

$$\begin{aligned}
& +K(z) \left\} + \frac{1}{2\rho(z)} \frac{\partial}{\partial z} \left\{ \frac{E(z)}{1+\sigma(z)} \left[\frac{\partial}{\partial z} \int_0^g \int_0^g u_{1y}(x, y, z, \tau) d\tau d\vartheta + \right. \right. \\
& \left. \left. + \frac{\partial}{\partial y} \int_0^g \int_0^g u_{1z}(x, y, z, \tau) d\tau d\vartheta \right] - K(z) \int_0^g \int_0^g T(x, y, z, \tau) d\tau d\vartheta \right\} \times \\
& \times \frac{\beta(z)}{\rho(z)} - \frac{\partial^2}{\partial y \partial z} \int_0^g \int_0^g u_{1y}(x, y, z, \tau) d\tau d\vartheta \left\{ \frac{E(z)}{6[1+\sigma(z)]} - K(z) \right\} \times \\
& \times \frac{1}{\rho(z)} - \frac{E(z)}{2\rho(z)[1+\sigma(z)]} \left[\frac{\partial^2}{\partial x^2} \int_0^g \int_0^g u_{1x}(x, y, z, \tau) d\tau d\vartheta + \right. \\
& \left. + \frac{\partial^2}{\partial x \partial y} \int_0^g \int_0^g u_{1x}(x, y, z, \tau) d\tau d\vartheta \right] - K(z) \int_0^g \int_0^g T(x, y, z, \tau) d\tau d\vartheta \times \\
& \times \frac{\beta(z)}{\rho(z)} - \frac{1}{\rho(z)} \frac{\partial^2}{\partial y^2} \int_0^g \int_0^g u_{1x}(x, y, z, \tau) d\tau d\vartheta \left\{ \frac{5E(z)}{12[1+\sigma(z)]} + \right. \\
& \left. + K(z) \right\} - \frac{K(z)}{\rho(z)} \frac{\partial^2}{\partial x \partial y} \int_0^g \int_0^g u_{1y}(x, y, z, \tau) d\tau d\vartheta - \frac{\partial}{\partial z} \left\{ \frac{E(z)}{1+\sigma(z)} \times \right. \\
& \left. \times \left[\frac{\partial}{\partial z} \int_0^g \int_0^g u_{1y}(x, y, z, \tau) d\tau d\vartheta + \frac{\partial}{\partial y} \int_0^g \int_0^g u_{1z}(x, y, z, \tau) d\tau d\vartheta \right] \right\} \times \\
& \times \frac{1}{2\rho(z)} + \frac{1}{\rho(z)} \frac{\partial^2}{\partial y \partial z} \int_0^g \int_0^g u_{1y}(x, y, z, \tau) d\tau d\vartheta \left\{ \frac{E(z)}{6[1+\sigma(z)]} - \right. \\
& \left. - K(z) \right\} + u_{0y} \\
& u_z(x, y, z, t) = \frac{E(z)}{2[1+\sigma(z)]} \left[\frac{\partial^2}{\partial x^2} \int_0^g \int_0^g u_{1z}(x, y, z, \tau) d\tau d\vartheta + \right. \\
& \left. + \frac{\partial^2}{\partial y^2} \int_0^g \int_0^g u_{1z}(x, y, z, \tau) d\tau d\vartheta + \frac{\partial^2}{\partial x \partial z} \int_0^g \int_0^g u_{1x}(x, y, z, \tau) d\tau d\vartheta + \right. \\
& \left. + \frac{\partial^2}{\partial y \partial z} \int_0^g \int_0^g u_{1y}(x, y, z, \tau) d\tau d\vartheta \right] \frac{1}{\rho(z)} + \frac{1}{\rho(z)} \frac{\partial}{\partial z} \left\{ K(z) \times \right. \\
& \times \left[\frac{\partial}{\partial x} \int_0^g \int_0^g u_{1x}(x, y, z, \tau) d\tau d\vartheta + \frac{\partial}{\partial y} \int_0^g \int_0^g u_{1x}(x, y, z, \tau) d\tau d\vartheta + \right. \\
& \left. + \frac{\partial}{\partial z} \int_0^g \int_0^g u_{1x}(x, y, z, \tau) d\tau d\vartheta \right] \left\} + \frac{1}{6\rho(z)} \frac{\partial}{\partial z} \left\{ \frac{E(z)}{1+\sigma(z)} \times \right. \\
& \times \left[6 \frac{\partial}{\partial z} \int_0^g \int_0^g u_{1z}(x, y, z, \tau) d\tau d\vartheta - \frac{\partial}{\partial x} \int_0^g \int_0^g u_{1x}(x, y, z, \tau) d\tau d\vartheta - \right. \\
& \left. - \frac{\partial}{\partial y} \int_0^g \int_0^g u_{1y}(x, y, z, \tau) d\tau d\vartheta - \frac{\partial}{\partial z} \int_0^g \int_0^g u_{1z}(x, y, z, \tau) d\tau d\vartheta \right] \right\} - \\
& - K(z) \frac{\beta(z)}{\rho(z)} \frac{\partial}{\partial z} \int_0^g \int_0^g T(x, y, z, \tau) d\tau d\vartheta + u_{0z}.
\end{aligned}$$

Framework this paper we determine concentration of dopant, concentrations of radiation defects and components of displacement vector by using the second-order approximation framework method of averaging of function corrections. This approximation is usually enough good approximation to make qualitative analysis and to obtain some quantitative results. All obtained results have been checked by comparison with results of numerical simulations.

3. DISCUSSION

In this section we analyzed dynamics of redistributions of dopant and radiation defects during annealing and under

influence of mismatch-induced stress and modification of porosity. Typical distributions of concentrations of dopant in heterostructures are presented on Figures 2 and 3 in direction, which is perpendicular to interface between epitaxial layer substrate. Figure 2 corresponds to diffusion type of doping. Figure 3 corresponds to diffusion type of doping. Increasing of number of curve corresponds to increasing of difference between values of dopant diffusion coefficient in layers of heterostructure under condition, when value of dopant diffusion coefficient in epitaxial layer is larger, than value of dopant diffusion coefficient in substrate. These distributions have been calculated for the case, when value of dopant diffusion coefficient in doped area is larger, than in nearest areas. Curves 1 and 3 on Figure 3 corresponds to annealing time $\Theta = 0.0048(L_x^2 + L_y^2 + L_z^2)/D_0$. Curves 2 and 4 on Figure 3 corresponds to annealing time $\Theta = 0.0057(L_x^2 + L_y^2 + L_z^2)/D_0$. Curves 1 and 2 corresponds to homogenous sample. Curves 3 and 4 corresponds to heterostructure under condition, when value of dopant diffusion coefficient in epitaxial layer is larger, than value of dopant diffusion coefficient in substrate. Annealing time of dopant, which corresponds to Figure 2, is equal to $\Theta = 0.005(L_x^2 + L_y^2 + L_z^2)/D_0$. The figures show, that inhomogeneity of heterostructure gives us possibility to increase compactness of concentrations of dopants and at the same time to increase homogeneity of dopant distribution in doped part of epitaxial layer. However, this manufacturing approach of biopolar transistor cannot effectively anneal the dopants and/or radiation defects. The annealing process should be optimized for the following reasons. If annealing time is small, the dopant did not achieve any interfaces between materials of heterostructure. In this situation one cannot find any modifications of distribution of concentration of dopant. If annealing time is large, distribution of concentration of dopant is too homogenous. We optimize annealing time framework recently introduces approach [29-37]. Framework this criterion we approximate real distribution of concentration of dopant by step-wise function (see Figures 4 and 5). These figures show spatial distributions of dopant in heterostructure after infusion (for Figure 4) or implantation of dopant (for Figure 5). Curve 1 is idealized distribution of dopant. Curves 2-4 are real distributions of dopant for different values of annealing time. Increasing of number of curve corresponds to increasing of annealing time. Farther we determine optimal values of annealing time by minimization of the following mean-squared error.

$$U = \frac{1}{L_x L_y L_z} \int_0^{L_x} \int_0^{L_y} \int_0^{L_z} [C(x, y, z, \Theta) - \psi(x, y, z)]^2 dz dy dx \quad (22)$$

where, $\psi(x, y, z)$ is the approximation function. Dependences of optimal values of annealing time on parameters are presented on Figures 6 and 7 for diffusion and ion types of doping, respectively. Curve 1 is the dependence of dimensionless optimal annealing time on the relation a/L and $\xi = \gamma = 0$ for equal to each other values of dopant diffusion coefficient in all parts of heterostructure. Curve 2 is the dependence of dimensionless optimal annealing time on value of parameter ε for $a/L=1/2$ and $\xi = \gamma = 0$. Curve 3 is the dependence of dimensionless optimal annealing time on value of parameter ξ for $a/L=1/2$ and $\varepsilon = \gamma = 0$. Curve 4 is the dependence of dimensionless optimal annealing time on value of parameter γ for $a/L=1/2$ and $\varepsilon = \xi = 0$. It should be noted, that

it is necessary to anneal radiation defects after ion implantation. One could find spreading of concentration of distribution of dopant during this annealing. In the ideal case distribution of dopant achieves appropriate interfaces between materials of heterostructure during annealing of radiation defects. If dopant did not achieve any interfaces during annealing of radiation defects, it is practically to additionally anneal the dopant. In this situation optimal value of additional annealing time of implanted dopant is smaller, than annealing time of infused dopant.

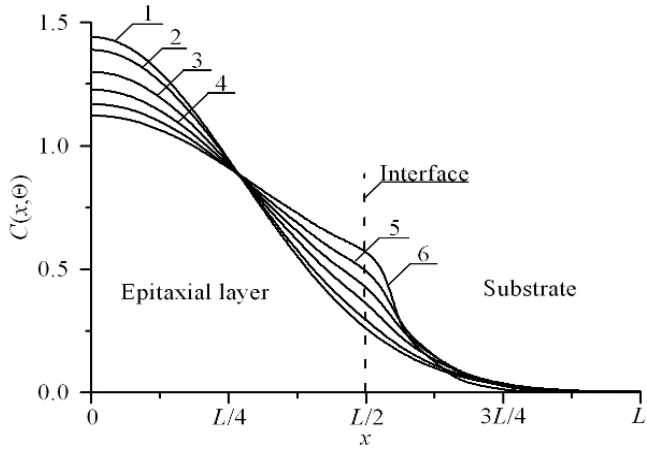


Figure 2. Distributions of concentration of infused dopant

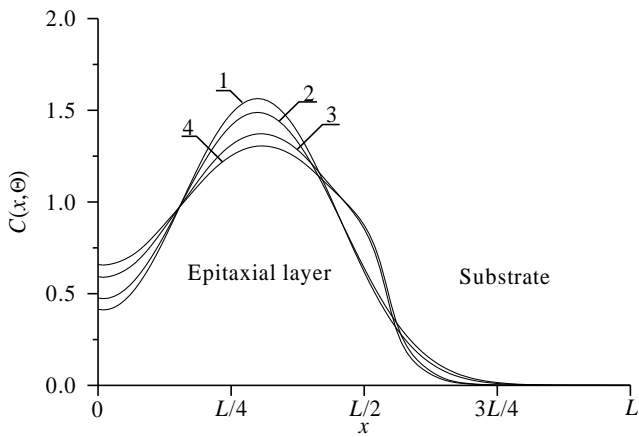


Figure 3. Distributions of concentration of implanted dopant

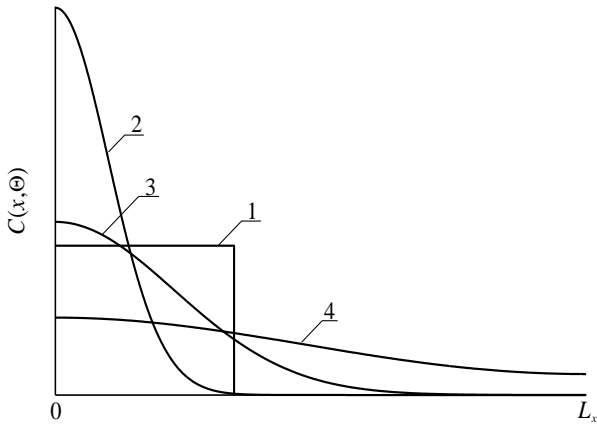


Figure 4. Typical spatial distributions of dopant in heterostructure after dopant infusion

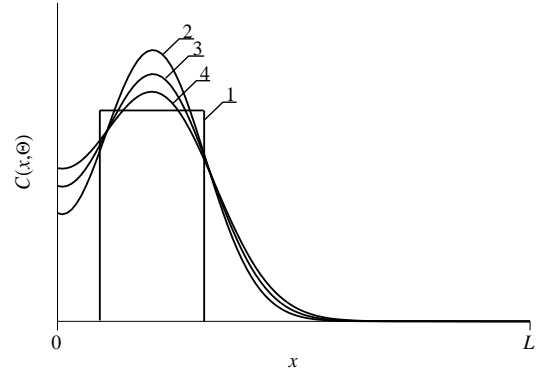


Figure 5. Typical spatial distributions of dopant in heterostructure after ion implantation

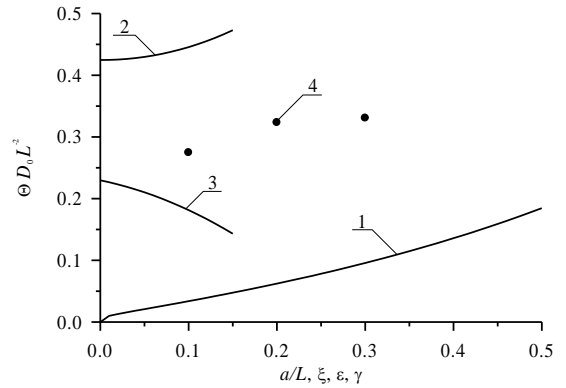


Figure 6. Dependences of dimensionless optimal annealing time for doping by diffusion

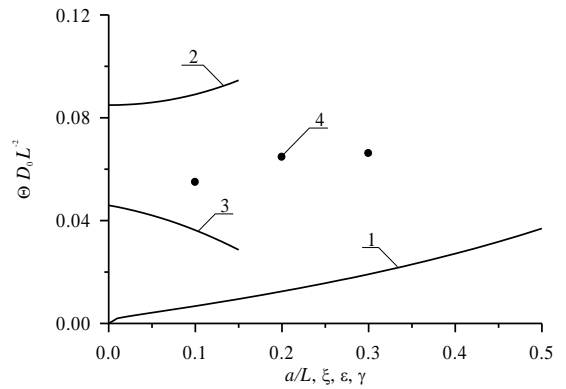


Figure 7. Dependences of dimensionless optimal annealing time for doping by ion implantation

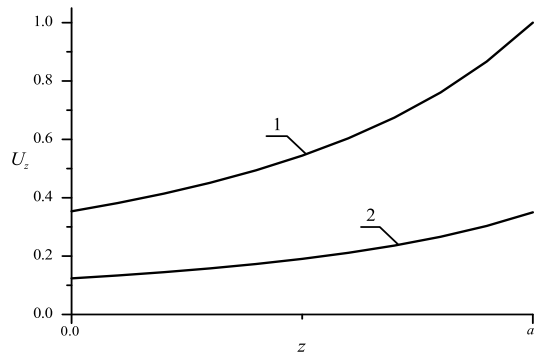


Figure 8. Normalized dependences of component u_z of displacement vector on coordinate z for nonporous (curve 1) and porous (curve 2) epitaxial layers

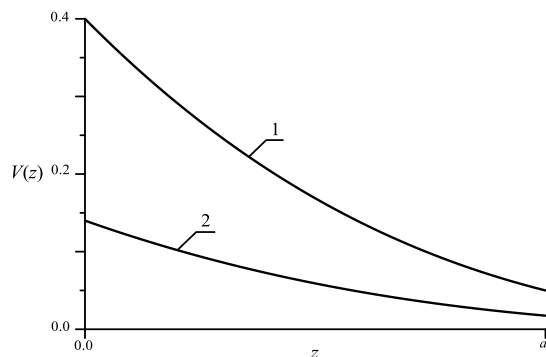


Figure 9. Normalized dependences of vacancy concentrations on coordinate z in unstressed (curve 1) and stressed (curve 2) epitaxial layers

Farther we analyzed influence of relaxation of mechanical stress on distribution of dopant in doped areas of heterostructure. Under following condition $\varepsilon_0 < 0$ one can find compression of distribution of concentration of dopant near interface between materials of heterostructure. Contrary (at $\varepsilon_0 > 0$) one can find spreading of distribution of concentration of dopant in this area. This changing of distribution of concentration of dopant could be at least partially compensated by using laser annealing [37]. This type of annealing gives us possibility to accelerate diffusion of dopant and other processes in annealed area due to inhomogenous distribution of temperature and Arrhenius law. Accounting relaxation of mismatch-induced stress in heterostructure could leads to changing of optimal values of annealing time. At the same time modification of porosity gives us possibility to decrease value of mechanical stress. On the one hand mismatch-induced stress could be used to increase density of elements of integrated circuits. On the other hand, it could leads to generation dislocations of the discrepancy. Figures 8 and 9 show distributions of concentration of vacancies in porous materials and component of displacement vector, which is perpendicular to interface between layers of heterostructure.

4. CONCLUSION

In this paper we model redistribution of infused and implanted dopants with account relaxation mismatch-induced stress during manufacturing field-effect heterotransistors framework a two-level current-mode logic gates in a multiplexer. We obtain, that using difference between materials of heterostructure and optimization of annealing of dopant and/or radiation defects gives a possibility to decrease dimensions of transistors and to increase their density. We obtain, that using ion implantation gives a possibility to decrease mismatch-induced stress. At the same time, it is necessary to choose materials with higher charge carrier motility and minimal mismatch-induced stress. Minimization of mismatch-induced stress gives a possibility to use diffusion type of doping without radiation damage of materials of heterostructure. Increasing of charge carrier motility gives a possibility to accelerate of transport of charge carriers. We also introduce an analytical approach to model diffusion and ion types of doping with account concurrent changing of parameters in space and time. At the same time the approach gives us possibility to take into account nonlinearity of considered processes.

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