

## Measurement of Base Transit Time and Minority Electron Mobility in GaAsSb-Base InP DHBTs

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GaAsSb/InP Type-II DHBTs are used extensively in a variety of instrument products from mmWave network analyzers to high performance oscilloscopes. The underlying DHBT IC technology requires a favorable combination of high current gain and bandwidth, low turn-on voltage, and high breakdown voltage [1] which are controlled mostly by epitaxial design. To achieve even higher performance DHBT ICs, the epi design must be further optimized. In this work, optimization of lattice matched GaAsSb bases is discussed. RF data on discrete DHBT devices and a methodology for extracting base transit time and electron mobility in the p+GaAsSb base are presented. The results of this work advance the design of the base region in p+GaAsSb base InP DHBTs, and the experimental method presented here can also be used to characterize the base transit time and electron mobility of any bipolar transistor.

A lot of seven wafers ("Lot A") was grown and processed identically. The epi structures used in "Lot A" all had lattice matched, p-doped, GaAsSb bases, but the base thickness was varied from  $W_B = 250\text{-}750 \text{ \AA}$ . Small-signal microwave measurements were performed on each wafer, and the emitter-collector delay time  $\tau_{EC} = 1/2\pi f_T$  was extracted from a single-pole approximation of  $h_{21}(f)$  for each device operating at constant  $V_{CB}$  and multiple collector currents. Using a single-pole approximation of  $h_{21}(f)$  of a small-signal T-model, the total delay time  $\tau_{EC}$  is estimated as a sum of internal delay times  $\tau_{EC} = \tau_T + (C_{JE} + C_{JC})\eta kT/qI_C$ , with  $\tau_T = \tau_B + \tau_C + (R_C + R_E)C_{JC}$ . The delay time  $\tau_T$  was extracted for each epi design by extrapolating  $\tau_{EC}$  vs.  $1/I_C$  to  $1/I_C = 0$ , as shown in Fig. 1 [2]. These delay times  $\tau_T$  followed a quadratic relationship versus base thickness  $W_B$ , as shown in Fig. 2. This quadratic dependence can be understood because the delays  $\tau_C$  and  $(R_C + R_E)C_{JC}$  remain the same for all the wafers, but the base delay varies with base thickness as  $\tau_B = W_B^2/2D_{nB} + W_B/V_{exit}$  [3]. The electron diffusion coefficient in the base  $D_n$  was then extracted from the RF data (see Fig. 2), producing a value for  $D_n$  of  $61 \text{ cm}^2/\text{s}$ . This electron diffusion coefficient together with the Einstein relation and our estimated junction temperature  $T_j$  were then used to calculate the minority electron mobility in the base, which gave a mobility value of  $\mu_n = 1970 \text{ cm}^2/(\text{V}\cdot\text{s})$ . For comparison, the value quoted for p-InGaAs at the same doping value was about  $1500 \text{ cm}^2/(\text{V}\cdot\text{s})$  [4].

A second lot ("Lot B") was grown and processed in the same way with a variety of base doping levels. All "Lot B" wafers were lattice matched GaAsSb with the same base thickness of  $336 \text{ \AA}$  and with base doping values of  $1.04, 1.30, \text{ and } 1.60 \times 10^{20} \text{ cm}^{-3}$ . The base transit times were extracted for each wafer (by finding  $t_T$  as before, and estimating the values of  $C_{JC}$ ,  $R_C$ ,  $R_E$ , and  $\tau_C$  from microwave measurements), and the electron diffusion coefficients  $D_n$  were calculated. The plot in Fig. 3 of  $D_n$  versus base doping for these bases shows a significant decrease in  $D_n$  for doping levels  $N_a$  above  $0.6 \times 10^{20} \text{ cm}^{-3}$  and approximately constant  $D_n$  value for higher values of  $N_a$ . This may be due to a strong dependence on doping concentration above  $6 \times 10^{19} \text{ cm}^{-3}$ , but growth reproducibility is also being investigated as a potential cause of this significant decrease.

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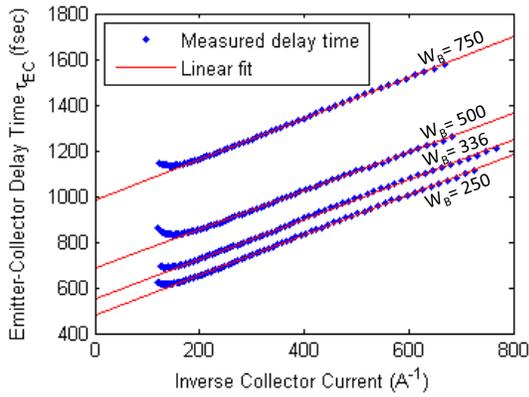


Fig. 1. Extrapolation of  $\tau_{EC} = 1/2\pi f_T$  to determine the delay time  $\tau_T$  according to the method in ref [2].

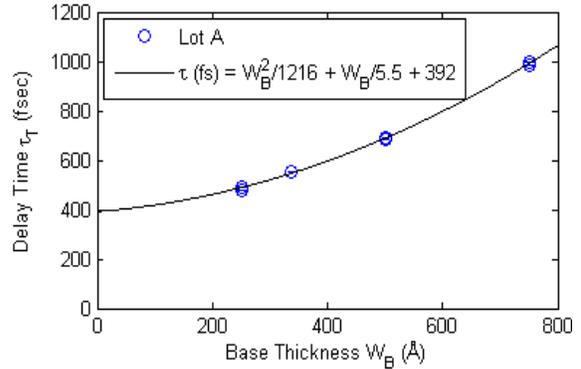


Fig. 2. Extrapolated delay time  $\tau_T$  versus base thickness  $W_B$ , fitted to the quadratic function that is expected from refs [2] and [3].

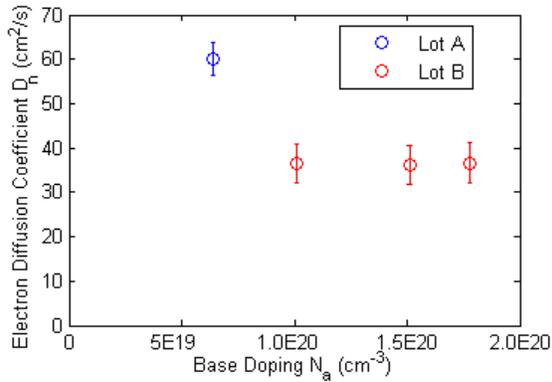


Fig. 3. Electron diffusion coefficient  $D_n$  versus base doping concentration for wafers in Lot A and Lot B