Acceptor Levels in GaSe:In Crystals Investigated by Deep-Level Transient Spectroscopy and Photoluminescence

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Optically and thermally detected deep levels in n-type Schottky and p + -n GaN diodes
Acceptor levels in GaSe:In crystals investigated by deep-level transient spectroscopy and photoluminescence

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Deep-acceptor levels associated with indium in indium-doped GaSe crystals have been measured. High-quality Schottky diodes of GaSe:In have been fabricated and characterized using current-voltage, capacitance-voltage, and deep-level transient spectroscopy (DLTS). Four DLTS peaks at 127, 160, 248, and 319 K, corresponding to 0.21, 0.22, 0.44, and 0.74 eV above the valence band, were well resolved and assigned to be an indium-on-gallium antisite (In_{Ga}), a gallium vacancy (V_{Ga}), an indium gallium vacancy complex (V_{Ga-In}), and a native defect associated with stacking fault or dislocation, respectively. Low-temperature photoluminescence (PL) spectroscopy measurements were performed on GaSe and GaSe:In crystals. The ground and the first excited states of the free exciton emissions were identified and the band-gap energies were determined. The results that the peak of exciton bound to acceptor (A',X) disappeared and the peak of donor-acceptor pair appeared in GaSe crystals have been consistent with the DLTS acceptor assignments.

I. INTRODUCTION

Gallium selenide (GaSe) has been studied for decades as a layered structure nonlinear optical material.1 Recently, it has become a material of choice for terahertz wave generation and detection.2–4 The electrical and optical properties of GaSe doped with different elements could differ significantly compared to the properties of unintentionally doped GaSe. The identification of the donor and acceptor levels between the valence and conduction bands of GaSe is important in view of its further applications in photoelectronic devices, such as room temperature radiation detectors.5

GaSe:In, GaSe doped with indium, is a promising semiconductor compound. It is well known that the mechanical softness of GaSe is the main flaw that limits GaSe applications. However, the microhardness of GaSe crystals can be improved noticeably after doping with indium.6,7 Not only the second harmonic generation efficiency, but also the hole mobility of GaSe:In as compared with GaSe have been improved. The highest hole mobility of GaSe:In at 300 K has been reported to be 1.08 × 10^3 cm^2/V s (see Table 2 in Ref. 6). The value is high enough to make GaSe:In potential candidates for room temperature radiation detectors although the dependences of hole mobility on temperature and indium concentration need further investigation to clarify the mechanism of hole scattering by the dopants and phonons.8 The carrier mobility-lifetime product is an important figure of merit in the evaluation of x-ray and gamma-ray semiconductor detectors; the carrier lifetime is determined by the trap concentrations and the capture cross sections.

Deep-level transient spectroscopy (DLTS) and low-temperature photoluminescence (PL) techniques are two sensitive techniques to identify defects. Acceptor levels in undoped p-GaSe (Ref. 9) and GaSe doped with elements, such as Sn,9 Cd,10 Mn,11,12 Cu,13,14 Er,15 N,16 and Co,17 have been investigated. However, to our knowledge, the acceptor levels of GaSe:In have not been yet reported. In this article, we have studied the acceptor levels of GaSe:In employing DLTS and PL techniques. Four well resolved acceptor levels have been observed in the DLTS spectrum, two of them associated with indium. The PL measurements are consistent with the DLTS assignments.

II. EXPERIMENT

GaSe crystals and GaSe:In crystals doped with 1000 ppm indium were grown in the EIC lab by the temperature gradient solution growth method. The typical size of a GaSe:In sample is 7.0 × 5.0 × 0.5 mm^3. For DLTS measurements, samples were cleaned in acetone using an ultrasonic bath. After drying in the air, they were cleaved and Au metal spots with diameter of 1.1 mm were sputtered using 50 W radio frequency power on the fresh surfaces to create Schottky contacts;18 high purity indium were then soldered to the other surfaces to form Ohmic contacts. The crystallographic c axis is perpendicular to the contact surfaces [(001) plane].
Current-voltage, capacitance-voltage, and DLTS measurements were carried out on the Schottky diodes to analyze their characteristics. The fully automated LABVIEW controlled DLTS system was designed by SEMETROL. DLTS capacitance was measured at 1 MHz using a Boonton 7200 capacitance meter. A SiC sample was used as a standard. The temperature deviation tolerance at each temperature point was less than 1 K.

For the low-temperature PL measurements, the samples were cooled down to about 9 K using an APD Cryogenic Inc. dual HC-4MK I helium compressors. The crystals were illuminated with the 488 nm line of an ILT 5500A air-cooled argon-ion laser with density of 2.0 W/cm². The photoluminescence spectra were detected using a SPEX 1877D Triplemate spectrometer in conjunction with a liquid nitrogen cooled charge coupled device (CCD) detector. A 0.1 mm slit and a grating with 300 grooves/mm were employed for the spectrometer. The spectral resolution was 1 cm⁻¹.

II. RESULTS AND DISCUSSION

Figure 1 shows the current-voltage characteristics and the plot of 1/C² versus voltage for a GaSe:In Schottky diode measured at room temperature. The plot of 1/C² versus voltage deviates from a straight line when the applied bias is larger than 1.5 V, which may indicate nonuniformity of indium doping. However, one can see that excellent rectification is obtained. The diode shows forward direction when a negative bias is applied to the Au contact, the ideality factor of the diode is 1.3. The formation of a high-quality Schottky diode is not only necessary for DLTS measurements, but also useful to make an electronic device such as room-temperature radiation detector, since a low leakage current is useful to make an electronic device such as room-diode is not only necessary for DLTS measurements, but also of the diode is 1.3. The formation of a high-quality Schottky diode measured at room temperature. The In contact was grounded. The diameter of the Au contact was 1.1 mm.

DLTS spectrum of GaSe:In is shown in Fig. 2. The filling pulse was 0 V, the measurement bias was 1 V, and the rate window was 29/s. The DLTS spectrum was generated from the difference in capacitance at two points during the emission transient as a function of temperature. The emission rate was extracted from a fit of the capacitance transient at each temperature as follows:

\[ C(t) = C_0 + \Delta C \exp(-t/\tau_p), \]  

where \( C_0 \) is the capacitance prior to the filling pulse, \( \Delta C \) represents the difference between the capacitance at the beginning and at the end of the filling pulse, and \( \tau_p \) is the hole emission rate. Four well-defined peaks, labeled as A, B, C, and D, were observed. The peaks B and D have been reported in undoped p-GaSe single crystals using photoinduced current transient spectroscopy (PICTS). In the area of peak C, a broad peak has been reported in GaSe samples doped with Cd, Mn, and Cu, and recently, two peaks close to peaks C and D have been found in a GaSe:Er sample.

When the holes were emitted to the maximum of the valence band, the dependence of hole emission rate \( \tau_p \) on the temperature was given as follows:

\[ \tau_p = (\sigma_p \nu_p N_v) \exp(-E_v/kT), \]  

where \( \sigma_p \) is the hole capture cross section, \( \nu_p \) the hole average thermal velocity, \( N_v \) the density of states of the valence band, \( E_v \) the thermal activation energy, \( k \) the Boltzmann constant, and \( T \) the peak temperature. If one assumes that \( \nu_p \) varies as \( T^{3/2} \) and \( N_v \) as \( T^{1/2} \), the activation energy and capture cross section of the trap levels can be determined from the slope and intercept of Arrhenius plot of \( \ln(T^{3/2}/\tau_p) \) versus \( 1/kT \). Furthermore, the concentration of a trap can be estimated from the peak height of the DLTS rate window spectrum as

\[ N_t = 2N_v(\Delta C/C_0). \]  

Figure 3 shows an Arrhenius plot for traps in GaSe:In. The trap energies capture cross sections, and assignments are listed in Table I. Since peaks B and D have been reported in the same temperature region for undoped GaSe, and the...
corresponding activation energy and capture cross section are comparable, it is reasonable to believe that peak B is due to a gallium vacancy and peak D to a stacking fault or dislocation, as suggested in Ref. 9. Peaks A and C were absent in undoped GaSe, so they are attributed to defects associated with indium. At region of peak C, a broad peak was observed in GaSe: M (M = Cd, Mn, Cu), and its intensity was a function of the concentration of M; therefore, it was attributed to a defect complex associated with M. It should be emphasised that peak A has never been reported in doped GaSe. Indium and gallium belong to the same IIIA family and because the activation energies of the acceptor at peaks A and B are almost same, it is reasonable to believe that peak A is due to an occupation of a Ga vacancy with an indium atom, and formation of a antisite InGa. Peak C cannot associate with an indium interstitial In, since like a gallium interstitial, the In may serve as a donor, instead of an acceptor. Therefore, we assign tentatively the acceptor at peak C to a VGa-In complex.

Low-temperature photoluminescence spectra of GaSe and GaSe:In crystals are shown in Fig. 4. The predominant peak in GaSe PL spectrum was attributed to the exciton bound to an acceptor (A0,X); the ground state free exciton (X)0, one longitudinal optical phonon replica of the indirect free excitonic recombination (IFE), and the radiative recombination of indirect excitons bound to deep neutral acceptor center (IBE) were also identified. It is interesting to see that the (A0,X) peak almost disappears in GaSe:In, while a strong broad peak associated with indium appears. The broad peak is attributed to a donor-acceptor pair (DAP) transition mainly, and to a free electron bound to neutral acceptor transition partly. It is generally accepted that the major acceptors in undoped GaSe are gallium vacancies VGa. Doped with indium, most of the gallium vacancies are occupied by indium atoms, and the InGa defects may be nonradiative centers. Therefore, the (A0,X) peak of GaSe:In decreases significantly while the (X)0,1 peak becomes dominate compared with those of GaSe. It is worth noting that the acceptor level C observed by DLTS is deeper than that measured from DAP.

The GaSe and GaSe:In samples used in the research have a high level of crystallinity. Except the ground-state free excitons observed in GaSe and GaSe:In crystals at temperatures varied from 9 to 300 K, first-excited-state free excitons (X)1,2 in GaSe crystal were successfully observed in the temperature range of 20–100 K. Figure 5 shows temperature dependence of the free exciton peak energies of GaSe and GaSe:In crystals. From the position of (X)0,1 and (X)2, one can determine the exciton binding energy (Rydberg energy R) and the band-gap energies at different temperatures, En(T), using the relation

$$E_n(T) = E_0(T) - R/n^2,$$

where En(T) is the peak position of the nth free exciton. The value of R is obtained as 18.6 meV at 20 K. One can see that the band gaps of undoped GaSe and GaSe doped with 1000 ppm indium are almost the same.

In summary, we have fabricated high-quality Schottky diodes and identified new acceptor levels associated with indium by deep-level transient spectroscopy. The assignments of new acceptor levels are consistent with the low-

![Fig. 3](image1.png)  
**FIG. 3.** Arhenius plot for traps in GaSe:In. The trap energies capture cross sections, assignments and concentration are listed in Table I.

![Fig. 4](image2.png)  
**FIG. 4.** Low-temperature (9 K) photoluminescence spectra of GaSe and GaSe:In crystals.

**TABLE I.** The acceptor types, energies, capture cross sections, and concentrations determined by DLTS for the GaSe:In crystals.

<table>
<thead>
<tr>
<th>Peak</th>
<th>Acceptor type</th>
<th>Activation energy (eV)</th>
<th>Capture cross section (cm²)</th>
<th>Density (cm⁻³)</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>InGa</td>
<td>0.21</td>
<td>3.7 × 10⁻¹⁵</td>
<td>3.0 × 10¹⁵</td>
</tr>
<tr>
<td>B</td>
<td>VGa</td>
<td>0.22</td>
<td>5.3 × 10⁻¹⁷</td>
<td>1.2 × 10¹⁵</td>
</tr>
<tr>
<td>C</td>
<td>VGa-In</td>
<td>0.44</td>
<td>2.2 × 10⁻¹⁵</td>
<td>8.3 × 10¹⁴</td>
</tr>
<tr>
<td>D</td>
<td>Stacking fault</td>
<td>0.74</td>
<td>1.1 × 10⁻¹²</td>
<td>6.1 × 10¹⁴</td>
</tr>
</tbody>
</table>
temperature photoluminescence results. The identification of new acceptor levels and the fabrication of Schottky diodes will be helpful to the development of electronic device based on GaSe:In crystals.

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FIG. 5. Temperature dependences of free exciton peak energies of GaSe and GaSe:In crystals.

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