

Pulsed X-Ray Detector Based on an Unintentionally-Doped High Resistivity ϵ -Ga₂O₃ Film

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Abstract—Gallium Oxide (Ga₂O₃), an emerging ultra-wide band gap semiconductor, has drawn great attention for application in radiation detection. In this letter, ultrafast X-ray detectors have been fabricated using a high resistivity unintentionally-doped (UID) ϵ -Ga₂O₃ film grown on sapphire by metal-organic chemical vapor deposition (MOCVD). The detector featuring a lateral metal–semiconductor–metal (MSM) structure exhibited a low dark current < 2 nA at 100 V and its sensitivity was as high as 28.6 nC/Gy or $\sim 1.0 \times 10^6$ nC/(Gy · cm³) at 40 V and an X-ray dose rate of 0.383 Gy/s. A stable and repeatable transient response was observed for the detectors under switching X-ray illumination. Furthermore, the detector achieved a pulsed X-ray detection with 50 ns in full width and its time resolution was revealed to be ~ 7.1 ns. These results imply the great potential of the MOCVD-grown high-resistivity UID ϵ -Ga₂O₃ film for ultrafast X-ray detection.

Index Terms—Ga₂O₃ film, X-ray detection, semiconductor radiation detector, pulsed measurement.

I. INTRODUCTION

GALLIUM Oxide (Ga₂O₃), which possesses a wide direct band gap of ~ 4.9 eV, a high critical electric field of 8 MV/cm and an excellent tolerance to irradiation, is a promising material for X-ray and high-energy particle detection applications [1], [2], [3]. Previously, Lu et al. demonstrated the first β -Ga₂O₃ Schottky X-ray detectors based on

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a single crystalline bulk substrate [4]. Later, various types of Ga₂O₃ X-ray detectors have been developed based on either single crystals or polycrystalline/amorphous thin films [5], [6], [7], [8], [9], [10], [11], [12]. Compared to polycrystalline/amorphous material, the single crystals material with low defects will ensure both fast response and high sensitivity of the detector [13], [14].

Usually, the unintentionally-doped (UID) Ga₂O₃ single crystals show an n-type conductivity, which are not suitable for fabricating metal–semiconductor–metal (MSM) detectors due to a high dark current issue. Very recently, pulsed X-ray detectors with a nanosecond time resolution have been realized using a Fe doped high resistivity β -Ga₂O₃ single crystal substrate [9], in which Fe was adopted as deep acceptors to compensate the background n-type conductivity in the material. However, the compensation doping would inevitably deteriorate the crystalline quality and produce defects within the Ga₂O₃ lattice, bringing some adverse effects to the performance of the detector, for example decreasing its sensitivity. Moreover, the β -Ga₂O₃ single crystal substrates at current stage are of small size and still too expensive from a commercial point of view.

On the other hand, high quality single crystalline ϵ -Ga₂O₃ films have been successfully grown on low cost sapphire substrates by metal-organic chemical vapor deposition (MOCVD) [15], [16], [17], [18]. Featuring an intrinsic high resistivity, the MSM X-ray detectors based on a hetero-epitaxial UID ϵ -Ga₂O₃ film demonstrated a remarkably high sensitivity [12], which suggested a great potential in reducing the production cost while maintaining a high performance for future X-ray detection applications.

In this study, based on an MOCVD-grown high resistivity UID ϵ -Ga₂O₃ film, a MSM detector was fabricated and characterized for pulsed X-ray detection. The detector achieved a 50 ns in full width pulsed X-ray measurement and its time resolution was revealed to be ~ 7.1 ns.

II. EXPERIMENTS

The MSM detector was fabricated using UID ϵ -Ga₂O₃ thin film grown on sapphire substrate by MOCVD. More detailed information about the epitaxy process could be found in our previous reports [15], [19]. The Al interdigital electrodes were deposited on top of the ϵ -Ga₂O₃ film by E-beam evaporation, and the distance between the neighboring electrodes was 0.5 mm.

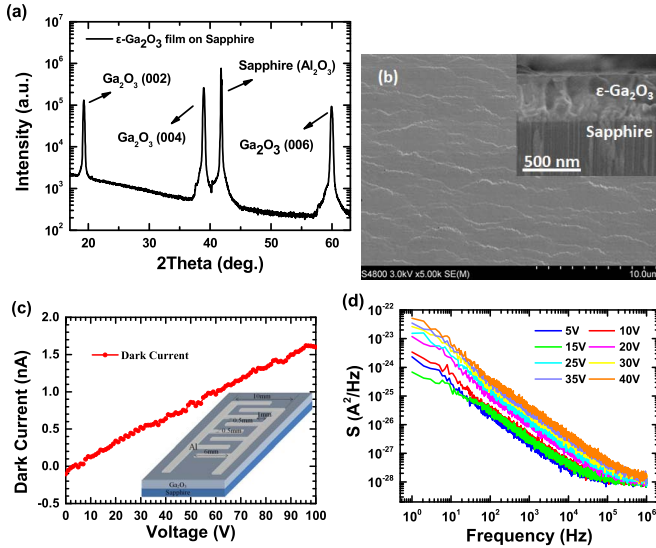


Fig. 1. (a) The XRD pattern and (b) The plan-view SEM image of the ϵ -Ga₂O₃ film grown on a sapphire substrate. (c) The typical dark I-V characteristics of the fabricated ϵ -Ga₂O₃ detector. Insert shows the structure of detector. (d) The LFN equivalent spectra at different bias voltages.

A test unit (Agilent B2902A) was used for measuring the I-V characteristics. The low frequency noise (LFN) analysis was performed to figure out the influence of defects on the carrier transport process in the detectors.

The X-ray source used in this study was an X-ray tube with a tungsten anode, which operated at the tube currents between 1 μ A and 400 μ A and a tube voltage of 30 kV with the corresponding X-ray dose rates ranging from 3.83 mGy/s to 1.532 Gy/s and proportional to the tube current. The maximum X-ray photon energy was 30 keV and the X-ray beam was perpendicularly illuminated to the ϵ -Ga₂O₃ detector in all the measurements. An XRS-4 pulsed X-ray source with 50 ns in full width was used for the pulsed X-ray detection test and the response signal was recorded by an oscilloscope from a 50- Ω resistor, which was connected in series with the detector.

III. RESULTS AND DISCUSSION

Fig.1 (a) shows the X-ray diffraction (XRD) pattern of the as-grown ϵ -Ga₂O₃ sample. Three peaks at 19.284°, 38.964° and 59.924° were attributed to the (002), (004) and (004) planes of the ϵ -Ga₂O₃ film, while the peak at 41.784° corresponded the sapphire (Al₂O₃) substrate. The full width at halfmaximum (FWHM) of the rocking curve of the (004) diffraction peak of ϵ -Ga₂O₃ was 0.1728° (622''), which confirmed that a high quality phase-pure single-crystalline ϵ -Ga₂O₃ film was achieved in our study [15], [19], [20], [21]. The plan-view scanning electron microscopy (SEM) image in Fig.1 (b) shows a high crystallinity of the heteroepitaxial ϵ -Ga₂O₃ film without obvious grain boundary. The inset shows the cross section of the fabricated film, and the thickness of the film was 380 nm. Fig.1 (c) shows the device structure of the fabricated ϵ -Ga₂O₃ MSM detector and its typical I-V characteristic under dark condition. The dark current of the detector remained below 2 nA at a bias of 100 V, indicating that the epitaxial UID ϵ -Ga₂O₃ film in our study had an

intrinsic high resistivity. Unlike the reported Ga₂O₃ detectors based on the Fe/Mg compensation doped material [6], [7], [8], [9], the intrinsic high resistivity property of the UID ϵ -Ga₂O₃ film would keep the carrier transport process of the detector far from the influence of defects. Fig.1 (d) shows the LFN spectra of the detector at various bias voltages. The measured curves can be represented as the function of $1/f$ and a weak generation-recombination (GR) noise appended at 10 Hz. When the bias voltage was increased from 5 V to 40 V, the measured noise of our detector kept in the same tendency and merely grew from 1×10^{-24} (A²/Hz) to 1×10^{-23} (A²/Hz), which were close to the system noise below 10^{-26} (A²/Hz). Thereby, the noise level of detector was quite low and weakly dependent on the bias voltage, and the device was not seriously affected by the defects [22], [23].

Fig.2 (a) shows the response net currents ($I_{X-ray} - I_{dark}$) of the detector to the X-ray illumination with different X-ray tube current as a function of bias voltage. Obviously, the net current of the detector increased with both the X-ray dose rates and the bias voltage. The unsaturated net current against bias voltage indicated an incomplete collection of the radiation-generated carriers in our detector. According to the Hecht equation [24], the efficiency of carrier collection could be increased by enlarging the electric field between the electrodes (E) for an MSM detector and it would eventually get saturated when E went strong enough. In this case, the response of the detector could be further enhanced.

The sensitivity of the detector is defined as the ratio between the net current and the incident X-ray dose rate reaching to the detector (D):

$$\text{Sensitivity} = \frac{\text{Net Current}}{D} = \frac{I_{X-ray} - I_{dark}}{D} \quad (1)$$

When the X-ray tube current was set at 100 μ A, the net current of the detector biased at 40 V was 11 nA (23 nA @100 V) and the X-ray dose rate reaching to the detector was calibrated to be 0.383 Gy/s. Therefore, the sensitivity was calculated to be 28.6 nC/Gy (60 nC/Gy @100 V). In order to make a comparison with our previous studies under the same X-ray illumination ($D = 0.383$ Gy/s) and bias condition ($E = 800$ V/cm @40 V), the normalized sensitivity to device active volume was calculated according to:

$$\text{Normalized Sensitivity} = \frac{\text{Sensitivity}}{\text{Device active Volume}} \quad (2)$$

The normalized sensitivity of the ϵ -Ga₂O₃ detector in this study was as high as $\sim 1.0 \times 10^6$ nC/(Gy \cdot cm³), which was much higher than that of a high-resistivity polycrystalline ZnO detector $\sim 1.1 \times 10^5$ nC/(Gy \cdot cm³) [25] and a Fe-doped high-resistivity single crystalline β -Ga₂O₃ detector $\sim 2.2 \times 10^1$ nC/(Gy \cdot cm³) [9]. The higher normalized sensitivity in this study could be mainly attributed to the low defects density in the UID ϵ -Ga₂O₃ epitaxial film.

The net current and sensitivity of the detector biased at -40 V and -100 V are plotted as a function of the X-ray dose rate in Fig. 2(b). The device showed a nonlinear output with a reduced sensitivity at higher X-ray dose rate, which indicated that the carrier collection efficiency degraded when the X-ray dose rate increased. According to the band-to-band

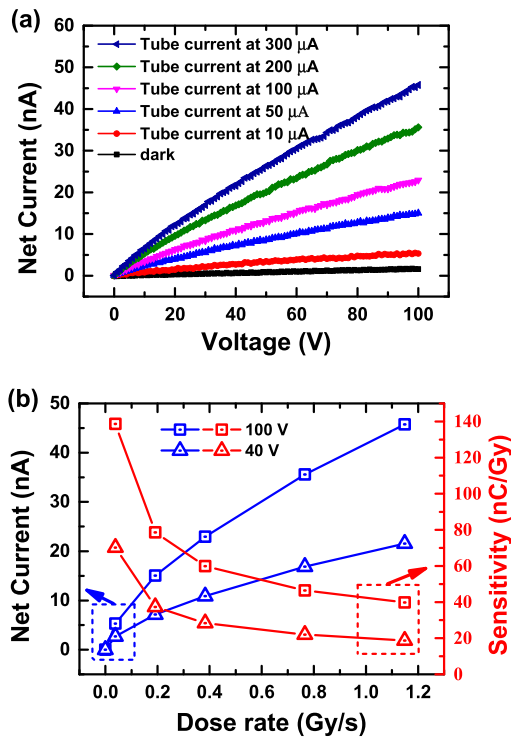


Fig. 2. (a) The net current of the detector at various X-ray tube currents as a function of the bias voltage. (b) The net current and sensitivity of the detector biased at -40 V and -100 V as a function of X-ray dose rate.

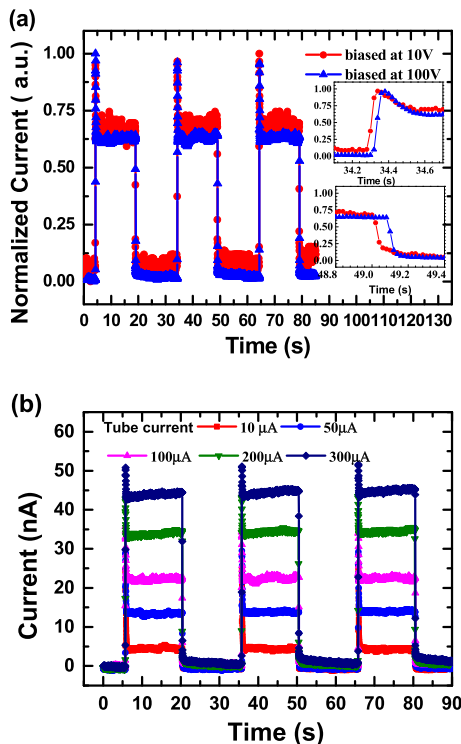


Fig. 3. The transient responses of the ϵ -Ga₂O₃ detector under (a) different bias voltages and (b) different X-ray dose rates.

recombination theory, the recombination rate of the radiation-generated carriers increased with the increase of the carrier density, which was directly proportional to the incident X-ray dose rate. However, the collection of the radiation-generated carriers in our detector was incomplete.

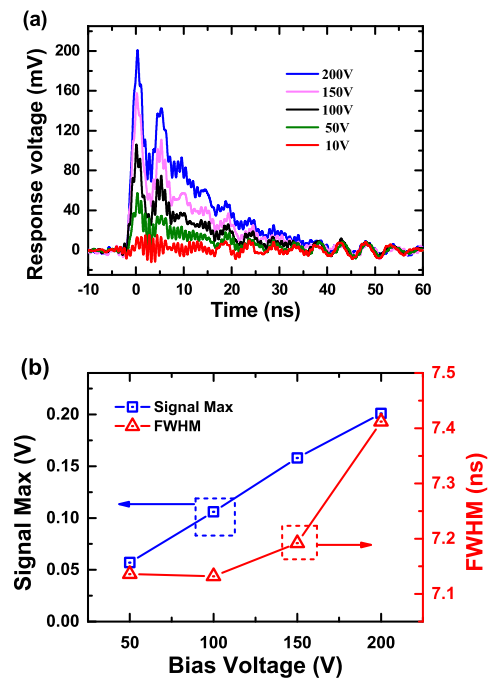


Fig. 4. (a) The pulsed X-ray responses of the fabricated ϵ -Ga₂O₃ detector at various bias voltages. (b) The maximal amplitude and FWHM of response signals.

Owing to the intrinsic high resistivity property of the UID ϵ -Ga₂O₃ film, the fabricated detector was expected to exhibit a fast response to the X-ray illumination. Fig.3 (a) and (b) show the transient responses of the ϵ -Ga₂O₃ detector under different bias voltages and different X-ray dose rates, respectively. The X-ray switching period was set to be 30 s and the sampling interval of the measurement was 20 ms. It was revealed that the detector exhibited a good and stable transient response with both the rise time (10% to 90%) and decay time (90% to 10%) of less than 50 ms, as shown by the enlarged plot of the rise and decay process in the inset of Fig. 3 (a).

The pulsed X-ray response of the fabricated ϵ -Ga₂O₃ detector was characterized using an XRS-4 pulsed X-ray source and the response signals at various bias voltages were shown in Fig. 4 (a). The ϵ -Ga₂O₃ detector exhibited an obvious pulsed response to the 50 ns in full width pulsed X-ray illumination when the bias voltage was larger than 50V, while it was hardly distinguished at a bias voltage of 10 V due the low signal-to-noise ratio (SNR). The maximum amplitude and the full width at half maxima (FWHM) of the response signals were plotted as a function of bias voltages in Fig. 4(b). The maximum value of the response signal increased linearly with the bias voltage, which was attributed to the incomplete carrier collection efficiency of the detector as discussed before. A time resolution of ~ 7.1 ns was determined by the FWHM of the pulsed response signal when the detector was biased at 100 V, which was in the same level with the previously reported Fe-doped β -Ga₂O₃ detector [9]. Compared to the polycrystalline ZnO detector in [23], which was in the same device topology but had a millisecond time resolution, the ϵ -Ga₂O₃ detector in this study showed a much faster response by three orders of magnitude. Owing to the single crystalline and intrinsic high resistivity properties of the epitaxial ϵ -Ga₂O₃

TABLE I
COMPARISON OF PERFORMANCE X-RAY DETECTORS
BASED ON WIDE GAP SEMICONDUCTORS

Detectors	Sensitivity / C·Gy ⁻¹ ·cm ⁻³	Rise/Fall time /s	Pulsed response
GaN p-i-n ^[26]	3.4×10 ⁻⁴ @0V	< 20 m	N/A
Ni/GaN:Fe/Ti (bulk) ^[27]	N/A	28	N/A
Al/ZnO/Al (p.film) ^[25]	1.1×10 ⁻⁴ @40V	< 0.2	1 ms
Pt/Diamond/Pt (film) ^[28]	5.8×10 ⁻² @150V	0.008	N/A
Ti/β-Ga ₂ O ₃ :Mg/Ti (bulk) ^[7]	3×10 ⁻³ @1000V	< 0.2	N/A
Ti/β-Ga ₂ O ₃ :Fe/Ti (bulk) ^[9]	2.2×10 ⁻⁸ @40V	0.02	< 2 ns
ITO/Ga ₂ O ₃ /ITO (a.film) ^[10]	0.271@50V	68.3/51.3	N/A
Au/α-Ga ₂ O ₃ /Au (n.film) ^[11]	0.1388@50V	< 0.035	N/A
Au/ε-Ga ₂ O ₃ /Au (film) ^[12]	4.75×10 ³ @20V	132/37 m	N/A
This work (film)	1.0×10 ³ @40V	< 0.05	< 7.1 ns

Notes: p.film means polycrystalline film; a.film means amorphous film; n.film means nanocrystalline film.

film, the carrier transport process in the detector would be less affected by defects.

The time resolution of the detector might be faster than 7.1 ns actually, limited by the pulsed source in the experiment. Because the detector exhibited the 50 ns pulsed waveform in the same width with the XRS-4 pulsed X-ray source and the FWHM had reached 7.1 ns beyond 50 V bias voltage. What's more, XRS-4 pulsed X-ray source was a multi-peak source, some pulsed peaks can be detectable with the increase of the sensitivity of the detector. Thereby, the FWHM of the envelopment of the pulsed response curve just slightly grew with the bias voltage beyond 100 V.

Table I shows the comparison of the normalized sensitivity and response speed of our device with various reported detectors using wide band gap semiconductors. The result clearly indicates that the low cost UID ε-Ga₂O₃ film based detector has great potential in fast X-ray radiation detection.

IV. CONCLUSION

In summary, a single crystalline UID ε-Ga₂O₃ film was grown on a sapphire substrate by MOCVD and fabricated into an MSM X-ray detector. Featuring an intrinsic high resistivity property of the ε-Ga₂O₃ epilayer, the detector exhibited a low dark current and a relatively high sensitivity. Pulsed X-ray detection with a time resolution of ~ 7.1 ns was demonstrated using the fabricated ε-Ga₂O₃ detector.

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