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Resonant inversion of the circular photogalvanic effect in *n*-doped quantum wells

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We show that the sign of the circular photogalvanic effect can be changed by tuning the radiation frequency of circularly polarized light. Here resonant inversion of the photogalvanic effect has been observed for direct intersubband transition in n-type GaAs quantum well structures. This inversion of the photon helicity driven current is a direct consequence of the lifting of the spin degeneracy due to k-linear terms in the Hamiltonian in combination with energy and momentum conservation and optical selection rules.

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I. INTRODUCTION

Effects caused by spin-orbit interaction in compound semiconductor heterojunctions have been the subject of a growing number of investigations recently.^{1,2} In twodimensional systems based on quantum wells (QW's) the electron spin couples to the electron motion and results, under optical orientation with circularly polarized light, in spin photocurrents.³ The direction and magnitude of this spin photocurrent depend on the degree of circular polarization of the incident light.⁴ This phenomenon belongs to the class of photogalvanic effects⁵ and represents here a circular photogalvanic effect (CPGE).

It was shown in Ref. 3 that the CPGE in zinc-blende structure based QW's is caused by spin orientation of carriers in systems where the spin degeneracy of the band structure is lifted by k-linear terms in the Hamiltonian.^{6,7} In this case homogeneous irradiation of QW's with circularly polarized light results in an asymmetric distribution of photoexcited carriers in k space which leads to the current. So far this effect has been observed only for indirect intrasubband transitions in n- and p-type QW's (Drude absorption) and for direct inter-subband heavy-hole–light-hole transitions in p-type QW's.^{3,8}

Here we report on an observation of a resonant inversion of the CPGE at direct transitions between size quantized subbands in n-type QWs. This effect demonstrates in a very direct way the spin splitting of subbands in k space in zero electric and magnetic field due to spin-orbit interaction. We show that the sign of the spin driven circular photogalvanic current can be reversed by tuning the radiation frequency. This inversion of the photon helicity driven current is a direct consequence of k-linear terms in the subband structure in combination with conservation laws and optical selection rules.

II. EXPERIMENTAL RESULTS

Direct intersubband transitions between the lowest (e1) and the second (e2) conduction subband in *n*-type GaAs QW's have been obtained by applying a line tunable pulsed transversely excited atmospheric pressure (TEA)-CO₂ laser.

The laser yields strong linearly polarized emission at wavelengths λ between 9.2 and 10.8 μ m corresponding to photon energies $\hbar \omega$ ranging from 135 to 114 meV. The quantum well widths were chosen to be ≈ 8 nm, so that the separation of the subbands e1 and e2 matches the photon energy range of the laser.^{9,10} Molecular-beam-epitaxy grown (001)- and (113)-oriented *n*-type GaAs/AlGaAs QW samples of 8.8, 8.2, and 7.6 nm width with free-carrier densities ranging between 2×10^{11} cm⁻² and 1×10^{12} cm⁻² were investigated at room temperature. The (113)-oriented samples have been grown on GaAs (113)A substrates employing growth conditions under which Si dopants are predominantly incorporated as donors¹¹ as confirmed by Hall measurements.

On each sample a pair of contacts along a line parallel to the x direction has been attached [see Fig. 1(c)]. We use Cartesian coordinates for (001)-oriented samples $x \parallel [1\overline{10}]$, $y \parallel [10]$, $z \parallel [001]$ and for (113)-oriented samples $x' = x \parallel [1\overline{10}]$, $y' \parallel [33\overline{2}]$, $z' \parallel [113]$. Right handed (σ_+) and left handed (σ_-) circularly polarized radiation was achieved by using a Fresnel rhomb. In order to correlate the spectral dependence of the CPGE current to the absorption of the QW's, optical transmission measurements were carried out using a Fourier transform infrared spectrometer. The current *j* generated by circularly polarized light in the unbiased devices was measured via the voltage drop across a 50 Ω load resistor in a closed circuit configuration [see Fig. 1(c)]. The voltage in response to a laser pulse was recorded with a storage oscilloscope.

Illuminating the unbiased QW structures with circularly polarized radiation results in a current signal due to CPGE which is proportional to the helicity $P_{\rm circ}$ of the radiation. The signal follows the temporal structure of the laser pulse and changes sign if the circular polarization is switched from σ_+ to σ_- . Typical signal traces are shown in Fig. 1 compared to records of a linear photon drag detector.¹² In (001)-oriented samples, belonging to the point group C_{2v} , the CPGE current is only observed under oblique incidence of radiation, as expected from symmetry.³ For illumination along the *y* direction the helicity dependent photocurrent flows in the *x* direction perpendicular to the wave vector of the incident light. This is observed in experiment. In Fig. 2



Time (10⁻⁷s)

FIG. 1. Oscillographic traces obtained through excitation with $\lambda = 10.6 \ \mu m$ radiation of (113)-grown *n*-GaAs QW's. (a) and (b) show CPGE signals obtained for σ_+ and σ_- -circular polarization, respectively. For comparison in (d) a signal pulse of a fast photon drag detector is plotted. In (c) the measurement arrangement is sketched. For (113)-grown samples being of C_s symmetry radiation was applied at normal incidence and the current was detected in the direction $x \parallel [1\overline{10}]$. For (001)-grown QW's oblique incidence with light propagating along [110] direction.

the photocurrent as a function of photon energy is plotted for σ_+ and σ_- polarized radiation together with the absorption spectrum. The data are presented for a (001)-grown *n*-GaAs QW of 8.2 nm width measured at room temperature. The current for both, left and right handed circular polarizations,



FIG. 2. Photocurrent in QW's normalized by the light power *P* as a function of the photon energy $\hbar \omega$. Measurements are presented for *n*-type (001)-grown GaAs/AlGaAs QW's of 8.2 nm width (symmetry class C_{2v}) at T=293 K. Oblique incidence of σ_+ (squares) and σ_- (circles) circular polarized radiation with an angle of incidence $\Theta_0 = 20^\circ$ was used. The current j_x was measured perpendicular to the direction of light propagation *y*. The dotted line shows the absorption measurement using a Fourier transform infrared spectrometer.



FIG. 3. Microscopic picture describing the origin of the inversion of the photocurrent in C_s point group samples. The essential ingredient is the splitting of the conduction band due to *k*-linear terms. Right handed circularly polarized radiation σ_+ induces direct spin-flip transitions (vertical arrows) from e1 subband with s = -1/2 to e2 subband with s = +1/2. As a result an unbalanced occupation of the k_x states occurs yielding a spin polarized photocurrent. (a) For transitions with k_x^- left to the minimum of e1 (s = -1/2) subband the current indicated by j_x is negative. (b) At smaller $\hbar \omega$ the transition occurs at k_x^+ , now right to the subband minimum, and the current reverses its sign.

changes sign at a frequency $\omega = \omega_{inv}$. This inversion frequency ω_{inv} coincides with the frequency of the absorption peak (see Fig. 2). The absorption peak frequency and ω_{inv} depend on the sample width according to the variation of the subband energy separation. This has been verified by measuring QW's of different widths. Spin orientation induced CPGE and its spectral sign inversion have also been detected in a (113)-oriented *n*-GaAs QW which belongs to the point group C_s . In this case the helicity dependent signal is observed in the *x* direction at normal incidence of radiation along z'.

III. MICROSCOPIC MODEL

The physical origin of the effect is sketched in Fig. 3 for C_s symmetry and in Fig. 4 for C_{2v} symmetry. For both symmetries the degeneracy in k space is lifted. First we consider the simplest case of C_s symmetry relevant for (113)-oriented samples. The $\sigma_{z'}k_x$ contribution to the Hamiltonian, responsible for the effect under normal incidence, splits the electron spectrum into spin sublevels with the spin components s = $\pm 1/2$ along the growth direction z'. As a result of optical selection rules right-handed circular polarization under normal incidence induces direct optical transitions between the subband e1 with spin s = -1/2 and e2 with spin s = +1/2. For monochromatic radiation optical transitions occur only at a fixed k_x where the energy of the incident light matches the transition energy as is indicated by the arrow in Fig. 3(a). Therefore optical transitions induce an imbalance of momentum distribution in both subbands yielding an electric current. However, a nonequilibrium distribution of carriers in the upper subband rapidly relaxes due to the very effective relaxation channel of LO-phonon emission, because the energy separation ε_{21} between e1 and e2 at $k_x = 0$ is well above the energy of LO phonons in *n*-GaAs QW's (ε_{LO}



FIG. 4. Microscopic picture describing the origin of the inversion of the photocurrent in C_{2v} point group samples. (a) Excitation at oblique incidence with σ_+ radiation of $\hbar \omega$ less than the energy subband separation ε_{21} induces direct spin-conserving transitions (vertical arrows) at k_x^- and k_x^+ . The rates of these transitions are different as illustrated by different thickness of the arrows. This leads to a photocurrent due to an asymmetrical distribution of carriers in k space if the splittings of the e1 and e2 subbands are nonequal. (b) Increase of the photon energy shifts more intensive transitions to the right and less intensive to the left resulting in a current sign change.

= 35.4 meV). Therefore the contribution of the e2 subband to the electric current vanishes and the magnitude and direction of electron flow is determined by the momentum distribution of carriers in the lowest subband.

Figures 3(a) and 3(b) show what happens when, as in our experiment, the energy of the incident light is varied from energies above ε_{21} to values below ε_{21} . Here ε_{21} is the subbands' energy separation at k=0. At large photon energy, $\hbar \omega > \varepsilon_{21}$, excitation occurs at negative k_x resulting in a current j_x shown by arrow in Fig. 3(a). A reduced photon frequency shifts the transition towards positive k_x and reverses the direction of the current [Fig. 3(b)]. The inversion of the current's sign occurs at a photon frequency ω_{inv} corresponding to the transition at the minimum of e1 (s = -1/2). The model suggests that the magnitude of the spin splitting described by the Rashba and Dresselhaus terms^{6,7} could easily be derived from the energy shift $\hbar \omega_{inv} - \varepsilon_{21}$. However, our measurements show, that the broadening of the optical transition in real QW's is too large to obtain this energy shift. On the other hand the model clearly shows that without k-linear terms in the band structure neither an inversion nor a current would exist. Similar arguments hold for C_{2v} symmetry [relevant for (001)-oriented samples] under oblique incidence (see Fig. 4) although the simple selection rules are no longer valid.¹³ This is pointed out in more detail at the end of the next section.

IV. MICROSCOPIC THEORY

The theory of the circular photogalvanic effect is developed by using the spin density matrix technique.⁵ Generally the total electric current that appears in a structure under intersubband excitation consists of the contributions from the e1 and e2 subbands which in the relaxation time approximation are given by standard expressions

$$j^{(\nu)} = e \sum_{k} \tau_{p}^{(\nu)} \operatorname{Tr}[\hat{v}^{(\nu)}(k) \dot{\rho}^{(\nu)}(k)].$$
(2)

Here *e* is the electron charge, $\nu = 1,2$ labels the subband $e\nu$, $\tau_p^{(\nu)}$ is the momentum relaxation time in the subband $e\nu$, $\dot{\rho}^{(\nu)}(\mathbf{k})$ is the generation of the density matrix, and $\hat{\mathbf{v}}^{(\nu)}$ is the velocity operator in the subband given by

$$\hat{\boldsymbol{v}}^{(\nu)} = \hbar^{-1} \nabla_{\boldsymbol{k}} \hat{H}^{(\nu)}. \tag{3}$$

For the sake of simplicity we will consider a parabolic electron spectrum for the subbands and take a Hamiltonian of the form

$$\hat{H}^{(\nu)} = \varepsilon^{(\nu)} + \frac{\hbar^2 k^2}{2m^*} + \hat{\mathcal{H}}_{1k}^{(\nu)}, \qquad (4)$$

where $\varepsilon^{(\nu)}$ is the energy of size quantization, $\hat{\mathcal{H}}_{1k}^{(\nu)}$ is the spin-dependent *k*-linear contribution, and m^* is the effective mass equal for both subbands.

It is convenient to write the generation matrices in the basis of spin eigenstates $\chi_{\nu ks}$ of the Hamiltonians $\hat{H}^{(\nu)}$. For the case of intersubband transitions $e_1 \rightarrow e_2$ the corresponding equations have the form (see Ref. 14)

$$\dot{\rho}_{ss'}^{(1)}(\mathbf{k}) = -\frac{\pi}{\hbar} \sum_{s''} \mathcal{M}_{s'',s'}(\mathbf{k}) \mathcal{M}_{s'',s}^{*}(\mathbf{k}) [f_{ks} \delta(\varepsilon_{2ks''} - \varepsilon_{1ks} - \hbar\omega) + f_{ks'} \delta(\varepsilon_{2ks''} - \varepsilon_{1ks'} - \hbar\omega)], \qquad (5)$$

$$\dot{\rho}_{ss'}^{(2)}(\mathbf{k}) = \frac{\pi}{\hbar} \sum_{s''} f_{ks''} \mathcal{M}_{s,s''}(\mathbf{k}) \mathcal{M}_{s',s''}^*(\mathbf{k}) [\,\delta(\varepsilon_{2ks} - \varepsilon_{1ks''} - \hbar\,\omega) + \delta(\varepsilon_{2ks'} - \varepsilon_{1ks''} - \hbar\,\omega)],$$

Here *s*, *s'* and *s"* are the spin indices, f_{ks} is the equilibrium distribution function in the subband *e*1 (the subband *e*2 is empty in equilibrium), $\varepsilon_{\nu ks}$ is the electron energy, and $\mathcal{M}_{s'',s'}(\mathbf{k})$ is the matrix element of intersubband optical transitions $(e1,\mathbf{k},s') \rightarrow (e2,\mathbf{k},s'')$. The latter is given by $\mathcal{M}_{s'',s'}(\mathbf{k}) = \chi^{\dagger}_{2ks''} \hat{\mathcal{M}} \chi_{1ks'}$, where $\hat{\mathcal{M}}$ is a 2×2 matrix describing the intersubband transitions in the basis of fixed spin states $s_z = \pm 1/2$,

$$\hat{M} = -\frac{eA}{cm^*} p_{21} \begin{bmatrix} e_z & \Lambda(e_x - ie_y) \\ -\Lambda(e_x + ie_y) & e_z \end{bmatrix}, \quad (6)$$

A is the amplitude of the electro-magnetic wave related to light intensity by $I=A^2\omega^2 n_{\omega}/(2\pi c)$, e is the unit vector of the light polarization, n_{ω} is the refraction index of the media, c is the light velocity, and p_{21} is the momentum matrix element between the envelope functions of size quantization $\varphi_1(z)$ and $\varphi_2(z)$ in the subbands e1 and e2,

$$p_{21} = -i\hbar \int \varphi_2(z) \frac{\partial}{\partial z} \varphi_1(z) dz.$$
 (7)

The parameter Λ originates from $k \cdot p$ admixture of valence band states to the electron wave function and is given by

$$\Lambda = \frac{\varepsilon_{21}\Delta(2\varepsilon_g + \Delta)}{2\varepsilon_g(\varepsilon_g + \Delta)(3\varepsilon_g + 2\Delta)},\tag{8}$$

where ε_g is the energy of the band gap, and Δ is the energy of spin-orbit splitting of the valence band. As one can see from Eq. (6), the parameter Λ determines the absorbance for the light polarized in the interface plane.

In ideal QW's the circular photocurrent j may be obtained from Eqs. (1)–(5). However, in real structures the spectral width of the intersubband resonance is broadened due to fluctuation of the QW width and hence exceeds the spectral width of the absorption spectrum of an ideal structure. The broadening can be taken into account assuming that the energy separation between subbands ε_{21} varies in the QW plane. Then by convolution of the photocurrent $j(\varepsilon_{21})$ with the distribution function $F(\varepsilon_{21})$ we obtain

$$\overline{j} = \int j(\varepsilon_{21}) F(\varepsilon_{21}) d\varepsilon_{21}.$$
(9)

The function $F(\varepsilon_{21})$ for broadening may be expanded in powers of $\varepsilon_{21} - \hbar \omega$ and by considering only the first two terms we obtain

$$F(\varepsilon_{21}) \approx F(\hbar\omega) + F'(\hbar\omega)(\varepsilon_{21} - \hbar\omega). \tag{10}$$

Taking into account the Hamiltonian $\hat{\mathcal{H}}_{1k}^{(\nu)}$ to be linear in k, the averaged current is finally given by

$$\overline{j} = e n_e \frac{\pi}{\hbar^2} \left[\tau_p^{(2)} F(\hbar \omega) + (\tau_p^{(1)} - \tau_p^{(2)}) F'(\hbar \omega) \overline{\varepsilon} \right] \\ \times \operatorname{Tr} [\hat{M}^{\dagger} (\nabla_k \hat{\mathcal{H}}_{1k}^{(2)}) \hat{M} - \hat{M} (\nabla_k \hat{\mathcal{H}}_{1k}^{(1)}) \hat{M}^{\dagger}], \qquad (11)$$

where n_e is the 2D carrier density, and $\overline{\varepsilon}$ is the mean value of the electron energy. For a degenerate 2D electron gas $\bar{\varepsilon}$ $=\varepsilon_F/2$ and for a non-degenerate gas $\overline{\varepsilon}=k_BT$, where ε_F is the Fermi energy, k_B is the Boltzmann constant, and T is the temperature. We note, that the distribution function $F(\hbar \omega)$ describes the spectral behavior of the absorbance taking into account both inhomogeneous and homogeneous broadening. For the absorption spectrum in Fig. 2, the full width at half maximum (FWHM) is about 15 meV which corresponds to 1-2 monolayer fluctuations of the QW thickness. The homogeneous broadening at room temperature can be estimated as $\hbar W_{12}$ where W_{12} is the rate of optical phonon induced transitions $e2 \rightarrow e1$. By using the available estimations of W_{12} (Ref. 15) we can conclude that the homogeneous broadening is almost by an order of magnitude smaller than the observed FWHM. We also note that, due to the nonparabolicity of the 3D electron spectrum and the difference of effective masses in the well and barrier materials, the electron in-plane effective masses in the e1 and e2 subbands differ which leads to an additional broadening of the absorption spectrum. This may lead to an asymmetry of the absorption peak and a small redshift of the photocurrent inversion point.

A. C_s -symmetry and normal incidence

In (113)-grown QW structures of C_s symmetry the CPGE occurs under normal incidence of the radiation. In this case the *k*-linear contribution to the Hamiltonian responsible for the effect is given by $\beta_{z'x}^{(\nu)}\sigma_{z'}k_x$. Here $\beta_{z'x}^{(\nu)}$ are the coefficients being different for the *e*1 and *e*2 subbands. The *k*-linear term splits the electron spectrum into spin sublevels with the spin components $s = \pm 1/2$ along the growth direction z' (see Fig. 3). Thus the electron parabolic dispersion in the subbands *e*1 and *e*2 has the form

$$\varepsilon_{\nu,k,\pm 1/2} = \varepsilon^{(\nu)} + \frac{\hbar^2 (k_x^2 + k_{y'}^2)}{2m^*} \pm \beta_{z'x}^{(\nu)} k_x.$$
(12)

For direct intersubband transitions under normal incidence selection rules allow only the spin-flip transitions (e1, -1/2) \rightarrow (e2,1/2) for σ_+ photons and (e1,1/2) \rightarrow (e2, -1/2) for σ_- photons.¹³ Due to these selection rules together with energy and momentum conservation laws the optical intersubband transition under, for example, σ_+ photoexcitation is only allowed for the fixed wave vector k_x given by

$$k_{x} = \frac{\hbar\omega - \varepsilon_{21}}{\beta_{z'x}^{(2)} + \beta_{z'x}^{(1)}}.$$
(13)

Velocities of electrons in the e^2 subband and of "holes" in the e^1 subband generated by this transition are given by

$$v_x^{(1)} = \hbar k_x / m^* - \beta_{z'x}^{(1)} / \hbar, \quad v_x^{(2)} = \hbar k_x / m^* + \beta_{z'x}^{(2)} / \hbar.$$
(14)

This unbalanced distribution of carriers in k space induces an electric current

$$j_{x} = j_{x}^{(1)} + j_{x}^{(2)} = -e \frac{\eta \| I}{\hbar \omega} (v_{x}^{(1)} \tau_{p}^{(1)} - v_{x}^{(2)} \tau_{p}^{(2)}) P_{\text{circ}}, \quad (15)$$

where *I* is the light intensity and η_{\parallel} is the absorbance or the fraction of the energy flux absorbed in the QW due to the intersubband transitions under normal incidence. Note that the magnitude of the photocurrent $j_x^{(2)}$, corresponding to the second term in the bracket of Eq. (15) and stemming from photoelectrons in the *e*2 subband is smaller than $|j_x^{(1)}|$ because $\tau_p^{(2)} \ll \tau_p^{(1)}$ as described above. The resonant inversion of the circular photocurrent is clearly seen from Eqs. (13)–(15) because η_{\parallel} is positive and $v_x^{(\nu)}$ changes its sign at a particular frequency.

For a degenerate 2D electron gas at low temperature we find that the dependence of the absorbance η_{\parallel} on $\hbar \omega$ and $\beta_{\tau'x}^{(\nu)}$ is given by

$$\frac{\eta_{\parallel}}{\hbar\omega} \propto \frac{1}{|\beta_{z'x}^{(2)} + \beta_{z'x}^{(1)}|} \left[\tilde{\varepsilon}_F - \frac{\hbar^2}{2m^*} \left(\frac{\hbar\omega - \varepsilon_{21}}{\beta_{z'x}^{(2)} + \beta_{z'x}^{(1)}} \right)^2 \right]^{1/2}, \quad (16)$$

where $\tilde{\varepsilon}_F = \varepsilon_F - m^* [\beta_{z'x}^{(1)}/(\sqrt{2}\hbar)]^2$.

Taking into account the broadening we finally obtain for the averaged circular photocurrent

$$\begin{split} \overline{j}_{x} &= \frac{e}{\hbar} \left(\beta_{z'x}^{(2)} + \beta_{z'x}^{(1)} \right) \left[\tau_{p}^{(2)} \overline{\eta}_{\parallel}(\hbar \, \omega) \right. \\ &+ \left(\tau_{p}^{(1)} - \tau_{p}^{(2)} \right) \overline{\varepsilon} \frac{d \, \overline{\eta}_{\parallel}(\hbar \, \omega)}{d\hbar \, \omega} \right] \frac{IP_{\text{circ}}}{\hbar \, \omega}, \tag{17}$$

where $\bar{\eta}_{\parallel} \propto F(\hbar \omega)$ is the calculated absorbance neglecting *k*-linear terms but taking into account the broadening.

B. C_{2v} symmetry and oblique incidence

In the case of C_{2v} point symmetry which is relevant for (001)-oriented QW's the current flows only at oblique incidence and is caused by *k*-linear contributions to the electron effective Hamiltonian given by

$$\mathcal{H}_{1k}^{(\nu)} = \beta_{xy}^{(\nu)} \sigma_x k_y + \beta_{yx}^{(\nu)} \sigma_y k_x.$$
(18)

The coefficients $\beta_{xy}^{(\nu)}$ and $\beta_{yx}^{(\nu)}$ are related to the bulkinversion asymmetry (BIA) or Dresselhaus term^{7,16} and structure-inversion asymmetry (SIA) or Rashba term⁶ by

$$\beta_{xy}^{(\nu)} = \beta_{\text{BIA}}^{(\nu)} + \beta_{\text{SIA}}^{(\nu)}, \quad \beta_{yx}^{(\nu)} = \beta_{\text{BIA}}^{(\nu)} - \beta_{\text{SIA}}^{(\nu)}.$$
(19)

The circular photocurrent due to intersubband transition in (001)-grown QW's in the presence of broadening can be calculated following Eq. (11) yielding

$$j_{x} = -\Lambda \frac{e}{\hbar} \left(\beta_{yx}^{(2)} - \beta_{yx}^{(1)}\right) \left[\tau_{p}^{(2)} \eta_{\perp}(\hbar \omega) + \left(\tau_{p}^{(1)} - \tau_{p}^{(2)}\right) \bar{\varepsilon} \frac{d \eta_{\perp}(\hbar \omega)}{d\hbar \omega} \right] \frac{IP_{\text{circ}}}{\hbar \omega} \hat{e}_{y}, \qquad (20)$$

where \hat{e} is the unit vector directed along the light propagation and η_{\perp} is the absorbance for the polarization perpendicular to the QW plane. The current in the *y* direction can be obtained by interchanging the indices *x* and *y* in Eq. (20).

The origin of the spin orientation induced CPGE caused by direct intersubband transitions in C_{2v} -symmetry systems is illustrated in Fig. 4 for σ_+ radiation. In C_{2v} symmetry the $\sigma_y k_x$ contribution to the Hamiltonian splits the subbands in k_x direction in two spin branches with $s = \pm 1/2$ oriented along y (see Fig. 4). Due to selection rules the absorption of circularly polarized radiation, which yields an in-plane spin polarization, is spin conserving.⁵ The asymmetric distribution of photoexcited electrons resulting in a current is caused by these spin-conserving but spin-dependent transitions [see Eq. (20)]. This is in contrast to spin-flip processes occurring in (113)-grown QW's described above. It turns out that under oblique excitation by circularly polarized light the rates of intersubband transitions are different for electrons with the spin oriented coparallel and antiparallel to the in-plane direction of light propagation.¹⁷ The difference is proportional to the product $|M_{\parallel}M_{\perp}|$, where M_{\parallel} and M_{\perp} are the absorption matrix elements for in-plane and normal light polarization. This is depicted in Fig. 4 by vertical arrows of different thickness. In systems with k-linear spin splitting such processes lead to an asymmetrical distribution of carriers in kspace, i.e., to an electrical current. Similar to C_s symmetry the variation of the photon energy leads to the inversion of the current direction [see Figs. 4(a) and 4(b)]. Since the circular photogalvanic effect in QW structures of C_{2v} symmetry is caused by spin-dependent spin-conserving optical transitions, the photocurrent described by Eq. (20) in contrast to Eq. (17) is proportional to the difference of subband spin splittings.

V. CONCLUSIONS

Our experiments show that direct inter-subband transitions in *n*-type GaAs QW's result in a spin orientation induced CPGE. The central observation is a change of the sign of the CPGE current close to resonance of intersubband transitions. The theoretical results give a detailed description of all features observed in experiment. The sign inversion follows for C_s symmetry from Eq. (17) and for C_{2v} symmetry from Eq. (20). If $\tau_p^{(2)} \ll \tau_p^{(1)}$ the first term on the right hand side in square brackets of both equations is vanishingly small compared to the second one. Therefore the photocurrent is proportional to the derivative of the absorbance and is zero at the frequency of the absorption peak as observed in experiment. Thus the assumption that the momentum relaxation in the upper subband is much faster than in the lower subband is satisfied.

Comparing theory with experiment we also note that for C_{2v} symmetry the spin splitting in k space is different for the e1 and e2 subbands. Equation (20) shows that for equal spin splitting $\beta_{yx}^{(2)} = \beta_{yx}^{(1)}$, the current vanishes. For C_s symmetry in contrast, the spin-orientation induced CPGE current is proportional to the sum of the subband spin splittings and therefore exist for $\beta_{zx}^{(2)} = \beta_{zx}^{(1)}$. The observation of the CPGE for direct intersubband transitions allows one to extend the method of spin-sensitive bleaching, previously demonstrated for *p*-type QW's,¹⁸ and therefore to investigate electron spin relaxation times in *n*-type QW's for monopolar spin orientation.^{18,19}

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