

- <sup>7</sup>J. Gelbwachs and T. S. Hartwick, IEEE J. Quantum Electron. QE-11, 52 (1975).
- <sup>8</sup>T. W. Hansch and A. L. Schawlow, Opt. Commun. 13, 68 (1975).
- <sup>9</sup>M. C. Mery, D. Silhouette, and J. Conard, C.R. Acad. Sci. B 275, 693 (1972).
- <sup>10</sup>Purchased from Tyco Labs, Saphikon Division, Waltham, Mass.
- <sup>11</sup>C. A. Elyard and H. Rawson, in *Advances in Glass Technology* (Plenum, New York, 1962), p. 270.
- <sup>12</sup>A. R. Gordon, J. Chem. Phys. 4, 100 (1936).
- <sup>13</sup>E. A. J. Marcatili and R. A. Schmelzter, Bell. Syst. Tech. J. 43, 1783 (1964).
- <sup>14</sup>P. W. Smith and T. Hänsch, Phys. Rev. Lett. 26, 740 (1971).
- <sup>15</sup>The 3S ground state of sodium is split by the hyperfine interaction into two levels, with  $F=2$  and  $F=1$ , separated by 1.77 GHz. For the purposes of this letter, we define the resonant frequency  $\nu_0$  to correspond to the  $3S_{1/2}(F=2) \rightarrow 3P_{3/2}$  transition. The  $3S_{1/2}(F=1) \rightarrow 3P_{3/2}$  transition then corresponds to  $\Delta\nu = \nu - \nu_0 = +1.77$  GHz.
- <sup>16</sup>For example, see *Scientific Foundations of Vacuum Technology*, 2nd ed., edited by Saul Dushman (Wiley, New York, 1962).
- <sup>17</sup>This seems to justify our assumption of velocity cross relaxation.
- <sup>18</sup>B. P. Kibble, G. Copley, and L. Krause, Phys. Rev. 159, 11 (1967).

## Application of scanning electron microscopy to determination of surface recombination velocity: GaAs

L. Jastrzebski,\* J. Lagowski,\* and H. C. Gatos

Department of Materials Science and Engineering, and Department of Electrical Engineering and Computer Sciences, Massachusetts Institute of Technology, Cambridge, Massachusetts 02139  
(Received 16 June 1975; in final form 31 July 1975)

A method is reported for the determination of the surface recombination velocity by scanning electron microscopy; this method is based on an established relationship between the effective diffusion length of the minority carriers, the penetration depth of the electron beam, and the surface recombination velocity. Values of surface recombination velocity, up to about  $2.5 \times 10^6$  cm/sec, were determined in *n*-type GaAs with a bulk minority-carrier lifetime of the order of  $10^{-8}$ – $10^{-10}$  sec; in GaAs, with carrier concentrations exceeding  $10^{18}$  cm $^{-3}$ , recombination velocity of about  $3 \times 10^6$  cm/sec represents a saturation value.

PACS numbers: 72.20.K, 72.80.E, 73.30.

Employing virtually "point" excitation of excess carriers by an electron beam in conjunction with high-resolution scanning, a number of investigators have achieved the determination of the diffusion length of minority carriers having extremely short lifetimes<sup>1-5</sup>. Excess minority carriers generated at the desired distance from a Schottky barrier (or *p-n* junction) diffuse to the barrier and produce an electric current; for a constant accelerating voltage applied to the electron beam, the current  $I$  is a function of the distance  $x$  between the impinging beam and the barrier, i.e.,  $I \approx \exp(-x/L_{eff})$ , where  $L_{eff}$  is the effective diffusion length of the minority carriers. It has been recognized that with decreasing accelerating voltage (i.e., decreasing penetration depth of the electron beam)  $L_{eff}$  decreases, apparently due to the increasing influence of surface recombination.<sup>3</sup>

In the present investigation the effect of surface recombination on the effective diffusion length was quantitatively analyzed. Single-crystal samples of *n*-type GaAs, with carrier concentrations  $6 \times 10^{16}$ ,  $2.4 \times 10^{17}$ , and  $1.8 \times 10^{18}$  cm $^{-3}$ , were used. Schottky barriers were obtained by evaporating gold onto GaAs "real" surfaces (prepared by chemical etching, chemical polishing, or by cleaving). The experiments were carried out in the standard configuration for diffusion length measurements<sup>1,6</sup> (see Fig. 1), with no bias applied on the barrier contact.

For a given (constant) accelerating voltage, a typical linear region of  $\ln I$  vs  $x$  was obtained,<sup>1,6</sup> from which the effective diffusion length was evaluated. The measured current changes ranged over about 3 orders of magnitude. From a family of  $\ln I$  vs  $x$  plots measured at different accelerating voltage values (ranging from 5 to 30 kV) effective diffusion length was determined as a function of the accelerating voltage and, thus, as a function of the penetration depth of the electron beam. Typical results obtained on *n*-type samples with a carrier concentration of  $2.4 \times 10^{17}$  cm $^{-3}$  are shown in Fig. 1. It is seen that with increasing accelerating voltage  $V$  from 5 to 30 kV,  $L_{eff}$  increases by a factor of about 3 and then approaches the saturation value. These results reflect the increasing influence of surface recombination on  $L_{eff}$  with decreasing penetration depth  $\xi$  of the electron beam; it should be noted that  $\xi \sim V^{1/2}$ .<sup>5,7</sup>

The effect of surface recombination on the effective diffusion length will now be considered on a quantitative basis. In the first order of approximation the surface recombination velocity will be taken as a constant parameter, independent of the penetration depth of the electron beam. In this respect, it should be noted that, in general, the surface recombination velocity depends on the surface potential which varies with the excitation of excess carriers. In GaAs, however, the surface barrier is associated with a depletion region, and it essentially vanishes upon relatively low electron-hole

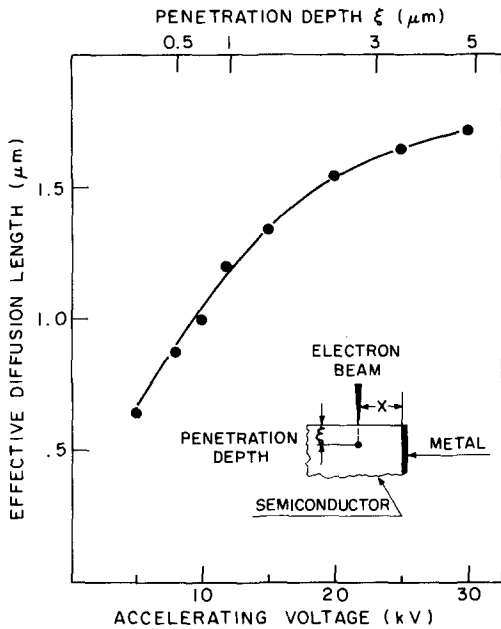


FIG. 1. Experimental configuration and typical results showing the dependence of the measured diffusion length on the accelerating voltage of the electron beam or penetration depth.

excitation.<sup>8</sup> Thus, in the present case of electron beam excitation—i.e., high level of excitation—there is virtually no surface barrier present.

On the basis of previous theoretical treatments of electron-hole pair excitation by an electron beam,<sup>5</sup> the electron beam is approximated by a steady-state point source  $G\delta(x, y, z-\xi)$ , with strength  $G$ , located at a distance  $\xi$  beneath the surface ( $z=0$ ) of an  $n$ -type semi-infinite semiconductor. The steady-state concentration of excess minority carriers,  $\Delta p(x, y, z)$ , can be obtained from the continuity equation with the surface boundary condition

$$D \frac{\partial \Delta p}{\partial x} \bigg|_{x=0} = S \Delta p \bigg|_{x=0}, \quad (1)$$

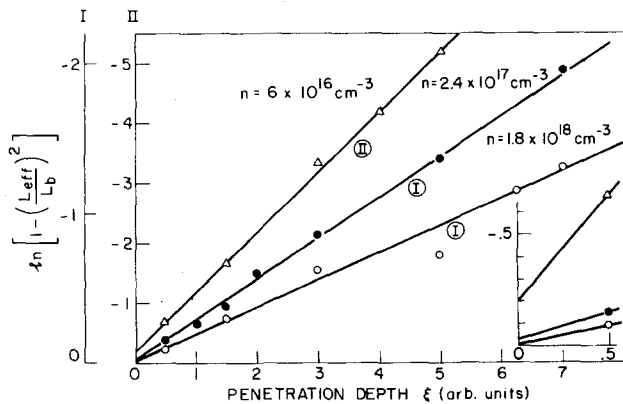


FIG. 2.  $\ln[1 - (L_{eff}/L_b)^2]$  as a function of the penetration depth of the electron beam, in arbitrary units. Curve I corresponds to the expanded scale (I), curves II correspond to the scale (II).

TABLE I. Measured surface recombination velocities of GaAs single crystals together with calculated values of  $\tau_b$  and  $L_b$  and values of  $L_{eff}$  measured for an electron-beam accelerating voltage of 25 kV.

$n$ (cm <sup>-3</sup> )	$S$ (cm/sec)	$\tau_b$ (sec)	$L_b$ (μm)	$L_{eff}$ (25 kV) (μm)	Material
$6 \times 10^{16}$	$5 \times 10^5$	$4 \times 10^{-10}$	0.5	0.45	boat grown, undoped
$2.4 \times 10^{17}$	$2 \times 10^6$	$5 \times 10^{-9}$	1.8	1.65	LPE
$1.8 \times 10^{18}$	$2.5 \times 10^6$	$1.5 \times 10^{-8}$	2.5	2.1	boat grown, Si doped

where  $D$  is the diffusion coefficient of the minority carriers and  $S$  is the surface recombination velocity. On the basis of Hackett's results<sup>5</sup> it can be readily shown that the total steady-state concentration of the excess holes,

$$\Delta p(\xi) \equiv \int_{-\infty}^{\infty} dx \int_{-\infty}^{\infty} dy \int_0^{\infty} dz \Delta p(x, y, z),$$

be reduced to

$$\Delta p(\xi) = (GL_b^2/D) \{1 - [s/(s+1)] \exp(-\xi/L_b)\}, \quad (2)$$

where  $L_b$  is the bulk diffusion length of the minority carriers and  $s = S\tau_b/L_b$ ;  $\tau_b$  is the bulk lifetime of the minority carriers.

Under steady-state conditions,  $\Delta p(\xi)$  can be related to the measured  $L_{eff}$  as follows:

$$\Delta p(\xi) = G\tau_{eff}, \quad (3)$$

$$L_{eff} = (D\tau_{eff})^{1/2},$$

where  $\tau_{eff}$  is the effective steady-state minority-carrier lifetime.

From Eqs. (2) and (3) one obtains

$$L_{eff}^2 = L_b^2 [1 - [s/(s+1)] \exp(-\xi/L_b)]. \quad (4)$$

According to Eq. (4), for  $\xi \leq L_b$ ,  $L_{eff}$  can be significantly reduced due to surface recombination.  $L_{eff}$  remains sensitive to  $S$  for an intermediate range of  $S$  values, depending on the values of  $\tau_b$  and  $L_b$ ; in GaAs this range is about  $10^4$ – $10^6$  cm/sec. For higher values of  $S$ ,  $L_{eff}$  approaches a saturation value

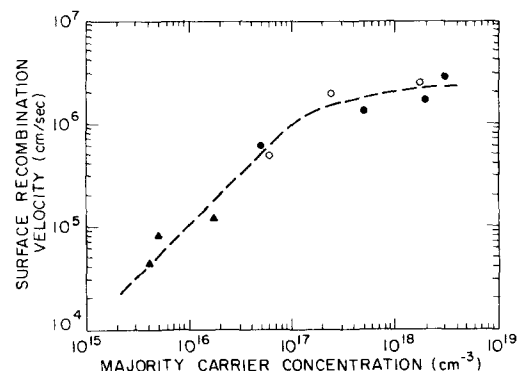


FIG. 3. Surface recombination velocity in GaAs vs majority-carrier concentration; ○, present results; ●, Ref. 7; ▲, Ref. 8.

$$L_{eff}^{sat} = L_b [1 - \exp(-\xi/L_b)]^{1/2}. \quad (5)$$

The experimental results will now be analyzed on the basis of Eq. (4). In Fig. 2,  $\ln[1 - (L_{eff}/L_b)^2]$  is plotted against  $\xi$  (or  $V^{1/2}$ ); consistent with Eq. (4) a straight line is obtained (using the least-squares method) for a given value of carrier concentration. From the intercept of the line with the ordinate one can determine  $s/(s+1)$ , and therefore the surface recombination velocity. The values of the surface recombination velocities obtained for the three GaAs crystals are shown in Table 1.

From Eq. (4) it is also possible to evaluate  $L_b$  (and  $\tau_b$ ). The calculated values of  $L_b$  and  $\tau_b$  are given in Table I together with the measured  $L_{eff}$  values for 25 kV. It should be noted that even for high values of accelerating voltage the measured  $L_{eff}$  is still noticeably reduced by surface recombination. The values of  $\tau_b$  cannot be satisfactorily related to the corresponding carrier concentrations; however, the GaAs crystals used in this study were grown by different techniques and at different laboratories, and thus they may contain different types and concentrations of active recombination centers.

The present surface recombination velocity together with earlier results obtained by other methods<sup>9,10</sup> are plotted in Fig. 3 as a function of carrier concentration. As seen in Fig. 3, the agreement between the present results and those previously reported is very satisfactory. It should be pointed out that all results shown in Fig. 3 were obtained at high excitation levels, and, thus, the uncertainty associated with possible effects of the excitation intensity on the measured values of surface recombination is minimized. It is further seen in Fig. 3 that the surface recombination velocity in GaAs increases with increasing carrier concentration and approaches a saturation value of about  $3 \times 10^6$  cm/sec for a carrier concentration of approximately  $10^{18}$  cm<sup>-3</sup>.

In summary, the scanning electron microscopy technique for determining the diffusion length of minority carriers in semiconductors was extended to the determination of surface recombination velocities. This new approach, based on the established quantitative relationship between the effective diffusion length and the penetration depth of the electron beam, is effectively applied to semiconductors with high surface recombination and short minority-carrier lifetime; standard methods for surface recombination measurements are not readily applicable to such materials.

The authors are grateful to the National Science Foundation for supporting the work; they are indebted to Professor A.G. Milnes for making available an LPE grown sample of GaAs and Ref. 1 prior to its publication; they also wish to acknowledge the skillful assistance of J. Adario in carrying out the SEM measurements.

\*On leave from Institute of Physics, Polish Academy of Sciences, Warsaw, Poland.

<sup>1</sup>A.M. Sekela, D.L. Feucht, and A.G. Milnes, in *Proceedings of the Symposium on GaAs and Related Compounds*, Deniville, France, 1974 (Institute of Physics, London, to be published).

<sup>2</sup>W.H. Hackett, Jr., R.H. Saul, R.W. Dixon, and G.W. Kammlot, *J. Appl. Phys.* **43**, 285 (1972).

<sup>3</sup>R.D. Ryan and J.E. Eberhardt, *Solid-State Electron.* **15**, 865 (1972).

<sup>4</sup>D.B. Wittry and D.F. Kyser, *J. Appl. Phys.* **36**, 1387 (1965).

<sup>5</sup>W.H. Hackett, *J. Appl. Phys.* **43**, 1649 (1972).

<sup>6</sup>C. van Opdorp, R.C. Peters, and M. Klerk, *Appl. Phys. Lett.* **24**, 125 (1974).

<sup>7</sup>J.S. Rao-Sahib and D.B. Wittry, *J. Appl. Phys.* **40**, 3745 (1969).

<sup>8</sup>J. Lagowski, I. Baltov, and H.C. Gatos, *Surf. Sci.* **40**, 216 (1973).

<sup>9</sup>D.B. Wittry, *J. Phys. Soc. Jpn. Suppl.* **21**, 312 (1966).

<sup>10</sup>N.L. Dymitriuk, V.I. Lyashenko, A.K. Tereshenko, and A.K. Spektor, *Phys. Status Solidi A* **20**, 53 (1973).

## Excitation of transversely excited CO<sub>2</sub> waveguide lasers

O. R. Wood II, P. W. Smith, C. R. Adams, and P. J. Maloney

*Bell Telephone Laboratories, Holmdel, New Jersey 07733*  
(Received 6 August 1975)

Using a preionization scheme based on the Malter effect, small-signal gains  $> 5\%/cm$  at  $10.6 \mu m$  have been produced in a 1-mm<sup>2</sup>-cross-section waveguide CO<sub>2</sub> amplifier at total operating pressures of 100–760 Torr. Comparisons are made between this preionization scheme and those using electron beams.

PACS numbers: 42.60.C, 52.80.H

A highly controllable transverse excitation technique for use with waveguide CO<sub>2</sub> lasers<sup>1,2</sup> has been developed. By applying this excitation technique to a 1-mm<sup>2</sup>-cross-section waveguide amplifier filled with CO<sub>2</sub>-N<sub>2</sub>-He mixtures, small-signal gains of peak value greater than

5%/cm on the P(20) line of CO<sub>2</sub> at  $10.6 \mu m$  have been produced at total operating pressures of 100–760 Torr. The performance achieved with this technique is comparable to previously published results using more complicated electron-beam preionization.<sup>3,4</sup> A 10-cm-long