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Structure, preparation and properties of liquid fluoroelastomers with different end groups

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In order to design and prepare liquid fluoroelastomers with different end groups, and reveal the relationship between the molecular chain structure and properties, we studied on the oxidation degradation method and functional group conversion method to prepare carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers, respectively. The reaction mechanisms were also deduced. Furthermore, the curing system was created for liquid fluoroelastomers, and systematically analyzed their properties. The sequence type and content of the $\text{C}=\text{C}$ and oxygen-containing groups in the samples were measured and characterized by attenuated total reflectance/Fourier transform infrared (ATR-FTIR) spectroscopy, ^1H nuclear magnetic resonance (^1H -NMR), ^{19}F -NMR spectroscopy and chemical titration, the molecular weights of liquid fluoroelastomers were measured by gel permeation chromatography (GPC). Differential scanning calorimetry (DSC) and thermogravimetric analysis (TGA) were used to examine the thermal properties, while a viscometer was used to measure the dynamic viscosity of the liquid fluoroelastomers. Then the mechanical and surface properties of the cured samples were examined by universal testing machine and contact angle measurement instrument, respectively. The results show that carboxyl-terminated liquid fluoroelastomer with 2.71 wt% carboxyl terminal groups can be prepared by oxidation degradation method. When lithium aluminium hydride (LiAlH_4) was used as the reducing agent, it can efficiently convert carboxyl group to hydroxyl group with a conversion rate of more than 95%. In addition, it can be seen that the dynamic viscosity of the liquid fluoroelastomers were all decreased with the increase of temperature, and it is similar to about 10 Pa s at 70 °C. Compared with carboxyl-terminated liquid fluoroelastomers, hydroxyl-terminated liquid fluoroelastomers has higher curing reactivity, higher glass transition temperature (T_g) and thermal decomposition temperature (T_d), and better mechanical properties of cured samples. The two types of liquid fluoroelastomers with distinct end groups presented distinct hydrophilicity.

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1. Introduction

Fluoroelastomers are synthetic polymer elastomers containing fluorine atoms on the main-chain or side-chain carbon atoms.¹ They possess exceptional thermal stability, oil resistance, oxidation resistance and chemical resistance due to the high bond energy (485 kJ mol^{-1}) of the C-F bond and the shielding effect of the fluorine atoms on the $\text{C}=\text{C}$ bond in its molecular structure.^{2–5} However, they do not fully meet the requirements of modern fluoroelastomer transport devices because of the poor processing performance, complicated processing and complex vulcanisation procedure.^{6,7} As a result, low-molecular-weight fluoroelastomers with improved fluidity and plasticity, known as liquid fluoroelastomers, were developed.^{8,9}

Liquid fluoroelastomers, also known as key materials, has a wide range of potential applications in transportation, medical protection, new energy, national defence and military industries due to their unique properties. They can also be utilised to make fluorine-containing functional polymers, novel battery-sealing materials, raw materials for additive manufacturing (3D printing), high-performance adhesives, caulking agents, coatings and processing complexes.

Liquid fluoroelastomers can be prepared not only by polymerisation methods^{10,11} but also through oxidation degradation method.¹² Because the oxidation degradation method is simpler than the polymerisation methods and produces carboxyl-terminated telechelic polymers *in situ*, it has attracted wide attention.¹³ Unfortunately, due to their low carboxyl-terminated activity and poor thermal stability, carboxyl-terminated liquid fluoroelastomers are difficult to cure at low temperatures, thus limiting their applicability in particular fields. As a result, carboxyl to hydroxyl group conversion is important for

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enhancing the thermal stability of liquid fluoroelastomers and for lowering the curing temperature.

In organic chemistry, carboxyl group reduction is a crucial process. Since Finholt¹⁴ discovered the preparation and application of LiAlH_4 in 1946, complex metal hydrides have been used as reducing agents in organic synthesis with considerable success. They were swiftly developed and extensively employed, and the number of reagents has gradually expanded in recent years. These reductants exist in the form of complex hydrogen anion salts, which are nucleophiles that can attack the positively charged atoms in unsaturated polar bonds, resulting in a reduction process via hydrogen anion transfer.^{15,16} The carboxyl group can be easily converted to a hydroxyl group simultaneously.

Furthermore, liquid fluoroelastomers themselves have no strength and cannot meet the current research and application requirements, so they must be cured. Epoxy,^{17,18} isocyanate,¹⁹ aziridine²⁰ and carbonised diamines are the most common curing agents used for liquid fluoroelastomers.²¹ Hexamethylene diisocyanate (HDI) is a particularly important aliphatic isocyanate monomer because its molecular chain contains a highly active isocyanate group ($-\text{NCO}$) that can react with active hydrogen groups, such as carboxyl, hydroxyl, amino, carbamate and urea, in the polymer to achieve crosslinking and obtain the desired network structure.^{22,23} Because HDI curing agents do not contain isocyanate groups directly attached to the carbon atoms on the benzene ring, the coatings and adhesives cured by HDI have a series of excellent properties, including weather resistance, chemical reagent resistance, yellowing resistance and powder resistance.^{24,25} The low surface energy of fluoroelastomers has made them popular hydrophobic materials.^{26,27}

At present, there are few reports on the preparation methods, reaction mechanism, structure and properties of liquid fluoroelastomers with carboxyl and hydroxyl end groups. Thus, in order to design and prepare liquid fluoroelastomers with different end groups, and reveal the relationship between the molecular chain structure and properties, we studied on the oxidation degradation method and functional group conversion method to prepare carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers, respectively. The poly(VDF-co-HFP) copolymer was chosen as the raw material, the fluoroelastomer formation was confirmed based on the results of spectral analyses and chemical quantification. Furthermore, property tests and comparative analysis of the relationship between the structure and properties of the liquid fluoroelastomers with different end groups were performed.

2. Experimental details

2.1 Materials and chemical agents

The fluoroelastomer that was made of the copolymer of vinylidene fluoride and hexafluoropropene was purchased from Chenguang Research Institute of Chemical Industry. Carboxyl-terminated liquid fluoroelastomer was prepared by the oxidation degradation method of the fluoroelastomer. The hydroxyl-terminated liquid fluoroelastomer was prepared by the

functional group conversion method of the carboxyl-terminated liquid fluoroelastomer. Benzyl triethyl ammonium chloride (BTEAC) and LiAlH_4 were purchased from Aladdin Industrial Corporation. HDI was purchased from Yantai Wanhua Chemical Group Co., Ltd. Potassium hydroxide (KOH), hydrogen peroxide (30 wt% H_2O_2), hydrochloric acid (HCl), anhydrous ethanol ($\text{C}_2\text{H}_5\text{OH}$) and tetrahydrofuran (THF) were purchased from Tianjin Kemiou Chemical Reagent Co., Ltd.

2.2 Characterization

2.2.1 Spectrum analyses. Attenuated total reflectance/Fourier transform infrared (ATR-FTIR) measurements were performed using the PerkinElmer spectrum one instrument. The FTIR spectra were obtained at the resolution of 4 cm^{-1} in the range $650\text{--}4000\text{ cm}^{-1}$.

The NMR spectra were recorded on a Bruker AC 80 spectrometer (500 MHz for ^1H , 470 MHz for ^{19}F) at room temperature using acetone- d_6 as the solvent and TMS (or CFCl_3) as the internal standard for the ^1H (or ^{19}F) nuclei.

2.2.2 Determination of molecular weight. The molecular weights (M_n) of the fluoroelastomers were estimated using an integrated gel permeation chromatography (GPC) system (PL-GPC50) from A Varian, Inc. Company. PS was employed as the standard, and tetrahydrofuran (HPLC) was used as the mobile phase. The flow rate was 1 mL min^{-1} , and the detector temperature was $30\text{ }^\circ\text{C}$.

2.2.3 Chemical titration. The carboxylic functions of the different liquid fluoroelastomers were titrated using a $\text{KOH}-\text{C}_2\text{H}_5\text{OH}$ ($C = 0.05\text{ mol L}^{-1}$) solution according to the China National Standards GB/T 603-2002 and GB/T601-2002.

The carboxyl-terminated or hydroxyl-terminated liquid fluoroelastomer ($m = 0.5\text{ g}$) and THF (50 mL) were added into an Erlenmeyer flask. Then, phenolphthalein was added (0.01 mL), and the $-\text{COOH}$ (ω) content was assessed by the following equation:

$$\omega = \frac{V \times C \times M_{\text{COOH}}}{m} \times 100\% \quad (1)$$

In eqn (1), “ V ” stands for the volume of the base employed for titration. Furthermore, the conversion rate of the carboxyl group was assessed by the following equation:

$$\alpha = \frac{\omega_0 - \omega_1}{\omega_0} \times 100\% \quad (2)$$

In eqn (2), ω_0 and ω_1 stand for the contents of $-\text{COOH}$ in the carboxyl-terminated liquid fluoroelastomer and the hydroxyl-terminated liquid fluoroelastomer, respectively.

2.2.4 Thermal properties. The glass transition temperatures of the fluoroelastomers were estimated by Differential Scanning Calorimetry (DSC). The DSC measurements were carried out on a NETZSCH DSC200. The samples were heated from -100 to $+30\text{ }^\circ\text{C}$ at a rate of $5\text{ }^\circ\text{C min}^{-1}$ to record the curves. The temperature at half-height of the corresponding heat capacity jump was defined as T_g .



The thermal decomposition temperatures of the fluoroelastomers were estimated by thermogravimetry analysis (TGA). The TGA measurements were carried out on a NETZSCH TG209. The samples were heated from 30 °C to 600 °C at a rate of 10 °C min⁻¹ to record the curves.

2.2.5 Dynamic viscosities. The dynamic viscosities of the fluoroelastomers were tested on a DV2T-RV Brookfield viscometer, according to GB/T 2794-2013. The test was carried out in the range of 30–70 °C, and the average value of 3 measurements was calculated for each sample. The absolute viscosity at 50 °C where it is not stated in the text.

2.2.6 Mechanical properties. Mechanical properties of the fluoroelastomers were estimated by universal testing machine which were carried out on INSTRON 3365. The tensile strength test was carried out in accordance with the national standard GB/T528-82 "Determination of the tensile properties of vulcanized rubber"; the test temperature was 23 ± 2 °C, and the tensile rate was 500 mm min⁻¹; for each group, 5 samples were tested, and the average was calculated. The hardness was tested with an LX-A digital Shore hardness tester.

2.2.7 Surface properties. The surface properties of the fluoroelastomers were estimated by FM40Mk2 for contact angle measurement instrument. The contact angle was measured on the cured product surfaces with a test water volume of 5 µL; 5 different points were selected for each sample tested, and the average value of the 5 points was considered.

2.3 Preparation

The carboxyl-terminated liquid fluoroelastomer was prepared by the oxidative degradation of the poly(VDF-co-HFP) copolymer by using an aqueous KOH solution and H₂O₂. The hydroxyl-terminated liquid fluoroelastomer was prepared by a reductive reaction, in which the carboxyl-terminated liquid fluoroelastomer was reacted with LiAlH₄. Then, these products were cured.

2.3.1 Carboxyl-terminated liquid fluoroelastomer. A 2000 mL flask was filled with the fluoroelastomer (100 g) and THF (700 mL). At 10 °C, benzyltriethylammonium chloride (BTEAC, 3 g), 30 wt% H₂O₂ aqueous solution and 45 wt% KOH aqueous solution were sequentially added to the flask and agitated for 10 h. After the reaction, the organic and inorganic phases were separated; concentrated hydrochloric acid and water were added to the organic phase, and the pH was corrected to 1–2. The product was collected and dried to a constant weight at 65–70 °C in a vacuum drying oven, finally producing a light-yellow viscous liquid.

2.3.2 Hydroxyl-terminated liquid fluoroelastomer. THF (100 mL) and the carboxyl-terminated liquid fluoroelastomer (10 g) were added to a 250 mL flask. LiAlH₄ was gently added at room temperature until the fluoropolymer dissolved fully; then, the temperature was elevated to the appropriate value, and the mixture reacted for 3 h. The reaction was then quenched with 20 mL hydrochloric acid at a concentration of 2.0 mol L⁻¹, and the product was rinsed several times with deionised water. Finally, the product was collected and vacuum-dried at 60–65 °C to a consistent weight, yielding a white viscous liquid.

2.3.3 Curing of the carboxyl-terminated liquid fluoroelastomer with HDI. HDI trimer with a COOH/NCO molar ratio of 1.0/1.0–1.0/1.5 was dissolved in acetone (2 mL), and the carboxyl-terminated liquid fluoroelastomer (10 g) was also dissolved in acetone (10 mL). To eliminate the solvent, they were evenly mixed and placed in a drying oven set to 60 °C. The solution was poured into a mould that had been preheated to 60 °C and placed in a vacuum drying box at 60 °C for 8–48 h to remove the residual solvent. The temperature was increased to the curing point temperature of 102 °C after the solvent was removed. A dark yellow cured product was formed after a 6 h reaction.

2.3.4 Curing of the hydroxyl-terminated liquid fluoroelastomer with HDI trimer. HDI trimer with a OH/NCO molar ratio of 1.0/1.0–1.0/1.5 was dissolved in acetone (2 mL), and hydroxyl-terminated liquid fluoroelastomer (10 g) was also dissolved in acetone (10 mL). To eliminate the solvent, they were evenly mixed and then placed in the drying oven set to 60 °C. When the remaining solvent was about 3–5 mL, the mixture was poured into a mould preheated at 60 °C and placed in a vacuum drying oven at 60 °C for 8–48 h to remove the remaining solvent. After removing the solvent, the temperature was raised to 90 °C, which was the solidification point, and the reaction was carried out for 4 h, yielding a light-yellow solidified product.

3. Results and discussion

3.1 Structure of the liquid fluoroelastomers

3.1.1 FTIR of the liquid fluoroelastomers. All the FTIR spectra revealed absorption peaks at about 870–890 cm⁻¹, 1155–1180 cm⁻¹ and 1395–1400 cm⁻¹, which could be ascribed to the stretching vibrations of CF₃, CF₂ and CF, respectively, as shown in Fig. 1.²⁸ There is a new peak belonging to the carboxyl group (–CF₂COOH) at 1760–1770 cm⁻¹ of carboxyl-terminated liquid fluoroelastomer (B) indicate that the oxidation degradation method can be used to successfully prepare the carboxyl-terminated liquid fluoroelastomer. From the characteristic peak of the carboxyl group at 1760–1770 cm⁻¹ weakened significantly with the progress of the reduction reaction and an obvious characteristic hydroxyl peak appeared at 3350–3450 cm⁻¹, demonstrating the noticeable reduction of –CF₂COOH when LiAlH₄ was used.

3.1.2 ¹H-NMR of liquid fluoroelastomers. As illustrated in Fig. 2, the characteristic structural peaks of the carboxyl-terminated liquid fluoroelastomer at δ = 1.55 ppm, 4.68 ppm and 7.70–7.50 ppm corresponded to –CF=C(CF₃)–CH₂–, –(CF₃)C=CH– and –CH=CF–, respectively. This demonstrates that the fluoroelastomer underwent an oxidation degradation reaction, with the dehydrofluorination reaction producing the –C=C– bond under alkaline conditions.²⁹

As shown in Fig. 3, compared with the carboxyl-terminated liquid fluoroelastomer, the hydroxyl-terminated liquid fluoroelastomer showed not only the characteristic structural peaks of –CH₂OH at δ = 3.63 ppm and 3.75 ppm but also peaks at δ = 1.55 ppm, 4.68 ppm and 7.70–7.50 ppm. The corresponding –C=C– characteristic structural peaks were significantly weakened, indicating that under the action of LiAlH₄, the carboxyl



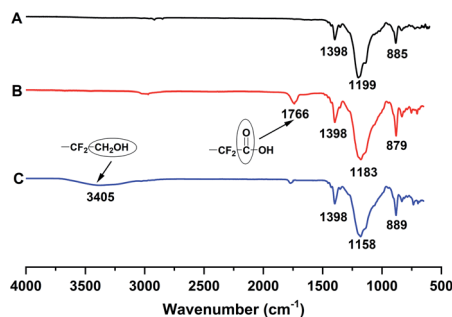


Fig. 1 FTIR spectra of the (A) poly(VDF-co-HFP) copolymer, (B) carboxyl-terminated liquid fluoroelastomer and (C) hydroxyl-terminated liquid fluoroelastomer.

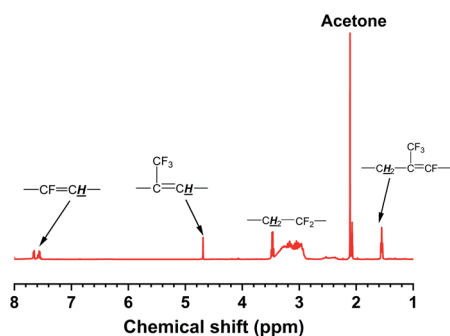


Fig. 2 ^1H -NMR spectrum of the carboxyl-terminated liquid fluoroelastomer.

groups and $\text{C}=\text{C}$ in the carboxyl-terminated liquid fluoroelastomer were all reduced.

3.1.3 ^{19}F -NMR of liquid fluoroelastomers. When ^{19}F -NMR spectroscopy was used for further characterisation, as shown in Fig. 4, a peak at -63.46 ppm ascribed to $-\text{CF}_2\text{COOH}$ could be seen. The multiplets at -70 to 76 ppm (b and d), -118 to 120 ppm (o) and -181 to 185 ppm (p and q) were assigned to $-\text{CF}_3$, $-\text{CF}_2$ and $-\text{CF}$ from the HFP sequences, respectively; the multiplets at -91 to 117 ppm (g–n) were assigned to $-\text{CF}_2$ from the VDF sequences.^{30–33} The detailed attribution is shown in Table 1.

Then, setting $\sum_1\text{C}=\text{C}$ as the integral of the $-\text{C}=\text{C}-$ sequences, it was calculated as $\sum_1\text{C}=\text{C} = 2/3(I_{-73.71} + I_{-80.66} + I_{-81.30})$; $\sum_1(\text{CF}_3 + \text{CF}_2)$ was the integral of the other VDF and HFP sequences, $\sum_1(\text{CF}_3 + \text{CF}_2) = 2/3(I_{-70.67} + I_{-75.19}) + I_{-63.46} + I_{-91.62} + I_{-93.56} + I_{-95.64} + I_{-103.62} + I_{-108.96} + I_{-110.51} + I_{-112.53} + I_{-113.95} + I_{-116.24}$. The molar fractions (mol%) of $-\text{C}=\text{C}-$ was calculated by the following equation.³⁴

mol% of $-\text{C}=\text{C}-$ in the copolymer:

$$X = \frac{\sum_1\text{C}=\text{C}}{\sum_1\text{C}=\text{C} + \sum_1(\text{CF}_3 + \text{CF}_2)} \times 100\% \quad (3)$$

According to the calculations, the molar fractions of $-\text{C}=\text{C}-$ in the carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers were about 1.12 mol% and 0.89 mol%, respectively.

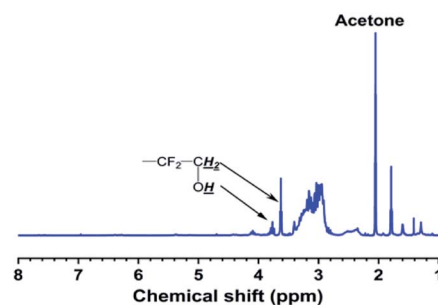


Fig. 3 ^1H -NMR spectrum of the hydroxyl-terminated liquid fluoroelastomer.

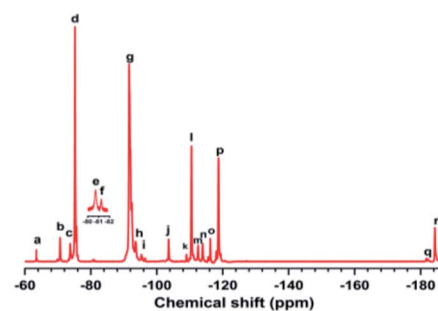


Fig. 4 ^{19}F -NMR spectrum of the carboxyl-terminated liquid fluoroelastomer.

When carboxyl was converted to hydroxyl, not only did the peak assigned to $-\text{CF}_2\text{COOH}$ (a) significantly decline and the peaks assigned to $-\text{C}=\text{C}-$ vanish, but the peak centred at -104.91 ppm(s) belonging to the $-\text{CF}_2\text{CH}_2\text{OH}$ structure also appeared, as seen in Fig. 5.³⁵ As a result, when carboxyl was converted to hydroxyl, the $-\text{C}=\text{C}-$ content decreased as well.

Thus, the successful preparation of the carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers was confirmed, and the characterisation results matched the FTIR and ^1H -NMR data.

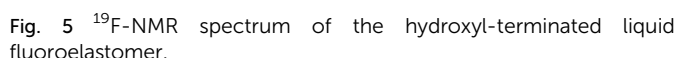
3.1.4 Reaction mechanism. According to the FTIR, ^1H -NMR and ^{19}F -NMR results, the reaction mechanisms of the formation of carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers were systematically analysed.

In the alkaline environment of the fluoroelastomer, the hydroxyl ion (OH^-) in KOH will attack the hydrogen ion (H^+) in the structure of $-\text{CF}_2\text{CH}_2-$, and a deprotonation reaction occurs, followed by a defluorination reaction to produce $-\text{C}=\text{C}-$, as illustrated in Scheme 1. Furthermore, when there is a high concentration of alkali or a high temperature in the reaction system, oxygen in the air, the alkali and fluorine continue to interact with $-\text{C}=\text{C}-$, resulting in an oxidation reaction. Subsequently, a hydroxyl group is introduced into the molecular chain, thus resulting in the carboxyl-terminated liquid fluoroelastomer.^{36,37}

There are two stages in the reduction of the carboxyl group by LiAlH_4 . The carboxyl group is reduced to an aldehyde in the first stage. The carboxyl group is first converted into a carboxylate lithium salt, and then when aluminium hydride (AlH_3) is



No.	δ (ppm)	Assignment	No.	δ (ppm)	Assignment
a	-63.46	-CF ₂ C _F ₂ COOH	j	-103.62	-CF ₂ CH ₂ C _F ₂ CF(CF ₃)CF ₂ -
b	-70.67	-CH ₂ CF ₂ CF(C _F ₃)CF ₂ CH ₂ -	k	-108.96	-CF(CF ₃)CH ₂ C _F ₂ CF(CF ₃)CF ₂ -
c	-73.71	-CF ₂ CH=C(C _F ₃)CF ₂ -	l	-110.51	-CF ₂ CH ₂ C _F ₂ CF ₂ CF(CF ₃)-
d	-75.19	-CF ₂ CH ₂ CF(C _F ₃)CF ₂ CF ₂ -	m	-112.53	-CF(CF ₃)CH ₂ C _F ₂ CF ₂ CH ₂ -
e	-80.66	-CH=CFCF(C _F ₃)-	n	-113.95	-CF ₂ CH ₂ C _F ₂ CF ₂ CH ₂ -
f	-81.30	-CF=CHCF(C _F ₃)CF ₂ -	o	-116.24	-CH ₂ CH ₂ C _F ₂ CF ₂ CF(CF ₃)-
g	-91.62	-CF ₂ CH ₂ C _F ₂ CH ₂ CF ₂ -	p	-118.72	-CH ₂ CF ₂ C _F ₂ CF(CF ₃)CH ₂ -
h	-93.56	-CF ₂ CH ₂ C _F ₂ CH ₂ CF(CF ₃)-	q	-181.74	-CH ₂ CF ₂ C _F (CF ₃)CF ₂ CH ₂ -
i	-95.64	-CH ₂ CH ₂ C _F ₂ CH ₂ CF ₂ -	r	-184.33	-CF ₂ CF ₂ C _F (CF ₃)CH ₂ CF ₂ -



End group	COOH (wt%)	Conversion ratio (%)
Carboxyl	2.71	—
Hydroxyl	0.12	95.57

The above spectral test results show that unlike hydrocarbons or polymers, when LiAlH_4 was utilised as the reducing agent in a fluorine-containing system, the carboxyl group was reduced, while only a portion of the $-\text{C}=\text{C}-$ was reduced. The reaction mechanism is shown in Scheme 2; the electron cloud density is high due to the weak polarity of $-\text{C}=\text{C}-$, which makes it a difficult target for attack by nucleophiles (AlH_4^-), and the electronegativity of the fluorine atom is relatively large, which causes the electron cloud density to shift and reduce, allowing nucleophiles to attack it. The hydride ions are then transferred to positively charged carbon atoms to form complexes, which combine with protons to complete the hydrogenation reduction process.³⁹

The end-group concentration in the carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers was then determined, and the conversion rate in the reduction reaction was nearly 96%. The results are shown in Table 2.



close by, it forms a complex with the carbonyl oxygen. Then, to convert LiAlH_2 to formaldehyde, the negative hydrogen from aluminium is transferred to the carbonyl carbon. The aldehyde



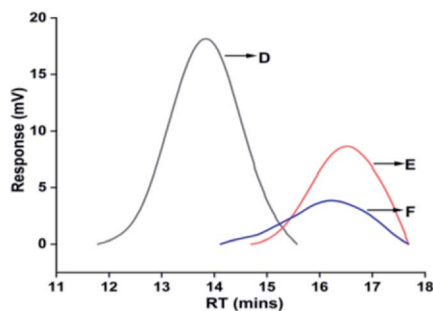


Fig. 6 GPC spectra of the (D) poly(VDF-co-HFP) copolymer, (E) carboxyl-terminated liquid fluoroelastomer and (F) hydroxyl-terminated liquid fluoroelastomer.

Table 3 Assignments of GPC in liquid fluoroelastomers

Sample	M_n	M_w	PD
Poly(VDF-co-HFP) copolymer	71 724	157 904	2.20
Carboxy-terminated fluoroelastomer	2458	4788	1.95
Hydroxy-terminated fluoroelastomer	2680	6904	2.57

3.1.5 GPC of the liquid fluoroelastomers. As demonstrated in Fig. 6, the M_n of the carboxyl-terminated liquid fluoroelastomer prepared by the oxidation degradation reaction was significantly low; however, when carboxyl was converted to hydroxyl, the M_n of the hydroxyl-terminated liquid fluoroelastomer increased, which is due to the simultaneous addition reaction at the $-C=C-$ groups in the molecular chain along with the reduction reaction. The detailed values are shown in Table 3.

3.2 Thermal properties of the liquid fluoroelastomers

3.2.1 Low-temperature property. The cohesive energy density of the fluoroelastomers was higher than those of other polymers due to the strong nature of their structural chemistry and strong polarity; hence, their low-temperature properties were not optimum.^{40,41} Meanwhile, the T_g of the raw material characterises the elastic capabilities of the product under low-temperature settings and reflects the thermal motion ability

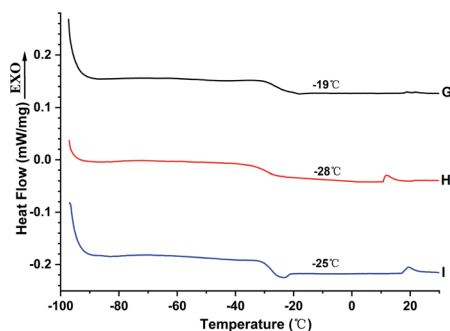


Fig. 7 DSC of the (G) poly(VDF-co-HFP) copolymer, (H) carboxyl-terminated liquid fluoroelastomer and (I) hydroxyl-terminated liquid fluoroelastomer.

of the macromolecular chains. DSC was used to study and analyse the T_g of the raw materials and the liquid fluoroelastomers.

The T_g of the liquid fluoroelastomers generated from the poly(VDF-co-HFP) copolymer fell dramatically from -19°C to -25°C , as illustrated in Fig. 7, with a temperature difference of about -6°C to -9°C . Due to the presence of isolated $-C=C-$ in the molecular chain, the carboxyl-terminated liquid fluoroelastomer generated by the oxidation degradation method showed increased flexibility of the molecular chain, resulting in a drop in T_g . The T_g of the hydroxyl-terminated liquid fluoroelastomer increased from -28°C to -25°C after the reduction reaction, and the temperature contrast was -3°C ; the $-C=C-$ in the molecular chain were reduced to single bonds. As a result, the number of isolated double bonds in the chain were reduced, limiting molecular chain flexibility and raising the T_g . The reason for this, based on the ^{19}F -NMR results in Fig. 4 and 5, is the addition of hydrogen atoms to $-C=C-$ during the reduction reaction, which would improve the tacticity of the molecular chains.^{42,43} The presence of $-C=C-$ in the molecular chains thus had a significant impact on the low-temperature property.

3.2.2 High-temperature stability. The T_d of the liquid fluoroelastomers is critical because the heat resistance of the raw material controls the heat resistance of the product.⁴⁴ In addition, we compared the raw materials with the liquid fluoroelastomers.

The TGA studies showed that the T_d of the liquid fluoroelastomers generated using the poly(VDF-co-HFP) copolymer fell dramatically from 442°C to 235°C , as shown in Fig. 8. It was found that contents of carboxyl and $-C=C-$ were the root causes of this phenomenon. The $-C=C-$ in the molecular chain were reduced to single bonds, and the content of isolated double bonds in the chain decreased, thereby limiting molecular chain flexibility and increasing the T_d . As a result, the T_d of the hydroxyl-terminated liquid fluoroelastomer was enhanced from 235°C to 241°C , according to TGA results. Hence, the content of $-C=C-$ had a significant impact on the T_d . Nevertheless, the TGA results showed that the material was stable and that no chemicals/residues could alter its thermal properties.

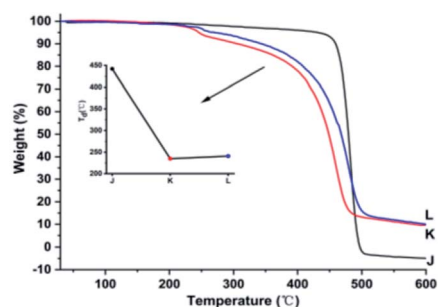


Fig. 8 TGA of the (J) poly(VDF-co-HFP) copolymer, (K) carboxyl-terminated liquid fluoroelastomer and (L) hydroxyl-terminated liquid fluoroelastomer.



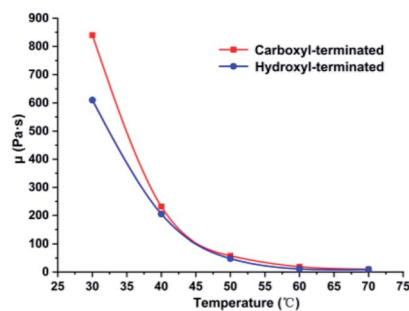


Fig. 9 Dynamic viscosities of the carboxyl-terminated liquid fluoroelastomer and the hydroxyl-terminated liquid fluoroelastomer.

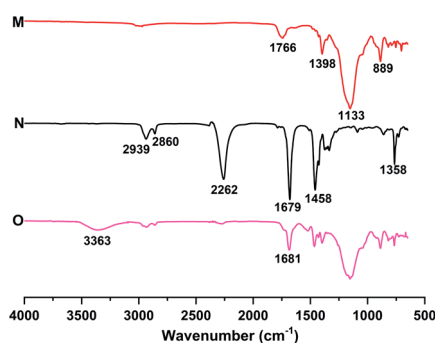


Fig. 10 FTIR spectra of the (M) carboxyl-terminated liquid fluoroelastomer, (N) HDI trimer and (O) cured fluoroelastomer.

3.3 Dynamic viscosities of the liquid fluoroelastomers

At 30–70 °C, the dynamic viscosity of the liquid fluoroelastomers with carboxyl-terminated and hydroxyl-terminated chains was measured, and the findings are presented in Fig. 9.

The dynamic viscosity of the hydroxyl-terminated liquid fluoroelastomer was lower than that of the carboxyl-terminated liquid fluoroelastomer at 30–50 °C, as shown in Fig. 9. Because the carboxyl group has stronger polarity than the hydroxyl group, the intermolecular force of a carboxyl-terminated liquid fluoroelastomer is larger, resulting in higher dynamic viscosity. The dynamic viscosity of the hydroxyl-terminated liquid fluoroelastomer was 48 Pa s at 50 °C, which was lower than the dynamic viscosity of carboxyl-terminated liquid fluoroelastomer, which was 58 Pa s. When the test temperature exceeded 50 °C, the dynamic viscosities of them are both decreases and tends to equalise, around 10 Pa s, and the end group effect on the dynamic viscosity of the liquid fluoroelastomer disappeared.

3.4 Curing of the liquid fluoroelastomers

At room temperature, the liquid fluoroelastomers were viscous liquids or semi-solids with no strength. Thus, they had to be cured before use as curing might impart outstanding mechanical properties, excellent thermal stability, and chemical stability. Therefore, the liquid fluoroelastomers were cured, and the structures of the cured products, as well as the influence of heat and the amount of curing agent on the mechanical and

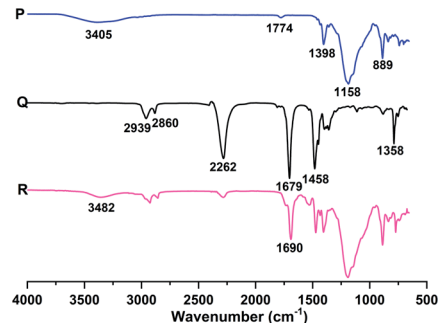
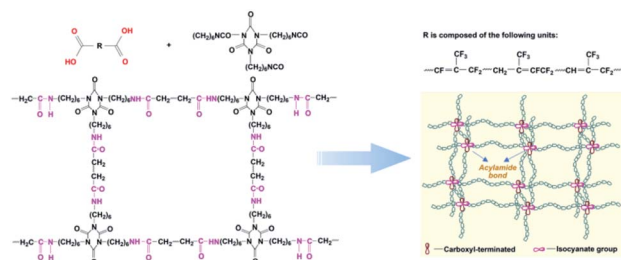


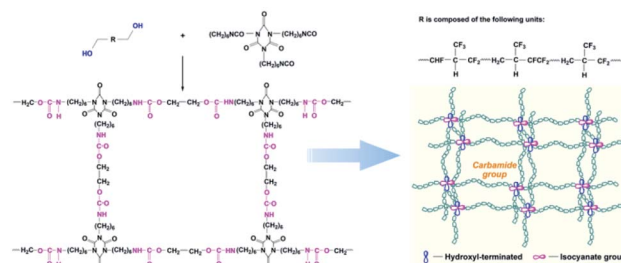
Fig. 11 FTIR spectra of the (P) hydroxyl-terminated liquid fluoroelastomer, (Q) HDI trimer and (R) cured fluoroelastomer.



Scheme 3 Mechanism of curing of the carboxyl-terminated liquid fluoroelastomer.

surface properties of the cured products, were investigated and analysed.

3.4.1 Structure of the cured liquid fluoroelastomers. In the FTIR spectrum, HDI (curve N) presented a stretching vibration peak and an opposite stretching vibration peak corresponding to the C–H at 2939 cm^{−1} and 2860 cm^{−1}, respectively, and an opposite stretching vibration peak corresponding to the –NCO at 2262 cm^{−1}, as illustrated in Fig. 10. The –NCO of the urea bond produced a stretching vibration peak at 1679 cm^{−1}, while the isocyanurate ring generated a stretching vibration peak at 1458 cm^{−1}. The cured carboxyl-terminated product (curve O) featured a new stretching vibration peak of N–H at 3363 cm^{−1}, strongly corresponding with the peak of the C=O at 1681 cm^{−1}. Thus, these demonstrated that HDI could cure the carboxyl-terminated liquid fluoroelastomer.



Scheme 4 Mechanism of curing of the hydroxyl-terminated liquid fluoroelastomers.



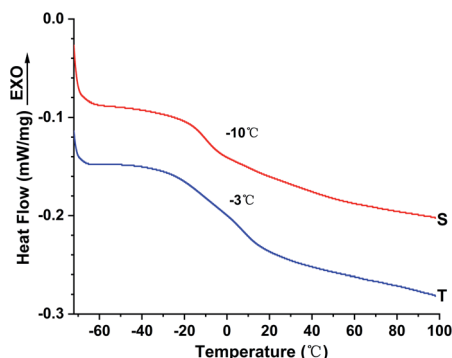


Fig. 12 DSC of the cured products of the (S) carboxyl-terminated liquid fluoroelastomer and (T) hydroxyl-terminated liquid fluoroelastomer.

As illustrated in Fig. 11, the characteristic peak of the hydroxyl group at 3405 cm^{-1} weakened significantly with the progress of the curing reaction, and the stretching vibration peaks corresponding to the N–H and C=O in the carbamide groups (–NHCOO–) structure appeared at 3482 cm^{-1} and 1690 cm^{-1} , respectively. Thus, HDI was shown to cure the hydroxyl-terminated liquid fluoroelastomer.

3.4.2 Mechanism of curing of the liquid fluoroelastomers.

Because of its highly unsaturated structure, the –NCO is highly reactive.⁴⁵ The isocyanate and carboxyl groups produce carboxyl anhydride with poor thermal stability during the curing procedure; it is then decomposed by heat to produce amide bonds and carbon dioxide, thus curing the carboxyl-terminated liquid fluoroelastomer, as shown in Scheme 3.

The –CH₂OH of the hydroxyl-terminated liquid fluoroelastomer comprises a nucleophilic active hydrogen group containing oxygen atoms, which attacks the –NCO carbon atoms, as indicated in Scheme 4. Meanwhile, because the carbon–nitrogen double bond has low bond energy, the active hydrogen may easily interact with the nitrogen atoms in –NCO and create stable –NHCOOH–, allowing the hydroxyl-terminated liquid fluoroelastomer to be cured.

3.4.3 Low-temperature property of the cured liquid fluoroelastomers. The T_g of the cured carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers dramatically

enhanced to -10 °C and -3 °C , respectively, as illustrated in Fig. 12. The molecular chain formed networks after curing, thus reducing the flexibility of the molecular chain of the liquid fluoroelastomer and raising the T_g . Because the hydroxyl-terminated liquid fluoroelastomer had a lower –C=C– content than the carboxyl-terminated liquid fluoroelastomer, its T_g was higher than that of the carboxyl-terminated liquid fluoroelastomer.

3.4.4 High-temperature stability of the cured liquid fluoroelastomers. The T_d of the cured carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers were 189 °C and 288 °C , respectively, as illustrated in Fig. 13. It is obvious that after the curing reaction, T_d shows an upward trend. Because of the low content of –C=C– in the molecular chains of the liquid fluoroelastomers with different end groups, the molecular weight increased after curing, and the molecular chains formed a network structure, which reduced the flexibility of the liquid fluoroelastomer molecular chains, thus increasing their T_d . Because the hydroxyl-terminated liquid fluoroelastomer had lesser –C=C– content than the carboxyl-terminated liquid fluoroelastomer, its T_d was higher than that of the carboxyl-terminated liquid fluoroelastomer.

3.4.5 Mechanical properties of the cured liquid fluoroelastomers. Fig. 14 shows the influence of HDI content on the mechanical properties of the carboxyl-terminated liquid fluoroelastomers at constant curing conditions.

The tensile strength of the carboxyl-terminated liquid fluoroelastomers after curing improved as the HDI proportion increased, and the elongation at break point decreased, as illustrated in Fig. 14. When the COOH/NCO molar ratio was 1.0/1.0, the tensile strength of the cured product was only 0.48 MPa, and the elongation at break was 198%; when the COOH/NCO molar ratio was 1.0/1.2, the tensile strength was 1.23 MPa, and the elongation at break was 125%; when the COOH/NCO molar ratio was further increased to 1.0/1.5, the tensile strength of the cured product increased to 2.26 MPa, and the elongation at break was the least at 73%. Because the cross-linking density in the cured product increases as the curing agents are increased, the mechanical properties of the cured product were also enhanced.

The tensile strength of the cured hydroxyl-terminated liquid fluoroelastomer increased when the amount of HDI trimer was

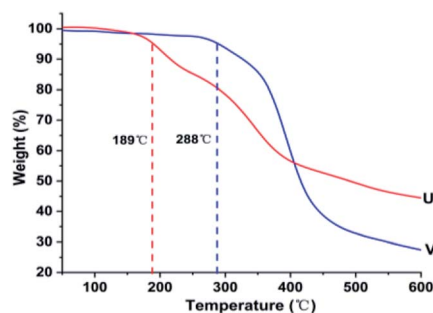


Fig. 13 TGA of the cured products of the (U) carboxyl-terminated liquid fluoroelastomer and (V) hydroxyl-terminated liquid fluoroelastomer.

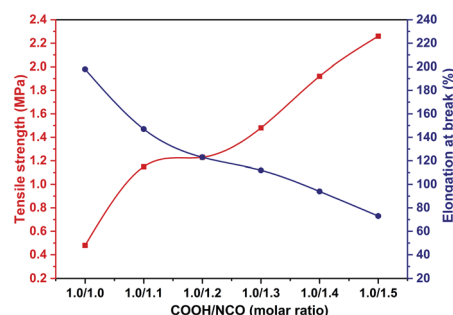


Fig. 14 Effect of HDI content on the mechanical properties of the cured carboxyl-terminated liquid fluoroelastomer.

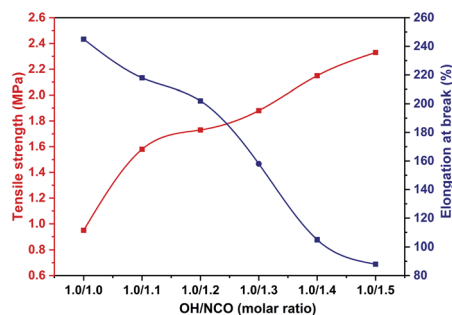


Fig. 15 Effect of HDI content on the mechanical properties of the cured hydroxyl-terminated liquid fluoroelastomer.

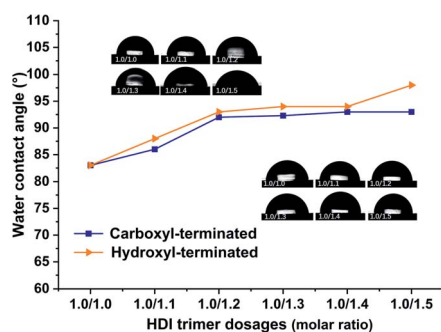


Fig. 16 Effect of HDI content on the surface properties of the cured carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers.

increased, and the elongation at break decreased, as shown in Fig. 15. The compressive mechanical properties of the cured product were higher when the OH/NCO molar ratio was 1.0/1.2, with a tensile strength of 1.73 MPa and 202% elongation at break. The tensile strength of the cured product was as high as 2.33 MPa when the COOH/NCO molar ratio was 1.0/1.5, and the elongation at break was the least at 88%. The mechanical properties of the cured hydroxyl-terminated liquid fluoroelastomer product were superior to those of the HDI-cured carboxyl-terminated liquid fluoroelastomer (Fig. 14). The reaction temperature was lower and the reaction time was shorter when the hydroxyl-terminated liquid fluoroelastomer was cured with HDI, preventing thermal oxygen agentisation of the liquid fluoroelastomers and oxidation degradation of the molecular chains. Further, there was no formation of small molecule products during the reaction, and there were no internal defects and holes in the cured product, which was also one of the reasons.

3.4.6 Surface properties of the cured liquid fluoroelastomers. The water contact angle of the cured carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers increased with the curing agent content, as illustrated in Fig. 16. The WCA tended to be stable at 93° and 94° when the ratio of the carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers to the curing agent was 1/1.4. The WCA reached its maxima at 93° and 98°, respectively, when the ratio of carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers to

Table 4 Effect of HDI content on the WCA

End group	1 / 1.0	1 / 1.1	1 / 1.2	1 / 1.3	1 / 1.4	1 / 1.5
Carboxyl-terminated	89°	86°	92°	92°	93°	93°
Hydroxyl-terminated	83°	88°	93°	94°	94°	98°

the curing agent was 1/1.5, and the hydrophilicity of the hydroxyl-terminated liquid fluoroelastomer was higher than that of the carboxyl-terminated liquid fluoroelastomer. The surface energy of the carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers decreased as the curing agent content increased, and a hydrophobic structure forms on the surface of the samples.

Furthermore, the hydroxyl-terminated liquid fluoroelastomer was more compatible with the curing agent than the carboxyl-terminated liquid fluoroelastomer. The range of surface energy loss was greater, and the samples were hydrophobic.^{46,47} The detailed results are shown in Table 4.

4. Conclusions

In summary, the molecular weight of the poly(VDF-co-HFP) copolymer reduced dramatically after the oxidation degradation reaction from 71 724 to 2458, so carboxyl-terminated liquid fluoroelastomer with 2.71 wt% carboxyl terminal groups can be prepared by oxidation degradation method, and there are about 1.12 mol% $-C=C-$ in the chains of carboxyl-terminated liquid fluoroelastomer were not completely been oxidative degraded. After reduction reaction, the concent of carboxyl group of carboxyl-terminated liquid fluoroelastomer was reduced to 0.12 wt%, with the end-group conversion rate about 96%. At the same time, the $-C=C-$ content was also been reduced. The M_n of the hydroxyl-terminated liquid fluoroelastomer increased to 2680 as a result of the hydrogen-addition reaction in the molecular chain.

Furthermore, the molecular weight and $-C=C-$ concentration had a direct impact on the carboxyl-terminated and hydroxyl-terminated liquid fluoroelastomers properties. The lower the molecular weight and the greater the $-C=C-$ concentration, the lower was the temperature resistance. Similarly, the lower the $-C=C-$ concentration and the higher the molecular weight, the better was the thermal stability. The T_g of the carboxyl-terminated liquid fluoroelastomer was -28°C , which was 3°C lower than that of the hydroxyl-terminated liquid fluoroelastomer, whereas the T_d of the hydroxyl counterpart was higher by 6°C at 241°C . The dynamic viscosity of the liquid fluoroelastomers with different end groups were all decreased with the increase of temperature and similar to about 10 Pa s at 70°C .

After curing, the different end groups of the liquid fluoroelastomers presented distinct curing efficiency and properties. Because the activity of the hydroxyl group is higher than that of the carboxyl group, it could be cured efficiently in a shorter time. The T_d of the cured hydroxyl-terminated liquid



fluoroelastomer reached 288 °C, which was about 100 °C higher than that of the cured carboxyl-terminated liquid fluoroelastomer. However, its low-temperature resistance property was poorer, with a T_g of −3 °C compared with −10 °C of its carboxyl counterpart. The tensile strength was 2.33 MPa and the elongation at break was 88%. In addition, they are both exhibited hydrophilic behaviour, the WCA of the hydroxyl-terminated liquid fluoroelastomer was 98°. And the higher the activity of the end group, the stronger was the hydrophilicity.

Conflicts of interest

The authors declare no competing financial interests.

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