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Eduard Kuhn

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Weierstrass Institute
Mohrenstr. 39
10117 Berlin
Germany
E-Mail: eduard.kuhn@wias-berlin.de

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Edited by
Weierstraß-Institut für Angewandte Analysis und Stochastik (WIAS)
Leibniz-Institut im Forschungsverbund Berlin e. V.
Mohrenstraße 39
10117 Berlin
Germany

Fax: +49 30 20372-303
E-Mail: preprint@wias-berlin.de
World Wide Web: <http://www.wias-berlin.de/>

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Abstract

In this publication a method to simulate the mode dynamics in broad-ridge laser diode is presented. These devices exhibit rich lateral mode dynamics in addition to longitudinal mode dynamics observed in narrow-ridge laser diodes. The mode dynamics are strongly influenced by higher order effects, which are described by effective interaction terms and can be derived from the band structure and the carrier scattering in the quantum well. The spatial dependency of pump current densities plays a crucial role in lateral mode dynamics, and thus, a Drift-Diffusion model is employed to calculate the current densities with an additional capturing term.

1 Introduction

Fabry-Pérot type laser diodes find use in a diverse range of applications such as laser displays [1–4] and projection [5–7], exhibiting phenomena related to mode competition [8, 9]. These lasers often demonstrate mode hopping, wherein the relative activity levels of different longitudinal modes change with respect to time due to an antisymmetric interaction among these modes. In a recent development, a similar effect has been noted in broad area laser diodes, where multiple lateral modes are present [10]. This mode interaction can be explained by the beating vibrations of carrier densities within the quantum well, which appear when several longitudinal modes are concurrently active [11].

The most common approach for simulating mode dynamics in these devices involves using rate equations, which entail formulating equations of motion for the photon numbers of the different optical modes and the quantum well carrier densities. An alternative method, the traveling wave method, involves solving a partial differential equation for the electrical field [12–18]. Nonetheless, thus far, simulations of mode dynamics using the traveling wave method have not succeeded in reproducing the experimental results.

As the mode dynamics are strongly influenced by higher order effects such as beating vibrations of the carrier densities or spectral hole burning, it is insufficient to consider a spatially constant carrier density in the rate equations. One possibility is to use spatial and energy-dependent distribution functions [19], but these calculations are very computationally expensive. Alternatively it is possible to add an effective mode interaction term to the rate equations for the photon numbers, which has been initially derived for maser devices by Lamb et al. [20]. In literature this term can be found for single lateral mode Fabry-Pérot laser diodes [11, 21–25] and also for devices with multiple lateral modes [26].

Another important aspect for the simulation of Fabry-Pérot laser diodes with multiple lateral modes is the spatial dependency of the pump current densities. A higher current density in a specific area can result in an increased carrier density, leading to higher optical gain. Consequently, the modal gain of lateral modes increases when their mode functions reach a maximum in that region. Thus, it is important to accurately describe these pump current distributions in order to describe the lateral mode dynamics. For this purpose the Drift-Diffusion equations are often used in literature to describe carrier transport and to study the properties of semiconductor laser devices [27–31].

The goal of the study is to combine the rate equations with an effective mode interaction term with an improved description of carrier transport using the Drift-Diffusion equations. As the time scale of the mode dynamics is in the region of 100 ns, using a time-dependent Drift-Diffusion model would be too computationally expensive. Thus the Drift-Diffusion equations are solved for a steady state. The resulting pump current densities are used in a subsequent mode dynamics simulation, where only the carriers in the quantum wells are considered in the dynamic equations. In section 3 simulation results using this method for a green nitride laser diode with a ridge width of 10 μm .

2 Theory

2.1 Mode Function

The goal of this publication is the simulation of the mode dynamics in Fabry-Pérot type laser diodes. For this purpose the optical field is expanded using mode functions:

$$\mathbf{E}(\mathbf{r}) = \sum_{mp} i \sqrt{\frac{\hbar\omega_{mp}}{2\epsilon_0}} [B_{mp} \mathbf{u}_{mp}(\mathbf{r}) - B_{mp}^* \mathbf{u}_{mp}^*(\mathbf{r})].$$

The goal of the mode dynamics simulation is to know how mode coefficients B_{mp} or the number of photons in each mode $S_{mp} = |B_{mp}|^2$ change with respect to time. It is assumed that the ridge width of the laser diode is small compared to the resonator length in order to be able to separate the longitudinal and transverse contributions of the mode functions $\mathbf{u}_{mp}(\mathbf{r}_{\parallel}, z)$:

$$\mathbf{u}_{mp}(\mathbf{r}) = \mathbf{t}_m(x, z) g_p(y).$$

Here the index m is used to number the transverse modes \mathbf{t}_m and p the longitudinal modes g_p . The coordinates are chosen so that the optical field propagates in y direction and the device is grown in z direction. For transverse electric (TE) modes the polarization primarily points in the x direction $\mathbf{t}_m(x, z) = \mathbf{e}_x t_m(x, z)$ and the transverse modes are given by the eigen equation [32]

$$\begin{aligned} \left(\frac{\partial^2}{\partial x^2} + \frac{\partial^2}{\partial z^2} \right) t_m(x, z) + \frac{\omega^2}{c^2} n^2(x, z, \omega) t_m(x, z) \\ = \frac{\omega^2}{c^2} (n_{\text{eff}}^m(\omega))^2 t_m(x, z) \end{aligned} \quad (1)$$

for a given refractive index profile $n(x, z, \omega)$ and the interface conditions for the electromagnetic field. In the simulations this equation is solved for a frequency ω_0 near the gain maximum in order to obtain the transverse mode functions $t_m(x, z)$, their respective effective refractive indices $n_{\text{eff}}^m(\omega_0)$ and the group refractive indices

$$n_{\text{gr}}^m(\omega_0) = n_{\text{eff}}^m(\omega_0) + \omega_0 \frac{\partial n_{\text{eff}}^m(\omega_0)}{\partial \omega_0}.$$

The group refractive index determines the longitudinal mode spacing and is used to calculate the mode frequencies

$$\begin{aligned} \omega_{mp} &= \frac{\pi c}{n_{\text{eff}}^m(\omega_0) L} p_0 + \frac{\pi c}{n_{\text{gr}}^m(\omega_0) L} (p - p_0^m), \\ \text{where } p_0^m &= \text{round} \left(\frac{n_{\text{eff}}^m(\omega_0) L \omega_0}{\pi c} \right) \end{aligned}$$

and L is the resonator length. The normalization of the mode functions is given by [33]

$$\int dx \int dz n^2(x, z, \omega) |t_m(x, z)|^2 = 1.$$

2.2 Mode Dynamics

The equations of motion for the carriers and photon numbers can be derived in the Heisenberg picture [33] and for the carrier densities n_e and n_h are approximately given by [26]

$$\begin{aligned} \frac{d}{dt} n_{e,h}(x) = & -B(n_e(x), n_h(x)) - \frac{n_{e,h}(x)}{\tau_{nr}} \\ & + D_{e,h} \frac{\partial^2}{\partial x^2} n_{e,h}(x) + \left. \frac{d}{dt} n_{e,h}(x) \right|_{\text{Pump}} \\ & + \sum_{mp} |t_m(x, z_{QW})|^2 \omega_{mp} \frac{S_{mp}}{L} \text{Im} \chi(\omega_{mp}, n_e(x), n_h(x)). \end{aligned} \quad (2)$$

The first two terms describe the losses due to spontaneous emission and nonradiative processes. The next two terms are used to describe the pumping and diffusion of carriers inside the quantum wells, while the last term gives the losses due to stimulated emission. The stimulated emission term is proportional to the photon numbers S_{mp} and the imaginary part of the susceptibility $\text{Im} \chi$ is evaluated at the respective mode frequencies. The losses due to spontaneous emission $B(n_e, n_h)$ and the susceptibility $\chi(\omega, n_e, n_h)$ depend on the carrier densities and are calculated using Fermi-Dirac distributions with a fixed temperature. The equations of motion for the photon numbers are given by [26, 33]

$$\begin{aligned} \frac{d}{dt} S_{mp} = & -\omega_{mp} S_{mp} \int dx |t_m(x, z_{QW})|^2 \text{Im} \chi(\omega_{mp}, n_e, n_h) \\ & + \int dx |t_m(x, z_{QW})|^2 I_{SE}(\omega_{mp}, n_e, n_h) \\ & - \frac{S_{mp}}{\tau_{\text{photon}}} + \left. \frac{d}{dt} S_{mp} \right|_{\text{Interaction}}. \end{aligned} \quad (3)$$

The first three terms denote changes of the photon numbers due to stimulated emission, spontaneous emission and losses respectively. The last term is used to describe an effective interaction between the optical modes due to third order effects. Without this last term there would be no mode competition effects and in a simulation a steady state would be reached after a few nano seconds, where only the longitudinal mode with the highest gain would be active. The effective mode interaction is derived in Ref. [26] and is given by the integral

$$\begin{aligned} \left. \frac{d}{dt} S_{mp} \right|_{\text{Interaction}} \approx & \sum_{nq} \frac{1}{2L} \frac{S_{mp} S_{nq}}{\omega_{mp} \omega_{nq}} \\ & \times \int dx |t_m(x, z_{QW})|^2 |t_n(x, z_{QW})|^2 \\ & \times G(\omega_{nq} - \omega_{mp}, n_e(x), n_h(x)), \end{aligned} \quad (4)$$

where the factor $G(\Delta\omega, n_e, n_h)$ determines the strength of the mode interaction, and depends on the carrier densities and the frequency difference of two modes.

A change of the carrier densities causes the susceptibility of the quantum wells to change. The resulting perturbations of the mode functions can also be included in the calculations by solving the one-dimensional Eigen equation [34]

$$\frac{\partial^2}{\partial x^2} \tilde{t}_p^x(x) + k_0^2 \epsilon_{1D}(x) \tilde{t}_p^x(x) = k_0^2 n_{\text{eff},p}^2 \tilde{t}_p^x(x),$$

where \tilde{t}_p is the corrected transverse mode function at the position of the quantum well and $n_{\text{eff},p}$ is the corrected refractive index. The effective dielectric function is given by

$$\begin{aligned} \epsilon_{1D}(x) = & \frac{\int dz |t_0(x, z)|^2 n^2(\omega_0, x, z)}{\int dz |t_0(x, z)|^2} \\ & + \delta\epsilon(x) \int_{\text{QW}} dx dz |t_0(x, z)|^2 \end{aligned}$$

where $t_0(x, z)$ is the solution of the two-dimensional Eigen equation (1) and $n^2(\omega_0, x, z)$ is the respective refractive index profile. The change of the dielectric function in the quantum well is given by

$$\delta\epsilon(x) = \frac{1}{d_{\text{QW}}} [\chi(\omega_0, n_e(x), n_h(x)) - \chi(\omega_0, n_e^{\text{ref}}, n_h^{\text{ref}})],$$

where the reference densities n_e^{ref} and n_h^{ref} are included in the two-dimensional Eigen equation and are chosen close to the threshold densities.

It is possible to solve the equations of motion for a given pump term, for example a constant pump current density:

$$\left. \frac{d}{dt} n_{e,h}(x) \right|_{\text{Pump}} = j_0.$$

For laser diodes with larger ridge widths multiple transverse modes participate in the mode dynamics. For an accurate simulation of the mode dynamics it is therefore important to know the spatial dependency of the pump term, in order to estimate which transverse modes benefit based on their respective lateral mode profiles. For example, if the pump term has a global maximum at a certain point then transverse modes will be important when their respective mode functions also have a maximum near that point.

2.3 Drift-Diffusion Model

For a more detailed calculation of the pump term we perform a steady state calculation, where the carriers in the bulk material are treated separately from the carriers in the quantum wells. For the bulk carriers we use the Drift-Diffusion equations which are given by [35, 36]

$$\begin{aligned} -\nabla \left(\frac{\mu_e^M}{e} n_e^{3D} \nabla \mu_e \right) &= R_e(n_e^{3D}, n_h^{3D}, \phi) \\ -\nabla \left(\frac{\mu_h^M}{e} n_h^{3D} \nabla \mu_h \right) &= R_h(n_e^{3D}, n_h^{3D}, \phi) \\ -\nabla (\epsilon_0 \epsilon_r \nabla \phi) &= \rho = e (n_h^{3D} - n_e^{3D} + C) + \rho_{\text{QW}}, \end{aligned} \quad (5)$$

where $\mu_{e,h}^M$ are the mobilities, $\mu_{e,h}$ are the chemical potentials, ϕ is the static electrical field and C is the charge density due to doping. The relationship between the bulk densities n^{3D} and the chemical potentials is given by the Boltzmann distribution as shown in Ref. [35], but is possible to use other statistical functions [27, 28].

The equations for quantum well carriers and the photon numbers are given by Eqs. (2) and (3) where the time derivatives are set to zero. There are no mode dynamics in the steady state calculation, therefore it is sufficient to only consider one longitudinal mode per transverse mode, here the mode with the frequency closest to the gain maximum ω_0 is used. The charge density of the quantum well carriers is included in poisson equation as an additional charge density:

$$\rho_{QW} = \begin{cases} \frac{e}{d_{QW}} (n_h - n_e) & \text{for } |z - z_{QW}| < \frac{d_{QW}}{2} \\ 0 & \text{elsewhere} \end{cases},$$

where d_{QW} is the quantum well thickness. The capture of the bulk carriers into the quantum wells is included in the recombination term:

$$R_{e,h} = R_{e,h}^{\text{capture}} - n_e^{3D} n_h^{3D} (1 - e^{\beta(\mu_e + \mu_h)}) \\ \times \left(\frac{1}{\tau_e (n_e^{3D} + n_e^0)} + \frac{1}{\tau_h (n_h^{3D} + n_h^0)} \right) \\ + c_e n_e^{3D} + c_h n_h^{3D} + r_{\text{spont}},$$

where the second term describes the losses due to spontaneous emission, SRH and Auger recombination [35]. The capture term is only considered for $|z - z_{QW}| < d_{QW}/2$, and is given by

$$R_{e,h}^{\text{capture}} = C_{e,h} n_{e,h} \eta_{e,h} (n_{e,h}, n_{e,h}^{3D}).$$

The constants $C_{e,h}$ determine the strength of the capture process and the efficiencies $\eta_{e,h}$ are given by

$$\eta(n, n^{3D}) = \frac{\sum_{\mathbf{k}, k_z} f_{\mathbf{k}, k_z}^{\text{Bulk}} (1 - f_{\mathbf{k}})}{\sum_{\mathbf{k}, k_z} f_{\mathbf{k}, k_z}^{\text{Bulk}}},$$

where both $f_{\mathbf{k}, k_z}^{\text{Bulk}}$ and $f_{\mathbf{k}}$ are Fermi-Dirac distributions that reproduce the densities n and n^{3D} for a fixed temperature. Similar capture terms can be found in literature [31, 37]. For the bulk carriers the $\mathbf{k} \cdot \mathbf{p}$ method is used to calculate the three-dimensional band structure using the parameters from the material surrounding the quantum well. For the carriers in the quantum well the $\mathbf{k} \cdot \mathbf{p}$ method is used as well, here a one-dimensional differential equation is solved in order to obtain the two-dimensional band structure.

The pump term in Eq. (2) is then given by

$$\left. \frac{d}{dt} n_{e,h} \right|_{\text{Pump}} = C_{e,h} \int_{QW} dz n_{e,h}^{3D} \eta_{e,h} (n_{e,h}, n_{e,h}^{3D}) = j_{e,h}^0. \quad (6)$$

This way the equilibrium current densities can be calculated for every voltage and can be included in the dynamic simulations. However, in order to include changes of the pump current densities due to changes of the quantum well densities in the dynamic simulations, the equation

$$C_{e,h} d_{QW} n_{e,h}^{3D,0} \eta_{e,h} (n_{e,h}, n_{e,h}^{3D,0}) = j_{e,h}^0,$$

is solved for an effective three-dimensional density $n_{e,h}^{3D,0}$ using the two-dimensional density and the current density from the equilibrium simulation. The pump term in Eq. (2) is then given by

$$\left. \frac{d}{dt} n_{e,h} \right|_{\text{Pump}} = C_{e,h} d_{QW} n_{e,h}^{3D,0} \eta_{e,h} \left(n_{e,h}, n_{e,h}^{3D,0} \right).$$

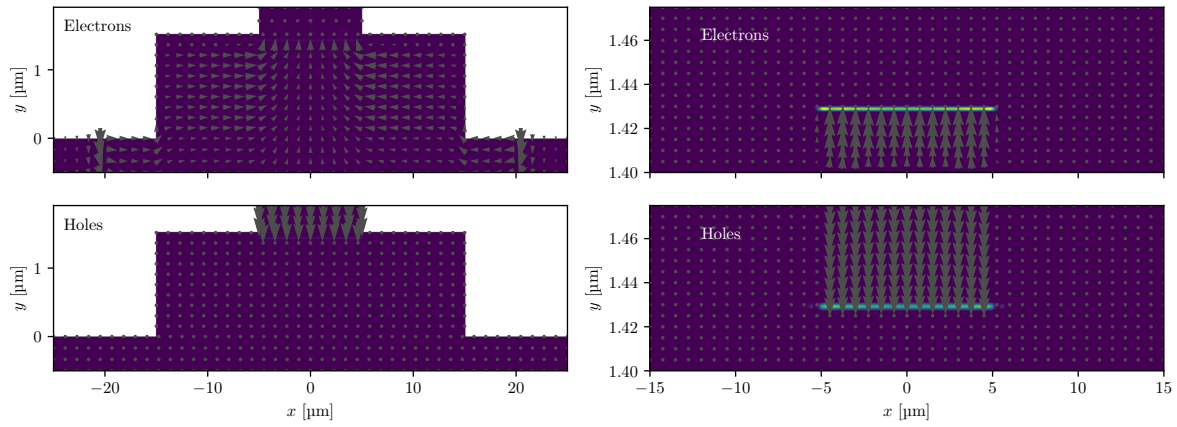


Figure 1: Solution of the stationary Drift-Diffusion model for a single QW laser diode with a ridge width of $10\ \mu\text{m}$ and a voltage of $4\ \text{V}$. The arrows denote the carrier current densities for electrons and holes, while the colors indicate the recombination term. On the right side the region near the quantum well is shown in more detail.

3 Results

In this section results for an example structure with a single quantum well and a ridge width of $10\ \mu\text{m}$ that exhibits multiple transverse modes are shown. The structure is described in more detail in Table 2 in the appendix. To calculate two-dimensional mode functions in Eq. (1) the dielectric function from Ref. [38] is used. Additionally the dielectric function of the quantum well is considered for a reference carrier density of $1 \times 10^{13}\ \text{cm}^{-2}$. As mentioned before, the band structure of the quantum well is required to compute the optical properties such as the susceptibility. For this purpose the $\mathbf{k} \cdot \mathbf{p}$ -Hamiltonian proposed by Chuang et al. is used [39–41], the method is described in more detail in Ref. [26]. The thickness of the InGaN quantum well used in the simulations is given by $2\ \text{nm}$, the indium concentration by $28\ \%$ and the material parameters are taken from Ref. [40]. The formulas for the susceptibility and the mode interaction term are given in the appendix.

For the charge carrier mobilities the model as described in Ref. [42] is used. The Drift-Diffusion equations are solved using the finite volume method with the Scharfetter-Gummel scheme for the carrier current densities [43]. The remaining simulation parameters for the simulations are shown in Table 1.

In Fig. 1 the current densities and the bulk carrier recombination are shown for a voltage above threshold. With the chosen parameters most of the carrier recombination is caused by the capture term at

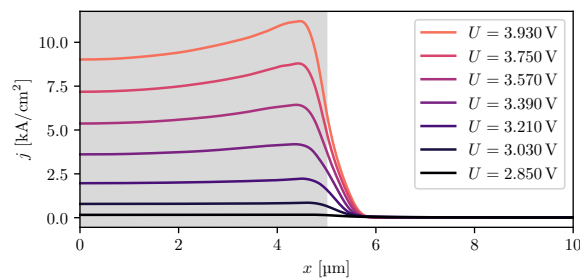


Figure 2: The electron pump current densities from Eq. (6) for different voltages near the threshold. The hole pump current densities look almost identical.

the position of the quantum well. The corresponding pump current densities are shown in Fig. 2 for different voltages. Due to the low hole mobility the pump current densities are high below the ridge and small in the other regions, but as the n -contacts are on the sides and far away from the ridge, the pump current densities have a maximum near the edge of the ridge. This maximum is more pronounced for larger voltages.

Table 1: Remaining simulation parameters

L	cavity length	600 μm
T	carrier temperature	300 K
τ_s	scattering time	300 fs
γ	dephasing constant	30 meV
τ_{nr}	nonradiative losses (QW)	5 ns
τ_{photon}	photon lifetime	5.9 ps
D_e	electron diffusion constant (QW)	151.9 $\text{nm}^2\text{ps}^{-1}$
D_h	hole diffusion constant (QW)	7.77 $\text{nm}^2\text{ps}^{-1}$
C_e	electron capture constant	1 ps^{-1}
C_h	hole capture constant	100 ps^{-1}
τ_e	electron nonradiative losses	1 ns
τ_h	hole nonradiative losses	1 ns
r_{spont}	bulk spontaneous emission losses	0.01 $\text{nm}^3\text{ps}^{-1}$
c_e	electron Auger losses	0.01 $\text{nm}^6\text{ps}^{-1}$
c_h	hole Auger losses	0.01 $\text{nm}^6\text{ps}^{-1}$
n_e^0	electron density for nonradiative losses	10^{19}cm^{-3}
n_h^0	hole density for nonradiative losses	10^{19}cm^{-3}

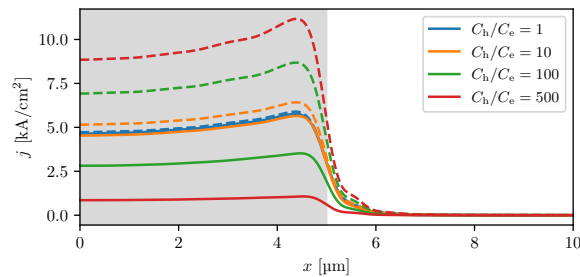


Figure 3: The electron pump current densities from Eq. (6) for the structure with two quantum wells are shown for different ratios C_h/C_e of the capture constants. The structure is the same as in Fig. 2 except for the additional QW, and is given in table 2 in the appendix.

The method can also be used for structures with multiple quantum wells. In this case the pump current densities depend strongly on the capture coefficients, which is shown in Fig. 3 for a structure with two quantum wells. For equal capture constants the current densities are split equally between the quantum wells. However, due to the lower hole mobility the hole capture constant is expected to be larger. As expected this results in a higher pump current density for the quantum well on the p -side.

Using these results it is possible to perform a mode dynamics simulation, as shown in Fig. 4 for different currents. The effect of mode hopping can be observed, where the laser output wavelength periodically changes from lower to higher values. As multiple transverse modes participate, the contributions of the

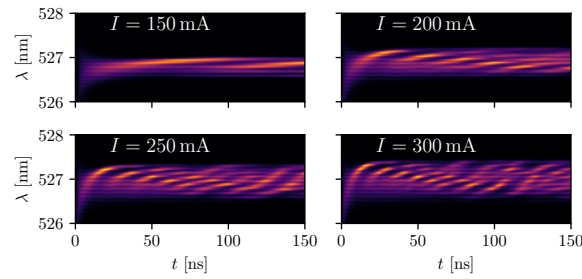


Figure 4: The longitudinal and transverse mode dynamics of a Fabry-Pérot laser diode with a single quantum well for four different currents. Here the output of the laser is shown as a function of wavelength and time. The data for this figure is calculated by multiplying the time-dependent photon numbers from the simulation with Gaussian functions that are centered at their respective vacuum wavelengths with a width of 0.02 nm.

respective longitudinal modes overlap, making it difficult to identify individual modes. This has been also observed experimentally for laser diodes with an even broader ridge of 40 μm [10].

Therefore it is of interest to look at the contributions of different transverse modes separately, which is shown in Fig. 5. In this case the different longitudinal modes can be observed, and the wavelength separation $\Delta\lambda = \frac{2L}{n_{\text{gr}}}$ is determined by the cavity length L . For the considered structure the fundamental mode is not active, because the pump current maximum is not in the middle of the structure but near the edge of the ridge, as shown in Fig. (2). If the current is increased further, more transverse modes will participate due to spatial hole burning.

In the model the time dependence of the mode wavelengths is also considered and is shown in Fig. 6 for different longitudinal and transverse modes. However after the relaxation oscillations are finished, the mode wavelengths remain constant in time and the changes do not play a role for the simulation of the mode dynamics.

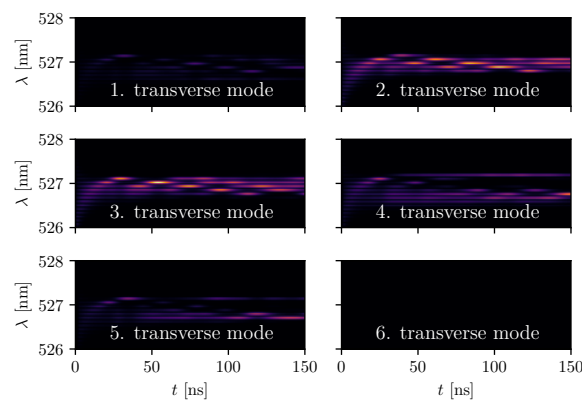


Figure 5: The contributions of different transverse modes to the mode dynamics for a laser diode with a single quantum well and a current of 200 mA.

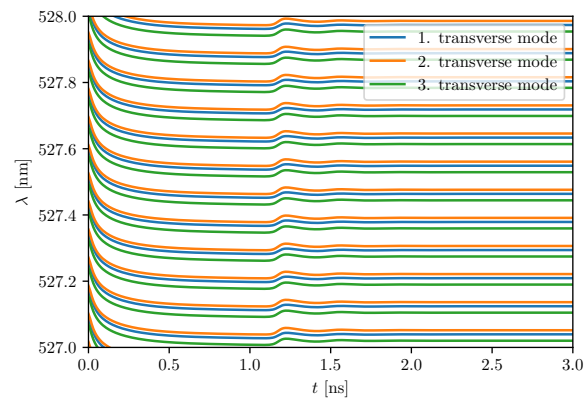


Figure 6: Changes of the mode wavelengths due to changes in the quantum well carrier densities for $I = 150$ mA.

4 Summary

In this publication a model for the simulation of mode dynamics in Fabry-Pérot type laser diodes with larger ridge widths and multiple lateral modes is presented. The carrier transport from the contacts to the active region is calculated for the steady state using the Drift-Diffusion model. The carriers in the quantum well are treated separately and are connected to bulk carriers using a capture term. The interaction between two different modes can be derived from third-order effects and is described in the simulation by an effective interaction term. This term strongly depends on the difference of the mode frequencies and the quantum well carrier densities.

Simulation results are shown for a green nitride laser diode with a ridge width of $10\ \mu\text{m}$, but in principle this model can be used for other materials and geometries. The effect of mode hopping can be observed in the resulting mode dynamics, which can also be observed experimentally for similar structures [10]. However, the experiments show multiple mode clusters. In order to obtain the same behaviour in simulations the presented model could be extended, for example fluctuations of the indium content in the quantum wells can be taken into account in future work.

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A Calculation of the susceptibility and the mode interaction

For the calculation of the susceptibility and the mode interaction the same formulas as in Ref. [26] are used. In order to keep the calculations simple for the example structure, a constant scattering time τ_s is used to describe the carrier scattering. It is of course possible to use more complicated scattering terms as described in Ref. [26] in more detail. Similarly a dephasing constant γ is used to describe the homogenous broadening and Hartree-Fock corrections are neglected. In this case the susceptibility is given by

$$\chi(\omega) = \frac{2}{A} \sum_{\lambda \mathbf{k}} \frac{e^2 |p_\lambda|^2}{\epsilon_0 m_0^2 \omega^2} \frac{1 - f_{\mathbf{k}}^e - f_{\mathbf{k}}^\lambda}{\hbar \omega + i\gamma - \epsilon_{\mathbf{k}}^e - \epsilon_{\mathbf{k}}^\lambda}.$$

Here λ is used as index for the hole bands and the carrier distributions are given by Fermi-Dirac distributions for a given temperature and carrier densities. The quantum well band structure $\epsilon_{\mathbf{k}}^\lambda$ and momentum matrix elements p_λ are calculated using the $\mathbf{k} \cdot \mathbf{p}$ method. The mode interaction in Eq. (4) can be split into symmetric and antisymmetric contributions with respect to the frequency difference of the two modes $\Delta\omega$:

$$G(\Delta\omega) = G_S(\Delta\omega) + G_A(\Delta\omega).$$

For a constant scattering time the symmetric mode interaction strength is given by

$$\begin{aligned} G_S(\Delta\omega) &= \omega_0^4 \text{Im} \chi(\omega_0) \text{Im} \chi'(\omega_0) \frac{\tau_s}{\Delta\omega^2 \tau_s^2 + 1} \\ &\quad - \sum_{\lambda \lambda' \mathbf{k}} \frac{e^4 |p_\lambda|^2 |p_{\lambda'}|^2 \tau_s (1 + \delta_{\lambda \lambda'})}{m_0^4 \epsilon_0^2 (\Delta\omega^2 \tau_s^2 + 1)} \\ &\quad \times \frac{2}{A} \frac{\gamma^2 (1 - f_{\mathbf{k}}^e - f_{\mathbf{k}}^{\lambda'})}{((\hbar\omega_0 - \epsilon_{\mathbf{k}}^e - \epsilon_{\mathbf{k}}^\lambda)^2 + \gamma^2) ((\hbar\omega_0 - \epsilon_{\mathbf{k}}^e - \epsilon_{\mathbf{k}}^{\lambda'})^2 + \gamma^2)} \end{aligned}$$

and the antisymmetric interaction term is given by

$$G_A(\Delta\omega) = \omega_0^4 \frac{\text{Im} \chi(\omega_0) \text{Re} \chi'(\omega_0)}{\Delta\omega}.$$

The carrier losses in the quantum well due to spontaneous emission are given by [26, 44]

$$B(n_e, n_h) = \frac{2}{A} \sum_{\lambda \mathbf{k}} \frac{2e^2 |p_\lambda|^2 n_{\text{eff}}^3}{3\pi m_0^2 \epsilon_0 c^3 \hbar^2} (\epsilon_{\mathbf{k}}^e + \epsilon_{\mathbf{k}}^\lambda) f_{\mathbf{k}}^e f_{\mathbf{k}}^\lambda.$$

The spontaneous emission spectrum I_{SE} in Eq. (3) is calculated like the susceptibility, except that the factor $1 - f_{\mathbf{k}}^e - f_{\mathbf{k}}^\lambda$ is replaced by $f_{\mathbf{k}}^e f_{\mathbf{k}}^\lambda$:

$$I_{\text{SE}}(\omega) = -\frac{2}{A} \sum_{\lambda \mathbf{k}} \frac{e^2 |p_\lambda|^2}{m_0^2 \epsilon_0 \omega} \text{Im} \left\{ \frac{f_{\mathbf{k}}^e f_{\mathbf{k}}^\lambda}{\hbar\omega + i\gamma - \epsilon_{\mathbf{k}}^e - \epsilon_{\mathbf{k}}^\lambda} \right\}.$$

B Structural parameters

The example structure with one quantum well used in the simulations has been obtained by stretching the structure in Ref. [45] in lateral direction. The structural parameters are given in Table 2. The example structure with two quantum wells is obtained by repeating the 2 nm and 8 nm layers. The p -contact is located on top of the ridge and the n -contact is on top of the substrate with a distance of 10 μm to the diode.

Table 2: Structural parameters for the laser diode used in the calculations, adapted from Ref. [45].

Thickness	Width	Material	Doping
55 nm	10 μm	GaN	$1 \times 10^{18} \text{ cm}^{-2}$
340 nm	10 μm	$\text{Al}_{0.08}\text{Ga}_{0.92}\text{N}$	$1 \times 10^{18} \text{ cm}^{-2}$
10 nm	30 μm	$\text{Al}_{0.15}\text{Ga}_{0.85}\text{N}$	$1 \times 10^{18} \text{ cm}^{-2}$
15 nm	30 μm	GaN	$1 \times 10^{18} \text{ cm}^{-2}$
50 nm	30 μm	GaN	0 cm^{-2}
13 nm	30 μm	$\text{In}_{0.05}\text{Ga}_{0.95}\text{N}$	0 cm^{-2}
2 nm	30 μm	$\text{In}_{0.22}\text{Ga}_{0.78}\text{N}$	0 cm^{-2}
8 nm	30 μm	$\text{In}_{0.05}\text{Ga}_{0.95}\text{N}$	0 cm^{-2}
2 nm	30 μm	$\text{In}_{0.22}\text{Ga}_{0.78}\text{N}$	0 cm^{-2}
13 nm	30 μm	$\text{In}_{0.05}\text{Ga}_{0.95}\text{N}$	0 cm^{-2}
115 nm	30 μm	GaN	$-1 \times 10^{18} \text{ cm}^{-2}$
1300 nm	30 μm	$\text{Al}_{0.06}\text{Ga}_{0.94}\text{N}$	$-1 \times 10^{18} \text{ cm}^{-2}$
4000 nm	50 μm	GaN	$-1 \times 10^{18} \text{ cm}^{-2}$