Rotational viscosity in spin resonance of hydrodynamic electrons

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In novel ultrapure materials electrons can form a viscous fluid, which is fundamentally different in its dynamics from the electron gas in ordinary conductors with a significant density of defects. The shape of the nonstationary flow of such an electron fluid is similar to the alternating flow of blood in large-radius arteries [J. R. Womersley, J. Physiol. 127, 553 (1955)]. The rotational viscosity effect is responsible for the interconnection between the dynamics of electron spins and flow inhomogeneities. In particular, it induces the spin polarization of electrons in a curled flow via an internal spin-orbit torque acting on electron spins. Here, we show that this effect in an electron fluid placed in a magnetic field leads to a correction to the ac sample impedance, which has a resonance at the Larmor frequency of electrons. In this way, via the electrically detected spin resonance, the Womersley flow of an electron fluid can be visualized and the rotational viscosity can be measured.

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Introduction. In ultrapure materials with a small density of defects, conduction electrons can form a viscous fluid at low temperatures. The electric transport in such a fluid occurs via the formation of inhomogeneous hydrodynamic flows, controlled by particular shapes of the samples. This idea was put forward many years ago for three-dimensional (3D) metals with strong electron-phonon coupling [1]. Recently, the hydrodynamic regime of electron transport has been realized in high-quality samples of graphene [2-7], the quasi-twodimensional metal PdCoO₂ [8], the Weyl semimetal WP₂ [9], and high-mobility GaAs quantum wells [10–19]. These experiments induced an avalanche of theoretical works (see, for example, Refs. [20-44]), which were aimed at the formulation and search for evidence of a hydrodynamic regime as well as to study various types and regimes of flows of the electron fluid.

An electric current in samples with a noticeable spin-orbit interaction induces various interesting spin-dependent transport effects. It was predicted nearly half a century ago for Ohmic conductors, where the scattering of electrons on disorder dominates, that the spin-orbit interaction results in the interconnection between electrical and spin currents [45,46]. An electric current **j** produces a transverse spin current $q_{ik} \sim \mathbf{j}$, leading to the generation of spin density near the sample edges (the direct spin Hall effect; this term was introduced in Ref. [47]) and, vice versa, inhomogeneous distribution of spin density results in corrections to the electric current (the inverse spin Hall effect). In these phenomena, the relaxation of the electron spin and the spin current are of large importance. The relaxation time of the first process is much longer than the relaxation time of the second one [48]. The action of both direct and inverse spin Hall effects induces a correction to the dc and the ac magnetoresistance in not too wide samples due

to the formation of near-edge layers with the perturbed spin density P [49,50].

In pure metals with a negligible density of defects, where the hydrodynamic regime of charge transport is realized, the effect of the rotational viscosity provides the interconnection between the electric current and spin density. Namely, the vorticity of the electron flow, $\nabla \times \mathbf{j}$, induces the difference in the nondiagonal components of the momentum flux density, $\Pi_{ik} \neq \Pi_{ki}$. Such a difference leads to the generation of the spin density **P** due to conservation of the total angular momentum in electron-electron collisions [51–55]. Similarly to spin Hall effects, the rotational viscosity effect is related to the band spin-orbit interaction of conduction electrons. In particular, an anomalous spin-orbit correction to the electron velocity leads to the mismatch of Π_{ik} and Π_{ki} . In defectless samples, the spin current relaxation rate is of the order of the interparticle scattering time $1/\tau_{ee}$, while the relaxation rate of the spin density is much lower, being typically proportional to $\Omega_{SO}^2 \tau_{ee}$ [56–59], similarly to the case of Ohmic conductors (here, Ω_{SO} is a small frequency of spin precession of conduction electrons due to the band spin-orbit interaction). As a reminder, let us note that in Ohmic materials the relaxation time due to scattering on disorder τ enters the precession spin relaxation rate $\Omega_{SO}^2 \tau$ instead of the time τ_{ee} in defectless materials [60].

Recently, a thorough study of spin generation in hydrodynamic flows of the two-dimensional electron fluid was performed [61]. Various microscopic contributions to the interconnection between the spin current, the spin density, and the electric current were calculated and compared. However, the relation between the rotational viscosity effect and microscopic mechanisms studied in Ref. [61] (for a particular geometry of the Poiseuille flow) remains unclear up to now. In this way, it is of interest to propose experimental methods to determine the rotational viscosity coefficient and/or other kinetic coefficients responsible for the mutual transformation

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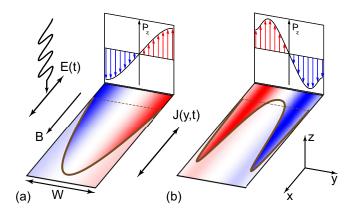


FIG. 1. Dynamic generation of the spin polarization in a hydrodynamic flow of two-dimensional electrons in the presence of a lateral magnetic field and ac electric fields. The polarization density (red and blue) and current density (brown) profiles are shown for (a) the quasistatic Poiseuille [62] and (b) the strongly nonstationary Womersley [63] regimes.

of the currents and the spin density in hydrodynamic conductors.

In this Letter we demonstrate that, under a high-frequency drive, a viscous flow of a two-dimensional electron fluid exhibits electron spin resonance related to the dynamics of a spin-polarized fluid in a weak magnetic field. The rotational viscosity effect leads to dynamic spin-vorticity coupling, namely, the spin density generated by a viscous flow with a nonzero vorticity affects the flow itself, resulting in a correction to the charge current density. In this way, we propose a purely electric method of observing the spin resonance in hydrodynamic conductors, for example, in high-mobility GaAs quantum wells and graphene. The studied effect may allow to measure the rotational viscosity of the electron fluid in these materials.

High-frequency hydrodynamics of spin-polarized electron fluid. We consider a flow of a two-dimensional electron fluid in a defectless sample (see Fig. 1) in an external radio-frequency electric field $\mathbf{E}_0(t)$ and magnetic field \mathbf{B} . We imply that the electrons form a Fermi gas. If the sample edges are rough and the scattering of electrons on them is diffusive, a nonstationary inhomogeneous viscous flow with the particle flow density $\mathbf{j}_0(\mathbf{r},t) \sim \mathbf{E}_0$ is formed in the sample due to the shear viscosity effect [32,40]. Our aim is to calculate the spin polarization of electrons $\mathbf{P}(\mathbf{r},t)$ in such a flow, induced by the rotational viscosity, and to find the spin-related correction to the current density, $\mathbf{J}_2(\mathbf{r},t) = e\,\mathbf{j}_2(\mathbf{r},t)$, and to the total electric current $\mathbf{I}_2(t) = \int_{\Sigma} dS_{\mathbf{r}}\,\mathbf{J}_2(\mathbf{r},t)$ (here, e < 0 is the electron charge and $dS_{\mathbf{r}}$ denotes the differential along the one-dimensional section of the sample Σ).

In such a system the interconnection between **J** and **P** is controlled, in particular, by the rotational viscosity effect. This effect is microscopically induced by the band spin-orbit interaction of conduction electrons, leading to spin-dependent interparticle scattering and an anomalous correction to the electron velocities [51,55,61]. These two processes establish the connection between the antisymmetric part of the momentum flux tensor Π_{ik} , $i \neq k$, the curl of the flow, $\nabla \times \mathbf{j}$, which can be interpreted as the local rotational frequency $\omega_{\text{orb}}(\mathbf{r})$

of the fluid, and the spin polarization density **P** [51,55]. The particular form of this connection is dictated by the symmetry of the system [64,65]. Rotational viscosity is the dissipative kinetic coefficient which interconnects the parts of the nonequilibrium thermodynamic flux (Π_{ik}) and forces (**P** and $\partial_i j_k$) transforming under symmetry operations according to the same one-dimensional irreducible representation [65]. For isotropic systems with O(3) symmetry this interconnection has the form [55,64,65]

$$\frac{\Pi_{ik}^a}{m} = \frac{\Pi_{ik} - \Pi_{ki}}{2m} = \eta_r \epsilon_{ikl} \left([\nabla \times \mathbf{j}] - \frac{\mu (\mathbf{P} - \mathbf{P}_0)}{\hbar} \right)_l. \quad (1)$$

For the two-dimensional fluid considered here we use the components of this equation describing the flows in the xy plane and three-dimensional spin polarization. Here, m is the electron mass, η_r is the kinematic rotational viscosity coefficient, ϵ_{ikl} is the unit antisymmetric tensor, μ is the total chemical potential of degenerate electrons, $\mathbf{P}_0 = \mathbf{S}_0/n_0 =$ $\chi \mathbf{B}/n_0$ is the equilibrium spin polarization, χ is the paramagnetic spin susceptibility of the electrons, and n_0 is the total electron density. In Eq. (1) the product μP is the so-called spin chemical potential [55] and thus $\mu P/\hbar = (\mu_+ - \mu_-)/\hbar$ is the effective frequency of the spin rotation of electrons at the spin-split Fermi surfaces (here, μ_{\pm} are their chemical potentials and the temperature is considered to be sufficiently low, $T \ll \mu_+ - \mu_-$). Microscopic calculations show [55] that in A_3B_5 quantum wells the rotational viscosity coefficient η_r is proportional to the squared Rashba coupling constant.

The hydrodynamic equations for the flow density \mathbf{j} , the spin polarization \mathbf{P} , and the spin current q_{ik} taking into account the shear viscosity and the rotational viscosity effects in a weak magnetic field can be written as

$$\frac{\partial \mathbf{j}}{\partial t} = \frac{e\mathbf{E}_0(t)}{m} n_0 + \eta \, \Delta \, \mathbf{j} - c_s^2 \frac{\eta_r}{\eta_0} [\nabla \times \mathbf{P}],$$

$$\frac{\partial P_k}{\partial t} + \frac{\partial q_{ik}}{\partial x_i} + [\mathbf{\Omega} \times \mathbf{P}]_k = -\frac{P_k - P_{0,k}}{\tau_s} + 2\frac{\eta_r}{\eta_0} [\nabla \times \mathbf{j}]_k,$$

$$\frac{\partial q_{ik}}{\partial t} = -c_s^2 \frac{\partial P_k}{\partial x_i} - \frac{q_{ik}}{\tau_{sc}},$$
(2)

where η is the kinematic shear viscosity coefficient, $c_s = v_F/\sqrt{2}$ is the velocity of sound, $\eta_0 = \hbar/(2m)$ is the characteristic value of the dimension of viscosity, the "quantum viscosity" [42], Ω is the Larmor frequency due to the applied magnetic field B, τ_s is the spin relaxation time related, for example, to the precession of the electron spins in the spin-orbit effective magnetic field during the electron motion between electron-electron collisions [56,57] (for this mechanism $1/\tau_s \sim \Omega_{SO}^2 \tau_{ee} \ll 1/\tau_{ee}$), and τ_{sc} is the spin current relaxation time in electron-electron collisions due to spin Coulomb drag [58,59] (it was shown that $\tau_{sc} \sim \tau_{ee}$ in those works). The form of Eqs. (2) follows from consideration of Refs. [51,55]

The first line in Eqs. (2) is the hydrodynamic Navier-Stokes equation for the evolution of the mean electron momentum, containing the shear viscosity term and the rotational viscosity term. That is, the momentum flux tensor term, $-\partial \Pi_{il}/\partial x_l$, accounts for the two contributions, $\Pi_{ik} = \prod_{ik}^s + \prod_{ik}^a$, where $\prod_{ik}^s = -m \eta(\partial j_i/\partial x_k + \partial j_k/\partial x_i)$ is the usual shear part of Π_{ik}

for incompressible flow, while Π_{ik}^a is the rotational viscosity contribution given by Eq. (1).

The second line of Eqs. (2) describes the evolution of the electron spin, namely, its transfer by the spin current q_{ik} , rotation described by the Larmor frequency Ω , relaxation to the equilibrium value $\mathbf{S}_0 = n_0 \mathbf{P}_0$, and the generation by the torque $T_k = \epsilon_{klq} \Pi_{lq}^a / m$ associated with the antisymmetric part of Π_{ik} (1). We imply that the relaxation term also contains the contribution from Π_{ik}^a

The third line of Eqs. (2) describes the evolution of the spin current q_{ik} . Provided that the spin current relaxation length $v_F \tau_{sc}$ is sufficiently short, one can neglect the dissipative transfer of the spin flux in this equation (unlike the dissipative transfer of electron momentum described by Π_{ik}). Additionally, we omit possible contributions to the local variation of the spin current, $\partial q_{ik}/\partial t$, from the particle current \mathbf{j} , which are crucial for the spin Hall effect in Ohmic conductors [45,45,48].

We consider slow flows with the frequencies $\omega \ll 1/\tau_{ee}$, thus the time dispersion of the viscosities η and η_r is insignificant. In this case, from the third line of Eqs. (2) we obtain the direct connection between the spin polarization and the spin current, $q_{ik} = -D_s \partial P_k/\partial x_i$, where $D_s = c_s^2 \tau_{sc}$ is the spin diffusion coefficient.

The boundary condition describing the absence of spin current at the closed part of the sample boundary ∂G takes the form $(\partial P_k/\partial n)|_{\partial G} = 0$, where $\mathbf{n} = \mathbf{n}(\mathbf{r})$ is the normal to the curve ∂G . The diffusive boundary condition at the closed part of the edges has the usual form, $\mathbf{j}|_{\partial G} = \mathbf{0}$.

Spin resonance due to rotational viscosity. First, we study the solutions of Eqs. (2) in a long sample. The magnetic and the electric field both are directed along the sample: $\mathbf{E}_0(t) = \mathbf{E}_0 e^{-i\omega t} + \text{c.c.}$, $\mathbf{E}_0 = E_0 \mathbf{e}_x$, $\mathbf{B} = B \mathbf{e}_x$. In this geometry, all values are homogeneous along the x coordinate and depend on the coordinate y along the sample section: $P_y(y, t)$, $P_z(y, t)$, $j = j_x(y, t)$ (see Fig. 1). The spin current tensor has two nonzero components: q_{xy} and q_{xz} .

The hydrodynamic Eqs. (2) take the form

$$\left(-i\omega - \eta \frac{d^2}{dy^2}\right)j = f_0 + r_1 \frac{dP_z}{dy},$$

$$\left(-i\omega + \frac{1}{\tau_s} - D_s \frac{d^2}{dy^2}\right)P_{\pm} \pm i\Omega P_{\pm} = r_2 \frac{dj}{dy},$$
(3)

where $P_{\pm}=P_z\pm iP_y$, $f_0=eE_0n_0/m$, $r_1=c_s^2\,\eta_r/\eta_0$, $r_2=2\eta_r/\eta_0$. The boundary conditions are formulated at the longitudinal edges: $j|_{y=\pm W/2}=0$ and $(dP_{\pm}/dy)|_{y=\pm W/2}=0$.

The first line of Eqs. (3) with $\eta_r = 0$ yields for the unperturbed particle flow $j(y, t) = j(y)e^{-i\omega t} + \text{c.c.}$, where

$$j(y) = f_0 \frac{i}{\omega} \left[1 - \frac{\cosh(\kappa y)}{\cosh(\kappa W/2)} \right], \quad \kappa = \sqrt{-i\frac{\omega}{\eta}}.$$
 (4)

Owing to the formula $\eta = v_F^2 \, \tau_{2,ee}/4$, where $\tau_{2,ee}$ is the shear stress relaxation time, we arrive at an estimate of the eigenvalue determining the characteristic width $L_\omega = 1/\text{Re}\,\kappa$ of the near-edge layers of nonstationary Womersley flow [63]: $\kappa \sim \sqrt{\omega/\tau_{ee}}/v_F$. In the static limit $\omega \to 0$, the particle flow profile (4) reduces to Poiseuille flow [62].

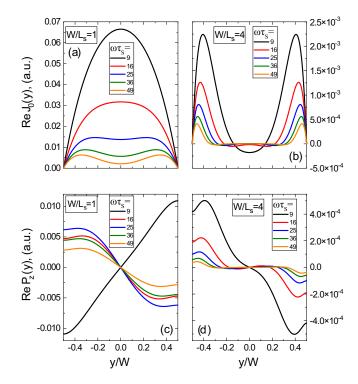


FIG. 2. (a), (b) Profiles of the amplitude $J_0(y)$ of the unperturbed electric current density $J_0(y,t)=J_0(y)e^{-i\omega t}+\text{c.c.}$ in a long sample in the parallel magnetic and electric field $\mathbf{E}_0(t)$ and \mathbf{B} for different parameters W/L and $\omega \tau_s$. Both values J_0 and P_z are plotted in arbitrary units. (c), (d) Profiles of the amplitude of the spin polarization $P_z(y,t)=P_z(y)e^{-i\omega t}+\text{c.c.}$ in the electron fluid, arising due to the rotational viscosity effect. The curves are plotted for the same sample at the same parameters as shown in (a) and (b).

The eigenvalues of the unperturbed equations for the spin polarization [the second line of Eqs. (3) with $r_2 = 0$] are

$$\lambda_{\pm} = (1/L_s)\sqrt{1 + i(\pm\Omega - \omega)\tau_s},\tag{5}$$

where $L_s = \sqrt{D_s \tau_s}$ is the spin diffusion length. The relation between the shear viscosity and the spin characteristic lengths L_{ω} and $L_{\lambda} = 1/\text{Re }\lambda_{\pm}$ is as follows: $L_{\omega} \sim L_{\lambda}$ far from the resonance frequency, $|\pm \Omega - \omega|\tau_s \gg 1$, and $L_{\omega} \ll L_{\lambda}$ near the resonance, $|\pm \Omega - \omega|\tau_s \sim 1$.

Circular components of the spin polarization generated by unperturbed flow (4) are first order by η_r and have the form

$$P_{\pm}(y) = \frac{i f_0}{\omega} r_2 \frac{t_{\lambda_{\pm}}(y) - t_{\kappa}(y)}{\lambda_{+}^2 - \kappa^2},$$
 (6)

where $t_{\xi}(y) = \sinh(\xi y)/[\xi \cosh(\xi W/2)]$. The profiles of the electric current $J_0(y) = ej_0(y)$ and the z component $P_z(y)$ of the spin polarization corresponding to Eq. (6) are drawn in Fig. 2.

Correction to the particle flow $j^{(2)}(y,t) = j^{(2)}(y)e^{-i\omega t} +$ c.c. generated by nonzero spin polarization (6) is second order by η_r and the equation for it is

$$\left(-i\omega - \eta \frac{d^2}{dy^2}\right)j^{(2)} = r_1 \frac{dP_z}{dy},\tag{7}$$

where $P_z = P_z^{(1)}$ is given by the sum of circular components P_{\pm} after (6). The result of its solution is $j^{(2)} = (j_+^{(2)} + j_-^{(2)})/2$,

where

$$j_{\pm}^{(2)}(y) = r_1 r_2 \frac{if_0}{\omega} \frac{1}{\lambda_{\pm}^2 - \kappa^2} \times \left\{ \frac{c_{\lambda_{\pm}}(y) - c_{\kappa}(y)}{\lambda_{\pm}^2 - \kappa^2} - \frac{F_{\kappa}}{8} \left[s_{\kappa}(y) \frac{y}{W/2} - c_{\kappa}(y) \right] \right\}.$$
(8)

Here, the notations $s_{\xi}(y) = \sinh(\xi y)/[\sinh(\xi W/2)]$, $c_{\xi}(y) = \cosh(\xi y)/[\cosh(\xi W/2)]$, and $F_{\xi} = \tanh(\xi W/2)/(\xi W/2)$ are used.

The main part of the total current $I(t)=e\int_{-W/2}^{W/2}dy\,j(y,t)$ associated with (4) is given by $I_0(t)=I_0e^{-i\omega t}+\text{c.c.}$, where the amplitude is $I_0=(i\,e\,f_0W/\omega)[1-F(\kappa)]$. The spin-orbit correction $I^{(2)}(t)$ to it corresponding to $j^{(2)}(y)$ (8) has the amplitude

$$\frac{I^{(2)}}{W} = r_1 r_2 \frac{ief_0}{\omega} \sum_{+} \frac{1}{\lambda_{\pm}^2 - \kappa^2} \left(\frac{F_{\lambda_{\pm}} - F_{\kappa}}{\lambda_{\pm}^2 - \kappa^2} - \frac{F_{\kappa}'}{2\kappa} \right). \tag{9}$$

It is seen from Eqs. (5) and (9) that Re $I^{(2)}$ as a function of magnetic field exhibits spin resonance for sufficiently narrow samples. This resonance originates from the formation of the near-edge layers with the perturbed spin polarization $\mathbf{P}(y, t)$, which induces the correction $I^{(2)}(y, t)$ to the current $I_0(t)$.

The real part of the correction to the mean sample impedance, $\Delta Z = E_0 W/(I_0 + I_2) - E_0 W/I_0$, is plotted in Fig. 3. This value is proportional to the squared rotational viscosity, $r_1 r_2 \sim \eta_r^2$ [see Eq. (9)]. We see from Fig. 3 that the higher is the ac frequency and the wider is the sample, the narrower are the resonances at $\Omega = \pm \omega$. The limiting half width at $W \gg L_s$ is estimated as $1/\tau_s$. The relative value of the correction to the impedance $\zeta = \text{Re } \Delta Z(\Omega, \omega)/\text{Re } Z(\omega) = -\text{Re } I_2/\text{Re } I_0$ in sufficiently wide samples, $W \gg L_s$ is independent of the sample width. The frequency dependence of ζ is determined by the values $\kappa(\omega)$ and $\lambda_{\pm}(\omega, \Omega)$ [see Eq. (9)].

The measurement of $\zeta(\Omega)$ at fixed ω , together with fitting of the whole curves $\zeta(\Omega)$ (see Fig. 3), may allow us to determine the rotational viscosity.

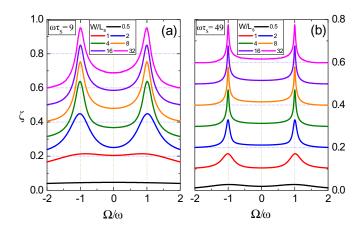


FIG. 3. The relative spin-orbit correction to the real part of the impedance, $\zeta = \text{Re }\Delta Z(\Omega,\omega)/\text{Re }Z(\omega)$, for the two-dimensional electron fluid in a long sample due to the rotational viscosity effect. This value is quadratic by the rotational viscosity coefficient: $\Delta Z \sim \eta_r^2$. The curves are plotted in arbitrary units for the different parameters W/L depicted on (a) for low and (b) high frequencies. For better visibility, the neighbor curves are shifted in the vertical direction by 0.1 at infinity, that leads to the almost same shifts at maximal values of Ω/ω , $\Omega/\omega = \pm 2$.

Conclusion. We have shown that the paramagnetic resonance of the conduction electrons can be observed in nonstationary viscous flow of two-dimensional electron fluid owing to the rotational viscosity effect. The resonance manifests itself in the spin-orbit correction to the sample impedance. Measurements of this resonance may allow us to determine the rotational viscosity coefficient of the viscous electron fluid.

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^[1] R. N. Gurzhi, Sov. Phys. Usp. 11, 255 (1968).

^[2] D. A. Bandurin, I. Torre, R. K. Kumar, M. Ben Shalom, A. Tomadin, A. Principi, G. H. Auton, E. Khestanova, K. S. Novoselov, I. V. Grigorieva, L. A. Ponomarenko, A. K. Geim, and M. Polini, Science 351, 1055 (2016).

^[3] R. Krishna Kumar, D. A. Bandurin, F. M. D. Pellegrino, Y. Cao, A. Principi, H. Guo, G. H. Auton, M. Ben Shalom, L. A. Ponomarenko, G. Falkovich, K. Watanabe, T. Taniguchi, I. V. Grigorieva, L. S. Levitov, M. Polini, and A. K. Geim, Nat. Phys. 13, 1182 (2017).

^[4] A. I. Berdyugin, S. G. Xu, F. M. D. Pellegrino, R. Krishna Kumar, A. Principi, I. Torre, M. Ben Shalom, T. Taniguchi, K. Watanabe, I. V. Grigorieva, M. Polini, A. K. Geim, and D. A. Bandurin, Science 364, 162 (2019).

^[5] L. Levitov and G. Falkovich, Nat. Phys. 12, 672 (2016).

^[6] J. A. Sulpizio, L. Ella, A. Rozen, J. Birkbeck, D. J. Perello, D. Dutta, M. Ben-Shalom, T. Taniguchi, K. Watanabe, T. Holder, R. Queiroz, A. Principi, A. Stern, T. Scaffidi, A. K. Geim, and S. Ilani, Nature (London) 576, 75 (2019).

^[7] M. J. H. Ku, T. X. Zhou, Q. Li, Y. J. Shin, J. K. Shi, C. Burch, L. E. Anderson, A. T. Pierce, Y. Xie, A. Hamo, U. Vool, H. Zhang, F. Casola, T. Taniguchi, K. Watanabe, M. M. Fogler, P. Kim, A. Yacoby, and R. L. Walsworth, Nature (London) 583, 537 (2019).

^[8] P. J. W. Moll, P. Kushwaha, N. Nandi, B. Schmidt, and A. P. Mackenzie, Science 351, 1061 (2016).

^[9] J. Gooth, F. Menges, N. Kumar, V. Süß, C. Shekhar, Y. Sun, U. Drechsler, R. Zierold, C. Felser, and B. Gotsmann, Nat. Commun. 9, 4093 (2018).

- [10] A. T. Hatke, M. A. Zudov, J. L. Reno, L. N. Pfeiffer, and K. W. West, Phys. Rev. B 85, 081304(R) (2012).
- [11] R. G. Mani, A. Kriisa, and W. Wegscheider, Sci. Rep. 3, 2747 (2013).
- [12] L. Bockhorn, P. Barthold, D. Schuh, W. Wegscheider, and R. J. Haug, Phys. Rev. B 83, 113301 (2011).
- [13] Q. Shi, P. D. Martin, Q. A. Ebner, M. A. Zudov, L. N. Pfeiffer, and K. W. West, Phys. Rev. B 89, 201301(R) (2014).
- [14] P. S. Alekseev, Phys. Rev. Lett. 117, 166601 (2016).
- [15] G. M. Gusev, A. D. Levin, E. V. Levinson, and A. K. Bakarov, AIP Adv. 8, 025318 (2018).
- [16] A. D. Levin, G. M. Gusev, E. V. Levinson, Z. D. Kvon, and A. K. Bakarov, Phys. Rev. B 97, 245308 (2018).
- [17] G. M. Gusev, A. D. Levin, E. V. Levinson, and A. K. Bakarov, Phys. Rev. B 98, 161303(R) (2018).
- [18] A. C. Keser, D. Q. Wang, O. Klochan, D. Y. H. Ho, O. A. Tkachenko, V. A. Tkachenko, D. Culcer, S. Adam, I. Farrer, D. A. Ritchie, O. P. Sushkov, and A. R. Hamilton, Phys. Rev. X 11, 031030 (2021).
- [19] A. Gupta, J. J. Heremans, G. Kataria, M. Chandra, S. Fallahi, G. C. Gardner, and M. J. Manfra, Phys. Rev. Lett. 126, 076803 (2021).
- [20] T. Scaffidi, N. Nandi, B. Schmidt, A. P. Mackenzie, and J. E. Moore, Phys. Rev. Lett. 118, 226601 (2017).
- [21] H. Guo, E. Ilseven, G. Falkovich, and L. S. Levitov, Proc. Natl. Acad. Sci. USA 114, 3068 (2017).
- [22] A. Lucas, Phys. Rev. B 95, 115425 (2017).
- [23] F. M. D. Pellegrino, I. Torre, and M. Polini, Phys. Rev. B 96, 195401 (2017).
- [24] P. S. Alekseev, I. V. Gornyi, A. P. Dmitriev, V. Y. Kachorovskii, and M. A. Semina, Semiconductors 51, 766 (2017).
- [25] A. Lucas and K. C. Fong, J. Phys.: Condens. Matter **30**, 053001 (2018).
- [26] P. S. Alekseev, A. P. Dmitriev, I. V. Gornyi, V. Y. Kachorovskii, B. N. Narozhny, M. Schütt, and M. Titov, Phys. Rev. B 95, 165410 (2017).
- [27] P. S. Alekseev, A. P. Dmitriev, I. V. Gornyi, V. Y. Kachorovskii, B. N. Narozhny, and M. Titov, Phys. Rev. B 97, 085109 (2018).
- [28] P. S. Alekseev, A. P. Dmitriev, I. V. Gornyi, V. Y. Kachorovskii, B. N. Narozhny, and M. Titov, Phys. Rev. B 98, 125111 (2018).
- [29] P. S. Alekseev and M. A. Semina, Phys. Rev. B 98, 165412 (2018).
- [30] P. S. Alekseev and M. A. Semina, Phys. Rev. B 100, 125419 (2019).
- [31] O. Kashuba, B. Trauzettel, and L. W. Molenkamp, Phys. Rev. B 97, 205129 (2018).
- [32] R. Moessner, P. Surówka, and P. Witkowski, Phys. Rev. B 97, 161112(R) (2018).
- [33] M. Semenyakin and G. Falkovich, Phys. Rev. B 97, 085127 (2018).

- [34] A. Lucas and S. Das Sarma, Phys. Rev. B 97, 115449 (2018).
- [35] R. Cohen and M. Goldstein, Phys. Rev. B 98, 235103 (2018).
- [36] P. S. Alekseev, Phys. Rev. B 98, 165440 (2018).
- [37] T. Holder, R. Queiroz, T. Scaffidi, N. Silberstein, A. Rozen, J. A. Sulpizio, L. Ella, S. Ilani, and A. Stern, Phys. Rev. B 100, 245305 (2019).
- [38] J. Y. Khoo and I. S. Villadiego, Phys. Rev. B 99, 075434 (2019).
- [39] P. S. Alekseev, Semiconductors **53**, 1367 (2019).
- [40] P. S. Alekseev and A. P. Alekseeva, Phys. Rev. Lett. 123, 236801 (2019).
- [41] P. S. Alekseev and A. P. Dmitriev, Phys. Rev. B 102, 241409(R) (2020).
- [42] K. Trachenko and V. V. Brazhkin, Sci. Adv. 6, eaba3747 (2020).
- [43] A. N. Afanasiev, P. S. Alekseev, A. A. Greshnov, and M. A. Semina, Semiconductors 55, 562 (2021).
- [44] A. N. Afanasiev, P. S. Alekseev, A. A. Greshnov, and M. A. Semina, Phys. Rev. B 104, 195415 (2021).
- [45] M. I. Dyakonov and V. I. Perel, JETP Lett. 13, 467 (1971).
- [46] M. I. Dyakonov and V. I. Perel, Phys. Lett. A 35, 459 (1971).
- [47] J. E. Hirsch, Phys. Rev. Lett. 83, 1834 (1999).
- [48] M. Dyakonov and A. Khaetskii, in *Spin Physics in Semiconductors*, edited by M. Dyakonov (Springer, Berlin, 2017).
- [49] M. I. Dyakonov, Phys. Rev. Lett. 99, 126601 (2007).
- [50] P. S. Alekseev and M. I. Dyakonov, Phys. Rev. B 100, 081301(R) (2019).
- [51] M. Matsuo, Y. Ohnuma, and S. Maekawa, Phys. Rev. B 96, 020401(R) (2017).
- [52] M. Matsuo, D. A. Bandurin, Y. Ohnuma, Y. Tsutsumi, and S. Maekawa, arXiv:2005.01493.
- [53] R. Takahashi, M. Matsuo, M. Ono, K. Harii, H. Chudo, S. Okayasu, J. Ieda, S. Takahashi, S. Maekawa, and E. Saitoh, Nat. Phys. 12, 52 (2016).
- [54] R. Takahashi, H. Chudo, M. Matsuo, K. Harii, Y. Ohnuma, S. Maekawa, and E. Saitoh, Nat. Commun. 11, 3009 (2020).
- [55] R. J. Doornenbal, M. Polini, and R. A. Duine, J. Phys.: Mater. 2, 015006 (2019).
- [56] M. M. Glazov and E. L. Ivchenko, JETP Lett. 75, 403 (2002).
- [57] M. M. Glazov and E. L. Ivchenko, JETP 99, 1279 (2004).
- [58] I. D'Amico and G. Vignale, Phys. Rev. B 62, 4853 (2000).
- [59] I. D'Amico and G. Vignale, Phys. Rev. B 68, 045307 (2003).
- [60] M. I. Dyakonov and V. I. Perel, Sov. Phys. Solid State 13, 3023 (1972).
- [61] M. M. Glazov, 2D Mater. 9, 015027 (2021).
- [62] J. L. M. Poiseuille, C. R. Acad. Sci. 11, 961 (1840).
- [63] J. R. Womersley, J. Physiol. **127**, 553 (1955).
- [64] S. R. De Groot and P. Mazur, *Non-Equilibrium Thermodynamics* (North-Holland, Amsterdam, 1962).
- [65] R. F. Snider and K. S. Lewchuk, J. Chem. Phys. 46, 3163 (1967).