Multi-Frequency Band Pyroelectric Sensors

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Abstract: A methodology is proposed for designing a multi-frequency band pyroelectric sensor which can detect subjects with various frequencies or velocities. A structure with dual pyroelectric layers, consisting of a thinner sputtered ZnO layer and a thicker aerosol ZnO layer, proved helpful in the development of the proposed sensor. The thinner sputtered ZnO layer with a small thermal capacity and a rapid response accomplishes a high-frequency sensing task, while the thicker aerosol ZnO layer with a large thermal capacity and a tardy response is responsible for low-frequency sensing tasks. A multi-frequency band pyroelectric sensor is successfully designed, analyzed and fabricated in the present study. The range of the multi-frequency sensing can be estimated by means of the proposed design and analysis to match the thicknesses of the sputtered and the aerosol ZnO layers. The fabricated multi-frequency band pyroelectric sensor with a 1 μm thick sputtered ZnO layer and a 20 μm thick aerosol ZnO layer can sense a frequency band from 4000 to 40,000 Hz without tardy response and low voltage responsivity.

Keywords: pyroelectricity; sensor; multi-frequency; zinc oxide; thin film
1. Introduction

Pyroelectricity, which is electric current generation from time-dependent temperature fluctuations, is useful in many applications, such as pollution monitoring, hot image detectors, intruder alarms, gas analysis and temperature sensors. A pyroelectric material exhibits a spontaneous polarization in the absence of an electric field. Thin-film pyroelectric sensors have many advantages, such as facile integration with on-chip circuitry, uncooled detection, room-temperature operation, speed, lower system costs, portability and a wide spectral response with high sensitivity [1–3]. Pyroelectric sensors have a pyroelectric layer sandwiched between top and bottom electrodes, which are built on thermally isolated structures or substrates to reduce heat loss. The principle of thin-film pyroelectric sensors is based on the pyroelectric effect, namely converting the heat transfer rates to the corresponding electrical signal. The pyroelectric effect is the property of selected dielectric materials with polar point symmetry, which show a spontaneous electrical polarization as a function of temperature. A change in the temperature of the material with respect to time (thermal fluctuations) causes a correspondent variation in the induced charge, thereby producing a pyroelectric current. The dynamic pyroelectric response current ($i_p$) of pyroelectric sensors can be described by the expression [1]:

$$i_p = \eta \times P \times A \times \frac{dT}{dt}$$

(1)

where $\eta$ is the absorption coefficient of radiation, $P$ is the pyroelectric coefficient of the pyroelectric film, $A$ is the electrode area and $\frac{dT}{dt}$ is the temperature variation rate of the pyroelectric film. When the materials and dimensions of the pyroelectric layers are defined, the absorption coefficient, the pyroelectric coefficient and the electrode area cannot be altered. However, the temperature variation rate has a huge maneuverability according to the design of the patterns, trenches, cavities and structures in the pyroelectric materials.

ZnO is a wide-band gap semiconductor of the II-VI semiconductor group. This semiconductor has several favorable properties, including good transparency, high electron mobility, wide band gap and strong room-temperature luminescence. These properties are used in emerging applications for transparent electrodes in liquid crystal displays, in energy-saving or heat-protecting windows and in electronics as thin-film transistors and light-emitting diodes. ZnO has a relatively large direct band gap of ~3.3 eV at room temperature. The advantages associated with a large band gap include higher breakdown voltages, the ability to sustain large electric fields, lower electronic noise, and high-temperature and high-power operation. ZnO films have been synthesized by numerous methods, such as metal organic chemical vapor deposition, molecular beam epitaxy, magnetron sputtering, pulsed laser deposition, atomic layer deposition, spray pyrolysis, filtered cathodic vacuum arc technique, sol-gel process and aerosol deposition. The quality of ZnO films obtained by the above methods depends on the specific growth methods and conditions. Thus, the preferential orientation of ZnO films depends on the growth conditions. The pyroelectricity of ZnO is attributable to non-centrosymmetrical crystals, and so it has a specific polar axis along the direction of spontaneous polarization [1–4]. The most densely packed and thermodynamically favorable growth orientation in a ZnO wurtzite structure is one in which the c-axis is perpendicular to the substrate. ZnO films with the c-axis normal to the substrate are preferred in many applications, such as ZnO pyroelectric devices [3,4] and film bulk acoustic resonators [5]. When ZnO is subjected to temperature variations, its internal polarization will produce an electric field. Therefore,
increasing the responsivity of a ZnO pyroelectric sensor depends on increasing the temperature variation rate of the ZnO layer, adopting a ZnO film with a strongly preferred orientation towards the c-axis, and using a high-performance thermal-isolation structure. The pyroelectric effect has been applied to environmental energy-harvesting systems. Pyroelectric energy conversion also offers a novel and direct way to convert time-dependent temperature fluctuations into electricity for micropower generators and low-energy-consumption systems [6,7].

The consideration of both the thermal and the electrical circuits is required in the analysis of pyroelectric devices. A radiation with power $W(t)$ sinusoidally modulated at a frequency $\omega$ is incident on the top surface of the pyroelectric element (area $A$ and thickness $d$), which has emissivity $\eta$. The pyroelectric element has a thermal capacity $H$ and a thermal conductance to the surroundings $G_T$; the thermal time constant can then be defined as $\tau_T = H/G_T$. The thermal capacity can be defined as $H = c'dA$, where $c'$ is the volume-specific heat. The electrical signal is further amplified by a high-input-impedance field effect transistor, which is used as an amplifier. $R_G$ is the gate resistor, and then the electrical time constant can be defined as $\tau_E = R_G(C_E + C_A)$, where $C_E$ is the capacitance of the pyroelectric element; $C_A$ is the capacitance of the amplifier; and $\tau_T$ and $\tau_E$ are the fundamental factors which determine the frequency response of the pyroelectric sensors. The voltage responsivity ($R_v$) can be calculated by the following equation [1]:

$$R_v = \frac{R_G\eta P A \omega}{G_T \sqrt{(1 + \omega^2 \tau_T^2) \sqrt{(1 + \omega^2 \tau_E^2)}}} \tag{2}$$

At a low frequency ($\omega << \tau_T^{-1}$), $R_v$ is proportional to the frequency, and is shown as the following equation:

$$R_v = \frac{R_G\eta P A \omega}{G_T} \tag{3}$$

Equation (3) can easily maximize $R_v$ by minimizing $G_T$ (i.e., by adding a thermal insulation layer between the pyroelectric film and the substrates, adopting a suspended structure fabricated by a bulk micromachining technique using anisotropic silicon etching or by using a substrate with a low thermal conductivity). At a high frequency ($\omega >> \tau_T^{-1}; \omega >> \tau_E^{-1}$), $R_v$ is inversely proportional to the frequency, and is shown as the following equation:

$$R_v = \frac{\eta P}{c'd(C_E + C_A)\omega} \tag{4}$$

Equation (4) can easily maximize $R_v$ by minimizing the thermal capacity of the pyroelectric element $H$ (i.e., decreasing the thickness of the pyroelectric element). Moreover, reducing the pyroelectric element’s thickness can retard the decline of the voltage responsivity at a high frequency. The frequency at $\tau_T^{-1}$ is a watershed to distinguish the ranges of low and high frequencies, and the pyroelectric element’s thickness determines the value of the thermal time constant ($\tau_T$) under the decided pyroelectric materials and electrode areas. Therefore, a thicker pyroelectric element increases the thermal time constant, which is suitable as the sensor for a low-frequency range. Unlike the thicker element, a thinner pyroelectric element reduces the thermal time constant, which is suitable as the sensor for a high-frequency range. It is difficult to apply a pyroelectric sensor with a single pyroelectric layer to multi-frequency sensing tasks when the materials and dimensions of the pyroelectric layers are already fixed. In other words, the
materials and dimensions of the pyroelectric layers directly determine the thermal time constant. Therefore, in this study, a structure with dual pyroelectric ZnO films was designed for a multi-frequency band pyroelectric sensor. The structure mainly comprised two ZnO pyroelectric layers: a thinner ZnO pyroelectric layer deposited by sputtering and a thicker ZnO pyroelectric layer deposited by aerosol deposition (AD). The thinner ZnO film was deposited by RF sputtering. Sputter deposition is a physical vapor deposition (PVD) method of depositing thin films by sputtering. Sputter deposited films have a composition close to that of the source material. Furthermore, sputtered films typically have a better adhesion on the substrate than evaporated films. The properties of ZnO films are affected by sputtering conditions such as the composition of mixed process gases, working pressure, substrate temperature, radio frequency (RF) power, the gap between the target and substrate, and the post-annealing temperature. Moreover, the thicker ZnO film was grown by the AD. In this technique, ceramic films are prepared by ejecting an aerosol consisting of a mixture of ultra-fine ceramic particles and gas from the nozzle to the substrate without vaporization of materials. Compared with the sputtering, screen-printing or sol–gel methods, the AD provides many advantages for producing films in the range of 1–100 μm thickness with a high deposition rate, low deposition temperature and low cost. The AD method can achieve fine patterning and fabricate a dense structure by the reduction of crystallite size by fracture or plastic deformation at room temperature [8–10]. Although ZnO film grown at a low temperature is available from the AD process, an annealing treatment is one of the most common methods for reducing the defects and improving the quality of as-grown thin film. Therefore, furnace annealing was adopted to improve the ZnO film quality in the present study. The annealing parameters for furnace annealing treatment included temperature, duration, atmosphere and pressure. In particular, annealing temperature and duration are the most effective factors for improving the film quality [11,12].

When a pyroelectric sensor is subjected to temperature variations, its internal polarization produces an electrical field that induces a voltage responsivity between the top and bottom electrodes. The responsivity is proportional to the temperature variation rates in the pyroelectric layers: no temperature variation in the pyroelectric layers results in no internal polarization change and, thus, no voltage response. Temperature variation rates in pyroelectric layers markedly affect the responsivity of pyroelectric sensors. In the present study, transient temperature fields in multi-frequency band pyroelectric sensors were simulated to probe into temperature variation rates at various pyroelectric layers and estimate the response of the sensors. Then, a multi-frequency band pyroelectric sensor was accomplished by designing a structure with dual pyroelectric layers, analyzing the temperature variation fields at various pyroelectric layers, fabricating the multi-frequency band pyroelectric sensor by MEMS process, measuring the responsivity of the sensors by a measurement system and treating and integrating the signals of the sensors by a LabVIEW system.

2. Materials and Methods

2.1. Design for the Multi-Frequency Band Pyroelectric Sensor

A general pyroelectric sensor, as shown in Figure 1, is in a sandwich structure, composed of a single pyroelectric layer sandwiched between top and bottom electrodes, and built on a substrate with an insulating layer for reducing heat and electric loss. As the thickness of the pyroelectric layers determines
the thermal time constant ($\tau_T$) under pyroelectric materials and electrode areas already fixed, the thermal time constant determines the optimal working frequency of the pyroelectric sensors. In the present study, a structure with dual pyroelectric film layers was adopted in the design of a pyroelectric sensor with an ability of multi-frequency band sensing; namely, a multi-frequency band pyroelectric sensor.

**Figure 1.** Schematic diagram of a general pyroelectric sensor with a single pyroelectric layer (unit: $\mu$m).

The present multi-frequency band pyroelectric sensor, consisting of a thinner and a thicker ZnO pyroelectric layer and top and bottom electrodes, was built on a silicon substrate with a thermal-insulation (silicon nitride) layer to reduce heat and electric loss. The thinner ZnO pyroelectric layer was named the sputtered ZnO layer, and the thicker ZnO pyroelectric layer was named the aerosol ZnO layer. Figure 2 shows the schematic diagram of the multi-frequency band pyroelectric sensor. The sputtered ZnO layer acted as producer of the responsivity at higher frequency bands, while the aerosol ZnO layer detected the signals of the sensors at lower frequency bands. Sputtering was used to deposit the thinner ZnO films with high quality. Furthermore, the thicker ZnO layer was to detect the signals of the sensors at lower frequency bands, and deposited by the AD. The starting powder was commercially available ZnO (Top Nano Technology Co. Ltd., New Taipei, Taiwan). The properties of the starting ZnO powder are shown in Table 1. Table 2 shows the deposition parameters used for the AD method. The powder was subjected to heat treatment at 150 °C for 1 h using an oven before ZnO films deposited. This could reduce moisture contents in the ZnO powder, and decrease agglomerated particles in ZnO films. Subsequently, furnace annealing was used to the ZnO films to improve the film quality. A furnace annealing system (SJ High Technology Company, Taipei, Taiwan), consisted of a single zone tube furnace with tube furnace model of T21-303, a single zone programmable temperature control console with controller model of SJ-C01, a cooling system and a vacuum system, was adopted for the ZnO film annealing in N$_2$ ambient. The parameters applied on furnace annealing of the ZnO films are shown in Table 3. The film thickness was further measured by a surface analyzer (ET-4000AK, Kosaka, Tokyo, Japan).
Figure 2. Schematic diagram of the multi-frequency band pyroelectric sensor (unit: μm).

Table 1. Material properties and geometry of commercially ZnO powder.

| Item                     | Data                                      |
|--------------------------|-------------------------------------------|
| Appearance               | White powder                              |
| Density                  | 5.61 g/cm³                                |
| Specific surface area    | 17 m²/g                                   |
| Particle form            | Sheet                                     |
| Particle size            | 300 nm (Diameter) × 20 nm (Thickness)     |

Table 2. Process parameters for ZnO films deposited by the AD method.

| Item                                      | Data                                      |
|-------------------------------------------|-------------------------------------------|
| Starting powder                          | ZnO                                       |
| Substrate                                | Silicon                                   |
| Pressure difference between deposition and aerosol chambers | 140 (Torr)                             |
| Carrier gas                              | Nitrogen                                  |
| Consumption of carrier gas               | 3 (L/min)                                 |
| Orifice size of nozzle                   | 0.4 × 10 (mm × mm)                        |
| Substrate temperature                    | 25 °C                                     |
| Deposition area                          | 70 × 70 (mm × mm)                         |
| Distance between nozzle and substrate    | 5 (mm)                                    |
| Scanning rate                            | 10 (mm/s)                                 |
| Deposition rate                          | 8.2 (nm/s)                                |
Table 3. Process parameters for ZnO films treated by furnace annealing.

| Item                        | Data                      |
|-----------------------------|---------------------------|
| Holding temperature        | 800 (°C)                 |
| Duration                    | 15 (min)                  |
| Speed for raising temperature | 10.4 (°C/min)            |
| Ambience                    | Nitrogen                  |
| Cooling type                | Cooling in furnace        |

2.2. Simulation

Temperature variation rates in the pyroelectric layers markedly affect the responsivity of pyroelectric sensors. Moreover, the temperature variation field is difficult to extract from thin films by experimental measurement. Li et al. [13] used a finite element model, built using the commercial software ANSYS, to explore the temperature variation rate in pyroelectric elements with various thermal properties and geometries. Hsiao et al. [4] also used a commercial multiphysics software package, COMSOL MULTIPHYSICS®, to explore the temperature variation rate in ZnO pyroelectric devices in the design and optimization of a partially covered mesh-type electrode. In the present study, a two-dimensional finite element model was constructed using the commercial multiphysics software COMSOL MULTIPHYSICS® 4.2 to explore the temperature variation rate in the multi-frequency band ZnO pyroelectric sensors. The material parameters of the films and substrate are shown in Table 4. There was an isotropic assumption for the films and substrate properties in this model. The model was meshed using a regular mesh, as shown in Figure 3. The thickness of the sputtered ZnO layer (Tr2) was fixed as 0.3 μm, while the thicknesses of the aerosol ZnO layer (TA2) were varied from 1, 2, 3, 4, to 5 μm. The incident irradiation power applied on the top side of the ZnO pyroelectric devices was approximately $1.228 \times 10^{-12}$ W/μm² [13]. The thermal isolation condition was applied to the rear side of the silicon substrate, and the symmetric condition was applied to the two lateral sides as boundary conditions.

Figure 3. Finite element model for the multi-frequency band ZnO pyroelectric sensor.
Table 4. Material parameters used for the simulation.

| Material                        | Thermal Conductivity (Wm\(^{-1}\cdot K^{-1}\)) | Specific Heat (Jg\(^{-1}\cdot K^{-1}\)) | Density (g·cm\(^{-3}\)) | Thickness (μm) |
|---------------------------------|-----------------------------------------------|----------------------------------------|--------------------------|----------------|
| Silicon substrate               | 163                                           | 0.703                                  | 2.330                    | 5              |
| Silicon nitride                 | 20                                            | 0.700                                  | 3.100                    | 1              |
| Electrodes                      | 317                                           | 0.129                                  | 19.300                   | 0.1            |
| sputtered and aerosol ZnO layers| 6                                             | 0.125                                  | 5.676                    | T\(_{PZ}\) and T\(_{AZ}\) |

2.3. Fabrication Process

The multi-frequency band pyroelectric sensor used both sputtering and AD to extract the advantages of the thin and the thick ZnO pyroelectric film to further integrate the electrical outputs. The process flow of the multi-frequency band ZnO-film pyroelectric sensor was divided into several steps, as follows.

A silicon wafer had specifications of (100) p-type, double-side polished, resistivity of 1–10 Ω-cm, and was used as a substrate to support the multi-frequency band pyroelectric sensor, as shown in Figure 4a. Low-stress silicon nitride layers with a thickness of 1 μm were deposited on both sides of the substrate by LPCVD, which was used to reduce or block the heat loss through the silicon substrate, as shown in Figure 4b. The bottom electrode was deposited on the substrate by electron beam (E-beam) evaporation and patterned by the shadow mask method, as shown in Figure 4c. The shadow mask method could simplify process steps and shorten process time. The bottom and top electrode were both composed of gold and chromium. The chromium was an adhesion layer to promote the adhesion between gold and the substrate or ZnO films. The electrodes comprised a 100 nm thick gold layer and a 10 nm thick chromium adhesion layer. The next step was to deposit the aerosol ZnO film with a thickness of about 20 μm by the AD, as shown in Figure 4d, and then those were promoted by furnace annealing, as shown in Figure 4e. Subsequently, a 1 μm thick ZnO layer (the sputtered ZnO layer) was deposited on the bottom electrode by RF magnetron sputter, as shown in Figure 4f. A ZnO target with 99.99% purity was adopted. Prior to the film deposited, the ZnO target was pre-sputtered for 15 min to remove any surface impurity. The chamber was pumped with base pressure up to \(8 \times 10^{-7}\) Torr before sputtering. The chamber was then filled with the mixture of argon and oxygen with the gas-mixing ratio of 5:3. The RF power was kept at 120 W. The chamber pressure was 2 mTorr during the film deposited. The substrate was heated up to 200 °C while deposition, which could help to make better ZnO film quality. The top electrode was deposited on the sputtered and the aerosol ZnO film by E-beam and patterned by the shadow mask method, as shown in Figure 4g. Finally, the wet etchant of CH\(_3\)COOH:H\(_3\)PO\(_4\):H\(_2\)O = 1:1:10 was used to pattern ZnO layers to open the bonding pads of the bottom electrode, as shown in Figure 4h. The fabricated multi-frequency band pyroelectric sensor is shown in Figure 5.

2.4. Signal Treatment and Measurement

The multi-frequency band pyroelectric sensor used the advantages of the thinner and the thicker ZnO pyroelectric film to integrate the electrical outputs yielded from those films into an all-round signal. The voltage responsivity of \(V_P\) was generated from the sputtered ZnO film for shouldering a high-frequency response, and the voltage responsivity of \(V_A\) was generated from the aerosol ZnO film for taking a low-frequency response. The integrated and treated electrical signal was \(V_T\).
Figure 4. Fabrication flow for the multi-frequency band pyroelectric sensor.

Figure 5. Fabricated multi-frequency band pyroelectric sensor.
A diagram of the flow of the signal treatment is shown in Figure 6. A voltage responsivity measurement system, as Figure 7, was adopted to appraise the performance of the multi-frequency band ZnO thin-film pyroelectric sensor.

**Figure 6.** Schematic diagram of signal treatment flow.

**Figure 7.** Schematic diagram of voltage responsivity measurement system.
The radiation source was a calibrated infrared (IR) laser of 900 nm wavelength and 7 mW maximum power. The laser beam was molded as a square wave with a modulated frequency (ω) by a programmable function generator. A prism was used to split the modulated beam into two beams, which had the same power: one was reflected on a photodiode as the reference signal, and the other was expanded and equalized via a beam expander and a beam equalizer such that the beam spot could uniformly cover the entire region of the top electrode of the sensors. Both the responsivities of $V_P$ and $V_A$ were filtered, amplified, modulated and combined as an integrated voltage responsivity by NI LabVIEW software. Moreover, both the integrated voltage responsivity of the sensors and the reference signal of the photodiode were acquired, recorded and displayed using NI LabVIEW system consisted of a case of NI PXIe-1082, a controller of NI PXIe-8135, a data acquisition card of NI PXIe-6366 and NI LabVIEW 2012 software.

3. Results and Discussion

The responsivities of pyroelectric devices are proportional to the temperature variation rate in the pyroelectric layers. Therefore, the transient temperature fields in multilayer pyroelectric devices with various geometric structures were simulated. Moreover, the response time was estimated. The points, as shown in Figure 8, were used to interpret the temperature variation rates in the ZnO pyroelectric devices with the single pyroelectric layer and the dual pyroelectric layers. Points SZ1, SZ3 and SZ5 were, respectively, located on the top, the middle and the bottom of the single ZnO layer. The thickness of the single ZnO layer was named as $T_{SZ}$. Points PZ1, PZ3 and PZ5 were, respectively, located on the top, the middle and the bottom of the sputtered ZnO layer. The thickness of the sputtered ZnO layer was named as $T_{PZ}$. Points AZ1, AZ3 and AZ5 were, respectively, located on the top, the middle and the bottom of the aerosol ZnO layer. The thickness of the aerosol ZnO layer was named as $T_{AZ}$.

**Figure 8.** The points defined for explaining the transient temperature fields in the ZnO pyroelectric devices for (a) the single layer type and (b) the dual layer type.
For the pyroelectric sensor with the single ZnO layer, Figure 9 shows the relationship between the temperature variation rate and time in the single ZnO layer along the thickness direction from points SZ₁ to SZ₅, when a single ZnO layer with a thickness of 0.3 μm was used to fabricate the ZnO pyroelectric sensor. The temperature variation rate increased when the point approached to the top side of the single ZnO layer, because incident radiation was applied on the top electrode. Moreover, the peak of the temperature variation rate moved leftward, and then the peak time of the maximum temperature variation rate also reduced, thus decreasing the response time. Figure 10 shows the peak temperature variation rate and the peak time of the maximum temperature variation rate at points SZ₁ to SZ₅, when the thickness of the single ZnO layer varied from 0.3 to 3 μm. A thinner ZnO layer was conducive to enhancing the temperature variation rate and improving the response speed in the pyroelectric sensor. However, the temperature variation rate at various locations in the ZnO film had different values and distributions at different time periods. Using the peak temperature variation rate and the peak time at point SZ₁ could not discriminate the properties of the pyroelectric sensor with various thicknesses of the ZnO films. Therefore, a conservative response was to adopt the properties at point SZ₅ for inspecting the sensors, because the peak temperature variation rate and the peak time at point SZ₅ were disappointing, as compared to points SZ₁, SZ₂, SZ₃, and SZ₄ under the specific thickness of ZnO film being used. Therefore, improvement in the temperature variation rate at the point SZ₅ possessed the lowest temperature variation rate certainly enhanced the responsivity of the pyroelectric devices.

Figure 9. Relationship between the temperature variation rate and time at points SZ₁ to SZ₅ under a 0.3 μm thick single ZnO layer.

For the multi-frequency band pyroelectric sensor, Figure 11 shows the relationship between the temperature variation rate and time at points PZ₅ and AZ₅ in the dual ZnO layers, when sputtered ZnO film with a constant thickness of 0.3 μm and aerosol ZnO film with various thicknesses from 1 to 5 μm were used to fabricate the ZnO pyroelectric sensors.
Figure 10. The peak temperature variation rate and the peak time of the maximum temperature variation rate at points SZ1 to SZ5 under a single ZnO layer 0.3 to 3 μm thick.

Figure 11. Relationship between the temperature variation rate and time at points PZ5 and AZ5 in the multi-frequency band pyroelectric sensor under sputtered ZnO film with a constant thickness of 0.3 μm and aerosol ZnO film with various thicknesses from 1 to 5 μm.

The sputtered ZnO film possessed a higher temperature variation rate and a shorter response time, while the aerosol ZnO film possessed a lower temperature variation rate and a longer response time. Increasing the thickness of the ZnO layer reduced the temperature variation rate and increased the peak time of the maximum temperature variation rate. Therefore, a ZnO film with a constant thickness in a pyroelectric device could not handle various multi-frequency sensing tasks. In fact, the pyroelectric devices use the voltage responsivity for presenting the performance of the sensors. Hence, the voltage responsivities were calculated for estimating the electrical outputs of the sensors. The voltage responsivity ($R_v$) is defined as the signal generated when the pyroelectric sensors are exposed to a
modulated radiation. Moreover, when the pyroelectric sensors are connected to a high impedance amplifier, the observed signal is equal to the voltage produced by the charge. \( R_v \) can be expressed as [1]:

\[
R_v = \frac{i_p}{Y \times W_0}
\]  

(5)

where \( i_p \) is the dynamic pyroelectric response current, \( W_0 \) is the magnitude of the incident radiation and \( Y \) is the electrical admittance, as:

\[
Y = R_G^{-1} + i\omega C_T
\]  

(6)

where \( R_G \) is the gate resistor, \( \omega \) is the modulated frequency of the incident radiation, \( C_T \) is the sum of \( C_E \) and \( C_A \), \( C_E \) is equal to \( \varepsilon_0 \varepsilon_r A/d \), \( \varepsilon_0 \) is the vacuum permittivity (8.85 \( \times \) 10\(^{-12} \) F/m) and \( \varepsilon_r \) is the relative permittivity or dielectric constant of the materials. \( R_v \) was estimated using the simulated data of the temperature variation rates and Equations (1), (5) and (6). The relative conditions for computing the voltage responsivity are shown in Table 5. Figure 12 shows the relationship between the voltage responsivities and time in the multi-frequency band pyroelectric sensor when sputtered ZnO film with a constant thickness of 0.3 \( \mu \)m and aerosol ZnO film with various thicknesses from 1 to 5 \( \mu \)m were used to fabricate the ZnO pyroelectric sensors. Furthermore, the temperature variation rates at points PZ5 and AZ5 were used to estimate the voltage responsivities generated by the sputtered and the aerosol ZnO film, respectively. The curve of the voltage responsivity generated by the sputtered ZnO film (VP) with a rapid response could be attributed to a thinner ZnO layer with a large temperature variation rate and a small thermal capacity, while that generated by the aerosol ZnO film (VA) with a tardy response could be attributed to a thicker ZnO layer with a large thermal capacity and a small temperature variation rate. Therefore, VP was responsible for shouldering a high-frequency response, and VA was responsible for shouldering a low-frequency response. VA was larger than VP at the low-frequency band, which could be attributed to the thicker ZnO layer with a small \( C_E \). The peak time of the maximum temperature variation rate increased when the thickness of the aerosol ZnO film increased due to the small temperature variation rate and large thermal capacity. The curves of VP and VA had an apparent intersection, which could be used for differentiating between the high- and the low-frequency bands. VT was the integrated voltage responsivity by combining VP with VA. However, integrating VP, generated by a 0.3 \( \mu \)m thick sputtered ZnO film, with VA, generated by a thicker aerosol ZnO film, was disadvantageous because the voltage responsivity did not produce a compensatory effect between the peak time of VP and VA, as shown in Figure 13a. Furthermore, integrating VP, generated by a 0.3 \( \mu \)m thick sputtered ZnO film, with VA, generated by a thinner aerosol ZnO film, was also unsuitable because the reinforced sensing frequency band was narrow, as shown in Figure 13b. An optimal VT needed a suitable combination, matching the thicknesses of the sputtered and the aerosol ZnO films. Obviously, as shown in Figure 13c, the voltage responsivity of VT could shoulder the sensing tasks of the pyroelectric devices in a multi-frequency band.

| \( R_G \) (M\( \Omega \)) | \( C_A \) (pF) | \( P \) (10\(^{-4} \) C/m\(^2\)·K) | \( A \) (mm\(^2\)) | \( \varepsilon_r \) (Unit-less) | \( W_0 \) (W/\( \mu \)m\(^2\)) |
|---|---|---|---|---|---|
| 22 | 6 | 0.1 | 9 | 11 | 1.228 \( \times \) 10\(^{-12} \) |
**Figure 12.** Relationship between the voltage responsivities and time in the multi-frequency band pyroelectric sensor under the sputtered ZnO film with a constant thickness of 0.3 μm and the aerosol ZnO film with various thicknesses from 1 to 5 μm.

**Figure 13.** The integrated voltage responsivities ($V_T$) generated by combining $V_P$, generated by a 0.3 μm thick sputtered ZnO film, with $V_A$, generated by various thicknesses of aerosol ZnO film: (a) $T_AZ = 5$ μm; (b) $T_AZ = 1$ μm; (c) $T_AZ = 2$ μm.
The transient temperature fields with various chopping frequencies were simulated for further calculating voltage responsivities when the multi-frequency band pyroelectric devices were exposed to irradiation power with various chopping periods. A square waveform was produced using a programmable function generator to modulate the irradiation power with a peak value of $1.228 \times 10^{-12}$ W/$\mu$m$^2$ and chopping frequencies of about 10,000, 33,000 and 250,000 Hz. Figure 14 shows the voltage responsivities of the multi-frequency band pyroelectric device with $TP_z = 0.3$ μm and $TA_z = 2$ μm when the incident irradiation power was modulated with various chopping frequencies of about 10,000, 33,000 and 250,000 Hz. The shape and amplitude of the integrated voltage responsivity of $V_T$ approached to that of $V_P$ generated by the sputtered ZnO film when the chopping frequency was near a high frequency about 250,000 Hz. Moreover, the shape and amplitude of the integrated voltage responsivity of $V_T$ approached to that of $V_A$ generated by the aerosol ZnO film when the chopping frequency was near a low frequency about 10,000 Hz. $V_T$ had a compensatory effect between $V_P$ and $V_A$. $V_T$ redeemed the drawbacks of $V_A$ with a tardy response at a high-frequency band and $V_P$ with a low voltage responsivity at a low-frequency band. This implied that the integrated voltage responsivity of $V_T$ generated by the
multi-frequency band pyroelectric devices could present the performances of various pyroelectric devices at various sensing frequency bands.

**Figure 15.** Voltage responsivities of the fabricated multi-frequency band pyroelectric device with $T_{PZ} = 1 \, \mu m$ and $T_{AZ} = 20 \, \mu m$ under the incident irradiation power modulated with various chopping frequencies of about (a) 4000 Hz; (b) 20,000 Hz and (c) 40,000 Hz.

An experimental setup was used to verify the above analytical results. The IR laser beam was chopped and molded as a square wave with various modulated frequencies (4000, 20,000 and 40,000 Hz) to obtain temperature variation rates in the multi-frequency band pyroelectric device by means of a programmable function generator. Figure 15 shows the voltage responsivities of the fabricated multi-frequency band pyroelectric device with $T_{PZ} = 1 \, \mu m$ and $T_{AZ} = 20 \, \mu m$ when the incident irradiation power was modulated with various chopping frequencies of about 4000, 20,000 and 40,000 KHz. The fabricated multi-frequency band pyroelectric device at the modulated frequency of 4000 presented a low-frequency property. $V_T$ presented a large temperature variation rate, but it did not possess a tardy response. Furthermore, the fabricated multi-frequency band pyroelectric device at the modulated frequency of 40,000 presented a high-frequency property. Although $V_T$ could not improve the temperature variation rate, the sensing frequency band increased. This implied that the voltage responsivity could hold on a
peak value with a larger sensing frequency band. Hence, the multi-frequency band pyroelectric device could enhance pyroelectric sensors detecting subjects with various speeds or frequencies.

4. Conclusions

A pyroelectric sensor with a single pyroelectric layer has only a single sensing frequency band. Multi-frequency sensing tasks could improve applications of pyroelectric sensors for detecting subjects with various speeds or frequencies. A multi-frequency band pyroelectric sensor was successfully designed, analyzed and fabricated in the present study. The combination of a thinner sputtered ZnO layer and a thicker aerosol ZnO layer proved useful in the present design. The ranges of the multi-frequency sensing tasks could be predicted by the proposed design and analysis to match the thicknesses of the sputtered and the aerosol ZnO layers for producing a compensatory effect and extending the reinforced sensing frequency band. The fabricated multi-frequency band pyroelectric sensor with a 1 μm thick sputtered ZnO layer and a 20 μm thick aerosol ZnO layer could cover the sensing frequency band from 4000 to 40,000 Hz without tardy response and low voltage responsivity.

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Author Contributions

Sheng-Yi Liu was involved in the data collection and experimental work under supervision of Chun-Ching Hsiao who carefully edited the paper, provided technical support and guidance, and directly contributed the results of this paper.

Conflicts of Interest

The authors declare no conflict of interest.

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