ABSTRACT: The eco-friendly insulating medium C₅F₁₀O gas has attracted the attention of many researchers due to its superior environmental protection and insulation properties. However, there are few studies on the compatibility of C₅F₁₀O gas with sealing materials for the gas-insulated equipment (GIE), which will bury hidden dangers for its engineering application. In this study, the compatibility of C₅F₁₀O gas with an ethylene propylene diene monomer (EPDM) sealing material which is most used in equipment was tested, and its reaction mechanism was simulated. The study found that the O element on the carbonyl group of C₅F₁₀O gas has high chemical reactivity and will interact with EPDM rubber. During the interaction process, EPDM rubber will also react with the C₃F₆ gas generated by the decomposition of C₅F₁₀O gas, and at the same time, it will also decompose C₅F₁₀O gas to produce more C₃F₆O and C₃HF₇. The surface of EPDM rubber will produce oily substances and a large number of crystal particles, which will cause its mechanical properties to deteriorate and shorten its service life. Therefore, it is necessary to carry out anti-corrosion treatment on the surface of EPDM rubber or replace the sealing material with better compatibility when designing and manufacturing GIE.

1. INTRODUCTION

Sulfur hexafluoride (SF₆) gas has been widely used in the gas-insulated equipment (GIE) of power systems due to its excellent insulation and arc extinguishing performance. However, SF₆ gas has a very high greenhouse effect. Its global warming potential (GWP) is 23,500 times that of CO₂, and its life span in the atmosphere is about 3200 years. As early as 1997, the “Kyoto Protocol” clearly listed it as a greenhouse gas that needs to be restricted. The “Paris Agreement” signed at the end of 2015 also pointed out that countries in the world should strive to achieve net-zero emissions of greenhouse gases by the end of this century while keeping the global warming value within 1.5–2%. In response to the global call for energy conservation and emission reduction, avoiding the greenhouse effect caused by the use of SF₆ gas and realizing the green and sustainable development of the power industry, it has become a research hotspot in related fields to find environmentally friendly insulating media for power systems.

The excellent insulation properties of C₅F₁₀O (1,1,1,3,4,4,4-heptafluoro-3-(trifluoromethyl)-2-butanone) gas have received extensive attention from many researchers in recent years. The insulation performance of pure C₅F₁₀O gas is about 1.4 times that of SF₆ gas, its GWP value is less than 1 (equivalent to CO₂), and its atmospheric lifetime is only 15 days. However, the liquefaction temperature of C₅F₁₀O gas of 26.9 °C under normal pressure limits its application. In engineering applications, pure C₅F₁₀O gas cannot be used alone as an insulating medium. It must be mixed with low-boiling buffer gases (such as N₂, synthetic air, CO₂, and so forth) to meet the requirements for liquefaction temperature in engineering practice.

Currently, many researchers have carried out a lot of research on the physical and chemical properties, insulation properties, decomposition characteristics, and arc extinguishing characteristics of C₅F₁₀O and its gas mixture and have achieved certain results, confirming that the C₅F₁₀O gas mixture has the potential to be used in various medium- and low-voltage GIE. It is necessary to carry out the compatibility study of C₅F₁₀O gas and the internal materials of the equipment before engineering applications. C₅F₁₀O gas will be in direct contact with the internal materials of GIE, so it is required that C₅F₁₀O gas and the internal materials of GIE can coexist for a long time and have good compatibility. Kessler et al. had carried out accelerated aging experiments of C₅F₁₀O gas and materials such as metals, alloys, and insulators inside GIE within the range of 75–225 °C. Studies had shown that C₅F₁₀O gas and metal oxides are still stable at high temperatures, whereas C₅F₁₀O gas will react with insulators or elastomers to

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produce \( \text{C}_3\text{HF}_7 \) and \( \text{C}_3\text{F}_6 \). Zhang et al. first studied the interaction mechanism between \( \text{C}_5\text{F}_{10}\text{O} \) and metal materials in electrical equipment through density functional theory (DFT). The calculation found that the oxygen atom in the \( \text{C}_5\text{F}_{10}\text{O} \) molecule has a strong interaction force with the \( \text{Cu}(1\ 1\ 1) \) surface, while the fluorine atom has a weak interaction force with the \( \text{Cu}(1\ 1\ 1) \) surface, which belongs to physical adsorption. The interaction force between \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Ag}(1\ 1\ 1) \) is stronger than that of \( \text{Ag}(1\ 1\ 1) \), which belongs to chemical adsorption, and the interaction mechanism between the \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Ag}(1\ 1\ 1) \) is physical interaction. Experimental research on the compatibility of \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Ag}(1\ 1\ 1) \) is physical adsorption. The interaction force between \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Cu}(1\ 1\ 1) \) surface, which belongs to physical adsorption, and the interaction mechanism between \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Cu}(1\ 1\ 1) \) is physical interaction. The interaction between \( \text{C}_5\text{F}_{10}\text{O} \) and \( \text{Al}(1\ 1\ 1) \) is stronger than that of \( \text{Ag}(1\ 1\ 1) \), which belongs to physical adsorption. The interaction mechanism between the \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Al}(1\ 1\ 1) \) is stronger than that of \( \text{Ag}(1\ 1\ 1) \), which belongs to physical adsorption. The interaction mechanism between \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Al}(1\ 1\ 1) \) is stronger than that of \( \text{Ag}(1\ 1\ 1) \), which belongs to physical adsorption.

Experimental research on the compatibility of \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Ag}(1\ 1\ 1) \) is physical adsorption. The interaction force between \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Cu}(1\ 1\ 1) \) surface, which belongs to physical adsorption, and the interaction mechanism between \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Cu}(1\ 1\ 1) \) is physical interaction. The interaction force between \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Al}(1\ 1\ 1) \) is stronger than that of \( \text{Ag}(1\ 1\ 1) \), which belongs to physical adsorption. The interaction mechanism between the \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Al}(1\ 1\ 1) \) is stronger than that of \( \text{Ag}(1\ 1\ 1) \), which belongs to physical adsorption. The interaction mechanism between \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Al}(1\ 1\ 1) \) is stronger than that of \( \text{Ag}(1\ 1\ 1) \), which belongs to physical adsorption. The interaction mechanism between \( \text{C}_5\text{F}_{10}\text{O} \) molecule and \( \text{Al}(1\ 1\ 1) \) is stronger than that of \( \text{Ag}(1\ 1\ 1) \), which belongs to physical adsorption.

The interaction between \( \text{C}_5\text{F}_{10}\text{O} \) gas and rubber sealing materials will have guiding significance for its engineering applications. In this study, \( \text{C}_5\text{F}_{10}\text{O} \) gas and rubber sealing material compatibility test platform was built, and ethylene propylene diene monomer (EPDM) rubber seal ring, the most commonly used sealing material in GIE, was used as the test object. The compatibility of the \( \text{C}_5\text{F}_{10}\text{O} \)−\( \text{CO}_2 \) gas mixture and EPDM rubber was studied by the thermal acceleration test.

### 2. RESULTS AND ANALYSIS

When using GCMS to detect the components of the gas mixture, first, one should use the SCAN mode to scan all possible decomposition products. The chromatographic peaks of the decomposition products are shown in Figure 1a. The mass-to-charge ratios \((m/z)\) of \( \text{C}_5\text{F}_{10}\text{O} \) gas and decomposition products are shown in Table 1. The decomposition products were found to be \( \text{C}_3\text{F}_6\text{O} \), \( \text{C}_3\text{F}_6 \), and \( \text{C}_3\text{HF}_7 \) through mass spectrometry identification. The reasons why the decomposition products of the control test group contained H-containing \( \text{C}_3\text{HF}_7 \) are as follows: In the test process of this research, the impurity inside the gas chamber was removed by vacuuming and using pure \( \text{CO}_2 \) gas to clean the gas chamber, which avoided the interference of impurity gas to the greatest extent, but there were still traces of \( \text{H}_2\text{O} \) remaining on the inside and inner wall of the gas chamber, and it participates in the decomposition reaction of \( \text{C}_5\text{F}_{10}\text{O} \) gas during the test. The safety data sheet of \( \text{C}_5\text{F}_{10}\text{O} \) gas shows that \( \text{C}_5\text{F}_{10}\text{O} \) reacts with water by violent hydrolysis, and the rate of reaction with water is fast at room temperature, and the half-life of hydrolysis is about 11.5 min. In the reaction process, \( \text{H}_2\text{O} \) will decompose to produce \( \text{H}^* \) and \( \text{OH}^* \) and react with \( \text{CF}_3^* \), \( \text{CF}_2^* \), and \( \text{F}^* \) particles generated by the decomposition of \( \text{C}_5\text{F}_{10}\text{O} \) gas and finally generate decomposition products such as \( \text{C}_3\text{F}_6 \) and \( \text{C}_3\text{HF}_7 \). From the chromatogram of the control group shown in Figure 1a, it can be seen that the \( \text{C}_5\text{F}_{10}\text{O} \)−\( \text{CO}_2 \) gas mixture will react with the stainless steel shell outside the heat source at high temperatures and produce three decomposition products of \( \text{C}_3\text{F}_6\text{O} \), \( \text{C}_3\text{F}_6 \), and \( \text{C}_3\text{HF}_7 \). In the test group, the main decomposition products produced by the contact of the \( \text{C}_5\text{F}_{10}\text{O} \)−\( \text{CO}_2 \) gas mixture with EPDM are also three decomposition products of \( \text{C}_5\text{F}_2\text{O} \), \( \text{C}_3\text{F}_6 \), and \( \text{C}_3\text{HF}_7 \). The difference is that the concentrations of the three decomposition products produced by them are different.

![Figure 1](https://pubs.acs.org/journal/acsodf)

**Figure 1.** Decomposition products of the \( \text{C}_5\text{F}_{10}\text{O} \)−\( \text{CO}_2 \) gas mixture and EPDM rubber after thermal aging. (a) Qualitative analysis of decomposition products of \( \text{C}_5\text{F}_{10}\text{O} \) gas. (b) Quantitative analysis results of decomposition products of \( \text{C}_5\text{F}_{10}\text{O} \) gas.

**Table 1.** \( \text{C}_5\text{F}_{10}\text{O} \) Gas and Its Decomposition Product Mass-to-Charge Ratios \((m/z)\) and Separation Time

| gas type   | mass-to-charge ratios \((m/z)\) | separation time \((\text{min})\) |
|------------|-------------------------------|-------------------------------|
| \( \text{C}_5\text{F}_{10}\text{O} \) | 69, 97, 169, 197, 266         | 6.04−6.90                    |
| \( \text{C}_3\text{F}_6\text{H} \)   | 69, 151                       | 5.55−6.01                    |
| \( \text{C}_3\text{F}_6 \)         | 69, 100, 131, 150             | 5.27−5.61                    |
| \( \text{C}_3\text{F}_2\text{O} \)  | 50, 69, 119                   | 4.97−5.56                    |
In order to study the effect of EPDM on the decomposition products of C$_5$F$_{10}$O gas under different temperature conditions, C$_3$F$_6$O and C$_4$H$_2$F$_7$ were quantitatively studied using C$_3$F$_6$ and C$_4$H$_2$F$_7$ as standard concentration gases. Since the standard concentration of C$_3$F$_6$O gas cannot be obtained temporarily, the peak area of C$_3$F$_6$O gas produced by different test groups can be obtained by integrating the mass spectrum peak area with the m/z of 119 in C$_3$F$_6$O gas, and the concentration change of C$_3$F$_6$O gas can be studied by comparing the peak area in a semi-quantitative way. The quantitative analysis diagrams of the three decomposition products of C$_3$F$_6$O, C$_3$F$_6$, and C$_3$HF$_7$ are shown in Figure 1b.

When the test temperature was 80 °C, the EPDM test group produced more C$_3$F$_6$O and C$_3$HF$_7$ than the C$_5$F$_{10}$O gas decomposition in the control group, while the content of C$_3$F$_6$ was greatly reduced. The concentrations of C$_3$F$_6$O and C$_3$HF$_7$ produced by the control test group were 83.70 and 232.33 ppm, respectively. The concentration of C$_3$F$_6$O produced by the 80 °C EPDM rubber test group decreased by 69.57%, and the concentration of C$_3$F$_6$O and C$_3$HF$_7$ increased by 156.55 and 26.98%, respectively. In the different temperature test groups, the concentrations of C$_3$F$_6$, C$_3$HF$_7$, and C$_3$HF$_7$ produced by the decomposition of the test group at 70 °C were 70.45 and 260.38 ppm, respectively, and the concentration of C$_3$F$_6$ produced by the EPDM rubber test group at 80 °C decreased by 64.41%, whereas the concentration of C$_3$F$_6$O and C$_3$HF$_7$ increased by 148.09 and 13.30%, respectively. The above results indicate that long-term contact between C$_5$F$_{10}$O gas and the surface of the stainless steel shell outside the heat source will decompose the trace of C$_5$F$_{10}$O gas and produce three main decomposition products: C$_3$F$_6$O, C$_3$F$_6$, and C$_3$HF$_7$.

After the test is over, the gas inside the gas chamber is extracted through the exhaust gas collection and recovery system, and the sample of the EPDM rubber sealing ring taken out is shown in Figure 2. From left to right, there are samples of EPDM rubber sealing rings before the test and when the test temperature is 70 and 80 °C, respectively. The EPDM rubber before the test showed a black matt appearance, and the rubber surface after the test had a layer of oily substance attached to the surface. In order to further explore the surface changes of EPDM rubber in contact with C$_5$F$_{10}$O gas under thermal aging conditions, it is necessary to characterize EPDM rubber samples before and after the test to reveal their interaction mechanism.

### 3. CHARACTERIZATION OF EPDM SAMPLES AND THEIR INTERACTION MECHANISM

#### 3.1. Rubber Tensile Test. The EPDM rubber sealing rings used in this study are the O-type rubber ring used in the GIE produced by the XJ Group and not the standard test rubber sample. One should refer to the rubber tensile standard test principle; the length of the tensile test sample is 65 mm and the tensile speed is 500 mm/min. The tensile stress and strain properties of EPDM rubber samples before and after the test are shown in Figure 3 and Table 2. The elongation at break of the EPDM rubber sample before the test was 111.60%, and the maximum tensile force was 250.00 MPa. However, the elongation at break of the EPDM sample at 70 and 80 °C was reduced by 19.47 and 16.98%, and the maximum tensile force was reduced by 10.88 and 1.92%, respectively. The changes in mechanical parameters indicate that the EPDM rubber has embrittled, and the mechanical properties of EPDM rubber samples are slightly reduced after the thermal aging test.

#### 3.2. SEM Test. Figure 4 shows the changes in the surface morphology of EPDM rubber before and after the test. The EPDM rubber before the test had some small ravines on the surface. Under high magnification, it can be observed that the surface of the rubber is covered with grit-like particles. These phenomena are caused by the rubber manufacturing process. When the test temperature is 70 °C, the small ravines on the surface of EPDM rubber are not as obvious as those in the control group. Most of the grit-like particles observed under high magnification have disappeared. Except for the precipitation of crystal particles (white particles in the picture) in some areas, the surface of the rubber has a relatively smooth morphology. The above results show that when the test temperature is 70 °C, the EPDM rubber has begun to appear slightly corroded after being thermally aged for 90 h in the C$_3$F$_6$O–CO$_2$ gas mixture. When the test temperature is 80

![Figure 2. Surface photos of EPDM rubber before and after the test.](image-url)
The results are shown in Table 3 and Figure 5.

To determine the chemical composition of the EPDM rubber sample, we first used XPS to perform energy spectrum scanning tests (the orbits of the tested elements are selected according to the international general test rules) on the rubber samples. This study used XPS to perform energy spectrum scanning tests on the EPDM rubber samples before and after the test. The different characteristic chemical bonds' information in the figure represents different material characteristics, which directly indicates that C,F10O gas interacts with EPDM rubber during the high-temperature thermal aging process. Figure 5a shows the photoelectron spectrum of the F 1s orbital on the surface of the rubber samples. The EPDM rubber before the test did not contain the F element, and the surface of the rubber after the test detected two characteristic peaks of the F element. The C−F bond and the C−F2 group were both detected at 684.50 eV and 688.15 eV, respectively. The C−F bond at 684.50 eV indicates that the chemical bond formed by the F atom fixed to the C atom was detected on the rubber surface. The reason may be that C,F10O gas or its decomposition product gas detected above was attached to the rubber surface. However, the −CF2−CH2=− group confirmed that the −CF2− group replaced the −CH2=− group in the rubber during the reaction of C,F10O gas with rubber and formed a long-chain structure containing organic fluorine groups. The photoelectron spectroscopy of the C 1s orbital on the surface of the rubber samples shown in Figure 5b detected the chemical bond formed by the C and O elements, which is a characteristic peak of the peroxide crosslinker. It is important to point out that the C element on the surface of EPDM rubber samples produced a new characteristic peak at 292.80 eV after the test, which corresponds to the −CF2−CF2− group. This is also consistent with the detection result of the F element shown in Figure 5a. Combined with the previous test results that the concentration of C,F10O produced by the decomposition of C,F10O is lower than that of the control group, it is shown that the −CF2−CF2− groups detected in Figure 5a,b are mainly produced by the reaction of C,F10O with EPDM. Although the photoelectron spectroscopy of the O 1s orbital on the surface of the rubber sample shown in Figure 5c

Table 3. Changes in Surface Element Content of EPDM Rubber Samples before and after the Test

| C (%)  | F (%)  | O (%) |
|-------|-------|-------|
| control group | 79.13 | 1.18 | 19.69 |
| 70 °C test group | 65.16 | 15.37 | 19.47 |
| 80 °C test group | 64.24 | 17.46 | 18.30 |

From the changes of the elements on the surface of EPDM rubber samples and after the test listed in Table 3, it can be seen that the content of the C element on the surface of the rubber samples is the highest, and its proportion of the surface of the EPDM rubber sample before the test is close to 80%. The reason that the rubber surface before the test contained 19.69% of the O element was that EPDM rubber was vulcanized with peroxide as a cross-linking agent to obtain more excellent mechanical and insulating properties, so its surface contained the O element. During the heat aging test, the proportion of O element decreased slightly, and the higher the test temperature, the more the proportion of the O element on the rubber surface decreased. This is due to the large increase in the content of the F element during the thermal aging test, which indirectly caused the relative decrease in the proportion of C and O. The increase in the content of the F element also indirectly indicates that C,F10O gas interacts with EPDM rubber during the thermal aging process.

Figure 4. Surface morphology of EPDM rubber before and after the test. (a) Control group. (b) 70 °C test group. (c) 80 °C test group. The EPDM rubber is a kind of copolymer formed by ethylene, propylene, and a small amount of non-conjugated diene. The chemical elements are mainly C and H elements. Since the photoionization interface of the H and He elements is small and the signal is too weak, XPS cannot detect these two elements, so the change of the H element is not considered. In order to further study the change law of elements (F 1s, C 1s, and O 1s) on the surface of EPDM rubber samples, this study used XPS to perform energy spectrum scanning tests (the orbits of the tested elements are selected according to the international general test rules) on the elements that may exist on the rubber surface. We first use XPS Peak software to perform a Shirley-type fitting deduction on the background of the energy spectrum and then use the Gaussian algorithm to perform peak fitting. Finally, refer to the National Institute of Standards and Technology XPS database and previous studies to determine the chemical state, molecular structure, and chemical bonds of the elements. The results are shown in Table 3 and Figure 5.
did not detect the formation of new characteristic chemical bonds, the two characteristic peak intensities of the O element also changed to varying degrees before and after the test.

The new characteristic peaks of F 1s and C 1s in the above results confirm that C₅F₁₀O gas and EPDM rubber have a chemical reaction under thermal aging conditions, and C₅F₁₀O gas will corrode the surface of EPDM rubber at high temperatures.

3.4. Reaction Mechanism of C₅F₁₀O and EPDM. In order to analyze the interaction mechanism between C₅F₁₀O gas and EPDM, density functional theory (DFT) was used. EPDM rubber is a copolymer formed of ethylene, propylene, and a small amount of non-conjugated diene. Ethylene norbornene is usually used as the diene. Its main advantages are fast vulcanization speed and high vulcanization efficiency. In this study, the ethylene-propylene-ethyldiene norbornene chain was selected as the model of EDPM rubber for DFT calculation. Yamada et al. also used this scheme to study the interaction mechanism between nitrile rubber and fuel species based on DFT. EPDM defects and particles have unsaturated chemical bonds and are more reactive than EPDM molecules. Therefore, the reaction between C₅F₁₀O and EPDM defects or particles can occur more easily than EPDM molecules. We used the Perdew–Burke–Ernzerhof-generalized gradient approximation method in our calculations to deal with electronic exchange and correlation. In the calculations, the double numerical plus polarization was used to add polarization functions to all atoms, and all electrons were processed using all electrons relativistically, and the relativistic effect was introduced in the processing of core electrons to make the calculation results more accurate and reliable. The convergence tolerance settings in the geometric optimization calculation process are as follows: 1.0 × 10⁻⁶ Ha for energy, (2) 0.005 Å for maximum displacement, and (3) 0.002 Ha/Å for maximum force. The reaction enthalpy of a chemical reaction is defined as follows:

\[ \text{enthalpy} = E_{\text{product}} - E_{\text{reactant}} \]  

where the negative value of the reaction enthalpy indicates that the reaction is thermodynamically spontaneous and the positive value of the reaction enthalpy indicates that the reaction is an endothermic process.

Because the EPDM molecule has a complex organic macromolecular structure, the interaction mechanism between EPDM rubber and C₅F₁₀O gas discussed in this study mainly focuses on the reaction of EPDM rubber defects or particles with C₅F₁₀O molecules. The defects and particles of EPDM rubber have unsaturated chemical bonds, and its chemical reaction activity is higher than that of EPDM rubber molecules. In other words, the reaction between C₅F₁₀O molecules and EPDM defects and particles is more likely to occur than EPDM molecules. The structure of the EPDM monomer is shown in Figure 6a. From its structural characteristics, it can be found that there are four possible defect formation paths for EPDM rubber (marked by the arrow in the figure).

The energy barrier and reaction enthalpy required by EPDM rubber to form the four defect paths are shown in Figure 6b–e. EPDM rubber requires 108.80 kcal/mol of energy to form defects through the process of the chemical bond breaking shown in reaction path 1, and the reaction enthalpy is 90.99 kcal/mol. EPDM rubber requires 184.74 kcal/mol of energy to form defects through the process of the chemical bond breaking shown in reaction path 2, and the reaction enthalpy is 89.18 kcal/mol. EPDM rubber requires 234.94 kcal/mol of energy to form defects through the process of the chemical
bond breaking shown in reaction path 3, and the reaction enthalpy is 114.41 kcal/mol. EPDM rubber requires 207.70 kcal/mol of energy to form defects through the process of the chemical bond breaking in the dehydrogenation process shown in reaction path 4, and the reaction enthalpy is 95.74 kcal/mol.

We calculated the possible interactions between the C5F10O molecule and four kinds of EPDM defects and considered the structural characteristics of the C5F10O molecule to construct the initial structure model of 24 possible reaction models. The calculation results show that C and F elements in the C5F10O molecule have low chemical reactivity and cannot react with EPDM defects, while the O element in the C5F10O molecule has high chemical reactivity and can react with EPDM defects to form chemical bonds. The adsorption models of the C5F10O molecule and EPDM defects obtained by calculation are shown in Figure 7.

The eight types of defects formed in the four reaction paths shown in Figure 6b–e interact with the O element in the C5F10O molecule to form eight adsorption models. When the O element in the C5F10O molecule is close to the defects of EPDM rubber, the optimized structures show that the distance between the O in the C5F10O molecule and the C element or the H element in the EPDM defect was shortened. When the O element in the C5F10O molecule is close to the C element in paths 1–3, the optimized structures show that the distance between the O and C elements in the EPDM defect was about 1.500 Å, the distances between the O and C elements in the two EPDM defects in reaction path 3 were 1.456 and 1.377 Å, respectively, and the distance between O in the C5F10O molecule and H elements in the EPDM defect in reaction path 4 was 0.979 Å.

Figure 6. Enthalpy of reaction formed by defects in EPDM rubber. (a) Monomer structure and defect formation path of EPDM. (b) Reaction path 1. (c) Reaction path 2. (d) Reaction path 3. (e) Reaction path 4.
In addition, the calculation results show that the enthalpy values of all reaction paths are negative, that is, the reaction process is exothermic. The reaction enthalpy of reaction paths 3A and 4B is significantly higher than that of the other six reaction paths (approximately $-20.00 \text{ kcal/mol}$). Among them, path 3A with the largest reaction enthalpy value is $-41.71 \text{ kcal/mol}$, and path 4A with the smallest reaction enthalpy value is $-17.61 \text{ kcal/mol}$. The two adsorption models formed by reaction paths 3A and 4B have the largest reaction enthalpy and the most heat released during the adsorption process. Therefore, the two reaction paths are most likely to undergo adsorption reaction between C$_5$F$_{10}$O molecules and EPDM defects. The chemical reaction that causes the fluorine element to accumulate on the rubber surface is also more likely to occur through the above two paths.

3.5. Discussion. According to the above results, the interaction between C$_5$F$_{10}$O and EPDM rubber defects and particles will cause the adsorption of C$_5$F$_{10}$O on the rubber surface. The O element in the carbonyl (C==O) group of C$_5$F$_{10}$O is chemically active and can easily react with EPDM defects or particles to form chemical bonds. C$_5$F$_{10}$O gas has poor compatibility with EPDM during high-temperature thermal aging. This shortcoming should be considered when designing and manufacturing GIE sealing materials. In order to prevent the corrosion of the sealing material from accelerating its aging process, resulting in the shortening of the service life of the equipment and even the insulation failure caused by gas leakage, we need to perform anti-corrosion treatment on the surface of EPDM rubber or use rubber with better compatibility with C$_5$F$_{10}$O gas as a sealing material.

4. CONCLUSIONS

In this study, the EPDM rubber sealing ring used in GIE by State Grid XJ Group Co., Ltd. was used as the test object to explore the compatibility of C$_5$F$_{10}$O gas and EPDM rubber under thermal aging. After the thermal aging test, the changes in gas composition, mechanical properties, surface morphology, and element changes of EPDM rubber were tested, and the following conclusions were obtained:
(1) Under thermal aging conditions, minor amounts of C₃F₆O gas will react with the outer shell of the stainless steel container to decompose and generate C₅F₁₀O, C₅F₆ and C₅HF₇. EPDM rubber will react with the C₅F₆ gas produced by the decomposition of C₅F₁₀O gas, reducing the content of C₅F₆ in the decomposition products.

(2) During the thermal aging test, oily substances will be formed on the surface of EPDM rubber, and a large number of crystal particles will be observed on the surface under the microscopic view. This change will make EPDM rubber brittle and decrease the mechanical properties.

(3) The O element on the carbonyl group in the C₅F₁₀O molecule has high reactivity and will form a stable adsorption structure and form new chemical bonds during the interaction process with EPDM rubber defects and particles.

(4) The reaction of EPDM and C₅F₁₀O gas will accelerate its aging and shorten its service life. In the process of designing and manufacturing GIE, the surface of EPDM rubber should be treated with anti-corrosion or replaced with rubber with better compatibility.

5. EXPERIMENTAL SECTION

The test platform is shown in Figure 8, which is mainly composed of a heating system, a temperature feedback control system, a detection system, and an exhaust gas collection and recovery system. The heating system includes a heat source and a 304 stainless steel shell outside the heat source. The heat source transfers heat to the surface through the stainless steel shell so that the gas is heated more uniformly. The outer wall of the gas chamber is also made of 304 stainless steel, which ensures that the gas chamber has high chemical stability. The temperature feedback monitoring and control system includes a temperature sensor and a proportion integration differentiation (PID) controller. The temperature sensor is used to monitor the temperature of the test gas and provide feedback of the temperature signal to the PID controller to control the test process. There is also a sensor for monitoring the gas temperature inside the gas chamber. The detection system includes a gas chromatograph–mass spectrometer, a field emission scanning electron microscope, and an X-ray photoelectron spectrometer. The gas chromatograph–mass spectrometer is the GCMS-QP2010 Ultra gas chromatography mass spectrometer produced by Shimadzu. The chromatographic column model is CP-SIL 5 CB, and the film thickness, length, and inner diameter of the chromatographic column are 8 μm, 60 m, and 0.32 mm, respectively. FESEM is produced by Carl Zeiss and is used to observe the changes in the surface morphology of EPDM rubber after heat aging. XPS is used to characterize the element composition, content, and characteristic chemical bonds of the EPDM rubber surface and can be used to analyze the chemical reactions that occur on the EPDM rubber surface.

The C₅F₁₀O gas used in the test was provided by Minnesota Mining and Manufacturing (3M) with a purity of 99.5%, and high-purity CO₂ was provided by Wuhan Niiude Special Gas Co., Ltd. with a purity of 99.999%. The EPDM rubber material is the sample of the EPDM seal ring in GIE produced by State Grid XJ Group Co., Ltd. The sample is a cylindrical shape with a diameter of 5 mm. The cleaning steps of EPDM rubber samples before the test were as follows: first, we should put the EPDM rubber sample in distilled water and use ultrasonic cleaning for 1 h to remove the impurities on the surface to prevent it from affecting the test results, and then we should put the sample in a drying oven to dry for 24 h to remove water.

The gas used in the experiment of this research was a 7.5% C₅F₁₀O−92.5% CO₂ gas mixture (volume fraction). Since the gas pressure in the gas chamber will rise during the thermal aging process, the gas pressure was set to 0.2 MPa (relative pressure) for safety reasons. The relevant regulations stipulate that the maximum ambient temperature for the safe and stable operation of GIE is 40 °C, and the allowable upper limit of temperature rise of metal products can reach 65−75 °C. EPDM is generally used as the sealing material of GIE, and the temperature of the equipment casing is usually lower. GB/T 11022-2011 stipulates that the maximum temperature rise of the part that can be touched in normal operation is 30 °C, and the maximum temperature rise of the part that does not need to be touched in normal operation is 40 °C. Therefore, considering the actual operating conditions of GIE, the test temperature was selected as 70 and 80 °C, and control groups with the temperature of 80 °C were set up. The control group had the same conditions as the test group except that no EPDM was placed. There is currently no compatibility test standard for non-SF₆ gas insulating media and rubber materials. In order to have sufficient reaction time between C₅F₁₀O gas and EPDM rubber, the thermal aging test time was 90 h. After the test, the gas sample in the gas chamber was collected and GCMS was used to detect the change in gas composition.

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Notes
The authors declare no competing financial interest.

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