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Generalized Wigner crystallization in moiré materials

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Recent experiments on the twisted transition metal dichalcogenide (TMD) material, WSe_2/WS_2 , have observed insulating states at fractional occupancy of the moiré bands. Such states were conceived as generalized Wigner crystals (GWCs). In this article, we investigate the problem of Wigner crystallization in the presence of an underlying (moiré) lattice. Based on the best estimates of the system parameters, we find a variety of homobilayer and heterobilayer TMDs to be excellent candidates for realizing GWCs. In particular, our analysis based on r_s indicates that MoSe_2 (among the homobilayers) and $\text{MoSe}_2/\text{WSe}_2$ or MoS_2/WS_2 (among the heterobilayers) are the best candidates for realizing GWCs. We also establish that due to larger effective mass of the valence bands, in general, hole-crystals are easier to realize than electron-crystals as seen experimentally. For completeness, we show that satisfying the Mott criterion $n_{\text{Mott}}^{1/2} a_* = 1$ requires densities nearly three orders of magnitude larger than the maximal density for GWC formation. This indicates that for the typical density of operation, bilayer moiré systems are far from the Mott insulating regime. These crystals realized on a moiré lattice, unlike the conventional Wigner crystals, are incompressible due to the gap arising from pinning with the lattice. Finally, we capture this many-body gap by variationally renormalizing the dispersion of the vibration modes. We show these low-energy modes, arising from coupling of the WC with the moiré lattice, can be effectively modeled as a Sine-Gordon theory of fluctuations.

I. INTRODUCTION

A strongly interacting dilute gas of electrons minimizes its energy by spontaneously breaking translation invariance to form a Wigner crystal (WC) [1]. Though this physics is a simple and intuitive manifestation of a strongly interacting many-body phase, experimental realizations of quantum Wigner crystals have been far and few between. Thus far, they have been seen in a two dimensional electron gas (2DEG) realized in semiconducting heterostructures [2] and liquid helium [3]. Recently, moiré materials, synthetic materials constituted from stacked monolayers with a mismatch in lattice size or orientation, have emerged as a highly tunable and experimentally accessible platform to study the physics of strong electronic correlations as well as topology [4–19].

In particular, homobilayer moiré (HoM) materials or heterobilayer moiré (HeM) materials based on transition metal dichalcogenides (TMD), see Fig. 1, have emerged as prime candidates for realizing WCs [13–15, 20, 21]. This can be largely attributed to the fact that the low energy moiré electrons in TMDs often reside in extremely narrow (quasi-flat) bands [18, 19, 22, 23] or have very large effective masses, even compared to the traditional 2DEG systems [3]. This makes them highly susceptible to charge localization. Such factors, coupled with the high controllability of TMDs for studying correlated phenomena [17–19], make them great candidates for studying Wigner crystallization. Given the plethora of TMDs, a primary goal in this article is to explore material characteristics—lattice constant (a), dielectric constant (ϵ), effective mass (m_*)—to characterize the ideal candidates for hosting a WC.

Typically, a pure WC formed in a 2DEG slides when

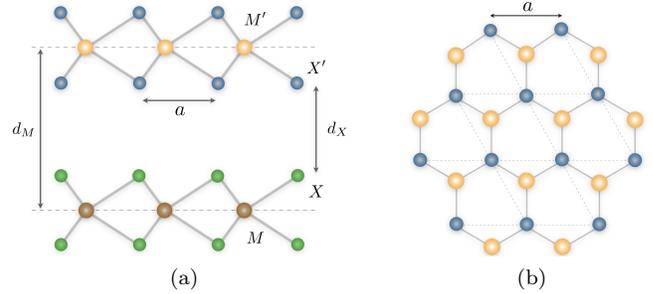


FIG. 1. Schematic of AA stacked TMD bilayer: (a) the side-view shows the MX_2 layer, with trigonal prismatic (H) coordination, stacked on top of the $\text{M}'\text{X}'_2$ layer. The large (yellow or brown) balls represent the metal ions, M or M' , and the small (blue or green) balls represent the chalcogens, X or X' . The distance between the metal ions and the chalcogens are, respectively, denoted by d_M and d_X . (b) The top-view is a honeycomb lattice of lattice constant a .

subjected to a nonzero electric field due to the lack of a momentum relaxation mechanism. A key signature of such a WC is its negative compressibility [24–28]. Disorder, however, pins the WC and renders it incompressible as a result of the activation or pinning gap. A WC realized in moiré materials [29, 30] is however, ineluctably influenced by the underlying moiré lattice, which provides a uniform periodic background potential as illustrated in Fig. 2. This provides a pinning mechanism distinct from that induced by disorder which will strongly influence its properties. Such a crystal is often referred to as a ‘generalized Wigner crystal’ (GWC) [31], see Fig. 2. Although disorder-pinned-WCs have been studied widely [2, 3, 32–36], an in-depth study of GWCs is still lacking.

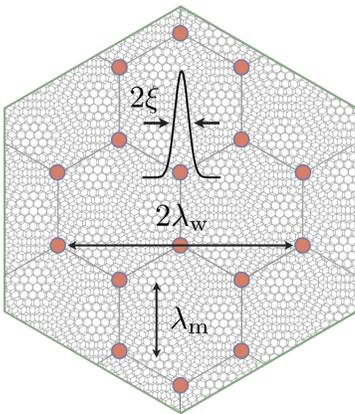


FIG. 2. A cartoon rendition of a WC and the relevant length scales: A Wigner lattice is realized on a moiré superlattice (gray background) at filling fraction $2/3$. The distance between two nearest dark (or bright) spots is the moiré periodicity, λ_m . The distance between the two nearest localized particles (red dots) is the Wigner lattice periodicity, λ_w . The bell-shaped curve, representing the wavefunction of a localized particle, has a width of 2ξ . Our discussion in this paper is confined to a crystal where the moiré electrons are highly localized, $\xi \ll \lambda_w$.

In light of the recent experiments in TMD platforms exploring the physics of strong correlations [13–16, 37], a study of the properties of the GWC is timely as it helps distinguish the GWC from other density ordered gapped states that a lattice system may host alongside a GWC [15, 37–40]. Insulating states observed in WSe_2/WS_2 are at fractional fillings, $\nu = 1/3, 2/3$, [13] and those in the twisted bilayer of graphene (TBLG) are at integer fillings [4–12]. Simple observables like compressibility (or capacitance) are often misleading and insufficient [41–44] to discern between a pinned WC and a Mott state as these exhibit similar capacitive signatures [43, 44]. However, in the presence of the moiré lattice, Mott states must preserve the underlying (moiré) lattice symmetry, and can only be observed at fillings for which a placement of the electrons preserves the underlying symmetry of the moiré lattice. While this is difficult for integer fillings exceeding unity [29, 30], it is impossible at fractional filling observed in WSe_2/WS_2 . Consequently, the nature of the insulating states at integer fillings remains ambiguous. In TBLG, our earlier works [29, 30], based on the Mott criterion, precluded the interpretation of the observed insulating states at integer fillings as Mott states. Wigner crystallization [29, 30] was envisaged to be more favorable than Mott insulation at low charge densities in TBLG.

In this paper, we explore from a materials perspective the viability of both homo- and hetero-bilayer TMDs for realizing WCs. Additionally, we study the impact of the moiré lattice on collective excitations [2, 3, 32] of GWCs and present estimates for the gap in the deep crystalline limit which can be directly accessed in transport experiments. Our results are directly of relevance to a slew of

recent experiments in these systems exploring the physics of the GWC [13, 16]. We organize this article as follows. In Sec. II, we analyze the material parameters of various HoM and HeM systems and assess their candidacy for crystal formation using several criteria. We identify a wide range of TMD materials that can support GWC phases and establish, broadly speaking, HeM to be better candidates than HoM for this purpose. In Sec. III, in the elastic limit [28, 45, 46], we obtain an effective Hamiltonian that describes harmonic fluctuations in a GWC pinned to a moiré lattice. We then move to obtaining the self-consistent equations for the pinning gaps corresponding to a GWC in Sec. IV. Finally, we conclude by connecting our results to the recent experiments in Sec. V. Technical details are relegated to various appendices.

II. TMD CANDIDACY FOR WIGNER CRYSTALLIZATION

In this section, we discuss the key criteria for assessing the candidacy of various TMD bilayers, both HoM and HeM for Wigner crystallization. Generally, a material with low carrier density and a high degree of correlation can be susceptible to forming a WC. A natural way to measure correlation is to compare the strength of electronic interaction (U) with the kinetic energy (W) of the relevant charge carriers. Denoting the mean separation between the moiré particles by r_e , we set the scale of the Coulomb repulsion to $U = e^2/\epsilon r_e$, where, e is electronic charge and ϵ is the dielectric constant. In principle, one can also use a more realistic interaction potential for TMDs that can account for the encapsulating environment (such as the hBN/SiO₂ surroundings) [47–49]. However, at long distances, such a potential distills to a Coulomb-type potential [50]. Therefore, our assumption remains useful for discussing the low energy physics of TMDs. Another simplifying assumption we make is to ignore the full details of the TMD bandstructure [18, 19]. We simply set $W = \hbar^2 k^2 / 2m_e^*$ with $k \sim 1/r_e$. m_e^* (m_h^*) is the effective mass of the electrons (holes) in the conduction (valence) band. For r_e of the order of the moiré lattice constant, λ_m , one gets, $W \sim \mathcal{O}(1 \text{ meV})$ and $U \sim \mathcal{O}(10 \text{ meV})$.

Here, we reiterate that the important (in-plane) length scales in the problem, as shown in Figs. 1 and 2, are – the monolayer lattice constant (a), the moiré periodicity (λ_m), the Wigner lattice periodicity (λ_w), and the localization length of the moiré particles (ξ). a is the smallest scale and can be neglected in a low energy theory. λ_m is a geometric scale which is fixed for a given TMD device. Unlike these two lengths, ξ and λ_w are dynamically generated. By working in the deep crystalline limit where $\xi \ll \lambda_m, \lambda_w$ we can drop ξ . Thus, the most important scale in our problem is λ_w , and its interplay with λ_m . Since $\lambda_w = 1/\sqrt{\pi n_e}$ is a function of electronic density n_e (or hole density n_h), it allows us to

study crystallization as a function of doping levels. Using this, the kinetic (W) and potential (U) energies can be recast as $W^{-1} = 2m/\pi n_e$ and $U^{-1} = \epsilon\sqrt{\pi n_e}/e^2$. The dimensionless ratio of these two parameters, also known as r_s , provides crucial insight into nature of a correlated state [51]. Ignoring the effect of the moiré potential on the energies, we obtain

$$r_s = \frac{g}{a_0 m_0} \frac{m_e^*}{\epsilon} \lambda_w, \quad \lambda_w = \frac{1}{\sqrt{\pi n_e}}, \quad (1)$$

where $a_0 = \hbar^2/m_0 e^2 = 0.529 \text{ \AA}$ is the Bohr radius with m_0 as the bare electron mass. And, $g = 2$ is a valley degeneracy factor for TMDs. This valley degree of freedom can significantly alter the correlation properties and the threshold for Wigner crystallization. In a 2-valley 2DEG the crystallization threshold drops to $r_s = 29.5$ [52] from $r_s = 37$ in a 1-valley system [51, 53]. The further r_s exceeds this threshold value, the easier it is to form a WC. For further discussion on a more fine-tuned definition of r_s , see [30]. Note however, due to the availability of a set of potential minima facilitated by the underlying moiré lattice, the threshold value for GWCs could be lower than $r_s = 29.5$.

Clearly, Eq. (1) shows that the material parameters that favor Wigner crystallization (or enhance r_s) are a high effective mass, reduced screening or a small dielectric constant and low carrier density. Firstly, low energy carriers in TMDs or twisted bilayers of TMDs are particularly heavy. Secondly, though the dielectric constant of a material is fixed, it can be altered by introducing a spacer layer [16], such as a hexagonal boron nitride (hBN) monolayer. Screening can then be reduced by a judicious choice of spacer material, thereby favoring Wigner crystallization.

Evidently however, the moiré scale dependence of r_s is not manifest in Eq. (1). This can be naturally restored by measuring the carrier density through the filling fraction of a moiré unit supercell. This can be understood as follows. The area of a (hexagonal) moiré unit supercell is given by $A_s = \sqrt{3}\lambda_m^2/2$. If the full occupancy of the relevant low energy band is N_0 , usually determined by the discrete symmetries of the system, then the supercell density is given by $n_s = N_0/A_s \sim 10^{11-12} \text{ cm}^{-2}$. A state consisting of N electrons in this band is observed at a filling fraction of $N/N_0 \equiv \nu$, or at a density $n_e = \nu n_s$. Inserting this in Eq. (1), we observe that, for a given material, there exists a critical density, n_e^{max} , or a maximal filling fraction, ν_{max} , above which a GWC cannot exist. Correspondingly, since $r_s \propto \lambda_m$ [replacing λ_w with λ_m in Eq. (1)], there also exists a critical moiré length below which a material cannot host a GWC. It is worth noting here that the true advantage of moiré materials in realizing WC is this availability of large length scales that govern most of the physics.

Before proceeding further, we note the above discussions are pertinent for zero temperature WC (or quantum WC) only. As the temperature increases, one needs to confront the problem of crystal melting. Although an

accurate estimation of this melting temperature can be a subtle issue [54–56], for simplicity, we estimate it using the classical Lindemann criterion, $k_B T_L \simeq 0.01U$. Our discussions in this paper will be confined to the physics of a GWC at $T \ll T_L$. In the subsections below, we will explicitly evaluate all the above mentioned parameters for several TMDs.

A. Homobilayers

In a HoM system, the top and the bottom layers consist of the same TMD where each layer projects to a 2D honeycomb lattice (see Fig. 1b). This, therefore, is geometrically equivalent to a twisted bilayer graphene system. The moiré periodicity in a HoM is thus given by [59] $\lambda_m(\theta) = \frac{a}{2 \sin(\theta/2)} \simeq a/\theta$. Here, θ is the twist angle between the two TMD layers. $(\epsilon_{\parallel}, \epsilon_{\perp})$ and $(\epsilon_{\parallel}^{(2)}, \epsilon_{\perp}^{(2)})$ denote the in-plane and out of plane dielectric constants of a monolayer and a homobilayer TMD respectively. We identify the geometric mean of these two constants, $\epsilon^{(2)} = \sqrt{\epsilon_{\perp}^{(2)} \epsilon_{\parallel}^{(2)}}$, as the dielectric constant of the bilayer system [60].

Using these parameters, we summarize our results for crystallization criteria in different candidate HoMs in Table I. For a typical twist angle $\theta = 1^\circ$, we find that, $U/W > 1$ for all the homobilayers in Table I, rendering them strongly interacting systems. The corresponding r_s computed using Eq. (1) shows that all the HoMs in

TABLE I. Wigner crystallization criteria for HoMs at $\theta = 1^\circ$: Due to the larger effective masses, the Mo-based compounds are generally better suited to forming GWCs as compared to the W-based compounds. In regard to r_s , or U/W , we conclude a twisted bilayer of MoSe₂ to be the best candidate for Wigner crystallization. The effective masses and dielectric constants are adapted from Ref. [57], and Ref. [58], respectively. Experimental lattice constant (a) data and the distance between the TMD layers (d_X) are compiled in Ref. [57]. Eq. (1) reduces to $r_s = 674m_e^*/m_0\epsilon$ for $n_e = 10^{11} \text{ cm}^{-2}$, which we use as the unit for densities mentioned here.

HoMs	MoS ₂	MoSe ₂	MoTe ₂	WS ₂	WSe ₂	WTe ₂
m_e^*/m_0	0.46	0.56	0.62	0.26	0.28	0.26
$\epsilon_{\perp}(\epsilon_{\parallel})$	4.8(3.0)	6.9(3.8)	8(4.4)	4.4(2.9)	4.5(2.9)	5.7(3.3)
$\epsilon_{\perp}^{(2)}(\epsilon_{\parallel}^{(2)})$	6.9(4.4)	7.9(4.6)	8.6(5.5)	6.1(4.2)	6.3(4.3)	8.4(5.2)
d_X [Å]	3.17	3.33	3.60	3.14	3.34	3.60
a [Å]	3.16	3.29	3.52	3.15	3.28	3.50
λ_m [nm]	18.1	18.8	20.2	18.0	18.8	20.0
U/W	5.0	5.8	6.0	3.1	3.3	2.6
$r_s _{10^{11} \text{ cm}^{-2}}$	56.3	62.6	60.8	34.7	36.3	26.5
n_e^{max}	3.6	4.5	4.2	1.4	1.5	0.8
ν_{max}	1.02	1.38	1.48	0.40	0.46	0.28
T_L [K]	1.7	1.5	1.2	1.8	1.7	1.3
$n_{\text{Mott}} 10^{-3}$	2.5	3.1	2.9	0.9	1.0	0.6

Table I are susceptible to forming GWCs since they all have r_s fairly above the crystallization threshold. The critical density for crystallization is found to be nearly the order of n_s . The critical density, or the closest rational filling fraction, $\nu_{\max} = n_e^{\max}/n_s$, below which a HoM system can host GWC is obtained by setting $r_s = 29.5$ in Eq. (1). Based on the Lindemann criteria, our results predict that the GWCs should be stable in the range of 1K-3K. Our simple analysis shows that Mo-based HoMs are more viable than W-based compounds for the realization of GWCs.

Finally, we evaluate the Mott criterion, $n_e^{1/2} a_0^* \approx O(1)$, which a system needs to satisfy in order to host Mott insulating states [61]. The effective Bohr radius, $a_0^* = \hbar^2/m_e^* e^2$ and $e_* = e^2/\epsilon$. Evaluating this for HoMs, we find that for experimentally relevant densities (that is near the fractional fillings of a moiré unit supercell) the Mott criterion is far from being met, $n_e^{1/2} a_0^* \sim O(10^{-2}) \ll 1$. Satisfying the Mott criterion, $n_{\text{Mott}}^{1/2} a_* = 1$ requires densities nearly three orders of magnitude larger than the maximal density for GWC formation. This indicates that for the typical density of operation, HoM systems are far from the Mott insulating regime.

B. Heterobilayers

In HeM materials, the top and bottom layers contain different TMDs. We now explore the potential for GWCs in HeMs in the manner done in the preceding section for homobilayers. Although the planar projection of each layer is a honeycomb lattice with different periodicities, a moiré pattern emerges even without introducing any twist angle ('near-aligned sample'). Twisting alters the moiré periodicity; in particular, it reduces with increasing twist angle and often approaches the original lattice constant at 'large-twist-angles'. For example, in a HeM with a small difference in lattice constants [72], the moiré periodicity is [63, 73]

$$\lambda_m \simeq \frac{a_{>}}{\sqrt{\delta_a^2 + 4 \sin^2(\theta/2)}} \quad , \quad \delta_a = 1 - \frac{a_{<}}{a_{>}}. \quad (2)$$

Here $a_{>(<)}$ is the largest (smallest) lattice constant among the two layers. We see that λ_m is strongly influenced by the twist angle for samples with small δ_a . As shown in Table I, this is the case of HeMs with differing metal ions [$MX_2/M'X_2$] which have $\delta_a \lesssim 0.1\%$. HeMs with differing chalcogens [MX_2/MX'_2] tend to have large δ_a , i.e. around 4% and are less sensitive to small angle twists. Motivated by the experiment of Ref. [13] which concern $\theta \lesssim 1^\circ$ [63], we confine our discussion to nearly-aligned heterobilayers.

The effective dielectric constant of the HeM system is obtained by treating the two layers as two dielectrics (or capacitors) in series,

$$\frac{d_1 + d_2}{\epsilon} = \frac{d_1}{\epsilon_1} + \frac{d_2}{\epsilon_2}, \quad (3)$$

where ϵ_i and d_i are the dielectric constants and the thickness of the top and bottom layers, respectively. We assume $d_1 = d_2$ and as the two layers are different and stacked along the direction that is normal to the dielectric plane, we set ϵ_i to be the in-plane monolayer dielectric constants, $\epsilon_{\parallel,i}$. For near-aligned samples with $\theta = 0.5^\circ$, using Eq. (2) and Eq. (1) we evaluate U/W and r_s for different HeMs. Our results are summarized in Table II (see also App. A for a similar table on $\theta = 5^\circ$). We find generically that hole carriers have larger r_s (or correlation) due to their larger effective masses. Almost all the HeMs considered in Table II can Wigner crystallize for a hole density of 10^{12} cm^{-2} or less. However, except for a few, most of the electronic carriers do not crystallize. Also note that the Lindemann temperature is the same for both electron and hole crystals since U , under our assumptions, simply depends on the geometry and not on the effective mass.

Since MoSe₂/WSe₂ and MoS₂/WS₂ share the same chalcogens, they are quite sensitive to twist angle. For $\theta \sim 0^\circ$, the moiré length can be as large as a micrometer and it gradually reduces to about a deca-nanometer by 5° of twisting. The correlation factor U/W , therefore, also reduces by nearly two orders of magnitude. For the remainder of the HoMs, though the above mentioned trend is still valid, however, quantitatively, no significant change is observed in the correlation factor since the moiré length scale remains largely insensitive to small changes in the twist angle. In particular, for WSe₂/WS₂, we find that at filling fraction $\nu = 1/3$, $r_s = 44.0$ (26.8) for holes (electrons), and at $\nu = 2/3$, it is 31.1 (19.0) for holes (electrons). This thus explains why Regan *et al.* [13] observe GWC states on the hole side but not on the electronic side. This is one of our key results as it bares directly on the experiments.

Lastly, we evaluate the critical density, or filling fraction, above which the heterostructure will be unable to host GWCs. In particular, for WSe₂/WS₂ we observe that no hole-crystal can exist above a filling fraction of 0.73 ($\approx 3/4$). States at any filling fraction below this, even other than those at 1/3 and 2/3 [39], are perfectly allowed. Similarly, on the electron side, GWC can exist up to $\nu = 0.28$ ($\approx 1/4$).

To summarize, based on the best estimates of the system parameters, we find a variety of homobilayer and heterobilayer TMDs to be excellent candidates for realizing WCs. In particular, our analysis based on r_s indicates that MoSe₂ (among the homobilayers) and MoSe₂/WSe₂ or MoS₂/WS₂ (among the heterobilayers) are the best candidates for realizing WCs. We also establish that due to larger effective masses of the valence bands, hole-crystals in general, are easier to realize than electron-crystals, an observation consistent with experiments. In the remainder of the paper, we focus on the properties of a GWC.

TABLE II. Wigner crystallization criteria for a nearly-aligned HeMs at $\theta = 0.5^\circ$: Among the HeMs listed here, MoSe₂/WSe₂ and MoS₂/WS₂ seem to be the most susceptible to forming an electronic and hole GWC, respectively. At least one experimental study of the pertinent HeM is referenced here. The effective masses are adapted from Ref. [62]. The unit of density is $n_e = 10^{12} \text{cm}^{-2}$. Asterisked values in the r_s row indicate values crossing the crystallization threshold.

HeMs	WSe ₂ /WS ₂	MoSe ₂ /MoS ₂	MoTe ₂ /MoSe ₂	MoSe ₂ /WS ₂	MoSe ₂ /WSe ₂	MoS ₂ /WS ₂	MoTe ₂ /WSe ₂
Refs.	[13, 63]	[64]	[65, 66]	[67]	[68, 69]	[70]	[71]
$m_{e(h)}^*/m_0$	0.28 (0.46)	0.42 (0.71)	0.46 (1.37)	0.28 (0.71)	0.54 (0.44)	0.46 (1.70)	0.30 (1.33)
$2/(\epsilon_1^{-1} + \epsilon_2^{-1})$	2.9	3.35	4.08	3.29	3.29	2.95	3.5
λ_m [nm]	8.1	8.1	5.3	7.6	35.6	34.0	5.1
$U/W _{e(h)}$	3.0 (4.9)	3.9 (6.6)	2.3 (6.9)	2.5 (6.3)	22.4 (18.2)	20.3 (75.1)	1.7 (7.5)
$r_s^{e(h)} _{10^{12} \text{cm}^{-2}}$	20.6 (33.8*)	26.7 (45.2*)	24.0 (71.6*)	18.2 (46.0*)	35.0* (28.5)	33.3* (122.9*)	18.3 (81.1*)
T_L [K]	7.1	6.1	7.6	6.7	1.4	1.7	9.3
$n_{e(h)}^{\max}$ [10^{12}cm^{-2}]	0.5 (1.3)	0.8 (2.3)	0.7 (5.9)	0.4 (2.4)	1.4 (0.9)	1.3 (17.4)	0.4 (7.5)
$\nu_{\max}^{e(h)}$	0.28 (0.73)	0.46 (1.31)	0.17 (1.46)	0.2 (1.19)	15.56 (10.0)	13.0 (174.0)	0.09 (1.7)

III. EFFECTIVE THEORY OF GWC

Understanding the collective excitations of a GWC is critical in distinguishing them from the other density ordered states observed in the lattice system. Here, we will focus on the vibrational modes of the GWC in absence of an external magnetic field. For analytical tractability we will confine our discussion to the limit when the GWC is deep in the crystalline regime. We represent the particle density of the system using a lattice of Gaussian wave-packets of size 2ξ (see Fig. 2),

$$\rho(\mathbf{x}) = \sum_i |\psi(\mathbf{x} - \mathbf{R}_i)|^2, \quad (4)$$

$$|\psi(\mathbf{x})|^2 = \frac{1}{2\pi\xi} \exp(-|\mathbf{x}|^2/4\xi^2).$$

where $\mathbf{R}_i = \mathbf{R}_i^0 + \mathbf{u}_i(t)$, describes fluctuations around the mean lattice sites \mathbf{R}_i^0 . The GWC we consider is far away from the phase boundary with the liquid phase so that we can treat the mean fluctuation in the position of the localized electrons, $\langle r^2 \rangle \sim \xi^2$, to be much smaller than the Wigner lattice periodicity, $\xi \ll \lambda_w$, as in Fig. 2. Since the field $\mathbf{u}_i(t)$ measures the fluctuation around the mean position of a particle, it is naturally $\mathcal{O}(\xi)$. For a GWC at $T = 0$, ξ (hence, \mathbf{u}_i) can be tuned by changing the density alone. A self-consistent solution of ξ as a function of density is discussed in Ref. [74]. Finally, since ξ increases with increasing temperature, we will restrict our discussion to low temperature, $T \ll T_L$.

In this regime, the above density functional can be written in terms of harmonics (see App. B for a derivation)

$$\rho(\mathbf{x}) \simeq \rho_0 \left[1 - \nabla \cdot \mathbf{u}(\mathbf{x}) + \sum_{l \neq 0} e^{i\mathbf{K}_l \cdot \mathbf{x}} \rho_l(\mathbf{x}) \right]. \quad (5)$$

Here, $\rho_l(\mathbf{x}) = e^{-i\mathbf{K}_l \cdot \mathbf{u}(\mathbf{x})}$ and ρ_0 is the average density (over the entire sample). The second term accounts for

long range density fluctuations over several λ_w and couples to couples to the long-range (or $q \sim 0$) component of the Coulomb interaction. The remaining terms take care of the density fluctuations at a length scale comparable to or smaller than λ_w and hence can be referred to as un-smearred density. The wave vectors $\mathbf{K}_l = \{\pm l\boldsymbol{\kappa}_n\}$ denote the Brillouin zone (BZ) vectors of the undeformed GWC. Here, $l = 1, 2, \dots$ are simply ‘size multipliers’ of the BZ. Formally, the $l = 0$ term is nothing other than ρ_0 in Eq. (5). The last term above also contains a summation over the index n appearing through \mathbf{K}_l . We perform this summation implicitly since it does not play any significant role in our analysis.

The long wavelength theory describing the fluctuations of the crystal is given by an elastic Hamiltonian

$$H_{\text{eff}} = \frac{1}{2} \sum_{\omega_n} \int \frac{d^2 \mathbf{q}}{(2\pi)^2} u_\alpha(\mathbf{q}, \omega_n) \Phi_{\alpha\beta}(\mathbf{q}, \omega_n) u_\beta(-\mathbf{q}, -\omega_n), \quad (6)$$

where $\alpha, \beta = x, y$ are summed over, momenta $\{\mathbf{q}\}$ form the Fourier basis, and the kernel $\Phi_{\alpha\beta}(\mathbf{q}, \omega_n)$ is the elastic matrix. Henceforth, we will express all the quantities after performing the frequency (ω_n) summation. In case of a classical ($\omega_n = 0$) free theory, this matrix is $\Phi_{\alpha\beta} = cq^2 \delta_{\alpha\beta}$, with the real space Hamiltonian $H_{\text{eff}} = \frac{c}{2} \int d^2 \mathbf{x} [\nabla \cdot \mathbf{u}(\mathbf{x})]^2$. Here c is an elastic modulus. The presence of the moiré potential and the Coulomb interaction between the particles generates the following terms in the hamiltonian

$$H_1 = H_{e-1} + H_{e-e}, \quad (7)$$

where electron-moiré lattice interaction and the electron-electron interaction terms, respectively, are

$$H_{e-1} = - \int_{\mathbf{x}} V(\mathbf{x}) \rho(\mathbf{x}), \quad (8a)$$

$$H_{e-e} = \frac{1}{2} \int_{\mathbf{x}, \mathbf{x}'} U(\mathbf{x} - \mathbf{x}') [\rho(\mathbf{x}) - \rho_0] [\rho(\mathbf{x}') - \rho_0]. \quad (8b)$$

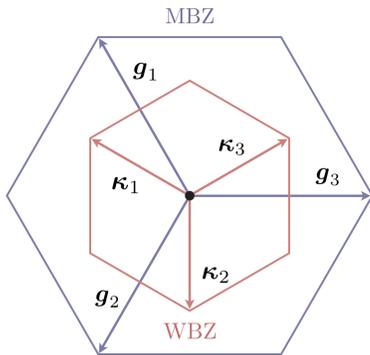


FIG. 3. Schematic of a MBZ (blue) and a WBZ (red). The BZ vectors are, $|\mathbf{g}_n| = \frac{4\pi}{\sqrt{3}\lambda_m}$ and $|\boldsymbol{\kappa}_n| = \frac{4\pi}{\sqrt{3}\lambda_w}$. In general, since $\lambda_w > \lambda_m$, the WBZ is smaller than the MBZ. For the particular case drawn above, $\lambda_w = 3\lambda_m$. In other words, the third WBZ is the same as the first MBZ ($|\boldsymbol{\kappa}_n| = 3|\mathbf{g}_n|$).

We will approximate the (triangular) moiré potential, $V(\mathbf{x})$, by [19, 75]

$$V(\mathbf{x}) = 2\tilde{V} \sum_{m=1}^3 \cos(\mathbf{x} \cdot \mathbf{g}_m + \phi), \quad (9)$$

where $\tilde{V} \sim \mathcal{O}(10 \text{ meV})$ sets the depth of the moiré potential and ϕ determines the shape of the potential. These two (intrinsic) parameters can be fixed for a given TMD using methods developed in Ref. [75]. Lastly, the unit vectors of the moiré Brillouin zone (MBZ) are given by $\mathbf{g}_m = \frac{4\pi}{\sqrt{3}\lambda_m} (\cos \frac{2\pi m}{3}, \sin \frac{2\pi m}{3})$. Similarly, the unit vectors of the Wigner Brillouin zone (WBZ) are $\boldsymbol{\kappa}_n = \frac{4\pi}{\sqrt{3}\lambda_w} (\cos \frac{2\pi n}{3}, \sin \frac{2\pi n}{3})$.

A. Interaction with the moiré potential

We now focus on the moiré potential given by the first term in Eq. (7). In terms of a reciprocal vector of the MBZ, $\mathbf{G}_m = \{\pm m\mathbf{g}_n\}$, see Fig. 3, the periodic moiré potential is [19, 63]

$$V(\mathbf{x}) = \sum_m V_m e^{i\mathbf{G}_m \cdot \mathbf{x}}. \quad (10)$$

As before, m is a size multiplier for the principal MBZ and a summation over the index n is made implicit. We assume the potential to be an even function in position space and set the $m = 0$ mode to zero. For the potential in Eq. (9), we obtain $V(\mathbf{G}_m) = \tilde{V} e^{i\text{sgn}(m)\phi}$. Substituting Eq. (10) in Eq. (8a), we obtain the following moiré term

$$H_{e-1} = -\rho_0 \sum_{l,m} V_m \int d\mathbf{x} e^{i(\mathbf{K}_l - \mathbf{G}_m) \cdot \mathbf{x}} \rho_l(\mathbf{x}). \quad (11)$$

In writing the above expression, we have set the energy of the moiré lattice, $\sim \int_{\mathbf{x}} V(\mathbf{x})$, to zero and neglected

the gradient term in the density as this term represents an external source term (linear in \mathbf{u}) and does not contribute to the physics of the pinning gap. Note that the integrand here involves both the WBZ and the MBZ vectors. This term plays a critical role in imposing a certain set of commensuration constraints. In general, a GWC need not conform to the lattice symmetries of a background (e.g., moiré) lattice. With changing density, one often anticipates the GWC to go through a large set of commensurate-incommensurate transitions, also known as the devil's staircase [76, 77], where the incommensurate structures may also have a completely different lattice symmetry [78] and associated stability issues. These states and the accompanying transitions cannot be described by the elastic (linear harmonic) theory developed here.

In this paper, we focus exclusively on the case where the GWC and the background lattice share the same lattice symmetry, such as in the experiment of Regan *et al.* [13]. As we will show below, this leads to a geometrical condition $r\mathbf{G}_1 = s\mathbf{K}_1$, where (r, s) are co-primes and the subscript 1 refers to the principal BZ vectors. Finally, since usually $\lambda_w \geq \lambda_m$, hence $|\mathbf{G}_1| \geq |\mathbf{K}_1|$. As a result, $r \leq s$. For instance, the WC observed in [13] at $1/3$ -filling, or a state at $\nu = 1/3^n$ in general, simply has its BZ shrunk (without any rotation) by a factor of 2^n . This state of affairs obtains because the GWC at $1/3^n$ -filling has a unit cell that is 2^n times larger than that of the moiré lattice. Therefore, for $\nu = 1/3^n$, $r = 1$ and $s = 2^n$.

B. Electronic interaction

Using the underlying translation invariance, we write the interaction term in, Eq. (7) as

$$H_{e-e} = \frac{\rho_0^2}{2} \int_{\mathbf{x}, \mathbf{x}'} U(\mathbf{x} - \mathbf{x}') [\nabla \cdot \mathbf{u}(\mathbf{x})] [\nabla \cdot \mathbf{u}(\mathbf{x}')] + \frac{\rho_0^2}{2} \int_{\mathbf{x}, \mathbf{x}'} \sum_l U(\mathbf{x} - \mathbf{x}') e^{i\mathbf{K}_l \cdot (\mathbf{x} - \mathbf{x}')} \rho_l(\mathbf{x} - \mathbf{x}'). \quad (12)$$

Note that terms with $K_l \neq K_m$ have been discarded as they are highly oscillatory.

We now switch from the cartesian basis u_x, u_y to one described by the longitudinal (u_{\parallel}) and transverse (u_{\perp}) components with respect to the momentum vectors (\mathbf{q})

$$u_{\alpha}(\mathbf{q}) = u_{\parallel}(\mathbf{q})\hat{\mathbf{q}}_{\alpha} + u_{\perp}(\mathbf{q})\epsilon_{\alpha\beta}\hat{\mathbf{q}}_{\beta}, \quad (13)$$

where $\alpha, \beta = \{x, y\}$, and $\epsilon_{\alpha\beta}$ is an antisymmetric tensor, $\epsilon_{xy} = 1 = -\epsilon_{yx}$. Note that u_{\parallel} and u_{\perp} are the bulk compression and shear modes respectively. In this basis,

the first term, $H_{e-e}^{(1)}$, in Eq. (8b) becomes

$$\begin{aligned} H_{e-e}^{(1)} &= \frac{\rho_0^2}{2} \int_{\mathbf{x}, \mathbf{x}'} U(\mathbf{x} - \mathbf{x}') [\nabla \cdot \mathbf{u}(\mathbf{x})] [\nabla \cdot \mathbf{u}(\mathbf{x}')] \\ &= \frac{d_{\parallel}}{2} \sum_{\mathbf{q}} q u_{\parallel}(\mathbf{q}) u_{\parallel}(-\mathbf{q}) \quad , \quad \frac{\rho_0^2 e^2}{\epsilon} \equiv d_{\parallel}. \end{aligned} \quad (14)$$

We see that the $q = |\mathbf{q}|$ term results from the long-range (in 2D) nature of the interaction, $U(\mathbf{q}) \sim 1/q$. Had we considered a shorter-range interaction of the form $U(\mathbf{q}) \sim 1/q^\gamma$, the proportionality above would have been modified to $q^{2-\gamma}$. The transverse modes do not change the local density and remain unaffected by the Coulomb interaction. Typically, long wavelength electrostatic fluctuations, namely the plasma modes, are always longitudinal in the absence of a magnetic field (since $\mathbf{q} \times \mathbf{E} = 0$, where \mathbf{E} is an electric field).

In the elastic limit $|\mathbf{u}(\mathbf{x}) - \mathbf{u}(\mathbf{x}')| \ll \lambda_w$, we Taylor expand the second term, $H_{e-e}^{(2)}$, in Eq. (12). The first-order term vanishes because the undeformed GWC has an energy minimum at $\mathbf{u} = 0$ and the second-order term gives the correction

$$\begin{aligned} H_{e-e}^{(2)} &\simeq \frac{\rho_0^2}{2} \sum_l \int_{\mathbf{x}, \mathbf{x}'} V(\mathbf{x} - \mathbf{x}') e^{i\mathbf{K}_l \cdot (\mathbf{x} - \mathbf{x}')} K_{l,\alpha} K_{l,\beta} \\ &\quad \times [u_\alpha(\mathbf{x}) - u_\alpha(\mathbf{x}')] [u_\beta(\mathbf{x}) - u_\beta(\mathbf{x}')] . \end{aligned} \quad (15)$$

Here, $\mathbf{K}_{l,\alpha}$ denote the $\alpha = x, y$ components of \mathbf{K}_l . Henceforth, unless mentioned, we will set $\rho_0 = 1$.

As shown in App. C, this term can be absorbed into a redefinition of the elastic coefficients [28, 79, 80]. We note that we have considered these elastic constants to be q -independent, which is a feature of the local elastic theory. One can also extend this analysis to non-local elastic theories where these constants can be considered to be q -dependent. Generalizing to an interaction of the form $U(\mathbf{x}) \sim 1/|\mathbf{x}|^\gamma$, we find that the full Hamiltonian defining the low energy fluctuations of the GWC can be expressed as

$$\begin{aligned} H_{\text{eff}} &= \int_{\mathbf{q}} u_{\parallel}(\mathbf{q}) \Omega_{\parallel} u_{\parallel}(-\mathbf{q}) + u_{\perp}(\mathbf{q}) \Omega_{\perp} u_{\perp}(-\mathbf{q}) + H_{e-1}; \\ \Omega_{\parallel}(\mathbf{q}) &= c_{\parallel} q^2 + d_{\parallel} q^{2-\gamma} \quad , \quad \Omega_{\perp}(\mathbf{q}) = c_{\perp} q^2. \end{aligned} \quad (16)$$

H_{e-1} is given by (11). Ω_a are the dispersions of the longitudinal and the transverse modes. As discussed previously, it is only the longitudinal mode whose dispersion is affected by γ , see Fig. 4. Secondly, as discussed in App. C, these elastic constants follow $c_a \propto \lambda_w^\gamma / \epsilon$. Notably, the elastic modulus d_{\parallel} is a density-independent constant only in the low density limit far away from WC melting. Also, as screening (ϵ) increases, the WC becomes loosely bound due to reduced interaction. This makes a WC less rigid, or c_a decreases with increasing ϵ .

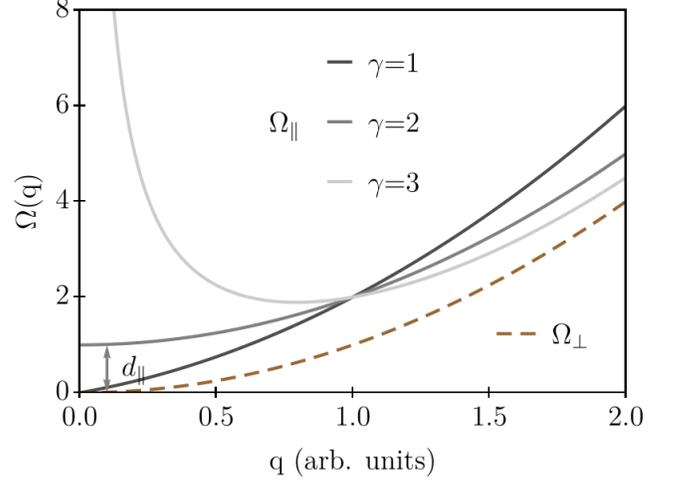


FIG. 4. Dispersion of the longitudinal (solid) and transverse (dashed) modes of a 2D WC (independent of its coupling to a moiré potential). Here, we have set $c_a = 1 = d_{\parallel}$. $\gamma = 1$ corresponds to the long-range Coulomb interaction. With increasing γ , the interaction becomes increasingly short-range. For $\gamma = 2$, as can be seen from Eq. (16), a gap of size d_{\parallel} appears in the longitudinal mode. With further increase in γ , this gap diverges.

IV. GAUSSIAN VARIATIONAL MINIMIZATION

In this section, we treat the effective Hamiltonian obtained in the previous section using the Gaussian variational method (GVM) developed in Refs. [45, 46, 81]. This allows us to obtain the dispersion of the vibrational modes of the GWC and the associated pinning gap arising from the interaction between the Wigner lattice and the moiré lattice. Motivated by the experiments, we assume the GWCs to be weakly coupled to the moiré lattice. This allows us to treat the vibrations of the localized particles as harmonic fluctuations. This is formalized by the GVM as follows. Consider a Hamiltonian $H = \frac{1}{2} \int_{\mathbf{q}} u(\mathbf{q}) \Omega(\mathbf{q}) u(-\mathbf{q}) + H'$, where the kernel $\Omega(\mathbf{q})$ is known *a priori* and H' can contain non-linear or polynomial terms in the field $u(\mathbf{q})$. For a vector field $\mathbf{u}(\mathbf{q})$, this kernel becomes a matrix. The goal is to approximate the Hamiltonian H by the following quadratic form,

$$H_0 = \frac{1}{2} \int_{\mathbf{q}} u(\mathbf{q}) \mathcal{G}^{-1}(\mathbf{q}) u(-\mathbf{q}). \quad (17)$$

The optimal function $\mathcal{G}(q)$ is then obtained by minimizing the variational free energy of the theory H , $F_{\text{var}} = F_0 + \langle H - H_0 \rangle_0$ where $\langle \dots \rangle_0$ is the expectation value evaluated with H_0 with respect to $\mathcal{G}(q)$. In App. D we provide a pedagogical discussion on using this GVM method for the simple case of a Sine-Gordon (SG) interaction as the hamiltonian in (16) closely resembles the SG problem.

A. Applying GVM to GWC

We use the GVM to obtain the gap opened by the moiré lattice. Since the displacement is a two component field we have both $\Omega_{\parallel}(\mathbf{q})$ and $\Omega_{\perp}(\mathbf{q})$. The variational free energy becomes

$$F_{\text{var}} = \frac{T}{2} \int_{\mathbf{q}} \sum_{a=\parallel,\perp} \left\{ \Omega_a(\mathbf{q}) \mathcal{G}_a(\mathbf{q}) - \log[T \mathcal{G}_a(\mathbf{q})] \right\} - \sum_{l,m} V_m \delta(\mathbf{K}_l - \mathbf{G}_m) \exp \left[-\frac{T}{2} \sum_{b=\parallel,\perp} K_{l,b}^2 \int_{\mathbf{q}} \mathcal{G}_b(\mathbf{q}) \right]. \quad (18)$$

Note that in the absence of a magnetic field there is no admixture of the longitudinal and transverse modes.

The Green function that minimizes the free energy in Eq. (18) can be approximated by $\mathcal{G}_a^{(0)}(\mathbf{q}) \simeq \frac{1}{\Omega_a(\mathbf{q}) + \Delta_a}$, where the gaps Δ_a satisfy the following self-consistent equations (SCE)

$$\Delta_a = \sum_{m \in \mathcal{M}} V_m G_{m,a}^2 \exp \left(-\frac{T}{2} \int_{\mathbf{q}} \sum_{b=\perp,\parallel} \frac{G_{m,b}^2}{\Omega_b(\mathbf{q}) + \Delta_b} \right). \quad (19)$$

Here a is not in the Cartesian basis but in the orthonormal basis discussed in Eq. (13). Though at first glance Eq. (19) seems independent of \mathbf{K}_l (or λ_w), we note that the conservation of momentum imposed through the delta function in Eq. (18), restricts the set of \mathbf{G}_m to those satisfying $r\mathbf{G}_1 = s\mathbf{K}_1$. The set of such restricted (momentum conserving) values of \mathbf{G}_m is denoted by \mathcal{M} . For instance, for the WC at 1/3-filling, since, as explained previously, $r = 1$, \mathcal{M} is trivially the first MBZ. After integrating, we find that the gap equations take the form

$$\Delta_a \simeq \sum_{m \in \mathcal{M}} V_m G_{m,a}^2 \left(\frac{\Delta_{\perp}}{c_{\perp} \Lambda^2} \right)^{\frac{T G_{m,\perp}^2}{8\pi c_{\perp}}} \left(\frac{\Delta_{\parallel}}{c_{\parallel} \Lambda^2} \right)^{\frac{T G_{m,\parallel}^2}{8\pi c_{\parallel}}} \times \exp \left[\frac{T G_{m,\parallel}^2}{8\pi c_{\parallel}} \tilde{\Delta}_{\parallel} \left(\pi + 2 \tan^{-1} \tilde{\Delta}_{\parallel} \right) \right]. \quad (20)$$

Here, $d_{\parallel} / \sqrt{4c_{\parallel} \Delta_{\parallel} - d_{\parallel}^2} \equiv \tilde{\Delta}_{\parallel}$ and Λ is a UV cutoff for the momentum space integration. The zero temperature limit for the gap above is $\sum_{m \in \mathcal{M}} V_m G_{m,a}^2 \equiv \Delta_a^0$. And, a low temperature expansion is obtained to be

$$\begin{aligned} \Delta_{\perp} &= A_{\perp} + B_{\perp} \log \Delta_{\perp}; \\ A_{\perp} &= \Delta_{\perp}^0 + T \sum_{m \in \mathcal{M}} V_m G_{m,\perp}^2 \left[D_m^{\parallel} - \frac{G_{m,\perp}^2}{8\pi c_{\perp}} \log c_{\perp} \Lambda^2 \right], \\ D_m^{\parallel} &= \frac{G_{m,\parallel}^2}{8\pi c_{\parallel}} \log \frac{\Delta_{\parallel}}{c_{\parallel} \Lambda^2} + \frac{G_{m,\parallel}^2}{8\pi c_{\parallel}} \tilde{\Delta}_{\parallel} \left(\pi + 2 \tan^{-1} \tilde{\Delta}_{\parallel} \right), \\ B_{\perp} &= T \sum_{m \in \mathcal{M}} V_m \frac{G_{m,\perp}^4}{8\pi c_{\perp}}. \end{aligned} \quad (21)$$

Here A_{\perp} is dependent on Δ_{\parallel} , and B_{\perp} is a geometric constant. From this, we obtain a closed-form expression for Δ_{\perp} in terms of Δ_{\parallel} . By bringing the above equation to the form $w e^w = z$, we obtain the solution $w = W_k(z)$, where $W_k(z)$ is the (multivalued) Lambert W function with its branch indexed by the integer k . In fact, when $w < 0$ (for us, $w = -\Delta_{\perp}/B_{\perp}$), the solution has two branches, $W_0(z)$ and $W_{-1}(z)$. We will drop the latter solution since it is not a regular function at $\Delta_{\perp} = 0$. Therefore,

$$\Delta_{\perp} = -B_{\perp} W_0 \left(-\frac{e^{-A_{\perp}/B_{\perp}}}{B_{\perp}} \right). \quad (22)$$

This is the explicit dependence of Δ_{\perp} on Δ_{\parallel} (through A_{\perp} only). Similarly, an SCE for the Δ_{\parallel} component is

$$\Delta_{\parallel} = \Delta_{\parallel}^0 + T \sum_{m \in \mathcal{M}} V_m G_{m,\parallel}^2 \frac{G_{m,\perp}^2}{8\pi c_{\perp}} \log \frac{\Delta_{\perp}}{c_{\perp} \Lambda^2}, \quad (23)$$

where Δ_{\perp} is given by Eq. (22). In the next subsection we discuss the solutions obtained here, especially in conjunction with the recent experiments.

B. Discussions

Note that the last term in Eq. (20) is an artifact of the long-range interaction which vanishes if $d_{\parallel} = 0$. This term, which is the compression term, purely accounts for the elastic contribution to the gap. For $d_{\parallel} = 0$, the gap equation is equivalent to the vector SG potential, see Eq. (D6).

Secondly, since Λ appears in the denominator of Eq. (20), the gap vanishes for temperatures larger than a characteristic temperature, $\min \left(\frac{8\pi c_{\parallel}}{G_{\parallel}^2}, \frac{8\pi c_{\perp}}{G_{\perp}^2} \right) \equiv T_*$. This is a feature of the equivalence of the effective interaction Hamiltonian to that with the SG potential, see discussions in the App. D. The analysis is valid only if T_* is much smaller than the melting temperature (such as T_L) of a GWC. Note that since $T_* \sim c_a \lambda_w^2$, this temperature scale can be controlled by means of the twist angle.

The pinning frequency is related to the zero temperature gap as [81] $\omega_p^a = \sqrt{\Delta_a^0 / \rho_0}$. Notably, since the pinning frequency scales with the size of the WBZ, $\omega_p^a \propto G_a$, it becomes increasingly difficult to de-pin a WC of smaller unit cell. This is since a WC with large unit cell (or small \mathbf{G}) will be loosely bound compared to one with smaller unit cell (as the particles are more tightly packed). Therefore, the former can be easily de-pinned by an external electric field. Similarly, for a deeper moiré potential the pinning frequency increases ($\omega_p^a \propto \tilde{V}^{1/2}$) since the particles get tightly bound to the potential minima. Introduction of a spacer layer can further modulate this frequency. Geometrical factors aside, the pinning gap thus becomes $\mathcal{O}(\lesssim \text{meV})$. With increasing temperature, as seen in Eq. (21), this gap softens as the increasing thermal fluctuation facilitates de-pinning. The extent to

which this gap decreases depends on various coefficients appearing in Eq. (21). Most notably, via the elastic constants, c_α , the logarithm term has a coefficient that is directly proportional to the dielectric term. Thus, the larger the screening, the smaller the pinning gap. Therefore, although the geometrical constants associated with various HoM or HeM TMDs may not affect the pinning gap of a GWC, the dielectric constant can however alter the physics. This gap translates into determining which state is a stronger insulator.

V. CONCLUSION

We have addressed the feasibility of realizing Wigner crystals in a host of HoM and HeM systems. Note however, that our results are based on estimated material parameters of the TMD moiré materials. Corrections to these results might arise principally from three sources. The first is from the full band structure of the TMD heterostructures [30]. Second, a material correction arising from twist-angle inhomogeneity across a sample [82, 83] which may cause additional pinning or de-pinning of the WC could also affect the physics. Similar effects may also arise from atomic relaxations [84, 85]. Third, the presence or absence of a spacer layer [16], such as a monolayer hBN, may also affect the correlation energy, thereby affecting Wigner crystallization. A first principles calculation of the elastic coefficients of the GWC is also important to obtain good qualitative and quantitative estimates for the pinning gap and the phonon spectrum. All of these aspects merit further studies as this will help narrow the density and temperature regimes where WC is feasible.

Due to the presence of a pinning gap, transport measurements to confirm the existence of WC states can be misleading as there can be many other kinds of insulating states with similar transport characteristics. Although observation of such states at fractional occupancy increases their likelihood of being Wigner states, especially for those observed at incommensurate fillings, however, the possibility of other density ordered states cannot be ruled out, particularly for commensurate fractional occupancies. Devising smoking gun evidence for various density ordered states may be an interesting task for theorists and experimentalists alike.

As was mentioned before, once a system meets the material constraints to realize a GWC, there exists a plethora of crystalline states below the filling fraction ν^{\max} . These states constitute a devil's staircase and have a rich physics of commensurate-incommensurate transitions [76, 86, 87]. Due to various stability criteria, only a few such states might display clear experimental signatures. However, with careful analysis or improvements in experimental conditions, one may gain insight into the other states as well. In fact, a theoretical framework to understand these commensurate-incommensurate transitions in presence of an underlying lattice is an interesting

theoretical task and is left for future work.

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APPENDIX

Appendix A: HeM at $\theta = 5^\circ$

In Table A1 we present crystallization parameters for HeMs for large twist angles. For large twist angles, doping levels need to be extremely low for obtaining GWCs.

Appendix B: Harmonic Expansion of Density

Following [45], we derive the elastic limit of the density written in Eq. (5). A continuum limit can be easily obtained if we treat the equilibrium GWC configuration, $\mathbf{R}_i^0 = \mathbf{R}_i - \mathbf{u}(\mathbf{R}_i^0)$, as a slowly varying smooth vector field, $\varphi(\mathbf{x})$, over the position of the particles

$$\varphi(\mathbf{x}) = \mathbf{x} - \mathbf{u}(\varphi(\mathbf{x})) . \quad (\text{B1})$$

Clearly a solution of $\varphi(\mathbf{x})$ is given by, $\varphi(\mathbf{R}_i) = \varphi(\mathbf{R}_i^0 + \mathbf{u}(\mathbf{R}_i^0)) = \mathbf{R}_i^0$. Using the above equality we can rewrite the density in terms of this new field as

$$\rho(\mathbf{x}) = \sum_i \delta^{(2)}[\mathbf{R}_i - \varphi(\mathbf{x}) - \mathbf{u}(\varphi(\mathbf{x}))] \quad (\text{B2a})$$

$$\simeq \det[\partial_\alpha \varphi_\beta(\mathbf{x})] \sum_i \delta^{(2)}(\mathbf{R}_i - \varphi(\mathbf{x})) \quad (\text{B2b})$$

$$= \det[\partial_\alpha \varphi_\beta(\mathbf{x})] \int \frac{d\mathbf{q}}{(2\pi)^2} \rho_0(\mathbf{q}) e^{i\mathbf{q} \cdot \varphi(\mathbf{x})} . \quad (\text{B2c})$$

The first simplification was done using the elastic limit, $\partial_\alpha \mathbf{u}_\beta \ll 1$. In the last line, we have used the integral representation of the delta function. In the presence of an undeformed GWC, we can introduce its reciprocal vectors, $e^{i\mathbf{K}_l \cdot \mathbf{R}_i} = 1$, to write

$$\rho_0(\mathbf{q}) = \sum_i e^{i\mathbf{q} \cdot \mathbf{R}_i} = \rho_0 (2\pi)^2 \sum_l \delta^{(2)}(\mathbf{q} - \mathbf{K}_l) . \quad (\text{B3})$$

Here ρ_0 is the average number density. Introducing the above simplification in Eq. (B2c) and using Eq. (B1), we obtain

$$\begin{aligned} \rho(\mathbf{x}) &= \rho_0 \det[\partial_\alpha \varphi_\beta(\mathbf{x})] \sum_l e^{i\mathbf{K}_l \cdot \varphi(\mathbf{x})} \\ &= \rho_0 \det[1 - \partial_\alpha \mathbf{u}_\beta(\varphi(\mathbf{x}))] \sum_l e^{i\mathbf{K}_l \cdot [\mathbf{x} - \mathbf{u}(\varphi(\mathbf{x}))]} \\ &\simeq \rho_0 - \rho_0 \nabla \cdot \mathbf{u}(\mathbf{x}) + \rho_0 \sum_l e^{i\mathbf{K}_l \cdot [\mathbf{x} - \mathbf{u}(\mathbf{x})]} . \end{aligned} \quad (\text{B4})$$

We again used the elastic limit by first Taylor-expanding the determinant operator, \det , and then substituting $\mathbf{u}(\varphi(\mathbf{x})) \approx \mathbf{u}(\mathbf{x})$ which works for \mathbf{x} close to the equilibrium position and in the elastic limit. This leads us

TABLE A1. Wigner crystallization criteria for a HeM at $\theta = 5^\circ$: The material parameters here are the same as those in Table II. Therefore, the r_s values remain the same for $n_e = 10^{12} \text{ cm}^{-2}$. Similarly, the $n_{e(h)}^{\max}$ values remain the same, however, $\nu_{\max}^{e(h)}$ is different since the supercell density has changed.

HeMs	WSe ₂ /WS ₂	MoSe ₂ /MoS ₂	MoTe ₂ /MoSe ₂	MoSe ₂ /WS ₂	MoSe ₂ /WSe ₂	MoS ₂ /WS ₂	MoTe ₂ /WSe ₂
λ_m [nm]	3.4	3.4	3.2	3.4	3.8	3.6	3.2
$U/W _{e(h)}$	1.3 (2.1)	1.6 (2.8)	1.4 (4.2)	1.1 (2.8)	2.4 (1.9)	2.2 (8.0)	1.0 (4.6)
T_L [K]	16.8	14.5	12.6	14.9	13.4	15.6	15.0
$\nu_{\max}^{e(h)}$	0.05 (0.13)	0.08 (0.24)	0.06 (0.53)	0.04 (0.24)	0.17 (0.11)	0.15 (1.98)	0.04 (0.66)

to Eq. (5). Note there is complete decoupling between the gradient term and the terms with \mathbf{K}_l . This occurs because $\mathbf{u}(\mathbf{x})$ has negligible Fourier components outside the WBZ.

Appendix C: Elastic Interaction Hamiltonian

In this Appendix, we clarify the derivation of Eq. (15). First, we Fourier transform the second part of Eq. (12),

$$H_{e-e}^{(2)} = \frac{1}{2} \sum_l \int_{\mathbf{x}, \mathbf{x}'} U(\mathbf{x} - \mathbf{x}') e^{i\mathbf{K}_l \cdot (\mathbf{x} - \mathbf{x}')} K_{l,\alpha} K_{l,\beta} \times \int_{\mathbf{q}, \mathbf{q}'} u_\alpha(\mathbf{q}) u_\beta(\mathbf{q}') \left(e^{i\mathbf{q} \cdot \mathbf{x}} - e^{i\mathbf{q} \cdot \mathbf{x}'} \right) \left(e^{i\mathbf{q}' \cdot \mathbf{x}} - e^{i\mathbf{q}' \cdot \mathbf{x}'} \right). \quad (\text{C1})$$

In order to simplify it further, we introduce the center of mass coordinate, $2\mathbf{X} = \mathbf{x} + \mathbf{x}'$ and the relative coordinate $2\boldsymbol{\delta} = \mathbf{x} - \mathbf{x}'$ to obtain

$$H_{e-e}^{(2)} = \sum_l K_{l,\alpha} K_{l,\beta} \int_{\mathbf{q}} u_\alpha(\mathbf{q}) u_\beta(-\mathbf{q}) \times \int_{\boldsymbol{\delta}} d\boldsymbol{\delta} U(\boldsymbol{\delta}) [1 - \cos(\mathbf{q} \cdot \boldsymbol{\delta})] e^{i\mathbf{K}_l \cdot \boldsymbol{\delta}}. \quad (\text{C2})$$

In coming to this line, we have also integrated out \mathbf{q}' , which introduced a delta function, $\delta^{(2)}(\mathbf{q} + \mathbf{q}')$. Next, we perform the last integration for a generic potential of the form, $U(\mathbf{x}) = e^2/\epsilon |\mathbf{x}|^\gamma$. One can obtain the long-range Coulomb potential by setting $\gamma = 1$, and with increasing γ the potential becomes increasingly short-range. For such a $U(\boldsymbol{\delta})$ we find that

$$H_{e-e}^{(2)} = \sum_l K_{l,\alpha} K_{l,\beta} \int_{\mathbf{q}} u_\alpha(\mathbf{q}) u_\beta(-\mathbf{q}) \times \frac{e^2}{\epsilon} \left(\frac{2}{|\mathbf{K}_l|^\gamma} - \frac{1}{|\mathbf{K}_l - \mathbf{q}|^\gamma} - \frac{1}{|\mathbf{K}_l + \mathbf{q}|^\gamma} \right). \quad (\text{C3})$$

For further simplification, we confine our discussion to the low-energy limit. This allows us to Taylor-expand the last term in Eq. (C3) for the limit $|\mathbf{q}| \ll |\mathbf{K}_l|$. The first term in this expansion, which is linear in \mathbf{q} , vanishes because it involves integrating over a $\cos \theta_l$ term. Here, θ_l

are the angles between the \mathbf{q} vector and \mathbf{K}_l . Therefore, retaining up to the $\mathcal{O}(q^2)$ term we obtain,

$$H_{e-e}^{(2)} \simeq \gamma \frac{e^2}{\epsilon} \sum_l \frac{K_{l,\alpha} K_{l,\beta}}{|\mathbf{K}_l|^{2+\gamma}} \int_{\mathbf{q}} q^2 u_\alpha(\mathbf{q}) u_\beta(-\mathbf{q}) \times [(2 + \gamma) \cos^2 \theta_l - 1]. \quad (\text{C4})$$

Note that unlike the long-distance term, $H_{e-e}^{(1)}$ in Eq. (14), the leading dispersion corresponding to $H_{e-e}^{(2)}$ remains quadratic regardless of the choice of γ .

$$H_{e-e}^{(2)} = \int_{\mathbf{q}} c_{\parallel} q^2 u_{\parallel}(\mathbf{q}) u_{\parallel}(-\mathbf{q}) + c_{\perp} q^2 u_{\perp}(\mathbf{q}) u_{\perp}(-\mathbf{q}). \quad (\text{C5})$$

Appendix D: GVM for Sine-Gordon Potential

In this Appendix, we demonstrate the GVM method discussed in the main text for a Sine-Gordon (SG) potential,

$$H = \frac{1}{2} c \int dx [\nabla \phi(x)]^2 - g \int dx \cos[2\phi(x)]. \quad (\text{D1})$$

Here c and g are free parameters. Using the simplifications discussed in the main text [and using $\Omega(q) = cq^2$], we obtain the variational free energy to be

$$F_{\text{var}} = -\frac{T}{2} \int_q \log[T\mathcal{G}(q)] + \frac{T}{2} \int_q cq^2 \mathcal{G}(q) - g \exp \left[\frac{T}{2} \int_q \mathcal{G}(q) \frac{\partial^2}{\partial \phi^2} \right] \int dx \cos(2\phi) \Big|_{\phi=0}. \quad (\text{D2})$$

Further simplifications of the last term leads us to

$$F_{\text{var}} = -\frac{T}{2} \int_q \log[T\mathcal{G}(q)] + \frac{T}{2} \int_q cq^2 \mathcal{G}(q) - ge^{-2 \int_q T\mathcal{G}(q)}. \quad (\text{D3})$$

In these equations, we fixed the sample area to $\int dx = 1$. The saddle point solution of the above free energy is

$$\mathcal{G}^{-1} = cq^2 + 4ge^{-2 \int_q T\mathcal{G}(q)}. \quad (\text{D4})$$

We now set $\mathcal{G}^{-1}(q) = cq^2 + m$ and solve m self-consistently,

$$m = 4ge^{-2T \int_q^\Lambda \frac{1}{cq^2 + m}} \simeq 4g \left(\frac{m}{c\Lambda^2} \right)^{T/2\pi c}. \quad (\text{D5})$$

Here, Λ is a UV cutoff in the momentum-space. A notable feature of this solution is that beyond a certain temperature maximum, $T > 2\pi c$, the SG mass must vanish simply due to the presence of the cutoff in the denominator above. Such a maximal temperature will also appear in our discussion in Sec. IV B. Additionally, from Eq. (D5) one can also deduce the scaling behavior of the mass, $m \sim g^{1/(1-\tau)}$, where $\tau = T/2\pi c$.

Pertaining to our discussion of GVM in the context of GWC, we extend the previous solutions for a SG potential to an n -component vector SG system. The interaction term here becomes $H' \sim \int dx \cos(\sum_n p_n \phi_n)$. The kernel corresponding to the field ϕ_n is $c_n q^2$. As before, we obtain the variational free energy

$$F_{\text{var}} = -\frac{T}{2} \sum_n \int_q \left\{ \log[T\mathcal{G}_n(q)] - c_n q^2 \mathcal{G}_n(q) \right\} - g e^{-\frac{T}{2} \sum_n a_n^2 \int_q \mathcal{G}(q)}. \quad (\text{D6a})$$

Since, due to the vanishing average of cosine functions, there are no cross terms such as $\cos \phi_m \cos \phi_n$ (with $m \neq n$), the saddle-point equation (setting $n = 1$ and $a_n = 2$ goes back to the original case)

$$\mathcal{G}_n^{-1} = c_n q^2 + g a_n^2 e^{-\frac{T}{2} \sum_n a_n^2 \int_q \mathcal{G}_n(q)}, \quad (\text{D6b})$$

$$\therefore m_n = g a_n^2 \exp \left[-\sum_n \frac{T a_n^2}{8\pi c_n} \log \left(\frac{c_n \Lambda^2}{m_n} \right) \right]. \quad (\text{D6c})$$

We can solve this SCE exactly and, in this case as well, there exists a similar temperature window where gap vanishes, $T a_n^2 / 8\pi c_n \equiv \tau_n > 1$. And, like before, the scaling of m_n with the coupling constant becomes, $m_n \propto g^{(1-\sum_n \tau_n)^{-1}}$. These solutions are not exactly transferable for our discussions in the main text since there the kernel has a $d_{\parallel} q^{2-\gamma}$ part. See Sec. IV B for the case when $d_{\parallel} = 0$, where the above results are perfectly applicable.

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