

## Dissipative and Hall Viscosity of a Disordered 2D Electron Gas

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Hydrodynamic charge transport is at the center of recent research efforts. Of particular interest is the nondissipative Hall viscosity, which conveys topological information in clean gapped systems. The prevalence of disorder in the real world calls for a study of its effect on viscosity. Here we address this question, both analytically and numerically, in the context of disordered noninteracting 2D electrons. Analytically, we employ the self-consistent Born approximation, explicitly taking into account the modification of the single-particle density of states and the elastic transport time due to the Landau quantization. The reported results interpolate smoothly between the limiting cases of a weak (strong) magnetic field and strong (weak) disorder. In the regime of a weak magnetic field our results describe the quantum (Shubnikov–de Haas type) oscillations of the dissipative and Hall viscosity. For strong magnetic fields we characterize the effects of the disorder-induced broadening of the Landau levels on the viscosity coefficients. This is supplemented by numerical calculations for a few filled Landau levels. Our results show that the Hall viscosity is surprisingly robust to disorder.

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*Introduction.*—Ordinary fluid motion is described by the theory of hydrodynamics, one of whose cornerstones is viscosity, which serves as the source of dissipation. Under certain conditions, charge transport in an electronic system can also be dominated by hydrodynamic viscous flow [1,2]. The discovery of graphene stimulated renewed theoretical [3–11] and experimental [12–18] interest in the hydrodynamic description of charge conduction.

In the absence of time-reversal symmetry the viscosity tensor has nondissipative antisymmetric components. In the presence of a magnetic field  $B$ , this nondissipative Hall viscosity ( $\eta_H$ ) was studied theoretically in the classical limit of high temperature plasmas [19–23], and for low temperature electron gas [24]. Later, interest in the Hall viscosity was rekindled in quantum systems with a gapped spectrum, due to the connection between  $\eta_H$  and the geometric response [25–31], and its expected quantization in the presence of translational and rotational symmetries [29]. It was understood that beyond the Hall conductivity and viscosity there are additional nondissipative electromagnetic and geometrical response functions in gapped quantum systems [32–44]. Within the hydrodynamic description of electron transport, nonzero  $\eta_H$  influences significantly the structure of the electron flow [45–50], which allows one to access  $\eta_H$  experimentally [51]. Also, it was argued that the dissipative and Hall viscosity affect the spectrum of edge magnetoplasmons [52–54].

For noninteracting electrons in the absence of disorder each filled Landau level (LL) gives a contribution to the Hall

viscosity equal  $\hbar(2n + 1)/(8\pi l_B^2)$  [25], where  $n$  denotes the LL index and  $l_B = \sqrt{\hbar c/(eB)}$  stands for the magnetic length. This result is stable against perturbations of the Hamiltonian which preserve translational and rotational invariance [29]. However, the fate of this result in the presence of disorder has not been studied yet. Therefore, it is not clear how the clean result obtained within the quantum treatment of the electron motion in a magnetic field connects to the result  $\eta_H = \nu_0 \mu^2 \omega_c \tau_{tr,2}^2 / [1 + 4\omega_c^2 \tau_{tr,2}^2]$  derived for a classical disordered electron gas [24]. Here  $\mu$  denotes the chemical potential,  $\tau_{tr,2}$  the second transport time,  $\omega_c = eB/(m_e c)$  the cyclotron frequency,  $\nu_0 = m_e/(2\pi\hbar^2)$  the density of states at  $B = 0$ , and  $m_e$  the effective electron mass.

In this Letter we report the results of an analytical and numerical study of the dissipative and Hall viscosities of noninteracting 2D electrons in the presence of disorder. Contrary to previous studies we explicitly take into account the Landau quantization of the electron spectrum. Analytically, within the self-consistent Born approximation (SCBA) [55] we derive expressions for the dissipative and Hall viscosities, which smoothly interpolate between the results known in the literature for a classical magnetic field [19–23,45] and for the strong magnetic field in the absence of disorder [25]. Since the SCBA is rigorously justified for high LLs only, we perform numerical calculation of  $\eta_H$  for a few lowest LLs. The obtained numerical results are in a perfect agreement with the SCBA predictions. They demonstrate a surprising resilience of the Hall viscosity to disorder.

*Model.*—Noninteracting electrons confined to a 2D plane are described by the following Hamiltonian:

$$H = (-i\nabla - e\mathbf{A})^2/2m_e + V(\mathbf{r}), \quad (1)$$

where  $V(\mathbf{r})$  stands for a random potential and  $\mathbf{A}$  for the vector potential corresponding to the static perpendicular magnetic field  $B$ . In this Letter we use the Landau gauge:  $A_y = -Bx$  and  $A_x = A_z = 0$ . We assume that the random potential has Gaussian distribution with a pair correlation function  $\overline{V(\mathbf{r})V(\mathbf{r}')} = W(|\mathbf{r} - \mathbf{r}'|)$  that decays with a typical length scale  $d$ . In what follows we use units with  $c = \hbar = 1$ .

*Kubo formula for the viscosity.*—The viscosity tensor can be computed by means of the Kubo formula [56–58]:

$$\eta_{jk,ps}(\omega) = -\frac{i\kappa^{-1}}{\omega} \delta_{jk} \delta_{ps} + \frac{1}{i\omega S} \int \frac{d\Omega d\epsilon}{\pi^2} \frac{f_\epsilon - f_{\epsilon+\Omega}}{\Omega - \omega - i0} \times \overline{\text{Tr} T_{jk} \text{Im} G_{\epsilon+\Omega}^R T_{ps} \text{Im} G_\epsilon^R}. \quad (2)$$

Here  $f_\epsilon = 1/\{1 + \exp[(\epsilon - \mu)/T]\}$  denotes the Fermi distribution function,  $G_\epsilon^R = 1/(\epsilon - H - i0)$  the retarded Green's function,  $S$  the system area, and  $\kappa$  the internal compressibility [59]. The form of the stress tensor  $T_{jk} = m_e(v_j v_k + v_k v_j)/2$  is not affected by a random potential (see Ref. [60] and the Supplemental Material for details [61]). Here  $\mathbf{v} = (-i\nabla - e\mathbf{A})/m_e$  stands for the velocity operator. Disorder averaging is denoted by an overbar.

*Self-consistent Born approximation.*—In order to compute the viscosity tensor from Eq. (2) we treat the disorder scattering using the SCBA [55]. This approximation holds under the following conditions [62,63]:

$$1/k_F, d \ll l_B, \quad d \ll v_F \tau_0. \quad (3)$$

Here,  $k_F = m_e v_F$  and  $v_F$  denote the Fermi momentum and velocity, respectively, and  $\tau_0$  is the total elastic relaxation time at zero magnetic field. It can be expressed in terms of the Fourier transform  $\tilde{W}(\mathbf{q})$  of the pair correlation function  $W(\mathbf{r})$ . Furthermore, it is convenient to generalize it to ( $m = 0, 1, 2, \dots$ ):

$$\frac{1}{\tau_m} = \nu_0 \int_0^{2\pi} d\phi \tilde{W}(2k_F \sin \phi/2) \cos m\phi. \quad (4)$$

The average density of states  $\nu(\epsilon)$  at nonzero  $B$  is determined by the average retarded Green's function  $\mathcal{G}_\epsilon^R$ . In the LL representation the average density of states is given as  $\nu(\epsilon) = -\sum_n \text{Im} \mathcal{G}_n^R(\epsilon)/(2\pi^2 l_B^2)$ . Within SCBA the retarded Green's function satisfies [55,62,63]

$$\mathcal{G}_n^R = (\epsilon - \epsilon_n - \Sigma_n^R)^{-1}, \quad \Sigma_n^R = \frac{\omega_c}{2\pi\tau_0} \sum_n \mathcal{G}_n^R, \quad (5)$$

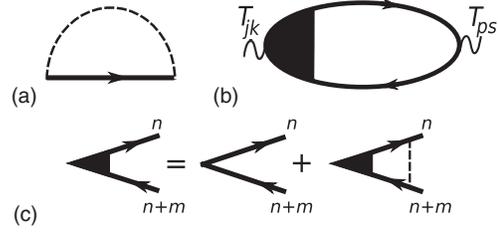


FIG. 1. (a) The self-energy diagram. (b) The diagram corresponding to the Kubo formula (2). (c) The equation for the vertex  $V_m$  in the ladder approximation. The solid line denotes the SCBA Green's function  $\mathcal{G}(\epsilon)$ . The dashed line denotes the disorder correlation function  $W(\mathbf{r})$ .

where  $\epsilon_n = \omega_c(n + 1/2)$ . There are two limiting cases in which the self-consistent Eq. (5) can be easily solved [55]. In the regime of overlapping LLs,  $\omega_c \tau_0 \ll 1$ , one can use the Poisson formula for summation over the LL index. The averaged density of states becomes  $\nu(\epsilon) = \nu_0[1 - 2\delta \cos(2\pi\epsilon/\omega_c)]$ . Here  $\delta = \exp(-\pi/\omega_c \tau_0) \ll 1$  is the Dingle parameter. In the opposite case, when the LLs are well separated, one can restrict the summation in Eq. (5) to the single LL which is closest to the energy of interest,  $|\epsilon - \epsilon_n| < \omega_c/2$ . Then the average density of states acquires the semicircle profile,  $\nu(\epsilon) = \nu_0 \tau_0 \sqrt{\Gamma^2 - (\epsilon - \epsilon_n)^2}$ , where  $\Gamma = \sqrt{2\omega_c/(\pi\tau_0)}$  determines the broadened LL width.

In the presence of long-range disorder correlations  $d \gg k_F^{-1}$  it is important to take into account the vertex corrections to the “bubble” contribution in the Kubo formula (2) (see Fig. 1). This implies that in addition to the average Green's function, one also needs to know the renormalized vertex, which is the stress tensor in the case of the viscosity. Within the SCBA  $T_{jk}$  can be approximated as a linear combination of operators which change the LL index by 2. Under conditions (3) one can show that an operator  $V_m$ , which transfers an electron from the  $(n + m)$ th LL to the  $n$ th LL, is renormalized by the ladder resummation of the disorder lines as follows [62–64] (see Ref. [61] for details):

$$V_m \rightarrow \frac{V_m}{1 - \tau_m^{-1} \Pi_m^{RA}}, \quad \Pi_m^{RA} = \frac{\omega_c}{2\pi} \sum_n \mathcal{G}_{n+m}^R(\epsilon) \mathcal{G}_n^A(\epsilon). \quad (6)$$

Here  $\Pi_m^{RA}$  is the contribution of the bubble without ladder insertions. Using Eq. (5), it can be rewritten as  $\Pi_m^{RA} = -i\nu(\epsilon)/[m\omega_c \nu_0 - i\nu(\epsilon)/\tau_0]$ . Therefore, within the SCBA the vertex corrections are expressed in terms of the average density of states only.

*Dissipative viscosity.*—Disorder averaging restores 2D rotational symmetry [65]. Hence, the viscosity tensor  $\eta_{jk,ps}$  is characterized by only three parameters:

$$\eta_{jk,ps} = \eta_s(\delta_{jp}\delta_{ks} + \delta_{js}\delta_{kp}) + (\zeta - \eta_s)\delta_{jk}\delta_{ps} + (\eta_H/2)(\epsilon_{jp}\delta_{ks} + \epsilon_{js}\delta_{kp} + \epsilon_{kp}\delta_{js} + \epsilon_{ks}\delta_{jp}), \quad (7)$$

where  $\zeta$  and  $\eta_s$  denote the bulk and shear viscosities, respectively. Within the SCBA the bulk viscosity vanishes,  $\zeta = 0$ . Using Eqs. (5) and (6), we find the following result for the shear viscosity at  $\omega = 0$  [61]:

$$\eta_s = \frac{1}{2} \int d\epsilon (-f'_\epsilon) \frac{\nu(\epsilon) \epsilon^2 \tau_{\text{tr},2}(\epsilon)}{1 + 4\omega_c^2 \tau_{\text{tr},2}^2(\epsilon)}, \quad (8)$$

where  $\tau_{\text{tr},2}(\epsilon) = \tau_{\text{tr},2} \nu_0 / \nu(\epsilon)$  is the renormalized second transport time and  $1/\tau_{\text{tr},2} = 1/\tau_0 - 1/\tau_2$  is the second transport rate at  $B = 0$ . We note that for  $k_F d \gg 1$  the second transport time becomes  $\tau_{\text{tr},2} = \tau_0 (k_F d / 2)^2 \gg \tau_0$ . We mention that Eq. (8) is analogous to the result for the dissipative conductivity [64].

In the regime of overlapping LLs,  $\omega_c \tau_0 \ll 1$ , the shear viscosity exhibits Shubnikov–de Haas-type oscillations:

$$\eta_s = \frac{1}{2} \frac{\nu_0 \mu^2 \tau_{\text{tr},2}}{1 + 4\alpha^2} \left( 1 - \frac{16\alpha^2 \delta}{1 + 4\alpha^2} \mathcal{F}_T \cos \frac{2\pi\mu}{\omega_c} \right), \quad (9)$$

where  $\alpha = \omega_c \tau_{\text{tr},2}$  and  $\mathcal{F}_T = (2\pi^2 T / \omega_c) / \sinh(2\pi^2 T / \omega_c)$ . The nonoscillatory term in  $\eta_s$  reproduces the classical result for the shear viscosity of an electron gas [24].

In the regime of well-separated LLs,  $\omega_c \tau_0 \gg 1$ , one finds from Eq. (8) that the shear viscosity is nonzero when the chemical potential is inside the  $N$ th broadened Landau level ( $|\mu - \epsilon_N| \leq \Gamma$ ):

$$\eta_s = (N^2 \tau_0) / (8\pi^2 l_B^2 \tau_{\text{tr},2}) [1 - (\mu - \epsilon_N)^2 / \Gamma^2]. \quad (10)$$

For chemical potential at the center of the LL, the shear viscosity is  $2\omega_c \tau_0 / \pi$  times larger than the one naively expected on the basis of purely classical expression. The dependence of the shear viscosity on the chemical potential in comparison with  $\nu(\epsilon)$  is shown in Fig. 2.

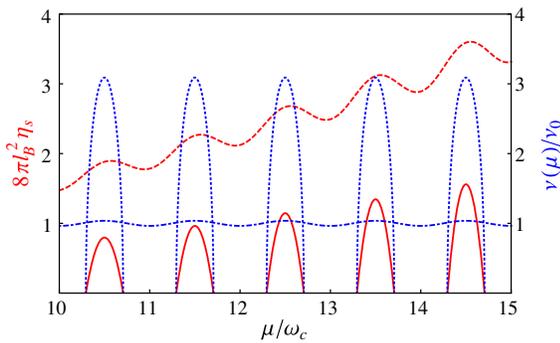


FIG. 2. The density of states (blue dotted and blue dash-dotted curves) and shear viscosity (red solid and red dashed curves) as functions of  $\mu/\omega_c$  for smooth disorder,  $\tau_{\text{tr},2}/\tau_0 = 40$ . Blue dotted and red solid curves correspond to well-separated LLs with  $\omega_c \tau_0 = 15$ . Blue dash-dotted and red dashed curves correspond to overlapping LLs with  $\omega_c \tau_0 = 0.8$ .

*Hall viscosity.*—The Hall viscosity can be extracted from the viscosity tensor as  $\eta_H = (\eta_{xy,xx} - \eta_{xy,yy})/2$ . Similarly to the Hall conductance, the evaluation of  $\eta_H$  from the Kubo formula (2) is complicated due to contributions that come from all the states below the chemical potential. Therefore, it is convenient to proceed in a way pioneered by Smrčka and Středa [67]: As for the Hall conductivity we split the expression for the Hall viscosity at  $\omega = 0$  into two parts,  $\eta_H = \eta_H^I + \eta_H^{II}$ , where [61]

$$\eta_H^I = \text{Re} \int \frac{d\epsilon}{4\pi S} (-f'_\epsilon) \overline{\text{Tr} T_{xy} G_\epsilon^R (T_{xx} - T_{yy}) G_\epsilon^A}, \quad (11)$$

$$\eta_H^{II} = \text{Im} \int \frac{d\epsilon}{2\pi S} (-f'_\epsilon) \overline{\text{Tr} J_{xy} (T_{yy} - T_{xx}) G_\epsilon^A} + \text{Im} \int \frac{d\epsilon}{2\pi S} f'_\epsilon \overline{\text{Tr} [J_{xy}, J_{yy} - J_{xx}] \text{Im} G_\epsilon^R}. \quad (12)$$

Here  $J_{jk}$  denotes the strain generators which are related with the stress tensor as  $T_{jk} = -i[H, J_{jk}]$  [56]. One can evaluate  $\eta_H^I$  in a similar way to  $\eta_s$  [61]:

$$\eta_H^I = \int d\epsilon (-f'_\epsilon) \frac{\nu(\epsilon) \epsilon^2 \omega_c \tau_{\text{tr},2}^2(\epsilon)}{1 + 4\omega_c^2 \tau_{\text{tr},2}^2(\epsilon)}. \quad (13)$$

The evaluation of  $\eta_H^{II}$  is more involved. Although one can write down the viscoelastic analog of the Smrčka and Středa formula for the Hall viscosity [68], it does not provide a suitable way for the calculation of  $\eta_H^{II}$  in the presence of disorder. In order to compute  $\eta_H^{II}$  one needs to know the expressions for the strain generators. In the absence of disorder they can be easily written down explicitly [56], e.g.,  $J_{xy}^{(0)} = (T_{xx} - T_{yy}) / (4\omega_c)$  and  $J_{yy}^{(0)} - J_{xx}^{(0)} = T_{xy} / \omega_c$ . In the presence of a random potential the strain generators can be constructed as a series in spatial derivatives of a random potential  $V$ . This allows us to evaluate  $\eta_H^{II}$  within the SCBA [61]:

$$\eta_H^{II} = \mathcal{E} / (2\omega_c) - \int d\epsilon (-f'_\epsilon) \nu(\epsilon) \epsilon^2 / (4\omega_c), \quad (14)$$

where  $\mathcal{E} = \int d\epsilon \nu(\epsilon) \epsilon f_\epsilon$  stands for the energy density. Combining Eqs. (13) and (14), we obtain

$$\eta_H = \frac{\mathcal{E}}{2\omega_c} - \frac{1}{4\omega_c} \int d\epsilon (-f'_\epsilon) \frac{\nu(\epsilon) \epsilon^2}{1 + 4\omega_c^2 \tau_{\text{tr},2}^2(\epsilon)}. \quad (15)$$

In the absence of disorder and for the chemical potential above the  $N$ th Landau level the energy density at  $T = 0$  can be computed as  $\mathcal{E} = \sum_{n=0}^N \epsilon_n / (2\pi l_B^2)$ , which yields the known result  $\eta_H = \sum_{n=0}^N (n + 1/2) / (4\pi l_B^2)$  [25]. Also, we mention that in the Boltzmann limit,  $T \gg E_F$ , the energy density is given by  $\mathcal{E} = n_e T$ , where  $n_e$  denotes the particle density, such that the Hall viscosity in the absence of

disorder and at  $T \gg E_F$  becomes  $\eta_H = n_e T / (2\omega_c)$ , in agreement with Eq. (59.38) of Ref. [69] in which the Hall viscosity is denoted by  $\eta_3$ . We note that the structure of Eq. (15) resembles the structure of the SCBA result for the Hall conductivity  $\sigma_H$  [70].

The appearance of the nonzero  $\eta_H$  can be explained on a pure classical level [21]. Hall viscosity describes the response of  $T_{xx} - T_{yy}$  to a shear velocity profile,  $U_x = uy$ . In the presence of a magnetic field this velocity can be considered as the result of a nonuniform electric field,  $E_y = -U_x B$ . This electric field results not only in a drift of the cyclotron orbit but in its deformation into an ellipse. To linear order in  $u$  the eccentricity of the ellipse is equal to  $u/(2\omega_c)$ . This asymmetry between motion in the  $x$  and  $y$  direction yields the nonzero ratio  $(T_{xx} - T_{yy})/u$  in the limit  $u \rightarrow 0$ . Hence, nonzero  $\eta_H$  arises, which is given by the first term in Eq. (15). An electron moving along an ellipse conserves its energy to the first order in  $u$ , in agreement with the nondissipative nature of  $\eta_H$ . In the presence of impurity scattering an electron experiences a friction force corresponding to an electric field  $E_x = -U_x/(e\tau_{tr,2})$ . This electric field leads to a velocity component  $U_y = U_x/(\omega_c\tau_{tr,2})$ . The nonuniformity of this velocity produces additional correction to the difference,  $T_{xx} - T_{yy} \sim -u\eta_s/(\omega_c\tau_{tr,2})$ . Thus, there is an additional correction to the Hall viscosity,  $\Delta\eta_H = -\eta_s/(\omega_c\tau_{tr,2})$ , which corresponds to the second term in Eq. (15) in the classical regime.

In the case of overlapping LLs,  $\omega_c\tau_0 \ll 1$ , from Eq. (15) we obtain the Shubnikov–de Haas oscillations of the Hall viscosity:

$$\eta_H = \frac{\nu_0\mu^2\omega_c\tau_{tr,2}^2}{1+4\alpha^2} \left( 1 + \frac{\delta}{2\alpha^2} \frac{1+12\alpha^2}{1+4\alpha^2} \mathcal{F}_T \cos \frac{2\pi\mu}{\omega_c} \right). \quad (16)$$

The nonoscillatory term in  $\eta_H$  coincides with the classical result for the Hall viscosity of electron gas [24].

In the case of well-separated LLs,  $\omega_c\tau_0 \gg 1$ , one finds from Eq. (15) that the Hall viscosity is reduced from the quantized value if the chemical potential lies within the broadened LL,  $|\mu - \epsilon_N| \leq \Gamma$ :

$$\eta_H = \frac{N^2}{8\pi l_B^2} \left[ 1 - \frac{\tau_0^2\Gamma}{2\pi\omega_c\tau_{tr,2}^2} \left( 1 - \frac{(\mu - \epsilon_N)^2}{\Gamma^2} \right)^{3/2} \right]. \quad (17)$$

We note that for the long-range-correlated random potential the Hall viscosity dominates the shear viscosity,  $\eta_H \gg \eta_s$  [cf., Eqs. (10) and (17)].

The deviation of the Hall viscosity from the clean value is controlled by the small parameter  $(\tau_0/\tau_{tr,2})^2/\sqrt{\omega_c\tau_0} \ll 1$ . In the case of short range random potential correlations  $\tau_0 = \tau_{tr,2}$  the deviation of  $\eta_H$  from its clean value is very small. For long-range-correlated random potential  $\tau_0 \ll \tau_{tr,2}$  the difference  $\eta_H - N^2/(8\pi l_B^2)$  is additionally suppressed (see Fig. 3).

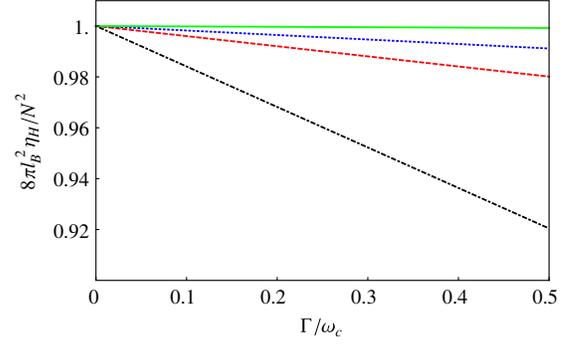


FIG. 3. The normalized Hall viscosity  $8\pi l_B^2 \eta_H / N^2$  as the function of  $\Gamma/\omega_c$  for different ranges of disorder in the case of well-separated LLs. The parameter  $\tau_{tr,2}/\tau_0$  is equal to 1, 2, 3, and 10 from the bottom to the top.

*Numerical results.*—We would now like to explore the quantum Hall regime, where the number of filled LLs is of order unity. Here the SCBA cannot be used anymore, and we resort to a numerical calculation. For this we discretize the system and employ the Hofstadter model with uncorrelated random potential, uniformly distributed between  $[-w/2, w/2]$  at each lattice site. We calculate the Hall viscosity at zero temperature using the retarded correlation function of discretized stress operators [71], and take both the continuum and thermodynamic limits to extrapolate to the behavior of our model (1) [61]. In the presence of disorder we can take these limits while keeping constant  $\omega_c\tau$ . The results for the Hall viscosity are plotted in Fig. 4, together with the behavior of the Hall conductivity ( $\sigma_H$ ) at zero wave vector. One sees that, somewhat surprisingly, the

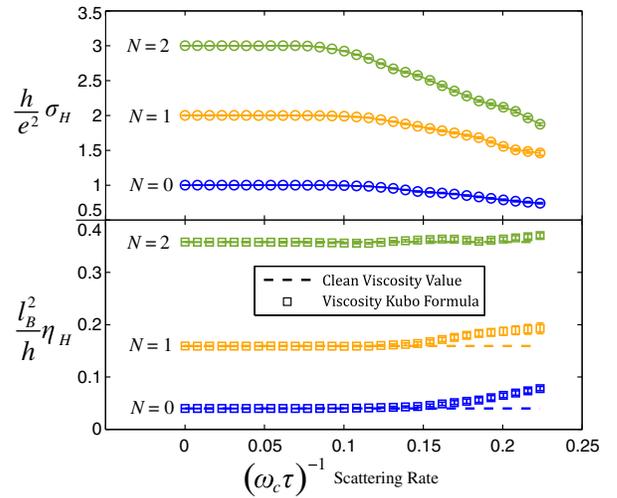


FIG. 4. The Hall conductivity (circles) and viscosity (squares) as a function of  $\omega_c\tau$  for  $N = 0, 1,$  and  $2$  filled Landau levels (blue, yellow, and green, respectively). The clean viscosity values are also indicated by dashed lines.  $\eta_H$  is seen to be robust to disorder to the same extent the  $\sigma_H$  is, within the numerical errors (error bars are smaller than the symbol sizes).

Hall viscosity maintains its quantization to the same extent as the Hall conductivity, that is, until the quantum Hall to insulator transition is approached.

*Conclusions.*—To summarize, we studied the dissipative and Hall viscosity of 2D electron system in the presence of a random potential. Within the self-consistent Born approximation we derived an expression for both the dissipative and Hall viscosities, which takes into account the modification of the single-particle density of states and the elastic transport time due to the Landau quantization. Our results smoothly interpolate between the case of weak magnetic field and strong disorder on the one hand and the case of strong magnetic field and vanishing disorder on the other hand. In the former regime, we derived the expressions for the quantum (Shubnikov–de Haas type) oscillations of the dissipative and Hall viscosities. In the case of strong magnetic field, we found that the disorder broadening of the Landau level does not lead to a significant change of the Hall viscosity in comparison with the clean result. Our numerical results for a few filled LLs support this striking conclusion.

There are various ways to extend our work. In Galilean invariant systems it was proven [36,56] that the viscosity tensor can be extracted from the nonlocal conductivity, that is, the conductivity tensor at finite wave vector  $\mathbf{q}$ . In the absence of Galilean invariance there is no reason to expect that  $\eta_H$  is related to  $\sigma_H(\mathbf{q})$  [72,73]. Also, the relation between  $\eta_H$  and  $\sigma_H(\mathbf{q})$  can be affected by the presence of a lattice [71,74] or disorder. However, if one treats disorder on the level of the Drude model with a classical magnetic field, the relation of Ref. [56] between  $\eta_H$  and  $\sigma_H(\mathbf{q})$  still holds [75]. This fact is not surprising since the Drude model does not properly take into account the LLs, which result in the energy dependence of the density of states and elastic scattering transport time. However, such a simplification can be dangerous since  $\eta_H$  and  $\sigma_H$  have contributions coming from the states well below the Fermi energy. It would therefore be worthwhile to extend the presented analytical and numerical approaches to the conductivity at finite wave vector [76]. We also note that our techniques can be applied to calculation of the dissipative and Hall viscosity in graphene, where only the result in the absence of disorder is known [77].

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