A Film Bulk Acoustic Resonator-Based Sensor with AlN Piezoelectric Material for Detecting Ethanol and Acetone

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Received 18 July 2022; Revised 17 September 2022; Accepted 21 October 2022; Published 21 November 2022

Measurement of a volatile solution is essential for laboratory safety and hospital clinic safety. In this paper, we present an ethanol-sensing and acetone-sensing device using an AlN piezoelectric material-based film bulk acoustic resonator (FBAR). In order to realize volatile solution sensing, the AlN-based FBAR was designed, fabricated, and characterized. In our sensor structure, the upper electrode is a Ti/Au (30nm/150nm) composite electrode, the bottom electrode is Mo material with 150nm thickness, and the piezoelectric sensing material is 0.8 μm thickness AlN. We conducted the experiment of ethanol measurement and acetone measurement by using this FBAR detector on the probe station within the vector network analyzer. The resonance frequency of the FBAR detector decreased as the concentration of ethanol increases, while under the circumstance of acetone concentration increasing, the detector’s response is the opposite. The sensing mechanisms of both ethanol measurement and acetone measurement are discussed in this paper, demonstrating that this FBAR detector could be able to distinguish acetone from ethanol due to different sensing mechanisms.

1. Introduction

During the last decade, the study of film bulk acoustic resonators (FBAR) has been used for the design of biochemical sensors. Compared to traditional detection method, including solution chromatography (GC) and liquid chromatography (LC) [1], FBAR detectors bring low fabrication cost and short detection time, together with the advantage of being compact and easy to carry [2]. Our study group in university focus on the design, fabrication, and measurement of micro biochemical sensors, thus, FBAR chips for the use of volatile solution sensing have come into our vision. For sensing principle, the detector mechanism of FBAR is similar as quartz crystal microbalance (QCM), all belonging to mass-sensitive detectors [3]. Under normal circumstance, the mass-sensitive detector is proportional to the fundamental resonance frequency. Once the biomedical target molecule binds to the piezoelectric area of the FBAR sensor, the increasing mass would lead to a natural resonance frequency decrease [4]. The film bulk acoustic resonator (FBAR) usually consists of a sputtered piezoelectric thin film (ZnO/AlN) sandwiched by two metal layers (top and bottom electrodes). The top and bottom electrode materials could be Au, Al, Mo, or composite electrodes such as Ti/Au, Ni/Au. Under the certain frequency, a resonance condition occurs of the thickness of the piezoelectric material thin film d is equal to an integer multiple of a half of the acoustic wavelength λ. Before the appearance of FBAR detectors, there have been other kinds of traditional filters like surface acoustic wave (SAW) and bulk acoustic surface (BAW). Different with the former traditional filters, the FBAR sensors are fabricated on silicon substrate by using MEMS technology and piezoelectric membrane [5]. Thus, the production of FBAR sensors would overcome the shortage of the former traditional filters, becoming vital cells in the next generation of wireless communication products. The FBAR would satisfy multifunction, multiband, and multiprotocol systems, which is the development direction of filters in the new century. For piezoelectric material choice, we usually have three selections: AlN, ZnO, and PZT [6]. Comparatively, ZnO FBAR would allow sensing characteristic to many
targets, while our sputtering membrane is more inclined to prepare AlN piezoelectric membrane. Although PZT brings higher piezoelectric properties, its fabrication process is much more difficult than that of AlN. And the cost of PZT is more expensive than that of AlN. Here, in this paper we report an AlN-based film bulk acoustic resonator (FBAR) detector for sensing ethanol and acetone, the bottom electrode is a 150 nm Mo electrodes, the top electrode is a Ti/Au composite electrode with a corresponding thickness of 30 nm/150 nm.

With the increasing importance of clinical diagnosis and medical technology, studies of biochemical sensors have been research hotspots in recent years [7]. Among numerous micro chemical sensors, FBAR has been used for important liquid and gas-phase chemical substance detection due to its significant performance and sensitivity [8]. For instance, Zhang et al. ’s paper introduced the design of an FBAR gas sensor based on ZnO film [9]. A. Kumar and Prajesh’s review also introduced the potential of FBAR for gas sensing applications [10]. For liquid detection, A. Soltan et al. ’s paper presented the FBAR microwave sensor for liquid classification [11]. Lin et al. ’s study has revealed an FBAR based explosive trace detection sensor for TNT detection [12]. FBAR sensor could also be utilized to predict frequency shift in viscous media due to Zhao et al. ’s study [13]. Furthermore, FBAR sensors could be modified and functionalized to be pesticide detectors for organophosphate measurement [14]. In our work, we finished the FBAR chip and produced this sensor to distinguish two common laboratory volatile gases. Ethanol and acetone are commonly used volatile chemicals in laboratory and medical occasion. Semiconductor sensors are widely used and accepted as ethanol and acetone detectors. There are many advantages for these kinds of semiconductor sensors, including low fabrication cost, sensitivity and sufficient detection limit. However, there is still one need to distinguish acetone vapor from ethanol vapors effectively. In our study, we fabricated the AlN-based FBAR sensor and conducted the experiment to measure ethanol and acetone, we have observed different responses. Then, the sensing mechanism of the FBAR sensor for these two volatile solutions is described and explained.

2. Sensor Structure

In our design, the sensor structure is one AlN-based FBAR sensor. The sputtered piezoelectric thin film of 0.8 μm AlN is sandwiched by two metal electrodes. The top electrode of this FBAR is Ni/Au with the thickness of 30 nm/150 nm; the bottom electrode is 150 nm Mo. We choose the p-doped Si substrate; one side of the substrate is deposited by the AlN/ Mo/AlN structure. The sputtering process of the 150 nm Mo film was conducted by the SPTS physical vapor deposition (PVD) system, the deposition is controlled by the argon flow. While the sputtering process of AlN piezoelectric material was also conducted by the SPTS physical vapor deposition (PVD) system, which allow the nitrogen solution and argon solution to flow to the Al target inside the equipment. The top electrode material, Ti/Au was conducted by the electron beam evaporation equipment, ULVAC Ei-5z.

Before the experiment, we need to fix the FBAR sensor on to the platform of the probe station by magnetic adsorption, under probe station’ microscope view, the FBAR arrays’ surface are shown in Figure 1. During the observation process, silica gel was coated around the detection area to form a chamber. The AlN piezoelectric material was inside the chamber. For each one-detection experiment, we dropped 10 μL solutions to be detected on to the AlN piezoelectric area inside the chamber. After about 10 s we recorded the resonance frequency of the chosen experimental module of the FBAR arrays. The resonance frequency of the FBAR was monitored with an Agilent E5071C network analyzer (Agilent, Santa Clara, CA). The probe station and Keithley source meter were also prepared to help record the experimental data. For sensing mechanism, the FBAR detection method is based on the inverse piezoelectric effect, when electrical signal is applied to the FBAR sensor, the electrical signal could be transformed the electrical signal into sound signal. The resonance frequency of this sound signal would represent the information of piezoelectric thin films and analysts. From the resonance frequency data, we could indirectly obtain the information of analyst to be detected.

In summary, in this paper, FBAR sensors are designed for sensing volatile solutions such as acetone and ethanol, by using the AlN piezoelectric material. FBAR technology could also be utilized to produce oscillators, duplexers, and other high-performance frequency devices. Compared to graphene sensors, the piezoelectric material AlN could be deposited by specific magnetic sputtering equipment within fewer fabrication hours, which is more efficient than the graphene fabrication period, avoiding graphene transfer craft by human labour. The sensor structure is shown in Figure 2.

3. Fabrication and Characterization

3.1. Fabrication of Microchip. In the beginning of sensor design, we finished our FBAR layout design using the software L-Edit. Along with the designed FBAR layout, we set the fabrication steps as the following procedure. The first step is the growth of AlN/Mo/AlN material on the silicon substrate by magnetron sputtering. The growth parameters of AlN and Mo during sputtering process are listed in Table 1. An AlN thin film was deposited in the magnetron sputtering process, which utilized nitrogen flow and argon flow to the Al target. The growth parameters of AlN and Mo are listed in Table 1. Second, the bottom electrode deposition was done. The photolithography process was conducted on the front side of the chip, and then, the step electron beam evaporation for 150 nm Mo was finished. Then we conducted the stripping by the ultrasound metal stripping method. For the third step, the sputtering and patterning of the piezoelectric thin films were accomplished. The SiO2 mask layer was deposited by the PECVD method, and the photolithography process was done. Afterwards, the SiO2 layer was etched by the RIE method, satisfying the condition of 50 nm over etched within the etching rate of 140 nm/min.
The chip was consequently deep etched in the BOE solution (1:7) for 5 s. Ultimately the AlN layer etching was done with a stable etching rate of 177 nm/min. For the fourth step, the upper electrode deposition was done. The front side of the chip was processed with the photolithography step. After postbake and removing the photoresist, the step electron beam evaporation for the 30 nm/150 nm thickness Ti/Au was finished. Then, we conducted the stripping by the ultrasound metal stripping method. The last fabrication step of the whole FBAR process is back etching. Then, the photolithography process was done. After that, the SiO₂ layer was over etched for 50 nm. The crucial point of the back etch is deep silicon etching for 25 min with 150 cycles. The back side of the chip needs to be observed to check whether the deep silicon etching is correctly accomplished. After the deep silicon etching, the chip surface was cleaned to remove the surface oil. When the deep silicon etching step has been correctly finished, the chip would be cleaned by deionized water and baked in the oven. Thus, one group of FBAR chips could be fabricated successfully. The fabrication steps are listed in Figure 3. Above all the parameters of each photolithography are listed in Table 2.

3.2. Characterization. After device fabrication, the AlN-based FBAR sensor needs to be conducted characterization for checking the sensing material. In order to observe the piezoelectric material AlN surface after magnetron sputtering, the SEM and XRD characterization were done to scan the sensor surface. We conducted the SEM characterization for the AlN thin film of our FBAR, Figure 4 represent the SEM results. We could observe that there had been many particles on the silicon substrate, which was the AlN crystal of the thin film after sputtering. Table 1 was listed for the mechanical parameters of the AlN, Mo, and silicon substrate within our sensor design. Second, we conducted XRD characterization for the AlN thin film, which is shown in Figure 5 below. The XRD results revealed that there is a (002) AlN preferential orientation (2θ ≈ 36°), besides the second peak is about 2θ ≈ 40°. The χ scan pattern shows that the c-axis of the AlN film is 18.5° inclined to the normal direction.

For the design of AlN-based FBAR sensor, we used several kinds of material, which are listed in Table 3 as below. The piezoelectric material AlN has stable physical properties and fine electrical characteristics. Besides, we conducted energy-dispersive X-ray spectroscopy (EDX) characterization and the EDX results have displayed the chemical element composition ratio of the AlN react area of the FBAR surface, which is shown in Figure 6 below. In Figure 6 the purple peak and the blue peak presents the Al chemical element and the N chemical element, respectively, which provides information about the AlN piezoelectric thin film.

4. Experiment and Sensor Signal

For sensor detection, we have measured the basic performance of our AlN-based FBAR sensor; the most important indexes of the FBAR include the resonance frequency and the quality factor (Q) [15]. After the index determination, we have found that the resonance frequency of our FBAR sensor is around 3.6 GHz and the quality factor (Q) is about 1050. The quality factor index figure of comparison between graphene and piezoelectric material AlN of this FBAR sensor is shown in Figure 7.

For the two target solutions, we collected the two samples from the laboratory within the common 100 ppm. During our measurement experiment, we secure our AlN-based FBAR chip on the surface of the probe station, and we drip the target solution onto electrode area of the FBAR sensor. Each time, we drip about 10 μL volume, and test for ethanol and acetone, respectively. The resonance frequency is recorded by the network analyzer. Before dropping the solution, we measure the initial resonance frequency with the FBAR sensor from the network analyzer, and after dropping the experimental solution, we measure the resonance frequency secondarily. Then, the two resonance frequency values are compared for both the ethanol sample and the acetone sample.
Figure 3: The illustration of fabrication protocol for FBAR chip. The upper electrode is Ni/Au, the bottom electrode is Mo. The piezoelectric material of the sensor is AlN. (b) Is for the photolithography and etching of the front side, (c) is for the SiO₂ deposition, (d) is for upper electrode deposition, (e) is for back etch.

Table 2: The photolithography parameters of the fabrication.

<table>
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<th>Which step</th>
<th>Photoreist</th>
<th>Exposure time</th>
</tr>
</thead>
<tbody>
<tr>
<td>Second step</td>
<td>Front side: AZ</td>
<td>Exposure 2 s—postbake 110°C</td>
</tr>
<tr>
<td>Third step</td>
<td>Front side: AZ</td>
<td>Exposure 6.5 s, 45 s in developer</td>
</tr>
<tr>
<td>Fourth step</td>
<td>Front side: AZ</td>
<td>Exposure 2 s—postbake 110°C</td>
</tr>
<tr>
<td>Fifth step</td>
<td>Both side: AZ4620</td>
<td>Exposure 22 s, 2 min in developer</td>
</tr>
</tbody>
</table>

During the measurement of acetone, we set the bias voltage ranging from −5 V to 5 V, with a 1 V increment for each one record. Before testing acetone, we measured the initial status of the FBAR sensor, then after adding acetone, the signal was recorded twice. For each one measurement, the peak frequency and the magnitude were recorded for the values. The two comparative figures below were the frequency and magnitude of the bias voltage +1 V. After acetone adding, the resonance frequency increased to 260 kHz when the bias voltage was kept at +1 V. For 100 ppm acetone, the resonance frequency increment would fluctuate as the bias voltage changed. For most bias voltage value, the resonance frequency increment was below 200 kHz. Before acetone adding the magnitude was about −14.5 dB, while the magnitude changed to about −8 dB after dropping the acetone solution.

In our experiment, the comparative two figures of resonance frequency were done for the condition before acetone addition and after acetone addition, which are shown in Figures 8 and 9, respectively. The detailed resonance frequency values with its magnitude values are listed in Table 4.

In order to figure out the FBAR sensors’ performance and sensitivity for different concentrations of the target analyt acetone, we prepared three acetone concentration samples and detected these three samples with the same module of our experiment FBAR sensor. For the experiment of different concentration measurements, we set the bias voltage near 0 V (about 0.1 V) to reduce the voltage’s effects. After we measured the resonance frequency for the 0 ppm, 50 ppm, 100 ppm, and 150 ppm, we conducted the concentration measurement curve for acetone, which is shown in Figure 10. From the concentration curve, we could find the resonance frequency would increase with the concentration increment, while the curve would become saturated near the 150 ppm concentration; the sensor has significant sensitivity from 0 to 100 ppm range.

For sensing mechanism, after the acetone addition the adsorbed oxygen ions on the AlN thin film surface react with the acetone, and release CO₂ as a reaction product [17]. Thus, the surface chemical reaction would lead to the density change of the AlN film, resulting in the resonance frequency increase [18]. The reaction between acetone and the adsorbed oxygen is shown as follows [19]:

\[ \text{CH}_3\text{COCH}_3 + \text{O}_2 \rightarrow \text{CH}_3^+ + \text{CH}_3\text{O}^- + \text{CO}_2 + e^- \]  

(1)

From the physical principle, it is demonstrated that the resonance frequency of the FBAR could be determined by the following equation [20]:

\[ \Delta f = \frac{\sqrt{E}}{2d} \cdot \left(1 - \sqrt{1 - \left(\frac{\Delta \rho}{\rho}\right)}\right) = \frac{\sqrt{E}}{4d\sqrt{\rho}} \cdot \Delta \rho, \]

(2)

where \( E \), \( \rho \) and \( d \) are the elastic modulus, density and thickness of the AlN thin film, respectively. The variable \( v \) represents the acoustic velocity within the ZnO film and \( f \) is the resonance frequency of the FBAR [21]. From the (3) we could find that \( \Delta f \) is positively proportional to \( \Delta \rho \), if the value of \( \Delta \rho \) is positive value, the value \( \Delta f \) is influenced to be positive. It could illustrate the phenomenon that after adding acetone the resonance frequency of this FBAR sensor gets increased.

In room temperature, the oxygen is adsorbed at the AlN thin film surface of the FBAR, which is shown in the chemical balance equation as follows:
The given equation is:
\[
\text{CO}_2 + \text{H}_2\text{O} + \text{O}_2 + e^- \overset{k_1}{\rightleftharpoons} k_1^{-1}\text{O}_2^{-1} + \text{H}_2\text{CO}_3. \tag{3}
\]

In this equation, \(e^-\) stands for a conduction electron at the surface, while \(k_1\) and \(k_1^{-1}\) in this equation stand for the two rate constants, \(k_1\) stands for the rate constant of forward reaction, \(k_1^{-1}\) stands for the rate constant of reverse reaction. When the oxygen gas upon the AlN surface becomes the dissolved oxygen within the AlN thin film, the target chemical acetone could be able to react with the dissolved oxygen \[22\]. The chemical reaction \(1\) would be influenced by the dissolved oxygen reaction \[23\], which is shown in equation \(3\).

From the relationship figure between gate voltage and resonance frequency in Figure 11, we could observe that the frequency shift keeps stable during the gate voltage sweep. During the whole detection experiment, the room temperature and room humidity were kept unchanged, and the experiment was conducted by the same person. From this point of view, we could deduce that the resonance frequency change is mainly due to the acetone addition, and other factors’ influence is excludes.

During the measurement of ethanol, we set the bias voltage ranging from \(-5\) V to \(5\) V, with a \(1\) V increment for each record. Before testing acetone, we measured the initial status of the FBAR sensor, and then, after adding ethanol the signal was recorded twice. The values of peak frequency and the magnitude were recorded at each measurement. The two comparative figures below were the frequency and magnitude of the bias voltage \(+1\) V. We could observe that the peak frequency decreased after ethanol addition. With 100 ppm ethanol, the peak frequency decreased by about \(1000\) kHz compared to the situation before ethanol addition when the bias voltage was set at \(+1\) V. For magnitude, the magnitude before ethanol addition was about \(-29.8\) averagely, while after ethanol addition, the magnitude was about \(-9.3\).

In our experiment, the comparative two figures of resonance frequency were done for the condition before ethanol addition and after ethanol addition, which are shown in Figures 12 and 13, respectively. The horizontal axis stands for the resonance frequency, and the vertical axis stands for the \(S_{11}\) parameter (unit: dB) in Figure 12 (before adding ethanol) and Figure 13 (after adding ethanol). The resonance frequency values and magnitude values were recorded during

![Figure 4: The Scanning electron microscopy (SEM) pictures of the AlN thin film, (a) is for AlN surface morphology, (b) is for cross section view.](image1)

![Figure 5: X-ray diffraction (XRD) patterns of the AlN thin film, (a) 2θ scan; (b) \(\chi\) scan of the AlN (002) peak.](image2)
the measurement. The detailed resonance frequency values with its magnitude values are listed in Table 5.

For the sensing mechanism of ethanol detection, after ethanol addition, the adsorbed oxygen ions on the AlN thin film surface react with the acetone, releasing H₂O as the reaction product which is absorbed by the AlN thin film [24]. Under this circumstance the mass per unit area on the AlN surface has been influenced by the chemical reaction, causing the density to increase of this AlN thin film [25]. The density change of the AlN thin film would lead to the frequency decrease [26]. The reaction between ethanol and the adsorbed oxygen is shown as follows [27]:

\[
\text{C}_2\text{H}_5\text{OH} + \text{O}_2^- \rightarrow \text{CH}_3\text{COOH} + \text{H}_2\text{O} + e^-.
\]  

From the (4) we could deduce that \(\Delta f\) is negatively proportional to \(\Delta \rho\) [28], if the value of \(\Delta \rho\) is positive value, the value \(\Delta f\) is influenced to be minus [29]. The (4) could explain the negative correlation relationship between the resonance frequency change and the ethanol addition [30].

From the relationship figure between gate voltage and resonance frequency in Figure 14, we could observe that except for the gate voltage of +2 V, the frequency shift keeps stable. During the whole detection experiment, the room temperature and room humidity were kept unchanged, and the experiment was conducted by the same person. From this point of view, we could deduce that the resonance frequency change is mainly due to the ethanol addition, and other factors' influence is excludes. Besides, we conducted the ethanol concentration measurement experiment and finished the ethanol

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Material</th>
<th>AlN</th>
<th>Molybdenum</th>
<th>Silicone</th>
</tr>
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<tr>
<td>Density (kg/m³)</td>
<td></td>
<td>3200</td>
<td>10000</td>
<td>2300</td>
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<tr>
<td>Young's modulus (GPa)</td>
<td></td>
<td>300</td>
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<td>130</td>
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<td>Poisson's ratio</td>
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<td>0.31</td>
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</table>

Figure 6: The energy-dispersive X-ray spectroscopy (EDX) characterization results for AlN area of FBAR sensor surface.

Figure 7: The quality factor (Q) figure comparison between graphene and AlN, black spot stands for graphene and blue spot stands for AlN in our sensor.

Figure 8: The resonance frequency figure of the AlN-based FBAR sensor before adding acetone, the horizontal axis is frequency while the vertical axis is \(S_{11}\) parameter.

Figure 9: The resonance frequency figure of the AlN-based FBAR sensor after adding acetone, the horizontal axis is frequency while the vertical axis is \(S_{11}\) parameter.

\[
\text{C}_2\text{H}_5\text{OH} + \text{O}_2^- \rightarrow \text{CH}_3\text{COOH} + \text{H}_2\text{O} + e^-.
\]  

Table 3: The physical material parameters of the FBAR sensor.
We prepared three ethanol concentration samples and detected these three samples with the same module of our experiment FBAR sensor. For the experiment of different concentration measurements, we set the bias voltage near 0 V (about 0.1 V) to reduce the voltage’s effects. After we measured the resonance frequency for the 0 ppm, 50 ppm, 100 ppm, and 150 ppm, the concentration figure of ethanol is finished in Figure 10: The concentration measurement results for the acetone samples.

<table>
<thead>
<tr>
<th>Voltage</th>
<th>Initial state (GHz)</th>
<th>Magnitude</th>
<th>After adding acetone (GHz)</th>
<th>Magnitude</th>
</tr>
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<tbody>
<tr>
<td>+5 V</td>
<td>3.61532</td>
<td>-14.29567</td>
<td>3.61559</td>
<td>-8.0435</td>
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<tr>
<td>+4 V</td>
<td>3.61536</td>
<td>-14.3501</td>
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<td>+3 V</td>
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<tr>
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<tr>
<td>+1 V</td>
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<tr>
<td>0</td>
<td>3.6151</td>
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<tr>
<td>-1 V</td>
<td>3.61518</td>
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<td>3.61509</td>
<td>-14.53756</td>
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<td>-5 V</td>
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<td>-14.70183</td>
<td>3.61525</td>
<td>-8.08964</td>
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</table>

Figure 10: The concentration measurement results for the acetone samples.

Figure 11: The figure for influence of gate voltage change on the resonance frequency shift value during acetone experiment. The black curve $f_1$ stands for the resonance frequency before acetone addition, the red curve $f_2$ stands for the resonance frequency after acetone addition.

Figure 12: The resonance frequency figure of the AlN-based FBAR sensor before adding ethanol, the horizontal axis is frequency while the vertical axis is $S_{11}$ parameter.

Figure 13: The resonance frequency figure of the AlN-based FBAR sensor after adding ethanol, the horizontal axis is frequency while the vertical axis is $S_{11}$ parameter.
Figure 15. From the concentration curve, we could find the sensor has significant sensitivity from 0 to 100 ppm. The real-time frequency shift figures are shown in Figure 16 below. For acetone and ethanol, the real-time curve was recorded under a 0 V voltage.

Table 5: The resonance frequency comparison of before and after adding ethanol solutions.

<table>
<thead>
<tr>
<th>Voltage</th>
<th>Initial state (GHz)</th>
<th>Magnitude</th>
<th>After adding acetone (GHz)</th>
<th>Magnitude</th>
</tr>
</thead>
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<tr>
<td>+5 V</td>
<td>3.617</td>
<td>−29.4286</td>
<td>3.61593</td>
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<td>+4 V</td>
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<tr>
<td>+2 V</td>
<td>3.61681</td>
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<tr>
<td>0</td>
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<td>−29.7705</td>
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<tr>
<td>−1 V</td>
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<td>−2 V</td>
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<td>−4 V</td>
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<td>−5 V</td>
<td>3.61646</td>
<td>−29.99142</td>
<td>3.61568</td>
<td>−9.29297</td>
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</table>

After the detection experiment, we have summarized some sensor reliability data of our FBAR sensor. The AlN-based FBAR could distinguish acetone and ethanol from 0 ppm to near 150 ppm in laboratory measurement. Among one fabrication batch of FBAR devices, about 95% of the devices could be able to detect sensor signals. For one FBAR chip, about 92% of the modules could effectively detect the analyst. The FBAR working temperature is from −30°C to 50°C, for general use and high precision testing the FBAR works under room temperature because extreme temperatures would lead to temperature drift for frequency measurement. The temperature coefficient of the FBAR sensor is usually 20 to −35 ppm/°C. But recently, the Suzhou Institute of Nano-tech and Nano-Bionics had conducted the research of temperature fluctuation compensation system within the FBAR sensor; thus, the temperature drift’s influence on frequency measurement would be weakened to some extent [31]. Above all, we calculated the sensitivity of this AlN FBAR sensor for acetone and ethanol detection, and compared it with the ZnO FBAR in previous literature [32]. The sensitivity indexes are listed in Table 6. The mean time to failure of our FBAR chips is about 28.5 days based on our failure analysis.
5. Conclusions

In summary, we have developed an AlN-based FBAR sensor and conducted acetone and ethanol sensing experiment using this FBAR sensor. From our experiment, we could find that the resonance frequency of our FBAR sensor increases after the acetone sample addition, while the FBAR sensor presents the opposite response after ethanol addition. For acetone detection’s sensing mechanism, the acetone reacts with the free oxygen within the AlN surface, the reaction products include CO₂ and other ions which would volatilize or dissolve. The AlN thin film’s entire thickness tends to be attenuated by this chemical reaction, and the film density tends to be increased. In this case, this property change is in charge of the frequency upshift. On the other side, for ethanol detection’s sensing mechanism, the ethanol reacts with the oxygen ions within the AlN surface; the reaction products include H₂O and acetic acid. The AlN thin film’s entire thickness tends to grow thicker due to this chemical reaction, because the reaction products are nonvolatile liquid. In that case the density of the AlN thin film tends to decrease, so the peak resonance frequency is attenuated. Furthermore, this AlN-based FBAR sensor could be used to detect other kinds of chemical volatile gases. However, our this work only conducted the FBAR sensor for detection of those two VOCs commonly used in laboratory, for more complex volatile organic compounds, the FBAR surface needs to be modified by the special chemical functional groups. In that case, with the concern of selectivity, more experiments would be scheduled to study the sensing mechanism of FBAR to different targets by different functionalization processes. The research significance of FBAR sensor needs to be deeply demonstrated and understood from laboratory experiments and sensor theoretical study, ultimately realizing microminiaturization and diversity detection.

Data Availability

The data are collected from Our own experiment.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

Acknowledgments

This paper’s project was supported by the research start-up funding No. GY-Z20045 of Fujian University of Technology. The authors are grateful to Professor Bao-Shun Zhang for his project supports, and Wen-Kui Lin for his experimental assistance from Suzhou Institute of Nano-tech and Nano-Bionics, Chinese Academy of Science. The authors also sincerely appreciate Professor Guan-Wen Zhang and Professor Jiang-Han Chen’s help from Guangdong Institute of Analysis (China National Analytical Guangzhou), the experiment was conducted with their technical guidance and suggestions.

References


