


Fig. 3 (a) ¹⁹F-NMR spectrum of the 0.5 wt% TMSPi containing LP47 electrolyte stored in glovebox for 45 days: 1 and 1' OPF₂(OH); 2 and 2' (OPF(OMe)₂); 3 and 3' (OPF(OEt)₂) (b) ³¹P-NMR spectrum of the same sample.

Therefore, the oxidation stability of the baseline LP47 electrolyte, the fresh TMSPi containing LP47 electrolyte and the aged TMSPi containing LP47 electrolyte was determined with a platinum working electrode against a lithium counter electrode (shown in Fig. 4). To get close to the equilibrium state of the reaction, a relatively low scan rate of 0.1 mV s^{−1} (50 times lower than in the work of Mai *et al.*) was used.

As shown in Fig. 4, the oxidative decomposition of all three electrolytes started at approximately the same onset potential. Similar to the results obtained by Mai *et al.*, the fresh TMSPi containing electrolyte clearly formed a passivating layer on the platinum electrode as the anodic current was much lower than the baseline electrolyte above 5.3 V *vs.* Li/Li⁺. However, the aged electrolyte containing TMSPi did not show this passivation behavior and the anodic current had a similar value as in the case of the baseline electrolyte. Obviously, the reaction products

Table 1 Water content in the contaminated LP47 electrolyte with and without the addition of 0.5 wt% TMSPi. The electrolytes were stored in a glovebox at 20 °C for 12 hours

Measurement	LP47 (H ₂ O contaminated)		LP47 (H ₂ O contaminated) + 0.5 wt% TMSPi	
	Sample mass (g)	H ₂ O (ppm)	Sample mass (g)	H ₂ O (ppm)
1	1.21579	44.00	1.32239	38.80
2	1.13990	50.90	1.35221	32.20
Average		47.45		35.50

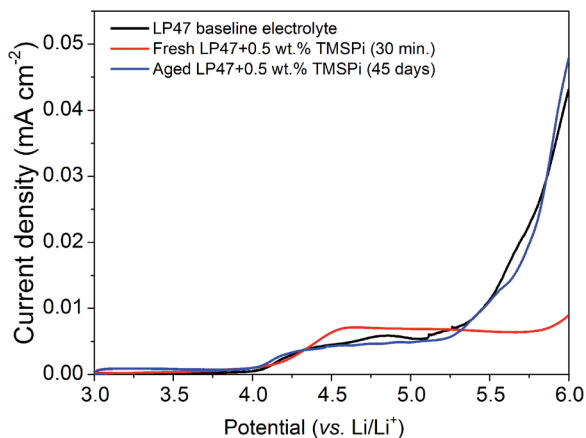


Fig. 4 Oxidation stability determination of the baseline LP47 electrolyte (black line), the fresh LP47 electrolyte containing TMSPi (red line) and the aged LP47 electrolyte containing TMSPi (blue line).

of TMSPi with HF or with other compounds, such as water, were not able to form a stable CEI. It is necessary to investigate the electrochemical performance of the aged TMSPi containing electrolyte in a full lithium ion cells to know whether the electrolyte additive still has a positive effect on the cycling stability.

The cycling stability of $\text{LiNi}_{0.5}\text{Co}_{0.2}\text{Mn}_{0.3}\text{O}_2/\text{graphite}$ full cells was investigated in the three electrolytes: the baseline LP47 electrolyte, the fresh TMSPi containing LP47 electrolyte and the aged TMSPi containing LP47 electrolyte (Fig. 5). The cut-off voltage range was set between 2.8 V and 4.6 V. To prove the reproducibility of the cell performance, two cells with the aged TMSPi containing electrolyte are shown in Fig. 5.

The reasons behind the fast capacity fading of the $\text{LiNi}_{0.5}\text{Co}_{0.2}\text{Mn}_{0.3}\text{O}_2/\text{graphite}$ full cells charged up to 4.6 V had been previously studied. On one hand, the irreversible transformation of the $\text{LiNi}_{0.5}\text{Co}_{0.2}\text{Mn}_{0.3}\text{O}_2$ from the pristine

rhombohedral phase to a mixture of spinel and rock salt phase²² significantly reduced the conductivity of the material. On the other hand, the carbonate based electrolytes could not resist the high cathode potential¹ and experienced a continuous oxidative decomposition. In this work, all investigated cells were charged/discharged within the same cut-off voltage window, so that the deterioration in the cycling stability caused by the phase transformation should be similar in each case. The difference in the cycling performance was thus determined by the electrolyte decomposition. During the first 70 cycles, the cells with fresh and aged TMSPi containing electrolytes showed similar specific discharge capacity (ca. 156 mA h g^{-1}) and capacity fading rate (Fig. 5). The cell with the baseline LP47 electrolyte showed a specific discharge capacity of 143 mA h g^{-1} and a faster capacity fading. This could be explained by the fact that the TMSPi helped to “purify” the electrolyte by removing the HF and H_2O contaminants. However, after 80 cycles, both cells with aged TMSPi containing electrolyte suffered from a faster capacity fading rate than the ones with the fresh TMSPi containing electrolyte. The better cycling stability of the cells with fresh TMSPi containing electrolyte could be attributed to the stable CEI layer formed in the presence of TMSPi. After 120 cycles, the discharge capacity of the cells with the aged TMSPi containing electrolyte decreased to the same level as the cell with the baseline electrolyte, showing that the aged TMSPi containing electrolyte failed to form an effective CEI to prevent further electrolyte oxidation.

Surface analysis by means of XPS was used to characterize the CEI formed by investigated electrolytes. In the P 2p spectra of the $\text{LiNi}_{0.5}\text{Co}_{0.2}\text{Mn}_{0.3}\text{O}_2$ cathode (Fig. 6), the peak at the higher binding energy (ca. 136.8 eV) corresponded to LiPF_x , whereas the broad peak at the lower binding energy (ca. 135 eV) originated from LiPF_xO_y .^{5,23,24}

Regarding the fluorophosphates compounds, the binding energy of the P 2p electrons is highly influenced by the attached elements or groups,²⁵ e.g. PF_xO_y^- has higher binding energy than PO_4^{3-} .^{23,24} Thus, the fluorophosphates showed a broad peak in the P 2p spectra. A comparison between Fig. 6a and b showed that the CEI film, which was formed by the oxidative decomposition of TMSPi in the freshly prepared TMSPi containing electrolyte, contained more fluorophosphates than the CEI obtained by the baseline LP47 and the aged TMSPi containing LP47 electrolytes. The contents of LiPF_xO_y (atomic ratio) in the CEI films formed in different electrolytes are listed in Table 2.

The thickness of the CEI films at the different cathode surfaces was calculated from the obtained XPS measurement results and by means of the theory developed by Niehoff *et al.*^{23,26} As shown in Table 2, the fresh TMSPi containing electrolyte enabled the formation of a CEI film of $10.5 \text{ \AA}$, which was thinner than the CEI formed by decomposition of the other two electrolytes. All three CEI films contained a similar amount of LiPF_x . However, the CEI formed by baseline LP47 had ca. 0.9 at% LiPF_xO_y , while the CEI formed by the fresh TMSPi containing electrolyte possessed as high as ca. 6.7 at% LiPF_xO_y . Based on the better electrochemical performance, the CEI film with more fluorophosphates could be assumed to be more

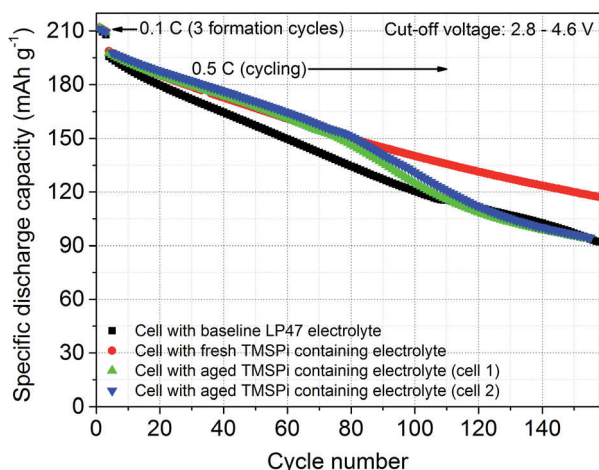


Fig. 5 Cycling stability of $\text{LiNi}_{0.5}\text{Co}_{0.2}\text{Mn}_{0.3}\text{O}_2/\text{graphite}$ full cells in the following electrolytes: baseline LP47 electrolyte (black square); freshly prepared TMSPi containing electrolyte (red dot); aged TMSPi containing electrolyte (green triangle and blue inverted triangle). Cut-off voltage ranging from 2.8 V to 4.6 V.

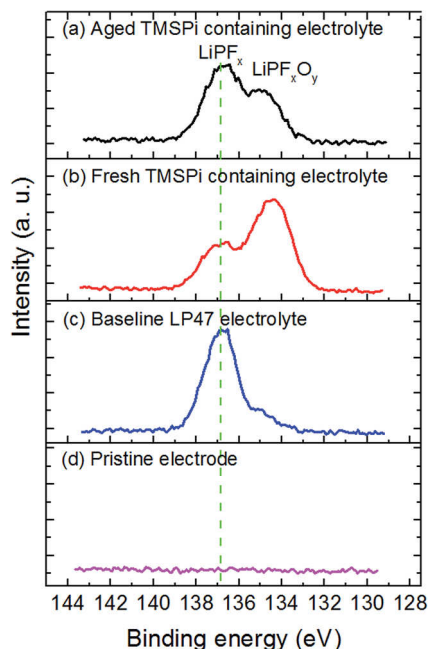


Fig. 6 P 2p spectra of the $\text{LiNi}_{0.5}\text{Co}_{0.2}\text{Mn}_{0.3}\text{O}_2$ cathode (after 150 cycles, discharged) with (a) aged TMSPi-containing electrolyte, (b) fresh TMSPi containing electrolyte, (c) baseline LP47 electrolyte and (d) a pristine electrode without exposure to the electrolyte.

robust and compact and thus effectively diminished the oxidation of the electrolytes.

The thickness of the CEI formed by the aged TMSPi containing electrolyte increased to 21.4 Å, while the amount of fluorophosphates decreased to 1.50 at%. The CEI built by the aged TMSPi containing LP47 electrolyte was less stable during long-term cycling.

Consequently, it is important to investigate the reaction between TMSPi and LiPF_6 to find out why less fluorophosphates remained after aging of the electrolyte. To avoid the side reaction with the carbonate solvents, the TMSPi was directly mixed with LiPF_6 in a molar ratio of 5 : 3, as it was suspected that the TMSPi reacted with PF_5 . The mixture was sealed in plastic vials (polypropylene, Shimadzu®) for the GC-MS measurement and stored in glovebox for two weeks at 20 °C. The product was investigated by NMR. The headspace gas was collected by a syringe and then injected into the GC-MS device. The mass spectrum is shown in Fig. 7. As the EI mode was applied to charge the molecule fragments, the obtained m/z values directly indicated the molecular weight of the gas species.

Table 2 Thickness of the CEI films on $\text{LiNi}_{0.5}\text{Co}_{0.2}\text{Mn}_{0.3}\text{O}_2$ cathodes cycled in different electrolytes, as well as the atomic ratio of the LiPF_x and LiPF_xO_y in each CEI film. "Dev." stands for "Mean deviation"

Electrolyte	Average thickness (Å)	LiPF_x (at%)	Dev. (at%)	LiPF_xO_y (at%)	Dev. (at%)
Baseline LP47	31.6	3.0	0.3	0.9	0.4
LP47 + TMSPi (fresh)	10.5	3.1	0.1	6.7	0.1
LP47 + TMSPi (aged)	21.4	2.6	0.2	1.50	0.05

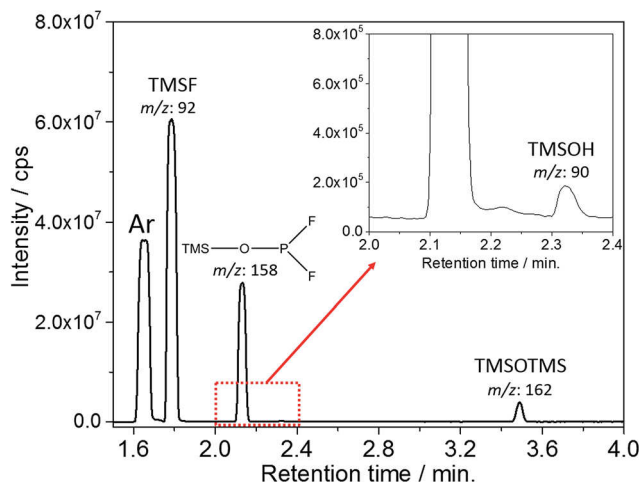


Fig. 7 GC-MS of the headspace gas generated from the reaction between LiPF_6 and TMSPi (molar ratio was 3 : 5).

Besides TMSF, TMSOH and TMSOTMS¹⁸ signals, one more peak with the m/z value of 158 was observed at the retention time of 2.1 min (Fig. 7). According to the possible reaction routines, this peak was assigned to trimethylsilyl difluorophosphinite ((TMSO)PF₂), which was firstly synthesized and reported by Cavell *et al.*²⁷ in 1973. In the reaction with LiPF_6 , the POF_2^- groups were obtained from the TMSPi *via* the exchange of the P–O[−] bond and the P–F bond (shown in Scheme 1). The POF_2^- was bound to the TMS groups and formed (TMSO)PF₂, which had a relatively low boiling point²⁷ and could be detected by GC-MS.

The existence of the (TMSO)PF₂ could also be confirmed by the ¹⁹F-NMR spectrum (Fig. 8a) of the reaction product of LiPF_6 and TMSPi. A doublet at −33.5 ppm in ¹⁹F-NMR spectrum and a triplet at 113.2 ppm in ³¹P-NMR with ¹J_{PF} coupling constant of 1032 Hz (Fig. 8b) were in agreement with previously reported NMR data.²⁷ Furthermore, two sets of doublets in ¹⁹F-NMR spectrum (Fig. 8a) at −84.0 ppm (broad signals, 1 and 1') and −73.3 ppm (2 and 2') with ¹J_{PF} coupling constant of 980 Hz and 925 Hz, respectively, could be assigned to the formation of (TMSO)POF₂ and (TMSO)₂POF₂²⁸ (structure shown in Fig. 8d). TMSF and a very small amount of LiPF_6 (3 and 3') were detectable in the ¹⁹F-NMR spectrum (Fig. 8a). The corresponding signals also appeared in the ³¹P-NMR spectrum (Fig. 8b) at −17.8 ppm for (TMSO)POF₂ as a broad triplet (1, 1' and 1'') and at −27.1 ppm for (TMSO)₂POF₂ as a doublet (2 and 2'). Besides, the ³¹P-NMR spectrum confirmed the formation of tris(trimethylsilyl) phosphate ((TMSO)₃PO) and tetrakis(trimethylsilyl) diphosphate ester ((TMSO)₂OPOPO(OTMS)₂), structure shown in Fig. 8d) as a singlet at −27.3 ppm (Fig. 8b, signal 3) and −30.5 ppm (Fig. 8b, signal 4), respectively.²⁹ A small amount of (TMSO)₂PHO (Fig. 8c, structure shown in Fig. 8d) was also produced in the reaction and could be assigned to a doublet in ¹H-NMR spectrum at 6.8 ppm (1 and 1') with a ¹J_{PH} coupling constant of 714 Hz which was in correlation with a doublet in the ³¹P-NMR spectrum at −13.6 ppm (5 and 5').²⁹

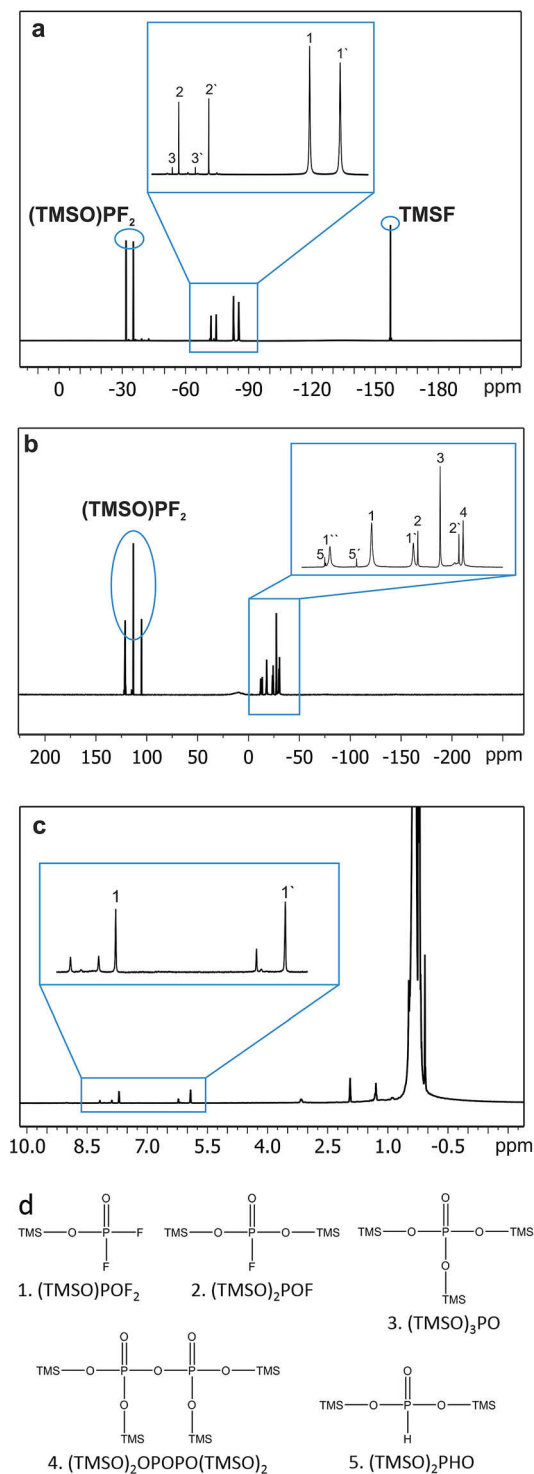


Fig. 8 NMR spectra of the reaction products of LiPF_6 and TMSPi. (a) ^{19}F -NMR: 1 and 1' $(\text{TMSO})\text{POF}_2$; 2 and 2' $(\text{TMSO})_2\text{POF}$. (b) ^{31}P -NMR: 1, 1' and 1'' $(\text{TMSO})\text{POF}_2$; 2 and 2' $(\text{TMSO})_2\text{POF}$; 3 $(\text{TMSO})_3\text{PO}$; 4 $(\text{TMSO})_2\text{OPOP}(\text{OTMS})_2$; 5 and 5' $(\text{TMSO})_2\text{PHO}$. (c) ^1H -NMR: 1 and 1' $(\text{OTMS})_2\text{PHO}$. (d) The schematic structure of the compounds found in the NMR analysis.

Based on the products observed by GC-MS and NMR, it is clearly shown that various TMS containing fluorophosphates have been formed. As a result, the polarity and reactivity of these

fluorophosphates dramatically decreased and could not take part in the formation of an effective CEI film. The mechanism of the reaction between TMSPi and LiPF_6 is proposed and presented in Scheme 1. In order to support the proposed mechanism and reaction steps, computational and theoretical chemistry approaches are in frame of our ongoing investigations. Not all the products are mentioned in Scheme 1, while only the formation of $(\text{TMSO})\text{PF}_2$ and $(\text{TMSO})\text{POF}_2$ as two key products are taken in consideration.

As the binding energy of Si-F bond is higher than the Si-O bond,¹⁸ the TMS groups in the TMSPi will react with the F^- from LiPF_6 , resulting in the formation of TMSF and P-O $^-$ bond. Afterwards, an exchange reaction between P-O $^-$ bond and P-F bond in PF_5 results in the formation of the highly reactive POF_3 . PF_2O_2^- (PF_xO_y^-) can be formed by a further exchange reaction between the P-O $^-$ bond and P-F (from POF_3). Due to the high anion affinity of the TMS groups, the TMSPi could react with various PF_xO_y^- moieties and generate the products illustrated in Fig. 8, e.g. the $(\text{TMSO})\text{POF}_2$. The major product of the reaction between TMSPi and LiPF_6 after two weeks is TMSOPF_2 , which has been detected by GC-MS analysis (Fig. 7). Therefore, the PO_3^{3-} group from TMSPi is consumed by the reaction with LiPF_6 and do not take part in the formation of the CEI. In the aged TMSPi containing electrolyte, the CEI film contains less fluorophosphate components than the CEI formed in the freshly prepared TMSPi containing electrolyte due to the formation of various TMS containing phosphates and fluorophosphates. Moreover, the concentration of LiPF_6 is also diminished by this side reaction with TMSPi, resulting in the loss of charge carriers in the electrolyte.

Conclusions

A freshly prepared TMSPi containing LP47 electrolyte (1 M LiPF_6 in a mixture of EC/DEC 3 : 7 (weight ratio)) shows a significant improvement to the cycling stability of the $\text{LiNi}_{0.5}\text{Co}_{0.2}\text{Mn}_{0.3}\text{O}_2$ /graphite lithium ion cells charged to 4.6 V. Due to its high anion affinity, the TMS group diminishes the HF and water content in the investigated electrolyte. A thin and robust CEI which contains more fluorophosphates can be formed in presence of TMSPi. From this point of view, TMSPi can be applied as an effective electrolyte additive.

However, after aging at 20 °C for 4 weeks, no TMSPi can be detected in the electrolyte anymore. It has been confirmed that TMSPi facilitates the decomposition of the electrolyte salt LiPF_6 , which reduces the content of the conducting lithium salt in the electrolyte. The fluorophosphates generated from the decomposition of LiPF_6 are captured by the highly electrophilic TMS groups, and thus cannot form a stable CEI anymore. Based on the reaction products between LiPF_6 and TMSPi, a failure mechanism of the TMSPi in the LiPF_6 containing carbonate electrolytes has been proposed. In conclusion, TMSPi has certain lifetime limit when being used as an additive in the LiPF_6 based carbonate electrolytes, which means the TMSPi containing electrolyte can only be used within a short time after the addition of TMSPi. Its beneficial effect of improving the cycling stability of the cells deteriorates upon aging.

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References

- 1 R. Wagner, N. Preschitschek, S. Passerini, J. Leker and M. Winter, *J. Appl. Electrochem.*, 2013, **43**, 481–496.
- 2 S. Y. Ha, J. G. Han, Y. M. Song, M. J. Chun, S. I. Han, W. C. Shin and N. S. Choi, *Electrochim. Acta*, 2013, **104**, 170–177.
- 3 E. Wang, D. Ofer, W. Bowden, N. Ilchev, R. Moses and K. Brandt, *J. Electrochem. Soc.*, 2000, **147**, 4023–4028.
- 4 M. Grutzke, V. Kraft, B. Hoffmann, S. Klamor, J. Diekmann, A. Kwade, M. Winter and S. Nowak, *J. Power Sources*, 2015, **273**, 83–88.
- 5 K. Edstrom, T. Gustafsson and J. O. Thomas, *Electrochim. Acta*, 2004, **50**, 397–403.
- 6 D. R. Gallus, R. Schmitz, R. Wagner, B. Hoffmann, S. Nowak, I. Cekic-Laskovic, R. W. Schmitz and M. Winter, *Electrochim. Acta*, 2014, **134**, 393–398.
- 7 M. Winter, *Z. Phys. Chem.*, 2009, **223**, 1395–1406.
- 8 D. R. Gallus, R. Wagner, S. Wiemers-Meyer, M. Winter and I. Cekic-Laskovic, *Electrochim. Acta*, 2015, **184**, 410–416.
- 9 X. X. Zuo, C. J. Fan, J. S. Liu, X. Xiao, J. H. Wu and J. M. Nan, *J. Electrochem. Soc.*, 2013, **160**, A1199–A1204.
- 10 Z. J. Cai, Y. B. Liu, J. H. Zhao, L. Li, Y. M. Zhang and J. Zhang, *J. Power Sources*, 2012, **202**, 341–346.
- 11 D. J. Lee, D. Im, Y. G. Ryu, S. Lee, J. Yoon, J. Lee, W. Choi, I. Jung, S. Lee and S. G. Doo, *J. Power Sources*, 2013, **243**, 831–835.
- 12 A. Wursig, H. Buqa, M. Holzapfel, F. Krumeich and P. Novak, *Electrochem. Solid-State Lett.*, 2005, **8**, A34–A37.
- 13 J. Demeaux, D. Lemordant, M. Caillon-Caravanier, H. Galiano and B. Claude-Montigny, *Electrochim. Acta*, 2013, **89**, 163–172.
- 14 Z. D. Li, Y. C. Zhang, H. F. Xiang, X. H. Ma, Q. F. Yuan, Q. S. Wang and C. H. Chen, *J. Power Sources*, 2013, **240**, 471–475.
- 15 S. W. Mai, M. Q. Xu, X. L. Liao, J. N. Hu, H. B. Lin, L. D. Xing, Y. P. Liao, X. P. Li and W. S. Li, *Electrochim. Acta*, 2014, **147**, 565–571.
- 16 Y. M. Song, J. G. Han, S. Park, K. T. Lee and N. S. Choi, *J. Mater. Chem. A*, 2014, **2**, 9506–9513.
- 17 N. N. Sinha, J. C. Burns and J. R. Dahn, *J. Electrochem. Soc.*, 2014, **161**, A1084–A1089.
- 18 Y. K. Han, J. Yoo and T. Yim, *J. Mater. Chem. A*, 2015, **3**, 10900–10909.
- 19 E. L. Myers, C. P. Butts and V. K. Aggarwal, *Chem. Commun.*, 2006, 4434–4436.
- 20 C. L. Campion, W. T. Li and B. L. Lucht, *J. Electrochem. Soc.*, 2005, **152**, A2327–A2334.
- 21 N. S. Choi, J. G. Han, S. Y. Ha, I. Park and C. K. Back, *RSC Adv.*, 2015, **5**, 2732–2748.
- 22 S. K. Jung, H. Gwon, J. Hong, K. Y. Park, D. H. Seo, H. Kim, J. Hyun, W. Yang and K. Kang, *Adv. Energy Mater.*, 2014, **4**, 1300787.
- 23 P. Niehoff and M. Winter, *Langmuir*, 2013, **29**, 15813–15821.
- 24 E. Fluck and D. Weber, *Z. Naturforsch., B: Anorg. Chem., Org. Chem.*, 1974, **29**, 603–607.
- 25 M. Pelavin, D. N. Hendrickson, J. M. Hollander and W. L. Jolly, *J. Phys. Chem.*, 1970, **74**, 1116–1121.
- 26 P. Niehoff, S. Passerini and M. Winter, *Langmuir*, 2013, **29**, 5806–5816.
- 27 R. G. Cavell, R. D. Leary, A. R. Sanger and A. j. Tomlinso, *Inorg. Chem.*, 1973, **12**, 1374–1380.
- 28 R. G. Cavell, A. j. Tomlinso and R. D. Leary, *Inorg. Chem.*, 1972, **11**, 2573–2578.
- 29 L. A. Wessjohann and M. A. Dessoy, *Polyhedron*, 2014, **70**, 133–137.