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To cite this article: Ching-Hao Chang and Carmine Ortix 2017 Nano Futures 1 035004

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Angle-dependent Weiss oscillations in a nanocorrugated two-dimensional electron gas

Ching-Hao Chang and Carmine Ortix

1 Institute for Theoretical Solid State Physics, IFW Dresden, Helmholtzstr. 20, D-01069 Dresden, Germany
2 Institute for Theoretical Physics, Center for Extreme Matter and Emergent Phenomena, Utrecht University, Princetonplein 5, 3584 CC, Utrecht, The Netherlands

E-mail: c.h.chang@ifw-dresden.de and c.ortix@uu.nl

Keywords: Weiss oscillations, guiding-center drift, snake orbits, perpendicular magnetic field

Abstract

We investigate the diffusive magnetotransport properties of a two-dimensional electron gas residing in a wrinkled nanostructure. The curved geometry of the nanostructure renders an effective inhomogeneous magnetic field which, in turn, yields Weiss oscillations. Since the relative strength of the effective inhomogeneous magnetic field can be tailored by changing the direction of the externally applied magnetic field, these Weiss oscillations exhibit a strong directional dependence. For large external magnetic fields we also find an anisotropic positive magnetoresistance.

1. Introduction

In very recent years, advanced nanostructuring techniques have enabled the synthesis of novel low-dimensional nanostructures in which thin film materials can be bent into curved deformable objects such as rolled-up nanotubes [1–3], nanohelices [4], and even complex nanarchitectures resembling structures that form naturally in the most basic forms of life [5]. On one hand, these next-generation nanomaterials carry an enormous potential in electronics, ranging from flexible displays to the integration of semiconductor electronics with the soft, curvilinear surfaces of the human body [6, 7]. On the other hand, the very fundamental quantum mechanical properties of charge carriers in these nanomaterials are strongly affected by the curved background in which they live [8, 9]. The recent theoretical advances in defining the quantum mechanics in constrained low-dimensional curved geometries have already brought to light a series of unique curvature-induced electronic and transport properties. Prime actors include the appearance of winding-generated bound states [10], a strongly anisotropic ballistic magnetoresistance (MR) in non-magnetic and spin–orbit–free semiconducting nanotubes [11], signatures of snake states in the transport properties of core–shell nanowires [12], peculiar Aharonov–Bohm oscillations in prismatic core–shell nanowires [13], the geometric shape control of spin–interference effects in quantum rings [14, 15], an all-geometrical driving of topological insulating states of matter in bent nanowires [16], and even reversal of thermoelectric currents in tubular nanowires [17].

Among the different geometrically deformed nano-objects that are nowadays produced, rippled nanostructures (see figure 1(a)) undoubtedly play a primary role. Wrinkling of thin solid films can occur over a wide range of length scales [18], and results from the presence of external strains. Controlled ripple texturing, for instance, has been achieved in suspended graphene sheets using spontaneously generated or thermally generated strains [19]. Another route involves the transfer of thin solid films onto pre-stressed elastomers in which strains are suddenly released. This method has enabled the fabrication of wrinkled silicon nanomembranes [20], and more recently of multilayer black phosphorous flakes with ripple textures [21]. Finally, controlled wrinkling of a nanomembrane with an embedded InGaAs quantum well has been also achieved by partial release and bond back of layers upon underetching [22].

From the theoretical point of view, the creation of ripples represents an avenue to modulate electronic properties on a local scale with the tremendous advantage that the system remains chemically pristine. Structural ripples can result in added periodic potentials, which can induce mini band gaps and Shockley end modes [23]. The aim of this paper is to show that the magnetotransport properties of a two-dimensional electron gas (2DEG)
in rippled nanostructures naturally display the so-called Weiss oscillations [24] occurring in conventional 2DEG when subject to additional periodic electric or magnetic field potentials [25]. This is because when subject to an external homogeneous magnetic field, electrons constrained to a wrinkled structure experience an effective periodically modulated magnetic field. More importantly, since the ratio between the effective homogeneous and inhomogeneous component of the magnetic field can be tuned on demand by simply rotating the externally applied homogeneous magnetic field, we find the Weiss oscillations to be strongly anisotropic, and consequently appear also in the angular dependence of the MR.

2. Results and discussion

2.1. Quantum theory of a nanocorrugated electron gas with an external magnetic field

To investigate the electronic properties of a gas of electrons constrained to a corrugated quasi-two-dimensional nanomembrane, we start out by approximating the corresponding curved surface \( S \) with a sinusoidal shape whose parametric equations in Euclidean coordinates read \( r = \{ x, y(x), z \} \equiv \{ x, a \cos(2\pi x / \lambda), z \} \), where \( a \) represents the maximum wrinkle amplitude while \( \lambda \) is its typical wavelength (see figure 1(a)). To proceed further, we use that the effective two-dimensional Schrödinger equation regulating the quantum motion of electrons moving along the curved surface \( S \) can be derived using the so-called thin-wall quantization procedure [26]: it treats the quantum motion on a generic curved surface as the limiting case of a quantum particle in three-dimensional space subject to a strong confining force acting in the direction normal to \( S \). Because of the lateral confinement, quantum excitation energies in the normal direction become much higher than in the tangential direction. Henceforth, one can safely ignore the particle motion in the direction normal to the surface, and, on the basis of this, deduce the effective two-dimensional Schrödinger equation. This procedure, which is the most rigorous and physically sound one for curved nanosystems embedded in the ordinary Euclidean three-dimensional space, simplifies the problem considerably since the surface curvature is eliminated from the Schrödinger equation at the expense of adding to it a potential, dubbed as quantum geometric potential (QGP). The QGP generally cause intriguing phenomena at the nanoscale, such as the emergence of winding-generated bound state in spirally rolled-up nanotubes [10]. Moreover, in strain-driven nanostructures the nanoscale variation of strain induced by curvature yields an additional strain-induced geometric potential (SGP) that is of the same functional form of the QGP, but (often strongly) boosting it [27]. With both the QGP and SGP taken into account, the effective Hamiltonian for a nanocorrugated electron gas takes a form, which, after employing the translational invariance along the \( \hat{z} \) direction and thereby separating the two-dimensional wavefunction as \( \Psi(x, z) = \psi(x) \times e^{ik_z z} \), reads

\[
\mathcal{H}^0 = -\frac{\hbar^2}{2m^*} \frac{\partial^2}{\partial x^2} + \frac{\partial_k}{\sqrt{g_{xx}}} + v_G \mathcal{V}_G + \frac{\hbar^2 k_z^2}{2m^*}
\]

In the equation above, \( m^* \) is the electron effective mass, \( g_{xx} = 1 + [\partial_y y(x)]^2 \) is the only non-trivial metric tensor component of the curved surface \( S \), whereas the functional form of the geometric potential is
\[
V_G = -\frac{\hbar^2}{8m^*g_{xz,z}} \left( \partial_x^2 y(x) \right)^2.
\]

The prefactor \( v_R \) characterizing the strength of the geometric potential in equation (1) measures the renormalization of the QGP due to strain effects. It is therefore a material-dependent parameter, which can be estimated to be \( v_R = 6 \) in GaAs heterostructures, and \( v_R = 600 \) in Si inversion layers assuming a 2 nm effective thickness [28].

It has been recently shown that the thin-wall quantization procedure can be safely formulated even in the presence of externally applied electric and magnetic fields [29, 30], which allows the investigation of unique curvature-induced effects on the magnetotransport properties of a 2DEG [11, 28]. We therefore consider the effect of an externally applied homogeneous magnetic field, which is orthogonal to the translationally invariant direction of the nanomembrane, and forms an angle \( \theta \) with the \( \hat{x} \) direction (see figure 1(a)), i.e. \( B = B(\cos \theta, \sin \theta, 0) \). We next fix the gauge in which the corresponding vector potential reads: \( A = (0, 0, A_z(\theta)) \) where \( A_z(\theta) = B(y \cos \theta - x \sin \theta) \). Since in this gauge the vector potential does not have a finite component in the direction normal to the curved surface \( S \), we can straightforwardly apply the framework of [29, 30], thereby obtaining the effective two-dimensional Hamiltonian:

\[
\mathcal{H} = -\frac{\hbar^2}{2m^*} \partial_x^2 + \frac{\hbar^2}{2m^*} (k_x - eA_z(\theta))^2 + v_R V_G.
\]

The Hamiltonian in the equation above can be interpreted as the effective one-dimensional Hamiltonian regulating the quantum motion of a particle constrained to a serpentine-shaped waveguide in the presence of both an attractive potential generated by the geometric curvature, and a \( k_x \)-dependent magnetic potential. Although both these two potential should be treated on an equal footing, we note that for corrugated nanomembranes with typical wavelengths of the order of a few hundreds of nanometers the geometric potential can be safely disregarded for \( v_R \approx 10^9 \). This is because the Landau magnetic length \( \approx \sqrt{26/\sqrt{B} \text{nm}} \) corresponding to an external magnetic field of a few Tesla is much smaller than the corrugation period, thereby implying that the amplitude of the QGP is more than two orders of magnitude smaller than the magnetic potential. In the remainder of this article, we will consider a corrugated 2DEGs formed at GaAs-based heterostructures where, as mentioned above, \( v_R \approx 6 \), and therefore the last term in equation (3) can be neglected.

To make further progress, we rewrite the effective Hamiltonian by introducing the arclength of the sinusoidal profile measured from \( x = 0 \), \( s = \int_0^x \sqrt{g_{xx}} \, dx \) [10], in terms of which equation (3) can be recast as

\[
\mathcal{H} = -\frac{\hbar^2}{2m^*} \left( \frac{1}{\sqrt{g_{xx}}} \frac{d}{ds} \right)^2 + \frac{\hbar^2}{2m^*} (k_x - eA_z(\theta))^2,
\]

with

\[
A_z(\theta) \approx B \left[ a \cos \frac{2\pi s}{\lambda} \cos \theta - s \sin \theta \right].
\]

In the equation above, we have used that in the shallow deformation limit \( a/\lambda \ll 1 \), \( s \approx x \). Most importantly, equations (4), (5) correspond to the Hamiltonian in the effective mass approximation for a 2DEG subject to a perpendicular magnetic field

\[
B(s) = B \left[ 2\pi a \sin \frac{2\pi s}{\lambda} \cos \theta + s \sin \theta \right].
\]

For a given angle \( \theta \), the effective field in equation (6) has an homogeneous component of strength \( B_0 = B \sin \theta \), as well as an inhomogeneous component \( B_1 = (2\pi B a / \lambda) \cos \theta \). As a result, depending on the direction of the externally applied homogeneous magnetic field, the effective field felt by electrons constrained to the corrugated nanomembrane changes from being purely homogeneous for \( \theta = \pi/2 \) to a zero-average periodic magnetic field for \( \theta = 0 \) (see figure 1(b)). In the next section, we will analyze the classical magnetic trajectories [31] that are thereby realized by rotating the external magnetic field.

2.2. Classical electron trajectories

We first analyze the classical electron trajectories in a nanocorrugated 2DEG subject to an homogeneous external magnetic field. Since, as demonstrated above, we can switch from the native curved system to a flat one in which the effective magnetic field both possesses an homogenous and an inhomogeneous component, the electron trajectories can be obtained, in perfect analogy with [32], by solving the classical equation of motion \( m^* \ddot{v} = -e v \times B(s) \) with \( B(s) \) given in equation (6). When the inhomogeneous component vanishes, i.e. assuming an external magnetic field that points along the \( \hat{y} \) axis (\( \theta = \pi/2 \)) in figure 1(a), the classical
trajectories simply corresponds to cyclotron orbits with a radius \( R_{\text{cycl}} = m^* v / (eB_0) \) that is inversely proportional to the magnetic field strength (see figures 2(a), (d)).

By rotating the magnetic field away from \( \hat{y} \) direction, the finite inhomogeneous component of the effective magnetic field leads to two different types of characteristic electron orbits depending on whether or not the angle-dependent mean cyclotron radius \( \bar{R}_{\text{cycl}} = m^* v / (eB_0 \sin \theta) \) exceeds the wrinkle wavelength \( \lambda \). In the weak-field regime, i.e. for \( \bar{R}_{\text{cycl}} > \lambda \), the inhomogeneous component of the magnetic field yields a drift of the electron orbit center which is simply superimposed to the unperturbed cyclotron motion due to the homogeneous part of the magnetic field. In complete analogy with the situation encountered in a 2DEG with an electrostatic potential grating \([33, 34]\), this guiding-center drift is oscillatory and can be neglected except when it resonates with the periodic cyclotron motion (see figure 2(b)). In the \( R_{\text{cycl}} < \lambda \) regime, instead, the homogeneous component of the magnetic field is strong enough to localize the electrons in different space regions of the wrinkles. More specifically, close to the regions where the gradient of the effective magnetic field is zero, and thus the local curvature of the wrinkle vanishes, the electronic trajectories form usual cyclotron orbits whose radius depends on the local magnetic field strength. Close to the valleys and crests of the wrinkles, instead, the gradient of the effective magnetic field is maximal and thus the cyclotron orbits acquire a strong drift along the \( \hat{z} \) and the \(-\hat{z}\) direction respectively (see figure 2(e)).

When the externally applied homogeneous magnetic field points exactly towards the \( \hat{x} \) direction in figure 1(a) the situation changes drastically. The homogeneous component of the effective magnetic field identically vanishes and thus in the weak-field regime, the electron orbits corresponds to slightly perturbed transverse orbits (see figure 2(c)). Cranking up the strength of the magnetic field to the regime in which the characteristic cyclotron radius \( \bar{R}_{\text{cycl}} = m^* v / (eB_1) \) is smaller than the wrinkle wavelength yields electronic orbits strongly resembling the ones encountered for \( \theta < \pi / 2 \) but with the following caveat: since the effective magnetic field averages to zero, the drifting cyclotron orbits of figure 2(e) are substituted by snake states with

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**Figure 2.** Top panels: effective magnetic fields felt by the electrons in a corrugated two-dimensional electron gas as obtained by changing the direction of an externally applied magnetic field. Middle panels: in the weak-field regime the usual cyclotron orbits obtained for a magnetic field direction \( \theta = \pi / 2 \) (a) acquire a guiding-center drift which is reinforced under resonance condition (b). For the magnetic field direction \( \theta = 0 \) the absence of an homogeneous component of the effective magnetic field yields instead slightly perturbed transverse orbits (c). Bottom panels: in the strong-field regime small radii cyclotron orbits (d) can coexist with drifting orbits (e) and snake states with reversed direction (f) localized at the crests and valleys of the wrinkles in the presence of an inhomogeneous component of the effective magnetic field.
opposite velocity similarly to the situation encountered in tubular nanostructures subject to a transversal magnetic field [12, 35].

2.3. Angle-dependent Weiss oscillations

Having established the nature of the classical electron trajectories, we now analyze their impact on the magnetotransport properties of a nanocorrugated 2DEG subject to an externally applied magnetic field. To do so, we first solve the magnetic spectra by diagonalizing the effective Hamiltonian equation (4) using as complete basis the Landau levels states due to the homogeneous field \( B_0 \) [36, 37]. Thereafter, we compute the dc conductivity reading [38]

\[
\sigma_{ij} = \frac{i\hbar e^2}{A} \sum_{\xi, \xi'} \int dE \int dE' P(E - E_\xi) P(E' - E_\xi') \times (f_{E'} - f_E) \frac{\langle \xi|v|\xi' \rangle \langle \xi'|v|\xi \rangle}{(E - E')^2},
\]

where \( i, j = s, z, |\xi\rangle = |n, k_z\rangle \) is an eigenstate with subband index \( n \) and longitudinal momentum \( k_z \), \( A \) is the area of the sample, \( v \) is the group velocity operator, and \( f_E \) is the Fermi–Dirac distribution function. In order to explicitly take into account the impurity broadening effect on the conductivity, which is important at low temperatures, we introduced an impurity broadening \( P(E) = (\Gamma / \pi) / (E^2 + \Gamma^2) \) with a constant width \( \Gamma = \hbar / \tau \), \( \tau \) being a constant relaxation time. Equation (7) explicitly accounts for both the diagonal and the non-diagonal diffusion contributions to the dc conduction but disregards the collisional conductivity describing transport through localized states due to impurity scattering. The latter, however, is mainly responsible for the Shubnikov–de Haas oscillations [38], and gives only a subleading contribution for the occurrence of the Weiss oscillations investigated in this study. Therefore, we will only account for the diffusional part of the dc conduction in our system. We emphasize that previous works on conventional 2DEG subject to additional modulated magnetic fields have accounted only for the diagonal diffusion contribution to the dc conductance [36, 37, 39]. This is a well justified assumption considering a magnetic modulation weak as compared to the external homogeneous magnetic field. In our case, instead, the off-diagonal contribution is essential in the small \( \theta \) regime, where the 2DEG is in a lateral magnetic superlattice [40] perturbed by a weak homogeneous magnetic field.

To obtain the conductivity tensor components from equation (7), we fix the temperature to \( T = 3.66K \), use the relaxation time \( \tau = 3.66 \times 10^{-11}s \) [38], and introduce a cutoff in the subband index \( n = 200 \), thereby reaching an accuracy within 1%. The resistivity tensor is thereafter found by inverting the conductivity tensor. Figure 3 shows the ensuing behavior of the MR in the direction orthogonal to the wrinkles \( \Delta \rho_{ij}/\rho_{ij}(B = 0) = \rho_\perp(B)/\rho_\parallel(0) - 1 \) assuming the homogeneous external magnetic field forms an angle \( \theta \approx 30^\circ \) with respect to the \( x \) direction (see figure 1(a)). The presence of both an homogeneous and an inhomogeneous component of the effective magnetic field felt by the electron gas leads, independent of the carrier density, to the presence of Weiss oscillations, which are the prime physical consequence of the guiding-center drift resonance discussed above. These Weiss oscillations, however, are clearly visible for magnetic fields smaller than a critical value \( B_c \propto \sqrt{n} \) with \( n \) the carrier density (see figure 3). In the \( B > B_c \) regime, indeed, we observe a strong positive diffusive MR (see the inset of figure 3), which is due to the fact that the inhomogeneous

![Figure 3](image-url)
component of the effective magnetic field is comparable in magnitude to the homogeneous one. We point out that this feature is unique of our system and is not generally realized in conventional 2DEG with additional ferromagnetic or superconducting strips where the characteristic strength of the modulated magnetic field is much smaller than the homogeneous one [38].

The tunability of the ratio between the effective inhomogeneous magnetic field strength and the homogeneous one, as obtained by simply rotating the direction of the external magnetic field, also yields a strong directional dependence of the magnetotransport properties. This is demonstrated in figures 4(a), (b) where we show the angle dependence of the magnetoresistance $\Delta \rho(\theta)/\rho(\pi/2) = 1$ in a nanocorrugated electron gas subject to an homogeneous external magnetic field with strength $B = 4 \, \text{T}$ for three carrier densities corresponding to $n = 0.5, 1, 2 \times 10^{11} \, \text{cm}^{-2}$.

![Figure 4. Angle dependence of the magnetoresistance $\rho_2(\theta)/\rho(\pi/2) = 1$ in a nanocorrugated electron gas subject to an homogeneous external magnetic field with strength $B = 0.4 \, \text{T}$ (a) and $B = 4 \, \text{T}$ (b) for three carrier densities corresponding to $n = 0.5, 1, 2 \times 10^{11} \, \text{cm}^{-2}$.](image)

To conclude, we have shown that the geometry of a nanocorrugated 2DEG can be used to engineer an effective periodically inhomogeneous magnetic field coexisting with an homogeneous one by simply applying a constant external magnetic field. As such, our proposed setup does not necessitate the usage of ferromagnetic or superconducting strips as routinely done in conventional semiconducting heterostructures. The magnetotransport properties of these patterned nanostructures have been extensively theoretically studied [36, 37, 39] in the regime of weak magnetic modulations. The geometrical engineering of modulated magnetic fields presented in this work, instead, can be conveniently used to access regimes where the modulated magnetic field is comparable in strength to the homogeneous one. Such a regime can be also realized in conventional flat nanostructures using magnetic barriers. However, in the latter situation the step-like character of the magnetic field inhomogeneity leads to localized charge densities [41], which are not expected to occur for the smooth magnetic field profile realized in our setup. Finally our setup also allows to mimic a lateral magnetic superlattice [40, 42] perturbed by an homogeneous magnetic field with tunable strength (tilting the magnetic field towards the $\hat{y}$ direction in figure 1(a)). The quadratic positive MR we find in this situation (see figure 4(a)) agrees with experimental observations [43]. We expect that the angular dependence of the Weiss oscillations at weak magnetic field and the strong anisotropic positive MR at strong external fields predicted in our study can be not only tested in wrinkled nanomembranes with embedded quantum wells as obtained by partial release and bond back of layers by selective underetching [22], but also in two-dimensional materials where ripples can be strain-engineered by depositing atomic crystals on prestretched elastomeric substrates [21].
Acknowledgments

We acknowledge the financial support of the Future and Emerging Technologies (FET) programme within the Seventh Framework Programme for Research of the European Commission under FET-Open grant number: 618083 (CNTQC). CO acknowledges support from the Deutsche Forschungsgemeinschaft (Grant No. OR 404/1-1), and from a VIDI grant (Project 680–47–543) financed by the Netherlands Organization for Scientific Research (NWO).

ORCID iDs

Ching-Hao Chang @ https://orcid.org/0000-0003-0878-9328

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