Electrochemical and Infrared Absorption Spectroscopy Detection of SF₆ Decomposition Products

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Abstract: Sulfur hexafluoride (SF₆) gas-insulated electrical equipment is widely used in high-voltage (HV) and extra-high-voltage (EHV) power systems. Partial discharge (PD) and local heating can occur in the electrical equipment because of insulation faults, which results in SF₆ decomposition and ultimately generates several types of decomposition products. These SF₆ decomposition products can be qualitatively and quantitatively detected with relevant detection methods, and such detection contributes to diagnosing the internal faults and evaluating the security risks of the equipment. At present, multiple detection methods exist for analyzing the SF₆ decomposition products, and electrochemical sensing (ES) and infrared (IR) spectroscopy are well suited for application in online detection. In this study, the combination of ES with IR spectroscopy is used to detect SF₆ gas decomposition. First, the characteristics of these two detection methods are studied, and the data analysis matrix is established. Then, a qualitative and quantitative analysis ES-IR model is established by adopting a two-step approach. A SF₆ decomposition detector is designed and manufactured by combining an electrochemical sensor and IR spectroscopy technology. The detector is used to detect SF₆ gas decomposition and is verified to reliably and accurately detect the gas components and concentrations.

Keywords: SF₆ gas decomposition products; electrochemical sensing (ES); infrared (IR) spectroscopy; detection characteristic; detector

1. Introduction

Electrical equipment is the fundamental basis of power systems, and its reliability is crucial to ensure grid security and stability [1]. With their many advantages, such as a compact structure, stable insulation performance, strong flowing-breaking ability, and reliable operation, SF₆ gas-insulated electrical equipment, especially gas-insulated-switchgear (GIS) and gas-insulated-transmission line (GIL), has been widely applied in high-voltage (HV) and extra-high-voltage (EHV) power systems worldwide [2]. However, partial discharge (PD) and local overheating can occur in the GIS because of insulation faults, causing SF₆ gas decomposition and ultimately generating several types of SF₆ decomposition products, including SOF₂, SO₂F₂, CO, SO₂, H₂S, HF, CF₄, and SiF₄ [3,4]. These decomposition products have a severe effect on the insulating materials, accelerate insulation deterioration in the GIS, and can even result in equipment damage. Once the GIS insulation fails, the
state-of-safe and reliable GIS operation faces a severe threat. The variation of these gases reflects the equipment internal PD and insulation status. Through quantitatively detecting and analyzing these gaseous components, potential equipment failure and security problems can be diagnosed to provide an early warning of degraded device status [5–8].

The analysis techniques for SF\(_6\) decomposition are an important method for diagnosing the insulation condition of SF\(_6\) gas-insulated electrical equipment. Relative to other detection methods, the SF\(_6\) decomposition detection methods have many advantages, including resistance against jamming, high-sensitivity and quantitative analysis. Furthermore, with the existence and development of equipment internal defects, the amount of SF\(_6\) decomposition products can gradually accumulate; thus, this method is particularly suitable for long-term monitoring. Substantial worldwide research has been directed to the detection of SF\(_6\) decomposition products. These detection methods are used to analyze the components and concentrations of such products and ultimately to deduce the partial discharge type and severity. The methods include gas chromatography, mass spectrometry, infrared (IR) spectroscopy, electrochemical sensing (ES), gas detector tubes, ion mobility spectrometry, and carbon nanotubes, among other methods [9–13]. However, these detection methods still generally depend on regular field sampling and complete quantitative analysis of SF\(_6\) decomposition products in the laboratory. Relative to other detection methods, electrochemical sensors and IR spectroscopy offer particularly favorable prospects for online detection [9,14,15]. Therefore, it is necessary a further investigation in electrochemical sensors and IR spectroscopy for detecting SF\(_6\) decomposition products, which is the focus of this study.

Here, by first establishing a testing platform of ES and IR spectroscopy, we obtain the response characteristics of the electrochemical sensor and the spectral characteristic of target detection gases (CO, SO\(_2\) and H\(_2\)S), including the temperature, temperature compensation and linearity. Then, a qualitative and quantitative analysis model of ES-IR is established by adopting a two-step approach. Furthermore, a SF\(_6\) decomposition detector has been designed by combining electrochemical sensors and IR spectroscopy to lay the foundation for SF\(_6\) decomposition product online monitoring with an ES-IR evaluation of SF\(_6\) gas-insulated electrical equipment.

2. Detection Principle

2.1. Electrochemical Sensing

With advantages such as a stable working performance, long-service life, low-power consumption, high-sensitivity and rapid-response speed, the electrometrical gas sensor is well-suited for field quantitative detection of gases that are toxic at low concentrations and combustible gases [9,16]. ES also offers advantages such as small volume, simple operation and low cost, and has been widely used to conduct online monitoring in multiple fields, such as petrochemical industry, home automation, air quality monitoring, etc.

Electrochemical sensors are electrochemical cells that operate in the amperometric mode. That is, sensors generate a current that is linearly proportional to the fractional volume of the target gas, such as CO, SO\(_2\) or H\(_2\)S. The structure of an electrochemical sensor is shown in Figure 1. The sensor typically consists of three electrodes immersed in an electrolyte: the working electrode (WE), the counter electrode (CE) and the reference electrode (RE). The working principle of an electrode sensor is that the target gas diffuses into the WE surface by crossing a gas diffusion barrier, the WE either oxidizes or reduces the target gas with the CE balancing the generated current, and an external circuit outputs a current linearly proportional to the measured gas concentration. Taking the reaction of CO electrochemical sensor as an example, when the sensor works, CO is oxidized on the WE and O\(_2\) is reduced on the CE. And the reaction equation is shown as follows:

\[
\text{Oxidation reaction (WE)}: \text{CO} + \text{H}_2\text{O} \rightarrow \text{CO}_2 + 2\text{H}^+ + 2\text{e}^- \\
\text{Reduction reaction (CE)}: \frac{1}{2}\text{O}_2 + 2\text{H}^+ + 2\text{e}^- \rightarrow \text{H}_2\text{O} \\
\text{Overall reaction: } \text{CO} + \frac{1}{2}\text{O}_2 \rightarrow \text{CO}_2 \tag{1}
\]
When the electrode potential and the catalytic activity are high enough, CO gas molecules penetrate the diffusion barrier close to the electrode and reacted rapidly. Then, the amplitude of the generated current is proportional to the CO gas concentration based on Fick’s law. Therefore, by measuring the current between WE and CE, the CO gas concentration can be obtained. Similarly, the total redox reaction of H₂S and SO₂ electrochemical sensors is expressed by the following equations:

\[
\begin{align*}
\text{SO}_2 + \text{H}_2\text{O} + \frac{1}{2}\text{O}_2 & \rightarrow \text{H}_2\text{SO}_4 \\
\text{H}_2\text{S} + 2\text{O}_2 & \rightarrow \text{H}_2\text{SO}_4
\end{align*}
\] (2)

In reality, the WE surface always undergoes a continuous electrochemical reaction, which may result in sensor performance degradation. Thus, the RE is added to improve sensor performance. The RE anchors the WE at the correct potential to ensure that the reaction stays in the transport-limited current plateau of the current-voltage curve.

![Figure 1. Schematic diagram of the electrochemical gas sensor.](image)

2.2. IR Spectroscopy

Material molecular movement follows the law of quantum mechanics. According to the Born-Oppenheimer approximation of quantum mechanics [17], the energy of molecular motion \( E \) is composed of translational \( (E_t) \), rotational \( (E_r) \), vibrational \( (E_v) \) and electronic energy \( (E_e) \). Thus, the energy of molecular motion can be expressed as follows:

\[
E = E_t + E_r + E_v + E_e
\] (3)

The energy difference of adjacent energy levels of a molecule’s translational kinetic energy is relatively small and can be regarded as changing continuously. The rotational, vibrational and electronic transitions are all quantized. When using IR to detect gas samples, the molecule undergoes an energy level transition from a lower energy level \( E_1 \) to a higher energy level \( E_2 \). The gas sample can selectively absorb the specific frequency of IR light that satisfies the following equation:

\[
\Delta E = E_2 - E_1 = h\nu
\] (4)

where \( h = 6.624 \times 10^{-34} \) J s and \( \nu \) is the frequency. IR absorption spectroscopy exploits the fact that molecules can selectively absorb electromagnetic (EM) radiation in the IR zone. Each type of molecule absorbs wavelengths in a unique IR spectrum corresponding to its molecular vibrational and rotational frequency. Thus, the IR absorption spectrum of a gaseous decomposition product features a peak at the gas absorption wavelength called the typical absorption peak for this gas. Through analyzing
the typical absorption position, value and shape of the gas sample IR spectrum, the decomposition products of SF6, for example, can be qualitatively and quantitatively detected [7]. Specifically, IR absorption spectroscopy for quantitative detection is based on the Lambert-Beer Law [18], which states that the gas sample absorbs a specific frequency of IR light when a beam of light passes through the gas sample; the absorption intensity is proportional to each component concentration and optical path length [19]. For the IR absorption spectra of a single gas sample, the absorbance at any wave number (v) is expressed as follows:

\[
A(v) = \lg \frac{1}{T(v)} = a(v)bc
\]

where the dimensionless parameter \(A(v)\) is the absorbance, the dimensionless parameter \(T(v)\) is the transmittance, \(a(v)\) is the absorbance coefficient at the wave number (v) and its units are \((\text{kPa} \times \mu\text{L}/\text{L} \times \text{m})^{-1}\), \(b\) is the optical path length (unit: m), and \(c\) is the gas concentration (units: \(\text{kPa} \times \mu\text{L}/\text{L}\)).

3. Testing Platform

To obtain the response of the electrochemical sensor and spectral characteristics of the target detection gas, an experimental platform is first configured, as shown in Figure 2. The platform comprises a gas configuration system and a measurement system, including the background SF6 gas, the sample gases (CO, SO2, H2S), a mass flow controller, a SF6 decomposition detection tank, electrochemical sensors, and a Fourier transform infrared spectroscopy (FTIR) detection system.

![Figure 2. Test platform for electrochemical sensing. The label in the picture is indicated respectively as follows: 1. Background gas SF6; 2. Sample gas CO, SO2, H2S; 3. Mass flow controller; 4. Thermocouple; 5. Heating plate; 6. SF6 decomposition detection tank; 7. Electrochemical sensor; 8. Digital ammeter; 9. Glass rotameter; 10. Interferometer; 11. IR source; 12. Gas pool; 13. Detector; 14. Barometer; 15. Vacuum pump.](image)

The experimental gas configuration system consists primarily of background SF6 gas, sample gas, a mass flow controller, and a SF6 decomposition detection tank. The gas flow controller can be used to accurately control the gas volume, and various species and concentrations of experimental gas can be configured in the SF6 decomposition detection tank.

The measurement system for sensor response contains the electrochemical sensors (see Figure 3), a digital ammeter, a heating plate and a thermocouple. The sensors that employ a threaded connection for ease of replacement is placed at the bottom of the tank and its main technical parameters is shown in Table A1. The heating plate is used to heat the experimental gas in the tank, and the thermocouple measures the ambient temperature inside the tank in real time, facilitating study of the sensor’s
temperature characteristics. Through configuring various components and concentrations of the experimental gas, the sensor linearity characteristics can also be obtained.

The FTIR detection system consists of an IR source, an interferometer module, a detector module and the gas pool, as shown in Figure 4 and Table A2. The IR source can emit 1–25 µm wavelength light, and the wavelength range can cover all the characteristic wavelengths for all measured gas components. The interferometer module is the key component and mediates the splitting and scanning of the IR light. The detector module is used to analyze the interference light containing the information from the measured gas; the model then obtains the gas component concentrations through IR spectroscopy.

4. Response Characteristics of the Sensors

4.1. Temperature Characteristics

The output signal of electrochemical sensors is closely related to the ambient temperature. To study the relationship between the sensor output signal and ambient temperature, the experiments are processed at different temperatures of 10 °C, 20 °C, 30 °C, 40 °C and 50 °C using sensors to detect the experimental gas. The temperature characteristics are shown in Figure 5.
The variation of ambient temperature affects the output signal of electrochemical sensors at a constant gas concentration. However, different gas sensors exhibit differences in their temperature-dependent response, as shown in Figure 5. Figure 5a shows that the output current signal of the CO electrochemical sensor decreases with increasing ambient temperature, although this temperature sensitivity is nearly absent over the range of 10–30 °C. Figure 5b,c show that the output signals of the SO₂ and H₂S sensors increase as the ambient temperature increases, and the higher the ambient temperature, the greater the rate of change for the output current signal. Figure 5d shows that the SO₂ sensor also responds to H₂S gas and that its output signal is positively correlated with ambient temperature. The sensor output signal is relatively stable at approximately 20 °C, which demonstrates that 20 °C may be an appropriate testing temperature for the sensors. When the ambient temperature is too high, the output signal exhibits a greater change. That is, a higher ambient temperature has a stronger effect on these electrochemical sensors. Therefore, such temperature-dependence sensors are difficult to be used directly for field detection.

The relationship between the sensor output signal and ambient temperature is influenced in two ways. First, the electrochemical sensors have an internal electrochemical reaction, and increasing the ambient temperature can enhance the reaction speed, thus increasing the output signal. Second, a gas diffusion barrier exists between the sensor and the external environment. With the use of various materials and fabrication processes, different sensor films exhibit differences in diffusion permeability, and the variation of ambient temperature influences the transport of gas molecules through the diffusion film. An increasing ambient temperature causes more energetic thermal motion of gas molecules, thereby affecting the gas contacting the working electrode surface, which may influence the
sensor output signal. As a result, an increasing ambient temperature may not necessarily lead to an increased sensor output signal. It is important when using electrochemical sensors to ensure an optimal operating temperature over long experimental durations, and temperature compensation is necessarily adopted to suppress the effect of ambient temperature insofar as it influence the sensor measurements.

4.2. Temperature Compensation

To further study the relationship between the ambient temperature and sensor output signal, the temperature compensation curves are fit using a quadratic fitting method, as shown in Figure 6. The x-axis represents the ambient temperature (°C), the sensor output signal at an ambient temperature of 20 °C is chosen as the reference signal, and the Q of the y-axis represents the ratio of output signals at the measuring temperature and at 20 °C, as shown in the following equation:

$$Q = \frac{I_x}{I_{20^\circ C}}$$

where $I_x$ is the measured output signals value; $I_{20^\circ C}$ is the measured output signals value at 20 °C.

![Figure 6. Cont.](image-url)
Figure 6. Temperature compensation curves of the electrochemical sensors. (a) CO sensor, (b) SO₂ sensor, (c) H₂S sensor.

The CO electrochemical sensor has a negative temperature sensitivity, whereas the SO₂ and H₂S sensors have a positive temperature sensitivity. The temperature compensation curves for the different sensors are as follows:

\[
Q_{CO} = f_1(T) = 1.00101 + 0.0006123T - 0.00002980T^2 \\
Q_{SO₂} = f_2(T) = 1.01002 - 0.0011700T + 0.00004893T^2 \\
Q_{H₂S} = f_3(T) = 0.99371 - 0.0006948T + 0.00004211T^2
\]

where \( T \) is the ambient temperature during the measurement. Therefore, after the data normalization preprocessing, the actual output signals (\( I_x \)) at the ambient temperature can be converted to the standard value (\( I_{20°C} \)) at 20 °C by the following equation:

\[
I_{20°C} = \frac{I_x}{Q} = \frac{I_x}{f_i(T)}
\]

4.3. Linearity Characteristics

The electrochemical sensor generates a current signal linearly proportional to the target gas concentration. Through configuring various components and concentrations of experimental gas in the detection tank at one atmosphere and 20 °C ambient temperature, the linearity characteristics of the electrochemical sensors are obtained, as shown in Figure 7.

The experimental results show that the output current signal of different types of electrochemical sensors responds linearly with the corresponding target gas concentration. In addition, the SO₂ sensor responds linearly to increasing H₂S gas concentration. The sensor response can be fit to a linear equation between the output signal (\( I \)) and the target gas concentration (\( C \)):

\[
I = mc + n
\]

where \( m, n \) is the slope and vertical intercept of the fitting line respectively. The sensor linearity characteristics are shown in Table 1.
Table 1. Linearity characteristics of the electrochemical sensors (at 20 °C).

| Sensor | Target Gas | Slope ($m$) | Vertical Intercept ($n$) | Linearly Dependent Coefficient ($R^2$) | Standard Error |
|--------|------------|-------------|-------------------------|--------------------------------------|----------------|
| CO     | CO         | 8.1310      | 4                       | 0.9934                               | 0.0834         |
| SO$_2$ | SO$_2$     | 3.5571      | 4                       | 0.9879                               | 0.0541         |
|        | H$_2$S     | 3.3049      | 4                       | 0.9925                               | 0.0875         |
| H$_2$S | H$_2$S     | 12.735      | 4                       | 0.9949                               | 0.0690         |

Based on the linearity characteristics of these electrochemical sensors, the theoretical detection precision and measurement uncertainty can be analyzed. The sensor detection precision is an important parameter for evaluating detection performance, and the target output current precision of the sensors is two valid digits after the decimal point in units of mA. Therefore, the ideal minimum gas concentration detection of sensors can be calculated by assuming a sensor output current value of 4.01 mA in the linear formulas. The measurement uncertainty is also an important parameter. The upper envelope is expressed by the following equation:

$$I_{max} = m_{max}C + n$$  \hspace{1cm} (10)

which can be fit by choosing the maximum measurement value at each gas concentration. The lower envelope is expressed by the following equation:

$$I_{min} = m_{min}C + n$$  \hspace{1cm} (11)
which can be fit by choosing the minimum measurement value at each gas concentration. Comparing the two envelope lines with the optimal linearity characteristic in Figure 7, the measurement uncertainty can be calculated, as shown in Table 2.

**Table 2.** Theoretical detection precision and measurement uncertainty of the electrochemical sensors.

| Sensors | Theoretical Precision (µL·L⁻¹) | Measurement Uncertainty (%) |
|---------|-------------------------------|-----------------------------|
|         | Upper Envelope                | Lower Envelope              |
| CO      | 1.2                           | 2.94                        | -3.21                      |
| SO₂     | 2.8                           | 5.14                        | -4.16                      |
| H₂S     | 0.8                           | 3.24                        | -3.85                      |

4.4. **Sensor Data Analysis Matrix**

The working principle of electrochemical sensors can cause the sensor to respond to several types of gas with similar chemical properties. We determined that the SO₂ sensor responds to SO₂ and H₂S gas. To accurately obtain the target gas concentration, the temperature compensation and crossing calculation are adopted, and the sensor data matrix is established [20]. The sensor data matrix between the real value of target gas concentration and the measured output current signal is as follows:

\[
I_1 = QI = AC + b \tag{12}
\]

where diagonal matrix \( Q = \text{diag}(Q_{CO}, Q_{SO₂}, Q_{H₂S}) \) represents the sensor temperature compensation coefficient matrix, \( I_1 = [I_{CO}, I_{SO₂}, I_{H₂S}]^T \) represents the sensor actual output signal matrix, \( I = [I_{CO}, I_{SO₂}, I_{H₂S}]^T \) is the sensor output signal matrix at 20 °C, \( A = [a_{ij}] (1 \leq i \leq 3, 1 \leq j \leq 3) \) represents the sensor crossing calculation matrix, \( C = [C_{CO}, C_{SO₂}, C_{H₂S}]^T \) is the actual target gas concentration matrix, and \( b = [4, 4, 4] \) is the sensor output signal in pure SF₆ gas.

Matrices \( Q \) and \( A \) can be obtained through a series of experiments, and then the actual target gas concentration can be obtained through matrix inversion:

\[
C = A^{-1}(I - b) = A^{-1}(Q^{-1}I_1 - b) \tag{13}
\]

A multi-component gas mixture is used to demonstrate the reliability and accuracy of the sensor data matrix. We configured the system to detect the multi-components with background SF₆ gas and the target gases of CO, SO₂ and H₂S, and the experimental results are shown in Table 3.

The experimental results show that the measurement uncertainty of the CO sensor is less than 2% and that the measurement uncertainty of the SO₂ and H₂S sensors is less than 3%. The results show that the sensor data matrix method offers high reliability and accuracy.

**Table 3.** The measurement results of SF₆ decomposition products using a sensor data analysis matrix.

| T (°C) | CO (µL·L⁻¹) | SO₂ (µL·L⁻¹) | H₂S (µL·L⁻¹) | CO (µL·L⁻¹) | SO₂ (µL·L⁻¹) | H₂S (µL·L⁻¹) | Uncertainty (%) |
|--------|-------------|---------------|---------------|-------------|---------------|---------------|----------------|
| 11.2   | 500         | 500           | 0             | 494         | 492           | 0             | -1.20          | -1.60          | 0             |
| 11.5   | 500         | 500           | 1000          | 500         | 514           | 1018          | 0.00           | 2.80           | 1.80          |
| 24.8   | 1000        | 500           | 0             | 985         | 495           | 0             | -1.50          | -1.00          | 0             |
| 25.1   | 1000        | 500           | 500           | 997         | 487           | 505           | -0.30          | -2.60          | 1.00          |
| 34.9   | 1000        | 1000          | 0             | 994         | 1027          | 0             | -0.60          | 2.70           | 0             |
| 35.0   | 1000        | 1000          | 1000          | 983         | 1009          | 1030          | -1.70          | 0.90           | 3.00          |
| 45.2   | 1500        | 1000          | 0             | 1480        | 983           | 0             | -1.33          | -1.70          | 0             |
| 45.4   | 1500        | 1000          | 500           | 1524        | 1016          | 503           | 1.60           | 1.60           | 0.60          |
5. Spectral Characteristic of the Target Gas

5.1. Spectral Simulation

Material studio is used to simulate the IR absorption spectrum of SF$_6$ and its decomposition products based on the molecular dynamics method [21]. The microstructure of different molecular molecules are first determined, as shown in Figure 8.

The calculated IR absorption spectroscopy of SF$_6$ and its decomposition products is simulated, as shown in Figure 9. The simulation results show that the IR absorption of SF$_6$ and its decomposition products is concentrated predominantly within the range of 300–4000 cm$^{-1}$. Different gas molecules have different IR absorption peaks, and their corresponding absorption intensities also exhibit a large difference. In addition, a single gas molecule can have multiple absorption peaks. Polyatomic molecules can have a variety of vibration modes that correspond to several IR absorption lines of different intensity; thus, it is necessary to determine the characteristic IR peaks to identify the SF$_6$ decomposition products. The calculated IR absorption spectroscopy is obtained by superimposing the IR absorption spectroscopy of SF$_6$ and its decomposition products, as shown in Figure 10. The position, intensity and shape of IR absorption characteristic peaks of these gas molecules differ substantially. The above characteristics of IR absorption peaks are used to qualitatively and quantitatively analyze the SF$_6$ decomposition products.

Through reviewing the relevant international standards and literature, we compared the results of calculated IR spectroscopy and measured IR spectroscopy, as shown in Table A3. The comparison shows that both calculated and measured results are consistent. It is possible to determine the attribution of IR absorption peaks for the SF$_6$ decomposition products.
5.2. Selection of a Typical Spectrum

The established FTIR detection system is intended to obtain the IR typical spectra of these three target gases. In the experiment, CO, SO2 and H2S are chosen as the target detection gases. Pure SF6 is the background gas, and the concentrations of CO/SF6, SO2/SF6, and H2S/SF6 are each 1000 µL/L. During the measurement, we ensure that the power supply and gas connection are safely and reliably...
connected. The gas pressure is adjusted to 0.1 MPa, and the measurement result is recorded after the
gas flows for approximately 15 min. The experimental results are shown in Figure 11, in which au is
the relative intensity of the IR spectrum.

The experimental results show that the IR spectra of CO, SO₂ and H₂S all contain prominent peaks.
The relative intensity of CO is 766.2 au at the 2169 cm⁻¹ characteristic peak, that of SO₂ is 3048.6 au
at the 1360 cm⁻¹ characteristic peak, and that of H₂S is 106.4 au at the 2625 cm⁻¹ characteristic peak.

The basic rules of characteristic IR line selection are as follows: (1) the characteristic IR line is typical for
a single component in the mixture; (2) a higher peak is correlated with improved detection sensitivity;
(3) the lines are independent of each other to reduce cross interference; and (4) the line is only minimally
influenced by external factors [22]. According to the above rules, the IR characteristic spectroscopy of
the three target gases (CO, SO₂ and H₂S) is obtained by comparing the calculated and measured IR
spectroscopic results, as shown in Table 4.

![Figure 11](image-url)
The quantitative analysis using this law is applied only in the absorbance spectrum. Therefore, the absorbance spectrum of these target detection gases must be pretreated. The collected IR spectrum data may contain irrelevant information, such as noise, instrument interference, and environmental factors. The absorbance spectrum pretreatment can obtain a high-quality IR spectrum by eliminating this irrelevant information, thus improving the qualitative and quantitative accuracy.

5.3. Pretreatment of IR Spectroscopy

Comparing the calculated and measured IR spectroscopy of these three target gases, the calculated spectral intensity of CO and SO$_2$ at their respective wavenumbers is 64.537 kJ/mol$^{-1}$ and 188.75 kJ/mol$^{-1}$, respectively; the calculated spectrum intensity of H$_2$S is only 0.3246 kJ/mol$^{-1}$ because the IR absorption capacity of CO and SO$_2$ is much stronger than that of H$_2$S gas. Due to these three target gases differing substantially in terms of IR absorption capacity, the experimental results show that CO and SO$_2$ have distinct IR characteristic peaks with higher relative intensity but that the IR spectrum of H$_2$S gas is noisier, with multiple IR peaks with lower relative intensity. It is important to ensure that the target detection gases exhibit strong and unambiguous IR characteristic peaks to enable reliable qualitative and quantitative detection of these target detection gases. Simulations based on the molecular dynamics calculated method are effective and help ensure the accuracy of the IR characteristic spectra of the target detection gases.

Table 4. Typical IR spectra of the three target detection gases.

| Detected Component | Characteristic Wavenumber (cm$^{-1}$) | Spectroscopic Attribution               | Calculated Intensity (kJ·mol$^{-1}$) |
|--------------------|----------------------------------------|-----------------------------------------|-------------------------------------|
| CO                 | 2168                                   | C–O stretching vibration                | 64.537                              |
| SO$_2$             | 1360                                   | S=O symmetrical stretching vibration    | 188.75                              |
| H$_2$S             | 2624                                   | S–H stretching vibration                | 0.3246                              |

Comparing the calculated and measured IR spectroscopy of these three target gases, the calculated spectral intensity of CO and SO$_2$ at their respective wavenumbers is 64.537 kJ/mol$^{-1}$ and 188.75 kJ/mol$^{-1}$, respectively; the calculated spectrum intensity of H$_2$S is only 0.3246 kJ/mol$^{-1}$ because the IR absorption capacity of CO and SO$_2$ is much stronger than that of H$_2$S gas. Due to these three target gases differing substantially in terms of IR absorption capacity, the experimental results show that CO and SO$_2$ have distinct IR characteristic peaks with higher relative intensity but that the IR spectrum of H$_2$S gas is noisier, with multiple IR peaks with lower relative intensity. It is important to ensure that the target detection gases exhibit strong and unambiguous IR characteristic peaks to enable reliable qualitative and quantitative detection of these target detection gases. Simulations based on the molecular dynamics calculated method are effective and help ensure the accuracy of the IR characteristic spectra of the target detection gases.

5.3. Pretreatment of IR Spectroscopy

IR absorption spectroscopy for quantitative detection is based on the Lambert-Beer Law. The quantitative analysis using this law is applied only in the absorbance spectrum. Therefore, the absorbance spectra of these target detection gases must be pretreated. The collected IR spectrum data may contain irrelevant information, such as noise, instrument interference, and environmental factors. The absorbance spectrum pretreatment can obtain a high-quality IR spectrum by eliminating this irrelevant information, thus improving the qualitative and quantitative accuracy.

Figure 12 shows that pretreatment of the target gas IR spectrum can fully retain the effective information by elimination of the interference noise and can further improve the quality of the absorbance spectrum of these target detection gases. Furthermore, the absorbance spectra of CO and SO$_2$ gases feature prominent IR peaks at their characteristic wavenumber; thus, these two gases

![Figure 11. Measured IR spectroscopic results for the target detection gas. (a) CO, (b) SO$_2$, (c) H$_2$S.](image-url)
can be quantitatively detected based on their absorbance spectrum. Relative to that of CO and 
SO$_2$, the IR absorption capacity of the H$_2$S is far lower; H$_2$S gas does not exhibit any prominent IR 
peak at its characteristic wavenumber and thus cannot be quantitatively detected using this FTIR 
detection system.

![Graphs showing absorbance spectra of CO, SO$_2$, and H$_2$S before and after pretreatment.](image)

**Figure 12.** Pretreatment of the target gas IR spectrum. (a) CO before pretreating, (b) CO after pretreating, (c) SO$_2$ before pretreating, (d) SO$_2$ after pretreating, (e) H$_2$S before pretreating, (f) H$_2$S after pretreating.
5.4. Response Characteristics of IR Spectroscopy

Through analyzing the pretreatment absorbance spectrum of these three target detection gases, a prominent difference emerges between qualitative and quantitative detection. In the qualitative analysis, these three target gases (CO, SO$_2$ and H$_2$S) can all be identified using IR spectroscopy. In the quantitative analysis, CO and SO$_2$ can be accurately calculated using the absorbance spectrum, but H$_2$S gas is challenging to quantitatively analyze with IR spectroscopy, and a supplemental electrochemical sensor is required to realize quantitative analysis. However, CO and SO$_2$ gas can be studied primarily in terms of the response characteristics IR absorption spectroscopy, as shown in Figure 13.

Figure 13. The response characteristics of the CO and SO$_2$ IR spectrum: (a) temperature characteristic of CO IR spectrum; (b) temperature characteristic of SO$_2$ IR spectrum; (c) temperature compensation of CO IR spectrum; (d) temperature compensation of SO$_2$ IR spectrum; (e) linearity characteristic of CO IR spectrum; (f) linearity characteristic of SO$_2$ IR spectrum.
5.4.1. Temperature Characteristics

The ambient temperature is an important factor influencing the measurement accuracy. To study the IR spectral characteristics of the target gas at different temperatures, the ambient temperature of the FTIR system is controlled using a vacuum oven. The temperature characteristics of the IR spectrum are obtained at 10 °C, 20 °C, 30 °C, 40 °C and 50 °C using the FTIR detection system to detect the experimental gas. The temperature characteristics are shown in Figure 13a,b.

Figure 13a,b shows that the absorbance value of CO and SO$_2$ increases with increasing ambient temperature. Due to the IR absorption capacity of SO$_2$ being stronger than that of CO, the SO$_2$ absorbance value is larger than that of CO at the same concentration and ambient temperature. In general, increasing the ambient temperature may strengthen absorption of IR light by gas molecules; for example, the absorbance values of CO and SO$_2$ increase with increasing temperature.

5.4.2. Temperature Compensation

To further study the relationship between the temperature and IR absorbance of CO and SO$_2$, the temperature compensation curves are fit by a quadratic fitting method, as shown in Figure 13c,d. The x-axis represents the ambient temperature (°C), and the Q of the y-axis represents the ratio of absorbance values at the measurement temperature and at 20 °C.

The absorbance spectra of CO and SO$_2$ have a positive temperature relationship, and the change of ambient temperature influencing the CO absorbance value is much higher than that for SO$_2$. The temperature compensation curves of CO and SO$_2$ absorbance are analyzed as follows:

$$Q_{CO} = 0.98554 - 0.0003167T + 0.00005351T^2$$
$$Q_{SO_2} = 0.98119 + 0.0005282T + 0.00002412T^2$$  (14)

where Q is the ratio of absorbance values at the measurement temperature and at 20 °C and T is the ambient temperature during a measurement.

5.4.3. Linearity Characteristics

The quantitative analysis of IR spectroscopy follows the Lambert-Beer Law, and the absorbance value of the target gas is proportional to the target gas concentration. To study the linearity characteristics of the CO and SO$_2$ IR spectra, through configuring various components and concentrations of experimental gas in the detection tank at one atmosphere and 20 °C ambient temperature, the linearity characteristics of CO and SO$_2$ IR spectra are obtained, as shown in Figure 13e,f.

The experimental results show that the absorbance values of CO and SO$_2$ follow a linear relationship with their gas concentration; the linearity characteristics of CO and SO$_2$ IR spectra are represented by the following formulas:

$$A_{CO} = 0.00015086C_{CO}$$
$$A_{SO_2} = 0.00059643C_{SO_2}$$  (15)

where A is the absorbance value of the target gas and C is the gas concentration.

Through analyzing the retreatment absorbance spectrum of CO and SO$_2$, the signal-to-noise ratio (SNR) of their IR typical spectra can be obtained, and the theoretical detection precision of the FTIR detection system can be calculated. In addition, the upper envelope $A_{max} = k_{max}C$ and the lower envelope $A_{min} = k_{min}C$ are linearly fit using the least squares method. Comparing these two envelope lines with the linearity characteristics in Figure 13e,f, the measurement uncertainty can be calculated, as shown in Table 5.
Table 5. Theoretical detection precision and measurement uncertainty of IR absorption spectroscopy.

| Electrochemical Sensor | Theoretical Precision (µL·L⁻¹) | Measurement Uncertainty (%) |
|------------------------|-------------------------------|----------------------------|
|                        |                               | Upper Envelope             | Lower Envelope             |
| CO                     | 4.8                           | 7.87                       | –8.10                      |
| SO₂                    | 1.3                           | 6.43                       | –7.24                      |

5.5. Spectral Analysis Matrix

The IR absorption spectrum can directly and quickly reflect the gas components of the detected gas sample, and the gas concentration of each component can be quantitatively calculated based on the Lambert-Beer Law. To accurately obtain the target gas component and concentration, the spectral analysis matrix is assembled. The gas component can be quickly obtained by analyzing the IR spectrum. The absorbance value of the target gas is converted to the value at 20 °C, and the gas concentration is calculated based on the linearity characteristics of the CO and SO₂ IR spectra:

\[ C = K^{-1}A = K^{-1}Q^{-1}A_1 \]  

(16)

where \( C = [C_{CO}, C_{SO₂}]^T \) is the actual target gas concentration matrix, \( K = [K_{ij}] (1 \leq i \leq 2, 1 \leq j \leq 2) \) represents the linearity characteristic matrices of the CO and SO₂ IR spectra, \( Q = diag(Q_{CO}, Q_{SO₂}) \) represents the temperature compensation coefficient matrices of CO and SO₂ IR spectra, \( A = [A_{CO}, A_{SO₂}]^T \) is the absorbance value matrix at 20 °C, and \( A_1 = [A_{1CO}, A_{1SO₂}]^T \) is the absorbance value matrix for the actual measurement.

The multi-component gas mixture is used to verify the data matrix. Through configuring and detecting the multi-components with SF₆ background gas with target gases of CO, SO₂ and H₂S, the experimental results of the FTIR detection system are shown in Table 6.

Table 6. The measurement results for SF₆ decomposition products using the spectral analysis matrix.

| T (°C) | Sample (µL·L⁻¹) | Measurement (µL·L⁻¹) | Uncertainty (%) |
|--------|-----------------|----------------------|----------------|
|        | CO   | SO₂  | H₂S | CO   | SO₂  | H₂S | CO   | SO₂  | H₂S |
| 11.2   | 500  | 500  | 0   | 479  | 482  | –   | –4.20 | –3.60 | –   |
| 11.5   | 500  | 500  | 1000 | 526  | 514  | –   | 5.20  | 2.80  | –   |
| 25.0   | 1000 | 500  | 0   | 1036 | 490  | –   | 3.60  | 2.00  | –   |
| 25.0   | 1000 | 500  | 500  | 1011 | 487  | –   | 1.10  | 2.60  | –   |
| 35.0   | 1000 | 1000 | 0   | 930  | 952  | –   | 7.00  | 4.80  | –   |
| 35.0   | 1000 | 1000 | 1000 | 972  | 1051 | –   | 2.80  | 5.10  | –   |
| 45.0   | 1500 | 1000 | 0   | 1450 | 979  | –   | 3.33  | 2.10  | –   |
| 45.4   | 1500 | 1000 | 500  | 1422 | 1016 | –   | 5.20  | 1.60  | –   |

The experimental results show that the measurement uncertainty of the CO absorbance spectrum is less than 7% and that the measurement uncertainty of the absorbance spectrum is less than 5%.

6. ES-IR Comprehensive Detection

6.1. Analysis Model Based on ES-IR

ES and IR spectroscopy are important for SF₆ decomposition analysis techniques for evaluating SF₆ gas-insulated electrical equipment. These two detection methods are both suitable for online detection. In this section, ES and IR spectroscopy are integrated to detect SF₆ gas decomposition through the response of electrochemical sensors and the spectral characteristics of target gases. A qualitative and quantitative analysis model for ES-IR has been established by adopting a two-step approach.

The first step is that the gas components of these three target gases (CO, SO₂ and H₂S) are quickly identified by analyzing the IR spectrum. Relative to the electrochemical sensor, IR spectroscopy
is superior for qualitative analysis, offering a faster response speed, more potential gas detection components and less cross-interference, as shown in Table 7. Therefore, IR spectroscopy is used to rapidly identify the target gas components.

Table 7. Comparison of electrochemical sensors and IR spectroscopy in qualitative analysis.

| Detection Method     | Response     | Gas Detection Components                                           | Cross Interference                          |
|----------------------|--------------|--------------------------------------------------------------------|---------------------------------------------|
| IR spectroscopy      | 1 spectrum/s | Most of the SF₆ decomposition products (e.g., SOF₂, SO₂, H₂S, HF, CF₄, and SF₆) | The typical spectrum of every component is independent |
| Electrochemical sensor | 25–60 s     | CO, SO₂, H₂S, and HF                                               | Some components exhibit cross-interference  |

The second step is that the gas concentrations of these three target gases (CO, SO₂ and H₂S) are accurately calculated by analyzing the measurement results of these two methods. The joint detection algorithm is used to obtain the gas concentration. It is challenging to use a single measurement to accurately reflect the gas concentration; thus, this algorithm selects the average of five measurement results as the detection result (CES, CIR) of these two detection methods. In addition, the relative standard deviation of these five measurement results is chosen as the evaluation criteria, and the final result for the target gas can be calculated using the following equation:

\[
CV = \sqrt{\frac{\sum (C_i - C_o)^2}{n-1}}
\]

where CV is the relative standard deviation of the five measurement results, \(C_o\) is the average of five measurement results, \(C_i\) is a single measurement result, and \(n\) is the number of measurements. In addition:

\[
C = \frac{C_{ES}}{C_{ES} + C_{IR}} \cdot C_o + \frac{C_{IR}}{C_{ES} + C_{IR}} \cdot C_{ES}
\]

where \(C\) is the final result of the target gas, \(C_{ES}\) is the relative standard deviation of the five measurement results using electrochemical sensors, \(C_{IR}\) is the relative standard deviation of the five measurement results using IR spectroscopy, \(C_{ES}\) is the average of five measurement results using electrochemical sensors, and \(C_{IR}\) is the average of five measurement results using IR spectroscopy.

Thus, the qualitative and quantitative analysis model of ES-IR is established by adopting a two-step approach. The qualitative analysis uses IR spectroscopy to rapidly identify the gas components. The quantitative analysis uses a joint detection algorithm to calculate the measurement results of these two methods and then obtains the gas concentration of each identified component.

6.2. SF₆ Decomposition Detector

A SF₆ decomposition detector was designed and manufactured by combining electrochemical sensors and IR spectroscopy, as shown in Figure 14. The volume of this detector is 460 mm × 380 mm × 132 mm, and the weight is 25 kg. Furthermore, there are several features for the developed SF₆ decomposition detector in this paper. First of all, using ES-IR comprehensive detection method and cross interference technique, the gas composition and concentration of SF₆ decomposition products could be analyzed accurately and fast, which meets the requirements of on-site monitoring. Secondly, temperature monitoring module is also installed inside the detector, then temperature compensation method is used to reduce the influence of ambient temperature on the measurement results and ensure the accuracy of the measurement results. Thirdly, pneumatic control module is applied to the detector to make the internal sensors work under the suitable condition of about 1 MPa gas pressure, which further ensures the veracity and reliability of test results.

The SF₆ decomposition detector was tested using a multi-component gas mixture to verify the data matrix. Through configuring and detecting the multi-components with a background gas of SF₆ and target gases of CO, SO₂ and H₂S, the results of the SF₆ decomposition detector are shown in Table 8.
After reviewing the complete research results, the main conclusions are as follows:

1. The SF$_6$ decomposition detection method based on electrochemical sensors was studied. The results show that the sensors can accurately detect the three target gases (CO, SO$_2$ and H$_2$S). Through adopting temperature compensation and a crossing calculation, the sensor data matrix is established to guarantee measurement accuracy. A multi-component mixed gas is configured and used to verify the sensor data matrix, and the results show that the method offers satisfactory reliability and accuracy.

2. Infrared spectroscopy is also an important detection method for analyzing SF$_6$ decomposition. The IR absorption spectrum of SF$_6$ and its decomposition products are simulated, and the typical IR spectra of the three target gases (CO, SO$_2$ and H$_2$S) were obtained. By assembling the spectral analysis matrix, the target gas can be rapidly identified, and CO and SO$_2$ can also be quantitatively detected.

The results show that the measurement uncertainty for CO, SO$_2$ and H$_2$S is less than 5%. The detector is used to detect SF$_6$ gas decomposition in the experiment, and the results show that the detector can reliably and accurately detect the gas components and concentrations.

### 7. Conclusions

ES and IR spectroscopy offer favorable application prospects for online detection. Furthermore, electrochemical sensors can be combined with IR spectroscopy to detect SF$_6$ decomposition products; following this concept, a SF$_6$ decomposition detector was also designed and manufactured. After reviewing the complete research results, the main conclusions are as follows:

1. The SF$_6$ decomposition detection method based on electrochemical sensors was studied. The results show that the sensors can accurately detect the three target gases (CO, SO$_2$ and H$_2$S). Through adopting temperature compensation and a crossing calculation, the sensor data matrix is established to guarantee measurement accuracy. A multi-component mixed gas is configured and used to verify the sensor data matrix, and the results show that the method offers satisfactory reliability and accuracy.

2. Infrared spectroscopy is also an important detection method for analyzing SF$_6$ decomposition. The IR absorption spectrum of SF$_6$ and its decomposition products are simulated, and the typical IR spectra of the three target gases (CO, SO$_2$ and H$_2$S) were obtained. By assembling the spectral analysis matrix, the target gas can be rapidly identified, and CO and SO$_2$ can also be quantitatively detected.

### Table 8. The results of the SF$_6$ decomposition detector.

| No. | Sample (µL·L$^{-1}$) | Measurement (µL·L$^{-1}$) | Uncertainty (%) |
|-----|----------------------|---------------------------|-----------------|
|     | CO       | SO$_2$     | H$_2$S     | CO       | SO$_2$     | H$_2$S     |                      |
| 1   | 6.20     | 6.72       | 4.84       | 6        | 7          | 5          | −3.23             | 4.17               | 3.31               |
| 2   | 11.99    | 12.41      | 9.55       | 12       | 13         | 10         | 0.08              | 4.75               | 4.71               |
| 3   | 24.34    | 19.03      | 20.21      | 24       | 19         | 20         | −1.40             | −0.16              | −1.04              |
| 4   | 48.81    | 50.51      | 39.52      | 48       | 51         | 39         | −1.66             | 0.97               | −1.32              |
| 5   | 85.20    | 82.36      | 59.64      | 84       | 84         | 59         | −1.41             | 2.00               | −1.07              |
| 6   | 100      | 100        | 100        | 1098     | 104        | 101        | −2.00             | 4.00               | 1.00               |
| 7   | 500      | 500        | 100        | 500      | 488        | 103        | 0.00              | −2.40              | 3.00               |
| 8   | 500      | 1000       | 500        | 514      | 1028       | 488        | 2.80              | 2.80               | −2.40              |
| 9   | 1000     | 500        | 500        | 1034     | 484        | 504        | 3.40              | −3.20              | 0.80               |
| 10  | 1000     | 1000       | 1000       | 1016     | 1047       | 1007       | 1.60              | 4.70               | 0.70               |

Figure 14. Structural design of the SF$_6$ decomposition detector.
The combination of electrochemical sensors with IR spectroscopy was used to detect SF$_6$ gas decomposition. The qualitative and quantitative analysis model of ES-IR was established by adopting a two-step approach. A SF$_6$ decomposition detector was designed and manufactured by combining electrochemical sensors and IR spectroscopy. The detector was used to detect SF$_6$ gas decomposition in the experiments, and the results show that the detector can reliably and accurately detect the gas components and concentrations.

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**Author Contributions:** Ming Dong and Ming Ren conceived and designed the experiments; Rixin Ye performed the experiments; Chongxing Zhang analyzed the data and wrote the paper; and Ricardo Albarracín gave substantive suggestions and guidance for the research.

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**Appendix A**

**Table A1.** Main technical parameters of electrochemical sensors.

| Sensor | Measuring Range (ppm) | Resolution Ratio (ppm) | Response Time (s) | Work Pressure (kPa) |
|--------|------------------------|------------------------|-------------------|---------------------|
| CO     | 0–2000                 | 0.5                    | <25               | 80–120              |
| SO$_2$ | 0–2000                 | 1                      | <20               | 90–110              |
| H$_2$S | 0–2000                 | –1                     | <60               | 90–110              |

**Table A2.** Main technical parameters of infrared spectrum detection system.

| Module Name | Interferometer | Detector | IR Source | Gas Pool |
|-------------|----------------|----------|-----------|----------|
| Type: Dual reflector design | Type: Photoelectric MCT | Type: ceramic lamp filament | Type: Gas Cell-2.4 m | Optical path length: 2.4 m |
| Lens diameter: 12.7 mm | Aperture diameter: 12.7 mm | Beam diameter: 12.7 mm | Spectrum range: 1–25 µm |
| Resolution ratio: 4 cm$^{-1}$ | Resolution ratio: 1 mm | | |
| Repetition rate of wave number: <10 ppm | Amplifier bandwidth: 0–20 kHz | Power source: 12/5 A | |
| Scanning frequency: 1 spectrum per second | Half-field angle: 28 mrad | Scanning frequency: 1 spectrum per second | |

**Table A3.** Comparison between calculated and measured IR spectroscopy.

| Decomposition Products | Simulation (cm$^{-1}$) | Measurement (cm$^{-1}$) | Relative Error (%) |
|------------------------|------------------------|------------------------|--------------------|
| SO$_2$F$_2$            | 523.15; 532.25; 540.90; 821.17; 1278.56; 1526.79 | 539; 544; 552; 848; 1269; 1502 | –2.57; –2.56; –2.01; –3.16; 0.75; 1.65 |
| SOF$_2$                | 373.25; 511.97; 780.89; 1355.55 | 390; 530; 806; 1333 | –4.29; –3.40; –3.12; 1.69 |
| CO                     | 2086.39 | 2169 | –3.81 |
| SO$_3$                 | 1163.69; 1368.85 | 1167; 1360 | –0.28; 0.65 |
| H$_2$S                 | 1380.97; 2657.83 | 1335; 2625 | 3.44; 1.25 |
| HF                     | 343.93 | 3644 | –0.61 |
| CF$_4$                 | 1231.598; 1232.343 | 1280; 1283 | –3.78; –3.95 |
| SiF$_4$                | 1044.98; 1045.93; 1047.79 | 1020; 1028; 1032 | 2.45; 1.74; 1.53 |

**References**

1. Sun, Q.; Dong, M.; Ren, Z.; Ren, M.; Li, Y.; Qiu, A. Field Test and Analysis of Partial Discharge on GIS under Impulse Voltage. *High Volt. Eng.* 2012, 38, 639–644.
2. Wang, F.; Qiu, Y. Recent Development Trend of Gas Insulated Switchgear. *Power Syst. Technol.* 2003, 27, 54–57.
3. Zhang, X.X.; Liu, W.T.; Tang, J. Study on PD detection in SF$_6$ using multi-wall carbon nanotube films sensor. *IEEE Trans. Dielectr. Electr. Insul.* 2010, 17, 838–844. [CrossRef]
4. Liu, C.; Palanisamy, S.; Chen, S.; Wu, P.; Yao, L.; Lou, B. Mechanism of Formation of SF₆ Decomposition Gas Products and its Identification by GC-MS and Electrochemical methods: A mini Review. *Int. J. Electrochem. Sci.* 2015, 10, 4223–4231.

5. Ji, Y.; Wang, C.; Yang, R. Measuring Technique of SF₆ Decomposition Products and Its Application to Fault Diagnosis of GIS. *High Volt. Appar.* 2011, 47, 100–103.

6. Hergli, R.; Casanovas, J.; Derdouri, A. Study of the Decomposition of SF₆ in the Presence of Water, Subjected to Gamma Irradiation or Corona Discharges. *IEEE Trans. Electr. Insul.* 1988, 23, 451–465. [CrossRef]

7. Nukaiyama, Y.; Takagi, I.; Ishihara, H. Principal Decomposition By-Products Generated at Various Abnormalities in Gas-Insulated Transformers. *IEEE Trans. Power Deliv.* 1994, 9, 1885–1891. [CrossRef]

8. Tang, J.; Yang, X.; Yang, D.; Yao, Q.; Miao, Y.; Zhang, C.; Zeng, F. Using SF₆ Decomposed Component Analysis for the Diagnosis of Partial Discharge Severity Initiated by Free Metal Particle Defect. *Energies* 2017, 10, 1119. [CrossRef]

9. Chen, X.; Peng, H.; Red, M. Detection Techniques of SF₆ Decomposition and Application. *High Volt. Appar.* 2010, 46, 34–38.

10. Sakthinathan, S.; Palanisamy, S.; Chen, S.; Wu, P.; Yao, L.; Lou, B. Electrochemical Detection of Phenol in Industrial Pollutant Absorbed Molecular Sieves by Electrochemically Activated Screen Printed Carbon Electrode. *Int. J. Electrochem. Sci.* 2015, 10, 3319–3328.

11. Zhang, X.; Tie, J.; Zhang, J. A Pt-Doped TiO₂ Nanotube Arrays Sensor for Detecting SF₆ Decomposition Products. *Sensors* 2013, 13, 14764–14776. [CrossRef] [PubMed]

12. Liu, H.; Zhou, Q.; Zhang, Q.; Hong, C.; Xu, L.; Jin, L.; Chen, W. Synthesis, Characterization and Enhanced Sensing Properties of a NiO/ZnO p–n Junctions Sensor for the SF₆ Decomposition Byproducts SO₂, SO₂F₂, and SOF₂. *Sensors* 2017, 17, 913. [CrossRef] [PubMed]

13. Thirumalraj, B.; Palanisamy, S.; Chen, S.; Wu, P.; Yao, L.; Lou, B. Electrochemical Sensing of SF₆ Decomposition Products Based on a Screen Printed Carbon Electrode. *Int. J. Electrochem. Sci.* 2015, 10, 3098–3105.

14. Chu, F. SF₆ Decomposition in Gas-insulated Equipment. *IEEE Trans. Electr. Insul.* 1986, 2, 693–725. [CrossRef]

15. Shi, H.; Qian, J.; Xiong, Z. Investigation on On-line Monitoring of SF₆ Decomposition in Electrical Device. *High Volt. Appar.* 2014, 50, 56–60.

16. Qi, B.; Li, C.; Luo, L. Experiment on the Correlation between Partial Discharge and Gas Decomposition Products in GIS. *High Volt. Eng.* 2010, 36, 957–963.

17. Cederbaum, L. Born-Oppenheimer Approximation and Beyond for Time-dependent Electronic Processes. *J. Chem. Phys.* 2008, 128, 124101. [CrossRef] [PubMed]

18. Joel, M.; Yi, X. Results on Gas Detection and Concentration Estimation via Mid-IR-Based Gas Detection System Analysis Model. *IEEE Sens. J.* 2012, 12, 2347–2354.

19. Ji, S.; Zhong, L.; Liu, K. Research Status and Development of SF₆ Decomposition Components Analysis under Discharge and Its Application. *Proc. CSEE* 2015, 35, 2318–2322.

20. Yang, D.; Tan, Q.; Xiong, J.; Tao, C. Design of Detencto foe SF₆ Gas Purity and Decomposition. *Instrum. Tech. Sens.* 2015, 2, 36–45.

21. Scott, A.; Radom, L. Harmonic Vibrational Frequencies: An Evaluation of Hartree-Fock, Møller-Plesset, Quadratic Configuration Interaction, Density Functional Theory, and Semiempirical Scale Factors. *J. Phys. Chem.* 1996, 100, 16502–16513. [CrossRef]

22. Chen, W.; Zhao, L.; Peng, S. Analysis of Dissolved Gas in Transformer Oil Based on Laser Raman Spectroscopy. *Proc. CSEE* 2014, 15, 2485–2492.

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