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## Research on the on-line determination of formaldehyde gas by the gas–liquid phase chemiluminescence method†

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Based on the gas–liquid phase chemiluminescence tester independently developed by our laboratory, a highly sensitive, fast and accurate on-line detection method of formaldehyde gas in ambient atmosphere is established. The chemiluminescence system and the trace formaldehyde gas in the air directly undergo an interface heterogeneous chemiluminescence reaction to obtain a strong chemiluminescence signal. Through the measurement of the chemiluminescence signal intensity, a highly sensitive, real-time and on-line method for the determination of formaldehyde in the air was established. Factors influencing the experimental results such as gallic acid, potassium dichromate, reaction medium concentration, surfactant type and concentration, pump speed, tube length, and interfering gas were discussed based on the single factor and orthogonal analysis results. Finally, the optimal detection conditions were collected, and the detection results were compared with the national standard phenol reagent method. The results show that when the concentration of the standard formaldehyde gas is in the range of 0–0.582 µg L<sup>-1</sup>, the linear equation of this method is  $y = 208x + 29.667$ , the linear coefficient is  $R^2 = 0.997$ , and the minimum detection concentration of formaldehyde is  $2.327 \times 10^{-3}$  µg L<sup>-1</sup>. Under the same external conditions, the comparison and analysis using the national standard phenol reagent method proved that this method has the advantages of fast detection speed, low detection limit, good sensitivity, and accurate results, which can be used for real-time and online determination of trace formaldehyde in ambient air.

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## 1 Introduction

With the rapid development of the social economy and industry, the use of formaldehyde (HCHO) as an important chemical organic raw material is steadily increasing. Formaldehyde is an irritant and toxic gas that is harmful to human health due to its mutagenicity and carcinogenicity.<sup>1</sup> The World Health Organization (WTO) and the U.S. Environmental Protection Agency (EPA) have long listed it as a class of carcinogens.<sup>2</sup> In an outdoor environment, formaldehyde will rapidly photodegrade under light exposure to generate hydroperoxy radicals (·HO<sub>2</sub>), and then react with nitric oxide (NO) to form active hydroxyl radicals (·OH), which ultimately leads to ozone (O<sub>3</sub>), peroxyacetyl nitrate (PAN) and other photochemical

pollution gas generation.<sup>3</sup> In an indoor environment, due to the large number of decorative materials containing formaldehyde and the slow air flow, the indoor concentration of formaldehyde is much higher than that in an outdoor environment. Formaldehyde can cause irritation and allergic reactions after being inhaled by the human body. A high concentration of formaldehyde has adverse effects on various organs and may even cause death.<sup>4–6</sup> Therefore, the detection of formaldehyde content is crucial in the field of environmental monitoring and human health.

Several detection methods of formaldehyde were reported<sup>7–9</sup> and they are generally divided into traditional analysis methods and modern analysis methods. Traditional analytical methods were developed in the 1990s, which mainly include spectrophotometry, chromatography, and chemiluminescence analysis.<sup>10–12</sup> Modern analytical methods are new analytical methods developed in the recent decade, which include the electrochemical method,<sup>13</sup> catalytic kinetic spectrophotometry (CKS),<sup>14</sup> sensor method,<sup>15–17</sup> gas–liquid chemiluminescence analysis (GLC)<sup>18</sup> and fluorescent probe (FP) method.<sup>8</sup> The gas–liquid heterogeneous chemiluminescence analysis (GLHC) is a new high-sensitivity detection method developed on the basis of chemiluminescence analysis. The reaction reagent directly

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reacts with formaldehyde gas, while the sampling and absorption process are not needed so that formaldehyde can be detected directly and quickly without an external light source. Moreover, this method can also avoid the interference of stray light, humidity and other external factors. Compared with the flow injection chemiluminescence (IC) analysis, GLHC analysis has significant advantages.

Based on the chemiluminescence analyzer of gas-liquid heterogeneous interface independently developed in the laboratory, a highly sensitive, fast and accurate on-line analysis and detection method of the formaldehyde gas in ambient atmosphere is established in this study. Alkaline conditions were employed for comparison with the classical gallic acid-hydrogen peroxide system<sup>19</sup> and luminol-hydrogen peroxide chemiluminescence system.<sup>20</sup> Potassium dichromate was selected as the oxidant to achieve more stable luminescence and less interference. The system provides a new idea for the on-line detection of formaldehyde gas by chemiluminescence.

## 2 Materials and methods

### 2.1 Apparatus and reagents

A gas-liquid phase chemiluminescence tester (self-made in the laboratory, 1000 V for negative high voltage of photocell) and a Lambda 35 ultraviolet visible spectrophotometer (PerkinElmer, USA) and MCS-2SLMP-D/5 M laminar flow mass flow controller (Alicat, USA) were used in this experiment. Formaldehyde standard gas (HCHO, 20  $\mu\text{g L}^{-1}$ , No. I63801091, filled with nitrogen, Wuhan New Reid Special Gas, CHINA), gallic acid ( $\text{C}_7\text{H}_6\text{O}_5$ , 99%, Aladdin, USA), potassium dichromate ( $\text{K}_2\text{Cr}_2\text{O}_7$ , 99.8%, Aladdin, USA), ammonium ferric sulfate, anhydrous ethanol ( $\text{NH}_4\text{Fe}(\text{SO}_4)_2$ ,  $\text{C}_2\text{H}_6\text{O}$ , AR, Chengdu Kelong Chemical Reagent, CHINA), phenol reagent ( $\text{C}_8\text{H}_9\text{N}_3\text{S}\cdot\text{HCl}\cdot\text{H}_2\text{O}$ , AR, Tianjin Guangfu Fine Chemical, CHINA), sodium hydroxide, sulfuric acid, formaldehyde ( $\text{NaOH}$ ,  $\text{H}_2\text{SO}_4$ , HCHO, AR, Chongqing Chuandong Chemical, CHINA), Tween 20, Tween 80, ethylene glycol, Brij58, and TritonX-100 ( $\text{C}_{26}\text{H}_{50}\text{O}_{10}$ ,  $\text{C}_{24}\text{H}_{44}\text{O}_6$ ,  $(\text{CH}_2\text{OH})_2$ ,  $\text{C}_{56}\text{H}_{114}\text{O}_{21}$ ,  $\text{C}_{18}\text{H}_{28}\text{O}_5$ , AR, Sigma, USA) were used as reagents.

### 2.2 CL assay for the detection of HCHO

The working principal diagram of the gas-liquid phase chemiluminescence tester is shown in Fig. 1. The reaction reagents (No. 1–3) and formaldehyde sample gas (No. 6) were pumped into the reactor (No. 8) through the peristaltic pump (No. 5) and vacuum pump (No. 9), respectively. The sample gas flow was controlled by a mass flowmeter (No. 7). The reaction reagent and the formaldehyde sample gas gave rise to a chemiluminescence reaction in the reactor, and a small amount of current was generated by an optical signal after being received by the photocell. It was received and processed by an industrial computer (No. 11) after the circuit was amplified. The obtained electrical signal was related to the intensity of the luminescence. Within a certain concentration range, the luminescence signal showed a linear relationship with the formaldehyde content in the sample gas. The reagent and waste gas after the

reaction was removed in time and collected in the waste liquid bottle (No. 4) and waste gas absorption bottle (No. 10).

A schematic diagram of the reactor is shown in Fig. 1c. An oval reaction zone (No. 5) in the middle of the reactor was paved with a layer of hydrophilic polymer materials. After the liquid reagent was pumped into the reaction zone, a uniform liquid film was formed. The gas sample into the reactor was used to keep the laminar flow state and to fully contact with the liquid membrane. The CL reaction occurred at the gas-liquid heterogeneous interface, and a luminescence signal was generated.

## 3 Results and discussion

### 3.1 Optimization of the chemiluminescence reagent system

In order to obtain a high detection sensitivity, the luminescent reagent system needs to be optimized. Each luminescent reagent and the formaldehyde standard gas at a certain concentration were pumped into the reactor at the same time for the chemiluminescence reaction and signal measurement. The variation curve of the luminescence intensity with the reagent concentration was recorded, and the optimal reagent concentration and luminescence intensity at this concentration were determined. The instrument parameters of the gas-liquid phase chemiluminescence tester set when optimizing the reagent system were: photocell negative high voltage of 1000 V, peristaltic pump speed of 2 (at the flow rate of 40  $\mu\text{L min}^{-1}$ ), vacuum pump flow rate of 2  $\text{L min}^{-1}$ , formaldehyde gas (entering the reactor) concentration was 2.3  $\mu\text{g L}^{-1}$ , and data filtering of 0.4.

**3.1.1 Optimization of the gallic acid solution.** The effect of the gallic acid concentration on the luminescence intensity was investigated. Gallic acid was used as the luminescent reagent in this experiment and its concentration directly affects the luminescence intensity of the reaction. The results are shown in Fig. S1a.<sup>†</sup> When formaldehyde was introduced into the system, the reaction luminescence intensity first increased and then decreased with the increase in the gallic acid concentration. The luminescence signal was the strongest at the gallic acid concentration of 0.03 mol  $\text{L}^{-1}$ . Therefore, the optimized concentration of gallic acid was 0.03 mol  $\text{L}^{-1}$ .

### 3.1.2 Optimization of the potassium dichromate solution.

The effect of the potassium dichromate concentration on the luminescence intensity was investigated at the concentration of 0.03 mol  $\text{L}^{-1}$  gallic acid while keeping other conditions unchanged. Potassium dichromate was an oxidant in the system, and its concentration also has a great influence on the luminescence signal. As shown in Fig. S1b,<sup>†</sup> with the increase in the concentration of the potassium dichromate solution, the chemiluminescence intensity of the system first increased significantly and then decreased gradually. The possible reason is that potassium dichromate is a colored solution, and the excessive concentration will lead to self-absorption. Therefore, the optimized concentration of the potassium dichromate solution was  $5.0 \times 10^{-4}$  mol  $\text{L}^{-1}$ .

**3.1.3 Optimization of the reaction medium.** The effects of an alkaline medium NaOH and acidic medium  $\text{H}_2\text{SO}_4$  on the chemiluminescence intensity of the system were investigated.



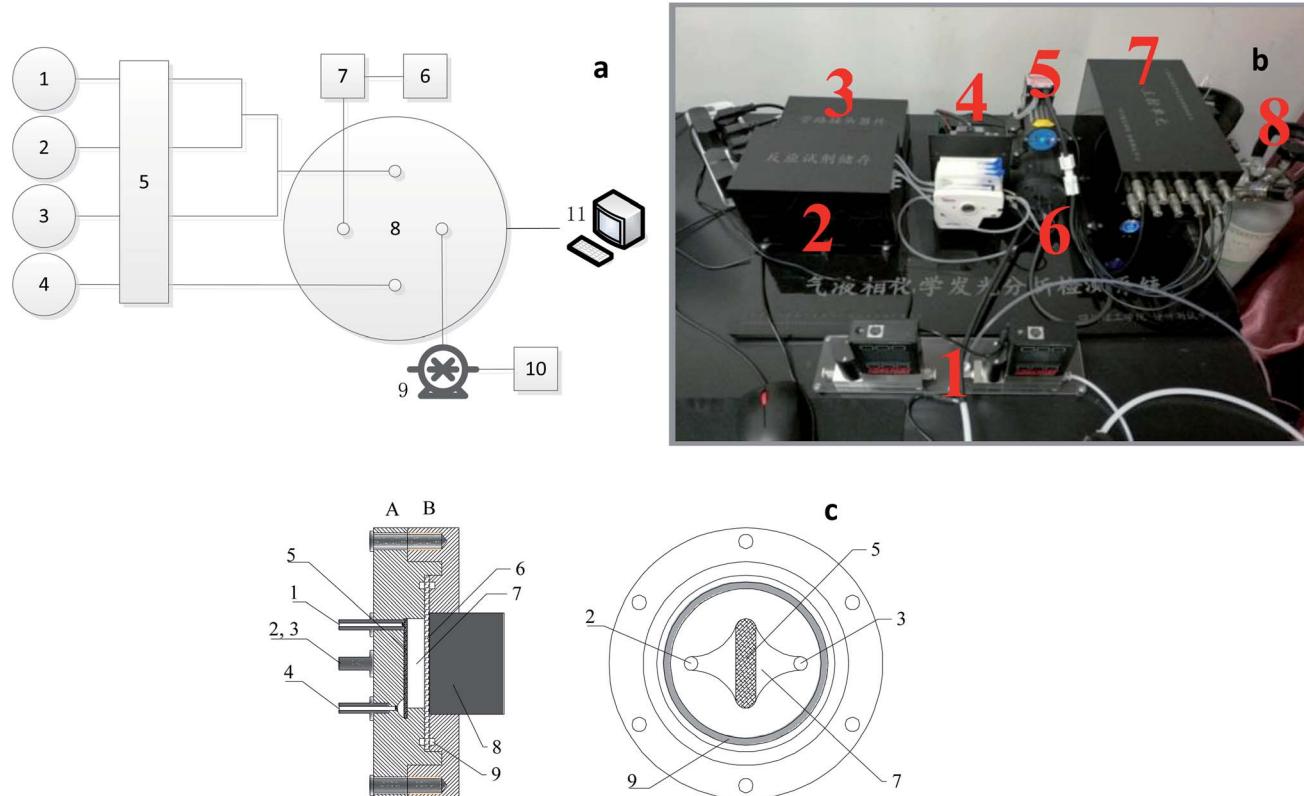


Fig. 1 (a) Schematic diagram of the chemiluminescence analyser: (1) gallic acid solution; (2) potassium dichromate solution; (3) sodium hydroxide solution; (4) waste liquid bottle; (5) multi-channel peristaltic pump; (6) formaldehyde like gas; (7) mass flow meter; (8) reactor; (9) vacuum pump; (10) waste gas absorption bottle; (11) control display system; (b) gas–liquid phase chemiluminescence analysis system; (1) mass flowmeter; (2) reaction reagent; (3) various joints; (4) veristaltic pump; (5) vacuum pump; (6) reactor; (7) control module; (8) standard gas; (c) (1 and 4) liquid inlet and outlet; (2 and 3) air inlet and outlet; (5) reaction interface; (6)quartz window; (7) reaction chamber; (8) photocell; (9) seal ring; (A) reactor front; (B) reactor rear.

Using the optimum potassium dichromate and gallic acid concentrations, the luminous intensity of the system reaction was tested when the concentrations of NaOH and  $\text{H}_2\text{SO}_4$  solutions were 0.01, 0.03, 0.05, 0.07, 0.09, 0.10, 0.30 and 0.50 mol L<sup>-1</sup>, respectively. As shown in Fig. S1c,† the luminous intensity of the system first increases rapidly with the increase in the NaOH concentration, reaches the maximum at 0.09 mol L<sup>-1</sup>, decreases rapidly after a short period of stability, and tends to be stable after 0.30 mol L<sup>-1</sup>. When  $\text{H}_2\text{SO}_4$  was selected as the reaction medium, the luminescence intensity of the system almost remained unchanged by the change in the  $\text{H}_2\text{SO}_4$  concentration. It was speculated that in the  $\text{H}_2\text{SO}_4$  medium, potassium dichromate was highly oxidizing and reacted quickly with reducing gallic acid, preventing formaldehyde gas from participating in the reaction between potassium dichromate and gallic acid. Therefore, 0.09 mol L<sup>-1</sup> NaOH alkali solution was selected as the reaction medium.

**3.1.4 Surfactant optimization.** Surfactants have the characteristics of adsorption, penetration, dispersion and solubilization, which can reduce the liquid surface tension, change the interface properties and greatly enhance the chemiluminescence intensity of the reaction system.<sup>21</sup> Under the above-optimized experimental conditions, six surfactants

(absolute ethanol, Tween 20, Tween 80, ethylene glycol and brij58) were added to the gallic acid solution at four different concentrations of 0.5%, 0.7%, 1.0% and 1.2%, and their chemical luminescence intensity was measured in the gas–liquid chemiluminescence system. As shown in Fig. 2a, in addition to Brij58, the other four surfactants caused significant enhancement on the luminous intensity of the system, but there enhancement effects had small differences. In order to further optimize the enhancement effect of the surfactants, the four surfactants were compounded in pairs with an additive, and their luminous intensity was measured. As shown in Fig. 2b, the chemiluminescence intensity of the system further enhanced when different surfactants were added. The combination of glycol and Tween 80 makes the luminescence intensity of the system reach 399.

### 3.2 System parameter selection

In order to obtain a stronger chemiluminescence signal, the effects of the liquid inlet speed and reagent mixing tube length on the intensity of the chemiluminescence signal were investigated under the condition of optimizing the luminescent reagent system. The reagent feeding speed was controlled by the peristaltic pump. The luminous instrument was equipped with



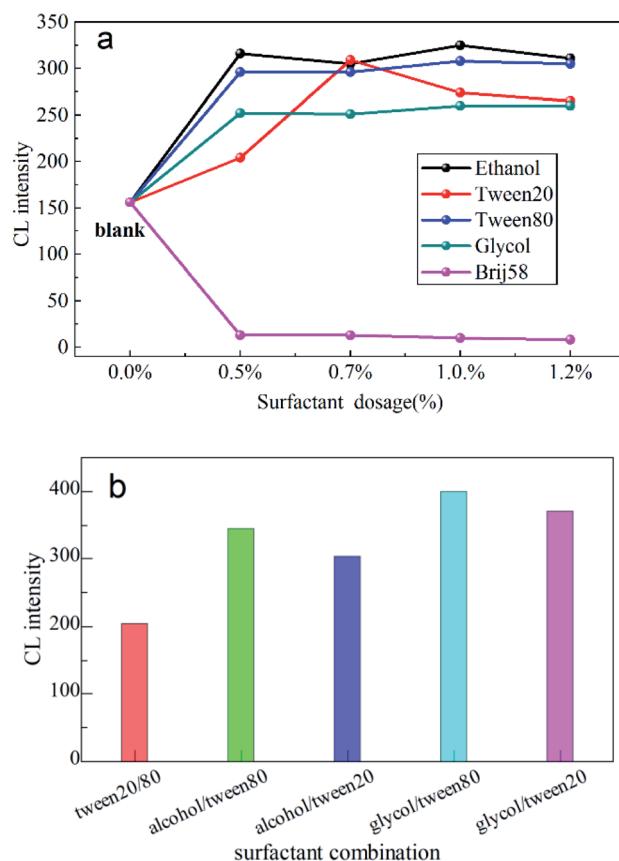


Fig. 2 (a) Different surfactant luminous intensity diagram; (b) optimized luminescence intensity map of different surfactants.

two pump speeds. When the pump speed was the first gear ( $20 \mu\text{L min}^{-1}$ ), the chemiluminescence intensity of the system was 257, and when the pump speed was the second gear ( $40 \mu\text{L min}^{-1}$ ), the chemiluminescence intensity of the system was 399. The results showed that when the pump speed was  $40 \mu\text{L min}^{-1}$ , the luminous intensity was high and the chemiluminescence intensity was stable and has a large signal-to-noise ratio.

Each reaction reagent was mixed before entering the reactor, and the length of the tube after mixing affected the mixing degree and reaction degree of the reagent before entering the reactor. The chemiluminescence intensity of the system was tested when the length of the pipeline from the mixing node to the reactor was 2, 4, 6, 8, 10, 12, 14, 16, 18 and 20 cm, and the optimal tube length was determined. As shown in Fig. 3, the luminous intensity of the system gradually increases first and then decreases with the increase in the tube length, but the luminous intensity reaches the maximum after 10 cm. Therefore, the optimal reagent mixing tube length was determined to be 10 cm.

### 3.3 Interference test

Under the optimized experimental conditions,  $10 \mu\text{g L}^{-1}$  mixture of acetaldehyde, nitrogen dioxide, sulfur dioxide, ozone, hydrogen sulfide and carbon monoxide diluted with zero

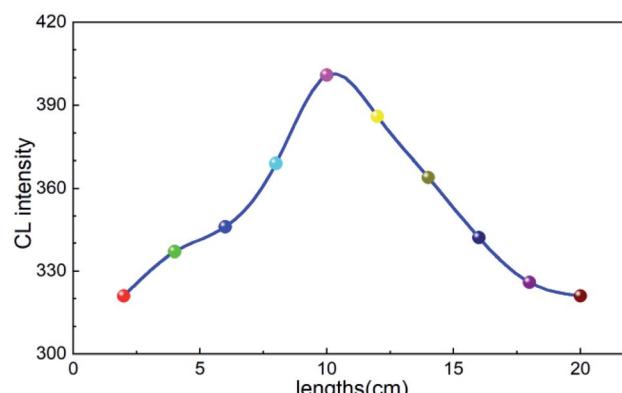


Fig. 3 Effect of tube lengths on chemiluminescence intensity.

air was pumped into the system for chemiluminescence measurement, and the luminous intensity was recorded. As shown in Fig. 4, After each interfering gas was introduced, the luminous intensity of the system showed no evident change and was negligible. Therefore, the interference gas of  $10 \mu\text{g L}^{-1}$  acetaldehyde, nitrogen dioxide, sulfur dioxide, ozone, hydrogen sulfide and carbon monoxide have no impact on the system detection.

### 3.4 Comparison between the phenol reagent method and instrument method

The phenol reagent method was a classical analytical method for the detection of formaldehyde in air,<sup>22</sup> which has the advantages of having a low detection limit, high sensitivity and good precision. However, it takes a long time to analyze, requires skilled testers and cannot test online. The phenol reagent method was compared with the instrument method, and the measured data were compared and analyzed. The detection limit was calculated as follows:  $D = 3S_b/S$ , wherein  $D$  is the detection limit,  $\mu\text{g L}^{-1}$ ;  $S_b$  is the standard deviation of

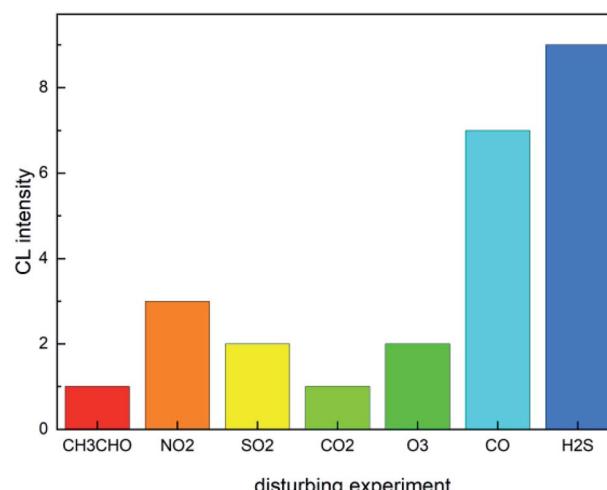


Fig. 4 Effect of interfering substances on the formaldehyde detection system.



blank signal, %;  $S$  is the equation sensitivity (the slope of the standard curve).

**3.4.1 Phenol reagent method.** The formaldehyde standard solution, color developer, and phenol reagent absorption solution were prepared. The 5 mL absorption solution was packed in a 10 mL bubble absorption tube, and the gas was collected at a flow rate of 0.5 L min<sup>-1</sup> for 20 minutes, and then the collected samples were analyzed. The standard curve was drawn using the formaldehyde content ( $x$ ) as the abscissa and the absorbance ( $y$ ) as the ordinate. As shown in Fig. S2(a),<sup>†</sup> the linear equation of the standard absorption curve of the phenol reagent was  $y = 0.0555x + 0.011$ , the linear correlation coefficient  $R^2 = 0.997$ , and the detection limit of the method was 0.066  $\mu\text{g L}^{-1}$ .

**3.4.2 Instrument method.** Under the optimized experimental conditions, the reagent system and different concentrations of formaldehyde were injected into the gas–liquid phase system for chemiluminescence measurement, and the luminescence intensity was recorded. The standard curve was drawn with the formaldehyde content ( $x$ ) as the abscissa and luminous intensity ( $y$ ) as the ordinate. As shown in Fig. S2(b),<sup>†</sup> the linear equation of the standard absorption curve of the gas–liquid chemiluminescence system was  $y = 208x + 29.667$ , the linear correlation coefficient  $R^2 = 0.997$ , and the detection limit of the method was 0.003  $\mu\text{g L}^{-1}$ . Motyka<sup>23</sup> reported a method for the determination of HCHO in the air, which belongs to Trautzsuch as Schorin (TSR) continuous CL. The detection limit was  $0.66 \times 10^{-3}$   $\mu\text{g L}^{-1}$  but there was no reaction interface in this reaction system and the  $\text{H}_2\text{O}_2$  is an unstable oxidant in the reagent system which cannot ensure the stable progress of luminescence reaction.

Comparing the test results of the phenol reagent method and the instrument method showed that the linear relationship of both the two methods was better, but the detection limit of the instrument method was lower. Through the above experiments, it was found that four formaldehyde samples were detected by the two methods, and the results are shown in Table 1. The experimental results showed that the measured results of 0.3  $\mu\text{g L}^{-1}$  and 0.5  $\mu\text{g L}^{-1}$  standard gas by the instrument method and phenol reagent method were basically consistent but for indoor and outdoor air samples in the laboratory, phenol reagent method was not detected. While the instrument method detects specific values and the operation of the instrument method was simpler than the phenol reagent method, there was no need to absorb the formaldehyde gas and the test time was shorter.

### 3.5 Chemiluminescence reaction mechanism

The gas–liquid chemiluminescence reaction system is obviously complex. As an element with variable valencies, chromium can

enter Haber–Weiss-type reactions leading to the generation of hydroxyl radicals ( $\cdot\text{OH}$ ).<sup>24</sup> The reaction of Cr(vi) with different reductants gives rise to Cr(v) complexes. The hydroxyl radical generation by Cr(vi) reduction has been demonstrated in a variety of model systems.<sup>25</sup> As a reducing agent, Ga is oxidized by Cr(vi) in the CL reaction. In order to explore the possible chemiluminescence mechanism, the luminescence wavelength of the reaction system was measured and studied by the BPCL-GP21Q-TGC ultra-weak chemiluminescence measuring instrument, and the chemiluminescence spectrum of  $\text{K}_2\text{Cr}_2\text{O}_7$ –Ga–KOH–HCHO was drawn.

As shown in Fig. 5, the luminescence signal was low when there was no formaldehyde in the reagent system. When formaldehyde was added, the strongest light was emitted by the chemiluminescence system at a wavelength of 665 nm. It was speculated that the luminescent body of the reaction was singlet oxygen ( $^1\text{O}_2$ ).<sup>26</sup>  $^1\text{O}_2$  is very active and can easily form a bimolecular singlet oxygen in the excited state ( $^1\text{O}_2\text{}_2^*$ ). CL occurs when the ( $^1\text{O}_2\text{}_2^*$ ) generated during the reaction transitions back to the ground state. It is speculated that the energy generated during the reaction between HCHO and  $\cdot\text{OH}$  is transferred to the excited bimolecular singlet oxygen,<sup>27</sup> and then transitions back to the ground state to produce CL. The light wavelength released after returning to the ground state is about 603–703 nm. The specific reaction mechanism remains to be explored. The CL emission in the Ga– $\text{K}_2\text{Cr}_2\text{O}_7$ –KOH system is very weak and thus, the HCHO content plays the main role in enhancing the CL intensity of this system. The possible

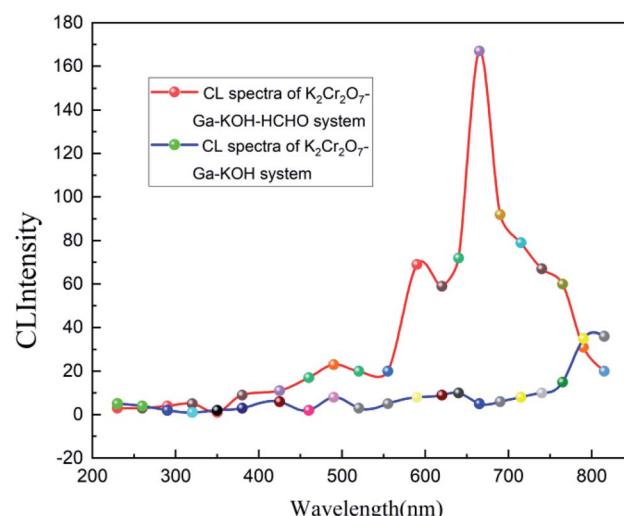


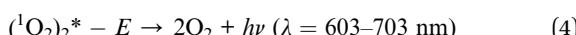
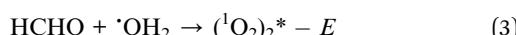
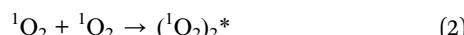
Fig. 5 CL spectra of  $\text{K}_2\text{Cr}_2\text{O}_7$ –Ga–KOH–HCHO and  $\text{K}_2\text{Cr}_2\text{O}_7$ –Ga–KOH systems.

Table 1 The results of four samples were compared between phenol reagent method and instrument method

Test methods	Indoor air	Outdoor air	0.30 $\mu\text{g L}^{-1}$ HCHO	0.50 $\mu\text{g L}^{-1}$ HCHO
Phenol reagent method	0.061 $\mu\text{g L}^{-1}$	ND	0.291 $\mu\text{g L}^{-1}$	0.488 $\mu\text{g L}^{-1}$
Instrument method	0.059 $\mu\text{g L}^{-1}$	0.011 $\mu\text{g L}^{-1}$	0.290 $\mu\text{g L}^{-1}$	0.480 $\mu\text{g L}^{-1}$



sequence of reactions occurring during the CL signal generation was as follows.



## 4 Conclusions

In this study, a formaldehyde-gallic acid-potassium dichromate chemiluminescence system under alkaline conditions was studied and a new method for the continuous determination of gaseous formaldehyde based on a gas-liquid chemiluminescence detection system was established. This method was based on the chemiluminescence reaction of the formaldehyde gas directly with gallic acid and potassium dichromate solution under alkaline conditions at the gas-liquid interface. The optimal concentration of each reagent and other reaction conditions were discussed, and compared with the phenol reagent method. The results showed that the chemiluminescence method has the advantages of providing accurate results, simple operation, low detection limit (as low as  $0.003 \mu\text{g L}^{-1}$ ), and nitrogen dioxide, sulfur dioxide, ozone and other gases have no obvious interference with the system. The method can be extended to the quantitative detection of other trace gases in ambient air.

In order to further optimize the formaldehyde gallic acid potassium dichromate chemiluminescence system, the material of the reaction interface can be optimized to improve the signal intensity. The liquid forms a liquid film on the surface of the reaction interface to increase the contact area of atmospheric liquid and reduce the detection limit. So far, the non-woven fabric used in this study is the best choice for the reaction interface. In addition, optimizing the sample gas path can increase the flow stability of the sample gas and reduce the freedom of system noise, so as to improve the optical stability of the gas-liquid chemiluminescence detection system. With the improvement of accuracy, the gas-liquid chemiluminescence detection system can not only be applied to the real-time detection of indoor formaldehyde but also to measure the spatial distribution of atmospheric HCHO concentration by carrying equipment.

## Conflicts of interest

There are no conflicts to declare.

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