

Material and Ultrafast Optoelectronic Properties of Furnace-Annealed Arsenic-Ion-Implanted GaAs

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Abstract—Structural, electrical, and ultrafast optical properties of furnace-annealed arsenic-ion-implanted GaAs (GaAs:As⁺) has been investigated for its applications in ultrafast optoelectronics. From these studies, we determine that GaAs substrates implanted with 200-keV arsenic ions at 10¹⁶ ions/cm² and furnace-annealed at 500 °C–600 °C would have recovered its crystallinity, be highly resistive, and exhibit picosecond photo-excited carrier lifetimes. The duration of the electrical pulses generated by photoconductive switches (PCS's) fabricated on the optimized material was ≈4 ps. The risetime (10%–90%) and 1/e falltime were, respectively, ≈2 and 3 ps. These results were measurement-system limited. We estimated the actual response to be ≈2 ps, consistent with a photo-excited carrier lifetime of ≈1.8 ps. The peak responsivity was ≥4×10⁻³ A/W. The dark current for the GaAs:As⁺ PCS biased at 40 V was as low as 5 nA. The break down field was higher than 150 kV/cm. These characteristics are comparable to those of state-of-the-art photoconductors such as LT-GaAs.

Index Terms—Arsenic-ion-implanted GaAs, photoconductive switch, ultrafast optoelectronics.

I. INTRODUCTION

PHOTOCONDUCTORS with ultrashort photo-excited carrier lifetime, good optical responsivity, high breakdown field, and low dark current are essential for ultrafast optoelectronic switching applications. Various classes of semiconductors, e.g., intrinsic, impurity-dominated, radiation-damaged, polycrystalline, and amorphous, have been explored as ultrafast photoconductors [1]. In particular, the ion implantation technique has been employed extensively [2]–[5]. Carrier lifetimes as short as 0.5–0.6 ps have been reported as the saturation limit for GaAs samples irradiated by protons (H⁺) [2]. Similar results were reported for oxygen-ion-implanted silicon-on-sapphire (SOS) materials [5]. On the other hand, nonstoichiometric arsenic-rich GaAs grown by molecular beam epitaxy (MBE) at low substrate tempera-

ture (LT-GaAs) have been the subject of intensive studies [6]. It exhibited nearly ideal electrical and optoelectronic properties for ultrafast optoelectronic applications. Recently, a new class of arsenic-rich material, arsenic-ion-implanted GaAs or GaAs:As⁺, has emerged as a potential alternative to LT-GaAs [7]. The structural and electrical characteristics of LT-GaAs and GaAs:As⁺ have been shown to be quite similar [8], [9]. The ultrafast optical characteristics of the as-implanted arsenic-rich GaAs materials have been reported [10]–[12]. Subpicosecond carrier lifetimes were observed. However, as-implanted or as-grown nonstoichiometric GaAs obtained by using either the arsenic-ion-implantation or the LTMBE process suffer from the following: the crystallinity of the material is degraded [13], [14]. As a result, the carrier mobility is substantially lower [12], [15]. This also leads to a defect-dominated hopping conducting mechanism, which destroys the highly resistive nature of the original substrate [8], [15]. Consequently, the as-implanted GaAs:As⁺ is not yet suitable for fabrication of ultrafast optoelectronic devices. To remedy this, rapid thermal annealing (RTA) has been successfully employed. The effects of RTA on photo-excited carrier lifetime and electrical properties of GaAs:As⁺ were reported [16]. Ultrafast photoconductive switches were demonstrated on RTA-annealed GaAs:As⁺ prepared by bombarding semi-insulating (SI) GaAs substrates with 200-keV arsenic ions at the dosage of 10¹⁶ ions/cm² [10], [16]. We have also reported picosecond photoconductive switches fabricated on low-dose-implanted GaAs:As⁺ [17]. An interesting alternative to RTA is furnace annealing. In this paper, we present a detailed study on the structural, electrical, and ultrafast optical properties of furnace-annealed GaAs:As⁺. The optimum processing condition is determined to be furnace-annealing at 600 °C for 30 min. Picosecond photoconductive switches (PCS's) were fabricated on this material. The devices exhibit characteristics better than or comparable to those of LT-GaAs.

II. SAMPLE PREPARATION

The substrates were liquid-encapsulated-Czochralski (LEC) grown SI GaAs wafers. For implantation of arsenic ions, we employ a commercial apparatus (Varian E220). The implanting dosage and energy were 10¹⁶ ions/cm² and 200 keV, respectively. Typical current density value was 1 mA/cm². The exposure time subject to the samples was 10–15 min. The postannealing process was performed *ex situ* by the encapsulated thermal annealing technique. We used a quartz-tube

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furnace with flowing nitrogen gas. The annealing temperature was varied from 200 °C to 800 °C. The annealing time was 30 min.

III. EXPERIMENTAL METHODS

For structural characterizations of the arsenic-ion-implanted GaAs substrates, we employ double-crystal X-ray diffractometry (DCXRD) and transmission electron microscopy (TEM) techniques. In DCXRD, a standard GaAs crystal and a Cu_α source were employed to check the crystallinity on (004) diffraction plane of the samples by measuring the rocking curve. For TEM analysis, the samples were carefully polished and ion-milled to reduce its thickness down to several nanometers. A JOEL model JEM-200CX electronic microscope with resolution of 2 Å was utilized to obtain the cross-sectional-viewed photograph of bright field and associated diffraction matrix patterns. For dark current measurements, Schottky diodes with thermally evaporated gold (0.5 μm thick) contacts were fabricated on GaAs:As⁺ samples. Their current voltage (I - V) characteristics were then measured by using an HP4145B semiconductor parametric analyzer.

For measurement of photo-excited carrier lifetime and transient index-of-refraction of the annealed samples, we employed a standard pump-probe setup [11], [18]. The light source was a home-made, passively mode-locked Ti:sapphire/HITCI laser [11]. It generated optical pulses at 82 MHz with a pulsewidth of about 130 fs. The polarization and power ratio of pump and probe beams were parallel and 50/1 in our experiment (400 mW/8 mW). The laser wavelength was 0.865 μm, which corresponds to an energy of only about 13 meV higher than the bandgap of GaAs (1.42 eV). This wavelength was chosen to prevent either the intervalley scattering of the photo-excited carriers from the Γ band to the satellite band (usually the L band, $\Delta E_{\Gamma-L} = 300$ meV) in k -space, or the emission of longitudinal optical phonon ($\Delta E_{LO} = 36$ meV). Thus the intraband relaxation and phonon scattering processes can be ignored in our experiments. The energy of the pump pulse mechanically chopped at 3.2 kHz was about 4 ± 0.5 nJ. This corresponds to an estimated injection density of photo-excited carriers of about 1×10^{19} cm⁻³. The fractional change of the reflected probe beam was detected by a pair of large area Si p-i-n diodes operated in differential-detection mode and subsequently processed in a lock-in amplifier operated in a (R, θ) mode. The detection sensitivity of our system is usually better than 10^{-5} . The transient change in reflectivity ΔR (in root mean square value) obtained from the lock-in amplifier was normalized to calculate fractional change in transient reflectivity, $\Delta R/R$. The transient refractive index change Δn was evaluated by using the Simplified Kramers-Kronig relation [19], i.e., $\Delta R/R = 4\Delta n/(n^2 - 1)$.

The ultrafast switching response of the switch was examined by using an external electrooptic sampling (EEOS) system. The wavelength was set at 0.8 μm. The pump power incident on the PCS was 140 mW while the probe power was 5 mW or less. The reflective-mode electrooptic sampling module consists of a polarization beam splitter (PBS), a quarter-

wave plate (QWP), a half-wave plate (HWP), and a LiTaO₃ electrooptic probing crystal. The crystal was thinned and polished down to 60 μm and placed upon the PCS. The probe point was about 2 mm away from the photoconductive gap. The temporal and spatial resolutions of the current system were estimated to be about 1 ps and 5 μm, respectively.

IV. STRUCTURAL AND ELECTRICAL PROPERTIES

A. DCXRD Diffractometry

The X-ray rocking curves for the as-implanted and annealed samples are shown in Fig. 1(a). The most significant side peaks at the lower angle side of the curves are attributed to the as-implanted layer. It is indicative of an expanded lattice constant for the implanted layer as compared with that of the SI GaAs substrate ($a = 5.6533$ Å). The angular separation between principal and side peaks is about 240 arc/s. By using the formula $t = \lambda \gamma_H / [\Delta\theta * \sin 2\theta_B]$, [20] we estimate the thickness of the as-implanted layer, $t \approx 90$ nm. In the formula, λ is the X-ray wavelength, γ_H is the direction cosine of the diffracted X-ray with respect to the surface normal, θ_B is the Bragg's angle, and $\Delta\theta$ is the angular separation between adjacent maxima of the rocking curve. The thickness is in good agreement with results of TRIM simulation of the implantation profile. In general, the maximum deviation of the lattice constant Δa and the vertical strain between the implanted region and the bulk $\Delta a/a$ can also be simply evaluated from the rocking curve by using the Bragg's law, $\Delta a/a = -\Delta\theta / \tan \theta_B$ [21]. A more accurate depth-dependent strain profile was obtained by fitting the X-ray rocking curves with calculated strain profiles. The latter was obtained by using a Phillips software package (HRD Simulation Software 1.0). The package is based on the usual kinematic theory for X-ray analysis [22]–[24]. The depth-dependent strain profile of the as-implanted GaAs:As⁺ is shown in the inset of Fig. 1(b). The maximum vertical strain is $\Delta a/a$ 0.35% at a depth of about 0.09 μm below the surface. In comparison, Fujioka *et al.* [25] reported a maximum lattice expansion of 0.36% for as-implanted GaAs:As⁺ in a numerical study. Similar X-ray side peaks with an estimated lattice expansion of 0.0053 Å was also observed in LT-GaAs grown at 200 °C. The corresponding maximum strain is only $\Delta a/a = 0.08 \sim 0.15\%$ [26], [27].

After annealing, the X-ray sidepeak was observed to gradually shift toward the central peak. This leads to a slightly broadened and asymmetric central peak with a slowly decreasing tail toward the negative angular direction. For GaAs:As⁺ annealed at and above $T_a = 500$ °C, the residual side peak essentially merged into the central peak. The asymmetry of the wings was less pronounced. For samples annealed at $T_a = 800$ °C, the X-ray rocking curve is identical to that of SI GaAs. A very similar trend was observed in LT-GaAs [28]. The average vertical strain $(\Delta a/a)_{\text{avg}}$ of GaAs:As⁺ annealed at different temperatures are shown in Fig. 1(b). These were obtained by integrating the strain profile over the implantation depth. Examining Fig. 1(a) and (b), we estimate that the critical annealing temperatures for the complete recovery of

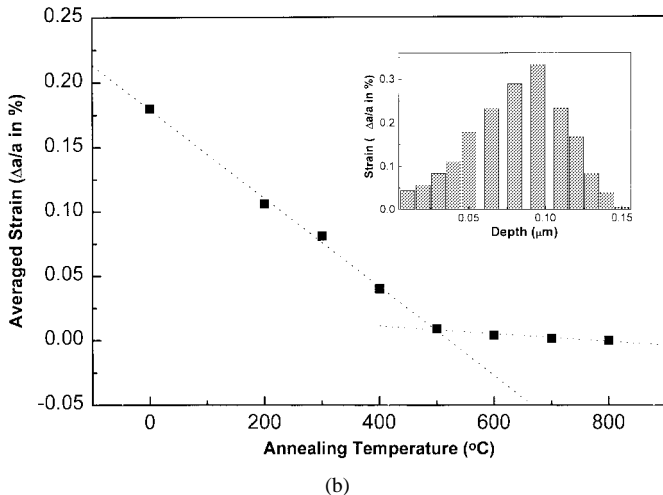
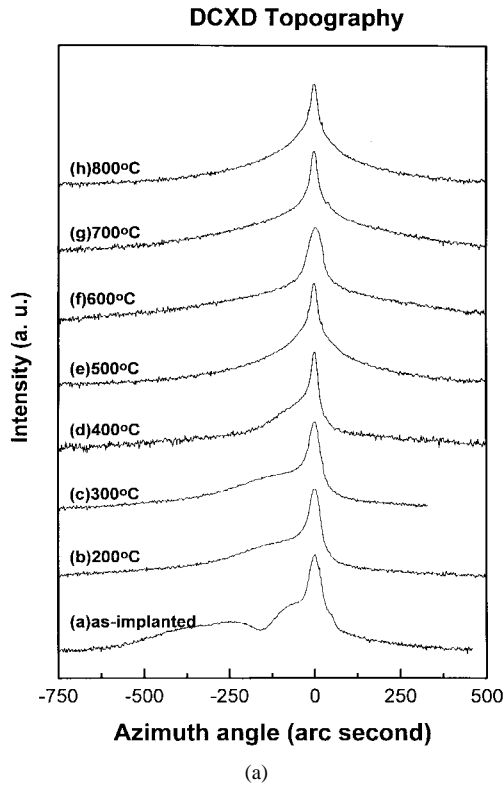


Fig. 1. (a) The double-crystal X-ray diffraction profiles of as-implanted and furnace-annealed GaAs:As⁺ samples. (b) The average vertical strain in the GaAs:As⁺ samples are plotted as a function of annealing temperature. The inset shows the depth-dependent strain profile of as-implanted GaAs:As⁺.

crystallinity is $T_a \approx 500$ °C as opposed to $T_a = 450$ °C for LT-GaAs [28].

The densities of the neutral As_{Ga} antisite ([As_{Ga}⁰]) defects in GaAs:As⁺ annealed at different temperatures can be evaluated from $(\Delta a/a)_{\text{avg}}$. These are plotted as a function of T_a in Fig. 2. The analysis is based on the following: it has been reported by several groups that three major kinds of defects—As_{Ga} antisite (including As_{Ga}⁰- and As_{Ga}⁺-type defects), interstitial (As_i), and vacancy (V_{ga}) defects are present in LT-GaAs [29]–[32]. The concentration of the antisite defects is higher than the other kind of defects by at least one order of magnitude. Typically, $N[V_{\text{Ga}}] \approx 3 \times 10^{17}$

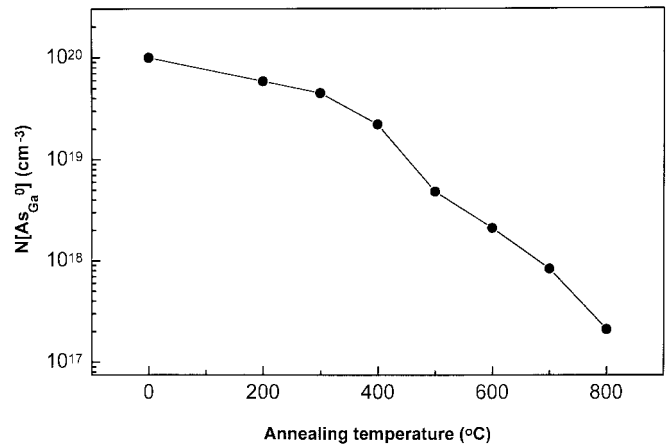


Fig. 2. The concentration of [As_{Ga}⁰] antisite defects in the GaAs:As⁺ samples plotted as a function of the annealing temperature.

cm⁻³ [29]. Furthermore, the concentration of As_{Ga}⁰ defects, $N[\text{As}_{\text{Ga}}^0] \approx 2\text{--}9.9 \times 10^{19}$ cm⁻³, [30], [31] is one order of magnitude higher than that of As_{Ga}⁺ defects, $N[\text{As}_{\text{Ga}}^+] \approx 2\text{--}8 \times 10^{18}$ cm⁻³ [32]. Now the structural and electrical characteristics of GaAs:As⁺ have been shown to be strikingly similar to that of LT-GaAs [33]. We thus argue that the As_{Ga}⁰-type defects would also be the most dominant traps for carriers in GaAs:As⁺. Next, we assume that $N[\text{As}_{\text{Ga}}^0]$ is linearly proportional to $(\Delta a/a)_{\text{avg}}$ in GaAs:As⁺. This is based on the empirical relation, $N[\text{As}_{\text{Ga}}^0] \approx 8.065 \times 10^{22} \cdot (\Delta a/a)_{\text{avg}}$ reported for LT-GaAs by Liu *et al.* [32]. In analogy to LT-GaAs, we suppose that $N[\text{As}_{\text{Ga}}^0]$ monotonically decreases from a maximum value of $\sim 10^{20}$ cm⁻³ for as-implanted GaAs:As⁺ [8] down to the minimum value of about 1×10^{16} cm⁻³ for the unprocessed SI GaAs substrate [34]. We can then write that $N[\text{As}_{\text{Ga}}^0] \approx 5.56 \times 10^{22} (\Delta a/a)_{\text{avg}}$ for GaAs:As⁺. The proportional constant in the expression is of the same order but slightly smaller than that for LT-GaAs. It correlated with the more severely damaged nature of the as-implanted GaAs:As⁺ layer as compared with that of as-grown LT-GaAs. As a result, GaAs:As⁺ must be annealed at a temperature higher than that for LT-GaAs to recover the crystallinity. As a supporting evidence, the concentration, $N[\text{As}_{\text{Ga}}^0]$ of the GaAs:As⁺ sample annealed at 600°C predicted by Fig. 2 was in reasonably good agreement with previously reported value of $\sim 3 \times 10^{18}$ cm⁻³ from the capacitance-voltage measurement [35].

The structural parameters, $\Delta a/a$ and $N[\text{As}_{\text{Ga}}^0]$, of as-implanted and annealed GaAs:As⁺ are summarized in Table I.

B. TEM Analysis

The cross-section views of the bright field TEM micrographs of as-implanted and furnace-annealed GaAs:As⁺ at $T_a = 600$ °C are shown in Fig. 3. The as-implanted sample is amorphous and reveals a nonstoichiometric and stress-rich structure. After annealing at $T_a = 200$ °C for 30 min, the 90-nm-thick implanted layer transformed into the polycrystalline phase with grain boundaries. If T_a was increased up to 400 °C, we found that the grain boundaries were almost not discernible.

TABLE I

	as-implant.	200 °C	300 °C	400 °C	500 °C	600 °C	700 °C	800 °C
$\Delta a/a$	1.8e-3	1.06e-3	8.1e-4	4e-4	8.7e-5	$\sim 3.79e-5$	$\sim 1.5e-5$	$< 3.8e-6$
$[As_{Ga}]$ (cm ⁻³)	$\cong 1e20$	5.9e19	4.5e19	2.2e19	4.8e18	2 \sim 3e18	8.5e17	$\sim 2e17$
D_{as} (nm)	–	–	–	–	–	2 \sim 3	5	20
I_{dark} (nA/ μm^2)	12	~ 0.6	~ 1	0.12	8.7e-5	5.2e5	0.05	~ 3.3
τ (ps)	0.26 \sim 0.3	0.4 \sim 0.42	0.47 \sim 0.5	0.58 \sim 0.6	0.85 \sim 0.9	1.6 \sim 1.9	2.8	4.2 \sim 4.6
Δn ($\times 10^{-2}$)	-1.1 \sim -1.3	-0.26 \sim -0.33	-0.1 \sim -0.21	-0.12 \sim -0.18	-0.14 \sim -0.21	-0.27 \sim -0.32	0.6 \sim 0.66	0.85 \sim 0.97
σ ($\times 10^{-14}$ cm ²)	~ 0.19	~ 0.21	~ 0.22	~ 0.39	~ 1.2	~ 1.5	~ 2.1	≥ 6

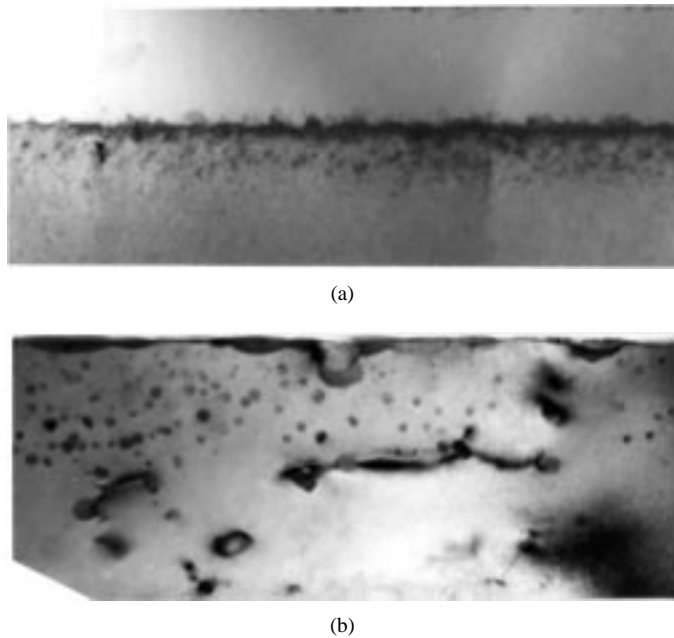


Fig. 3. The cross-sectional views of the bright-field TEM micrographs and associated electron diffraction patterns of (a) as-implanted and (b) furnace-annealed GaAs:As⁺ at $T_a = 600$ °C.

The implanted region exhibited a large number of dislocation defects in the GaAs matrix and stacking faults near the surface. No precipitates were observed, however. In comparison, the arsenic precipitating process was found to occur at T_a between 400 °C and 475 °C for LT-GaAs [28]. For samples annealed at $T_a = 500$ °C, the sample appeared to be single-crystalline in the TEM micrograph. A large number of cluster-type defects, mostly coffee-bean type in shape with a few As precipitates, were observed. The sizes of the precipitates, however, were still beyond the resolution limit of the TEM. The density of clusters was estimated to be $\geq 10^{17}$ cm⁻³. For samples annealed at 600 °C, we found the associated (110) diffraction pattern to be perfectly crystalline. Some weak spots in the diffraction pattern also indicate the existence of As precipitates in normal GaAs matrix. The crystal symmetry was assigned as the 3-m rhombohedral As (metallic) group. A significant number of arsenic precipitates and dislocation defects were also observed. The concentration and mean size of arsenic precipitates were about 2×10^{16} cm⁻³ and 2–3 nm, respectively. In comparison, the concentration and size of As clusters in the LT-GaAs annealed at the same temperature

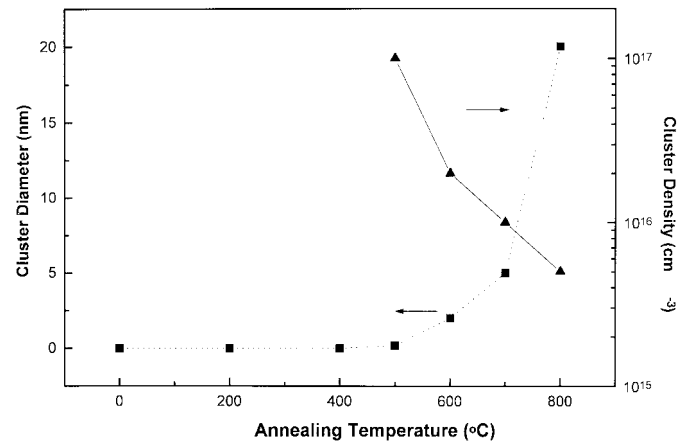


Fig. 4. The diameter and density of arsenic precipitates in GaAs:As⁺ are plotted as a function of annealing temperature.

was reported to be 5×10^{17} cm⁻³ and 5–8 nm [36], [37]. GaAs:As⁺ samples RTA-annealed at 600 °C for 30 s were still polycrystalline with a large amount of grain boundaries, dislocation lines, and defects [18]. If the RTA-annealing time was increased from 30 s up to 10 min at temperature of 600 °C, the crystal regrowth and arsenic precipitating process become more pronounced [18]. For samples annealed at $T_a = 700$ °C, we observed a large number of precipitates with larger diameters (~ 5 nm) and lower density ($\sim 1 \times 10^{16}$ cm⁻³) in the bright-field micrographs. Very few dislocation defects were observed. In the electron diffraction pattern of the same sample, we identified less distinct spots superimposed onto the principal diffracted spots. This is interpreted as the manifestation of a decrease in number of As precipitates from the GaAs matrix. Still larger arsenic clusters (~ 20 nm) with further reduced densities ($\sim 3 \times 10^{15}$ cm⁻³) were observed when the annealing temperature was increased up to 800 °C. The decreasing density may be attributed to both the precipitation into clusters of larger size and the out-diffusion of arsenic ions at high annealing temperatures. Our results illustrate that the arsenic precipitating phenomenon and the fully crystalline regrowth of the implanted region were more pronounced only at higher annealing temperatures. The trends in decreasing concentration but increasing size of As precipitates at higher annealing temperatures are shown in Fig. 4. The size and concentration of the arsenic precipitates as a function of annealing temperature are also summarized in Table I.

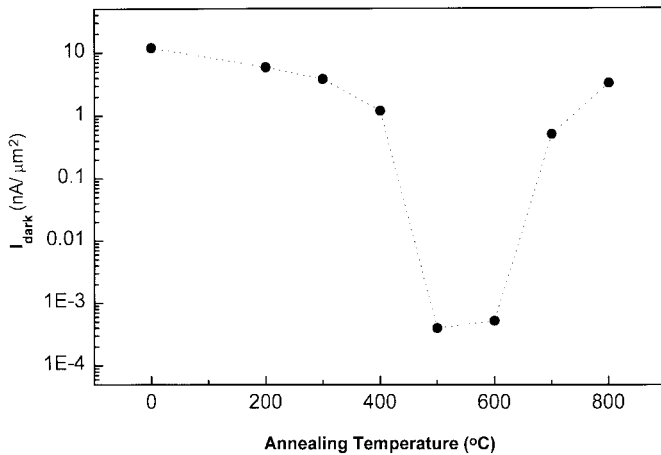


Fig. 5. The dark current densities of Schottky diodes made on the arsenic-ion-implanted GaAs substrates annealed at different temperatures.

C. Electrical Characteristics

The leakage current densities, I_{dark} 's, of Schottky diodes fabricated on GaAs:As⁺ as a function of T_a are shown in Fig. 5. The diodes were biased at -5 V. For example, the leakage current density for the as-implanted sample was -12 nA/ μm^2 . As we increased T_a from 200 °C to 400 °C, I_{dark} decreased from -6 to -1 nA/ μm^2 . Following our previous analysis [38], the change from hopping conduction to thermal emission mechanism, corresponding to the decline of density of trapping sites after annealing, is thought to be responsible for the recovery of the resistivity of the damaged surface layer. A reduction of four orders of magnitude of the leakage current from that of the as-implanted sample to a minimum of only about -3×10^{-4} nA/ μm^2 was observed for samples annealed at 500 °C and 600 °C. This is attributed to the onset of arsenic precipitation phenomenon presented in Section IV-B. It is consistent with the buried Schottky barrier model proposed by Warren *et al.* [39]. For $T_a > 600$ °C, I_{dark} increased drastically and was as high as -3.3 nA/ μm^2 for GaAs:As⁺ annealed at $T_a = 800$ °C. In Section IV-B, we have also shown that the density of arsenic precipitates decreased as the number of arsenic precipitates in the GaAs matrix dissolved for samples annealed at $T_a > 600$ °C. This behavior is also congruous with the buried Schottky barrier model. It likewise agrees with the work of Claverie *et al.* [40]. They showed that a large concentration of very small As precipitates were more efficient for increasing the resistivity of arsenic-implanted and furnace-annealed n-type GaAs substrates than a smaller number of larger precipitates. As a result, the implanted layer became more conductive as $T_a > 600$ °C. Furthermore, in our recent work [41], we have identified a shallow acceptor level located at 0.3 eV above the valance band maxima for samples annealed at high temperatures. This level is related to the gallium vacancy (usually denoted as V_{Ga})-related defects. These V_{Ga} -related defects will lead to incomplete compensation between deep donor and shallow acceptor defects. Consequently, the implanted layer becomes more leaky. In view of the electrical characteristics of GaAs:As⁺, the optimal annealing temperature for device applications is between 500 °C and 600 °C.

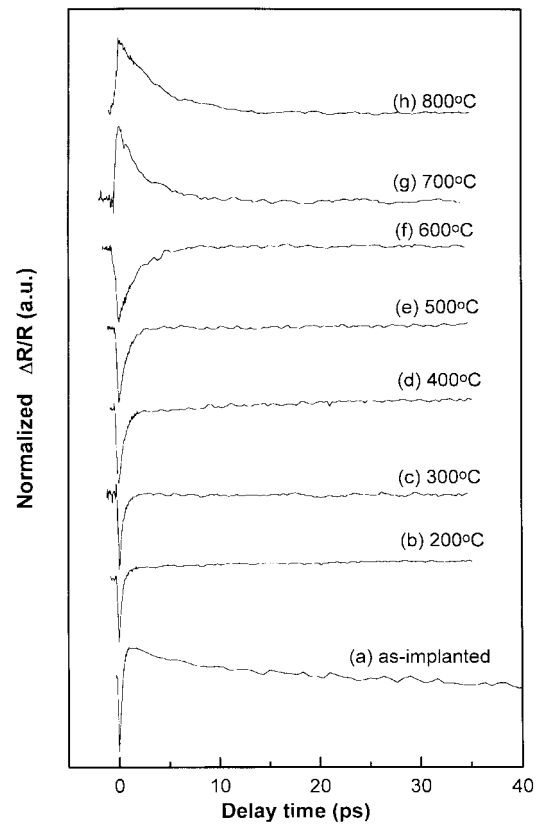


Fig. 6. The normalized time-resolved reflectivity $\Delta R/R$ of the arsenic-ion-implanted GaAs samples annealed at different annealing temperatures.

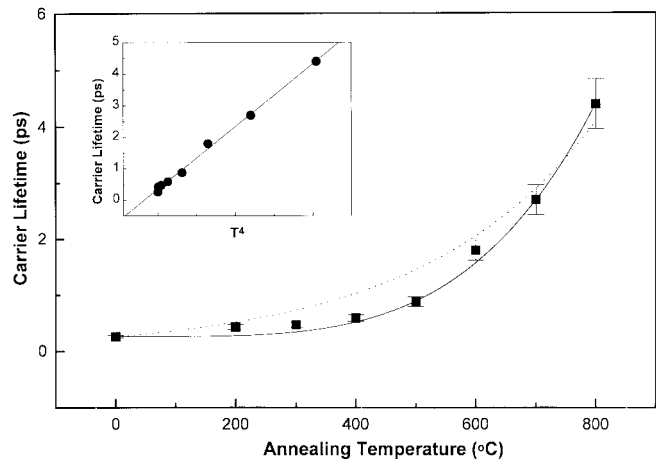


Fig. 7. The photo-excited carrier lifetimes of the arsenic-ion-implanted GaAs samples annealed at different temperatures. The solid and dotted line are fitted curves showing the T_a^4 and exponential dependence (see also the inset which is a semi-log plot).

V. ULTRAFAST OPTICAL PROPERTIES

Previously, we have reported subpicosecond photo-excited carrier lifetimes, τ_c 's, ranging from 0.19 to 0.28 ps in GaAs implanted with 200 keV arsenic ions at dosages from 10^{12} to 10^{16} ions/ cm^2 [11]. For RTA-annealed GaAs:As⁺ at 600 to 800°C, τ_c 's were observed to lengthen from 0.48 to 2.3 ps [18]. From results of femtosecond time-resolved reflectivity measurements (see Fig. 6) performed in this paper, we find that photo-excited carrier lifetimes of furnace-annealed GaAs:As⁺

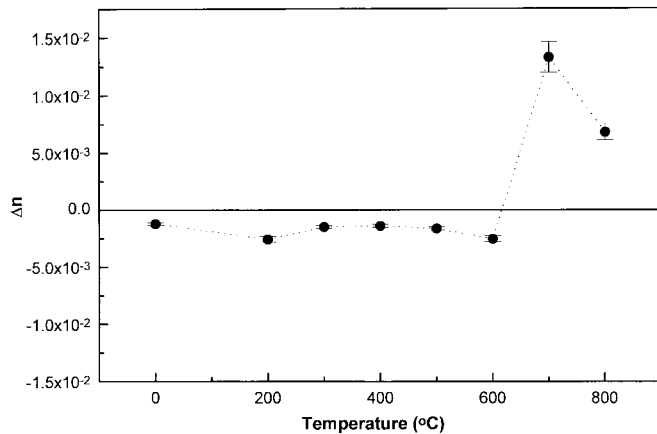


Fig. 8. The peak transient refractive index change Δn is plotted as a function of the annealing temperature.

are in the range of ≈ 0.4 to 4.6 ps (200 °C $\leq t_c$'s ≤ 800 °C) with an approximate $(T_a)^4$ dependence (inset of Fig. 7). The τ_c 's are plotted as a function of T_a in Fig. 7. For $T_a \leq 500$ °C, τ_c 's were found to evolve relatively slowly with increasing T_a and were all in the subpicosecond range. The corresponding changes of the refractive index, Δn , were of the order of -2 to -3×10^{-3} . This is shown in Fig. 8. The maximum Δn is comparable with the value reported in previous works on either arsenic-ion-implanted GaAs [42] or MBE grown LT-GaAs materials [43]. A sign reversal for Δn was observed for 600 °C $\leq T_a \leq 700$ °C. The peak magnitude of Δn was as large as 0.01 at $T_a = 700$ °C. This can be explained if the bandgap of GaAs:As⁺ shrinks due to bandgap renormalization toward shorter wavelength after annealing. This picture is in agreement with results of the CW photoreflectance (CWPR) experiment, as shown in Fig. 9. The signal-to-noise ratio (SNR) of the CWPR data for samples annealed at lower temperatures are relatively poor. This is due to the surface Fermi-level pinning effect, which strongly attenuates the modulated pump signal and the probe beam, reflected from the damaged surface. We observed a blue shift of the CWPR signal for samples annealed at 500 °C $\leq T_a \leq 700$ °C. This may indicate a recovery on bandgap energy of GaAs:As⁺ samples due to the reduction of defects near the band edge. A sign change of the CWPR amplitude was observed at probe energy of 1.433 eV for samples annealed between 600 °C and 700 °C. As we have shown in Section IV, the crystallinity of furnace-annealed GaAs:As⁺ recovers approximately after furnace-annealing at $T_a \geq 500$ °C for 30 min. Thus the sign reversal in both time-resolved reflectivity and CWPR may be explained by the bandgap shrinkage effect associated with the structural change. At $T_a = 800$ °C, a further sign reversal was observed in the CWPR signal. This may correspond to a type change of the material from n-type to p-type. The reduction of deep-donor defects and the presence of shallow acceptor defects in the samples may explain this result. The CWPR signal at 1.433 eV, however, is quite small for the sample annealed at $T_a = 800$ °C as compared with that of samples annealed at lower temperatures. This is not consistent with the trend for the transient reflectivity signal. Further study is required for clarification.

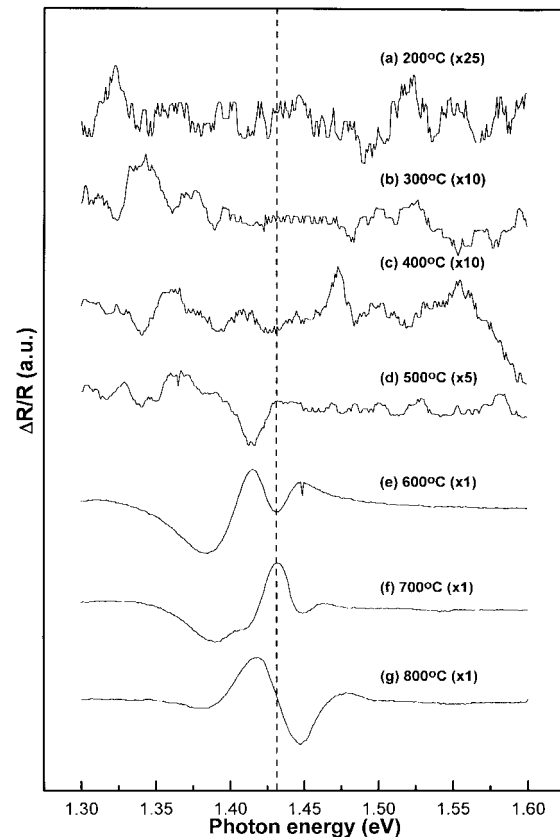


Fig. 9. Results of the CW photoreflectance (CWPR) measurement of GaAs:As⁺ samples annealed at different temperatures. The dashed line shows the pump wavelength for the time-resolved transient reflectivity experiments.

The ultrafast carrier lifetimes are consistent with structural and electrical characterization of GaAs:As⁺ that demonstrated ultrahigh densities of defects present in the samples. For example, the density of neutral As_{Ga} antisite defects remained $\approx 2 \times 10^{17}$ cm⁻³ at T_a as high as 800 °C. Arsenic precipitates, however, were not observed for $T_a \leq 600$ °C. This suggests that arsenic clusters were not as important for the ultrafast carrier trapping process in GaAs:As⁺ as for LT-GaAs. We have evaluated the capture cross section of photo-excited carriers in GaAs:As⁺ by using the Shockley-Read-Hall recombination theory [44]. In this model, $t_c = (N_t \sigma v_{th})^{-1}$, where N_t is the concentration of the carrier traps, σ is the capture cross section for the photo-excited carriers, and v_{th} is the thermal velocity of carriers in the photoconductor. We first assumed that $N_t \approx N[\text{As}_{\text{Ga}}^0]$. Using data in Section IV, we find $\sigma \approx 0.2\text{--}0.4 \times 10^{-15}$ cm² for samples annealed at $T_a \leq 400$ °C. The cross section increased up to $1\text{--}2 \times 10^{-14}$ cm² for annealing temperatures in the range of 500 °C– 700 °C. At $T_a = 800$ °C, $\sigma \approx 6\text{--}7 \times 10^{-14}$ cm², which is at least three times as large as σ 's at lower annealing temperatures. To check these values, samples annealed at 600 °C were also examined by using deep-level transient spectroscopy (DLTS) [36], [39]. Fitting from arrenius plot of the DLTS data, we obtain $\sigma \approx 7 \times 10^{-14}$ cm². On the other hand, we can estimate σ by using the diameters of the arsenic precipitates. That is, $s \approx \pi d^2/4$. These are $s \approx 2.8 \times 10^{-13}$, 2.0×10^{-13} , and 3.14×10^{-12} cm²,

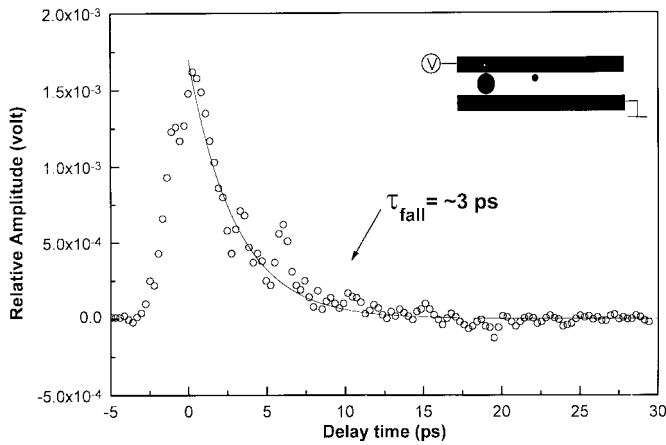


Fig. 10. The electrooptically sampled switching response of a photoconductive switch fabricated on arsenic-ion-implanted GaAs substrate furnace-annealed at 600 °C for 30 min.

respectively, for GaAs:As⁺ samples annealed at $T_a = 600$ °C, 700 °C, and 800 °C. The corresponding densities of the arsenic precipitates were for the GaAs:As⁺ samples annealed at different temperatures were $N_{As} \approx 2.0 \times 10^{16}$, 1.0×10^{16} , and 3.0×10^{15} cm⁻³, respectively. Accordingly to the Shockley–Read–Hall model, the expected lifetimes would be 5.1, 14.6, and 3.0 ps. These are not in agreement with the experimental data. We conclude that point and structural defects such as the arsenic antisite defects, instead of arsenic precipitates, are dominant as trapping centers of carriers in furnace-annealed GaAs:As⁺. The capture cross sections and ultrafast optical properties of GaAs:As⁺ are also summarized in Table I.

VI. ULTRAFAST OPTOELECTRONIC SWITCHING

From the structural, electrical, and ultrafast optical characterizations of GaAs:As⁺ in the previous sections, the optimum processing condition for switching applications were determined to be furnace-annealing at 600 °C and 30 min. A coplanar stripline (CPS) structure with its characteristic impedance of $\sim 75 \Omega$ was fabricated on this material. The 2-cm-long CPS consists of two 30- μ m-wide transmission lines separated by 15 μ m. The contact metal of the PCS was Au–Ti (0.5 μ m–0.03 μ m). The photoconductively generated electrical pulse as determined by the EEOS system exhibits a pulsewidth of ~ 4 ps (FWHM). This is shown in Fig. 10. The risetime and falltime of the impulse response were evaluated to be about 2.2 and 3–4 ps, respectively. These are instrument-limited. Dispersion effect may have broadened the pulse also, as the probe spot was located at 2 mm from the pump spot. We estimated that the actual response of the PCS to be ~ 2 ps. This is consistent with a $t_c \approx 1.8$ ps from transient reflectivity measurement. In comparison, Wang *et al.* [10] has reported that the response time of the same device fabricated on GaAs:As⁺ RTA-annealed at 600°C for 15 s was ≈ 1.23 ps (FWHM). Not long ago, we have also reported that ultrafast switching response of PCS fabricated on low-dosage (10^{13} ions/cm²) implanted GaAs:As⁺ substrate. The switching response of

RTA- and furnace-annealed low-dose-implanted PCS were ~ 3 and 8 ps, respectively [17].

The dark current of an Auston-type GaAs:As⁺ PCS with a gap size of 30 μ m fabricated on the same substrate was measured to be as low as 15 nA under a bias voltage of 40 V. The corresponding electric field biased across the gap is higher than 150 kV/cm. Such low dark current indicates that the annealed sample can sustain much higher bias voltage before breakdown. In comparison, the dark currents of PCS's fabricated on SI GaAs and RTA-annealed GaAs:As⁺ were > 10 nA and $\approx 100 \mu$ A at the same bias.

With a bias voltage of 20 V, the responsivity of the PCS operated in CW and pulse mode were 0.004 A/W and ≥ 0.03 A/W, respectively. The injected optical average powers were 100 mW (pulse mode) and 3 mW (CW mode), respectively. The responsivity is somewhat larger than the value of the devices made on different materials such as LT-GaAs (~ 0.001 A/W) [45]. The higher responsivity may be attributed to the contribution from the GaAs substrate since the thickness of the implanted layer was too thin to absorb the injected photons entirely.

VII. CONCLUSION

We have performed extensive structural, electrical, ultrafast optical, and optoelectronic characterization of furnace-annealed GaAs:As⁺. Structurally, we find the as-implanted GaAs:As⁺ to be amorphous. The crystallinity of the implanted region fully recovers for $T_a \geq 500$ °C. Arsenic precipitating process is observed for $T_a \geq 500$ °C only. Instead, arsenic antisite defects, As_{Ga}, are shown to be the dominant carrier trap for carriers for T_a as high as 800 °C. The trap densities are evaluated from the strain data and shown to decrease from 10^{20} for the as-implanted sample to $\sim 2 \times 10^{17}$ cm⁻³ at 800 °C. Increasing the annealing temperature from $T_a = 400$ °C to 500 °C, the dark current densities of the samples decrease drastically by two orders of magnitude to $\approx 3 \times 10^{-4}$ nA/mm². A sharp rise in I_{dc} was then observed for $T_a \geq 700$ °C. This is explained in terms of the density and size of the average As precipitates for samples annealed at different T_a 's. The photo-excited carrier lifetimes are found to evolve slowly but remain subpicosecond in samples annealed at $200 \text{ °C} \leq T_a \leq 500 \text{ °C}$. The τ_c 's then rise distinctly to 4.6 ps as T_a is increased to 800°C with an approximately T_a^4 dependence. If we assumed that As_{Ga} was the dominant trap, the capture cross sections of the deep-level trapping centers in furnace-annealed GaAs:As⁺ are in agreement with the Shockley–Read–Hall model and estimated to be in the range of $0.2\text{--}6 \times 10^{-14}$ cm². The transient change in refractive indices is of the same order as those of annealed LT-GaAs. The Δn is as large as 0.01 for GaAs:As⁺ furnace-annealed at $T_a \geq 600$ °C. A sign reversal in Dn is observed at this temperature. We have tentatively attributed this phenomenon to the bandgap shrinkage effect. From the above studies, we have determined that GaAs substrate implanted with 200-keV arsenic ions at 10^{16} ions/cm² and furnace-annealed at 500 °C–600 °C for 30 min would have recovered its crystallinity, be highly resistive, and exhibit picosecond photo-excited carrier lifetimes. Indeed, switching responses as fast as 3–4 ps (instrument limited) were

realized for coplanar-stripline type PCS's fabricated on the above material. We estimated the actual response to be ≈ 2 ps, consistent with a photo-excited carrier lifetime of ≈ 1.8 ps. The peak responsivity was $\geq 4 \times 10^{-3}$ A/W. The dark current for the GaAs:As⁺ PCS biased at 40 V was as low as 5 nA. The breakdown field was higher than 150 kV/cm. These characteristics are comparable to those of state-of-the-art photoconductors such as LT-GaAs. We thus conclude that the furnace-annealed GaAs:As⁺ is an attractive alternative to LT-GaAs for ultrafast optoelectronic applications.

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