Evidence of frustrated magnetic interactions in a Wigner–Mott insulator

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Electrons in two-dimensional semiconductor moiré materials are more delocalized around the lattice sites than those in conventional solids1,2. The non-local contributions to the magnetic interactions can therefore be as important as the Anderson superexchange3, which makes the materials a unique platform to study the effects of competing magnetic interactions3,4. Here we report evidence of strongly frustrated magnetic interactions in a Wigner–Mott insulator at a two-thirds (2/3) filling of the moiré lattice in angle-aligned WSe2/WS2 bilayers. Magneto-optical measurements show that the net exchange interaction is antiferromagnetic for filling factors below 1 with a strong suppression at a 2/3 filling. The suppression is lifted on screening of the long-range Coulomb interactions and melting of the Wigner–Mott insulators by a nearby metallic gate. The results can be qualitatively captured by a honeycomb-lattice spin model with an antiferromagnetic nearest-neighbour coupling and a ferromagnetic second-neighbour coupling. Our study establishes semiconductor moiré materials as a model system for lattice-spin physics and frustrated magnetism5.

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We examined the magnetic response of the moiré bilayers by performing magnetic circular dichroism (MCD) measurements near the fundamental moiré exciton resonance of WSe₂, under an out-of-plane magnetic field (Fig. 2). The MCD is the difference between the reflection of left and right circularly polarized incident light normalized by the total reflection. It provides a measure of the magnetization, M, of the moiré bilayer because of the valley-dependent optical selection rules and spin-valley locking in monolayer TMDs. The magnetic-field-dependent MCD spectrum in Fig. 2a is from device 1 at v = 2/3 and T = 1.7 K. (Extended Data Fig. 1 shows the raw reflectance contrast (RC) spectrum.) The signal is enhanced near the moiré exciton resonance (≈1.67 eV). We used the integrated MCD over a narrow spectral window around the resonance to represent M, as described in earlier studies. The results were insensitive to the exact choice of the spectral window (Extended Data Fig. 2).

Figure 2b shows the magnetic-field dependence of the integrated MCD of v = 2/3 at representative temperatures. The MCD increased linearly with magnetic field B for small fields and saturated at around 1 T at 1.7 K. With increasing temperature, the MCD decreased and no clear saturation was observed within the field range of 2 T at high temperatures. We extracted the magnetic susceptibility, the Curie constant, and the Curie–Weiss (CW) law is known to describe the high-temperature magnetic response of interacting local moments with a nearest-neighbour magnetic interaction J between the local moments. The CW law is known to describe the high-temperature magnetic response of interacting local moments with a nearest-neighbour magnetic interaction J between the local moments. The CW law is known to describe the high-temperature magnetic response of interacting local moments with a nearest-neighbour magnetic interaction J between the local moments. The CW law is known to describe the high-temperature magnetic response of interacting local moments with a nearest-neighbour magnetic interaction J between the local moments.

We limited the CW analysis to the spectral window (Extended Data Fig. 2). The susceptibility follows the Curie–Weiss (CW) law (solid line), ξ = 0.07 T cm where the Curie constant ξ is proportional to the saturation magnetization and θ (about −1.4 K) is the CW temperature. The CW law is known to describe the high-temperature magnetic response of interacting local moments with reflecting the net exchange interaction energy J between the local moments (we adopted the convention of expressing J in kelvin). A negative θ corresponds to an AF exchange interaction. We limited the CW analysis to the spectral window (Extended Data Fig. 2).
5–35 K (above $\theta$) but below the melting temperature of the Wigner–Mott insulator, in which the picture of local moments applies. The fitting results were, however, not sensitive to the precise choice of the temperature range. The picture of local moments is also consistent with Fig. 2b, which reflects the alignment and saturation of the local moments under an external magnetic field. In the low-temperature limit, the saturation field ($B_s = 1$ T) also reflected $\theta$; a higher Zeeman field was required to overcome a stronger AF exchange to achieve magnetic saturation. At temperatures above $\theta$, the magnetic response was strongly affected by thermal excitations.

We performed similar measurements and analysis for the entire filling range of $\nu \leq 1$ (Extended Data Figs. 3 and 4). Figure 2c shows the filling-dependent $\chi$ at varying temperatures. At high temperatures, a linear dependence was observed (Extended Data Fig. 5), which reflects the expected linear dependence of the local moment density on filling, $C \propto \nu$. As the temperature decreased below $\nu$, $\chi$ increased drastically and a non-monotonic filling dependence emerged (Figs. 2c and 3a). Concurrently, both the CW temperature (Fig. 3b) and the saturation field at 1.7 K (Fig. 3c) also exhibited a strongly non-monotonic filling dependence. For all $\nu$ values, the CW temperature was negative or close to 0. The three quantities, $\chi$, $\theta$ and $B_s$, were fully consistent with each other. For instance, a small $|\theta|$ was accompanied by a small $B_s$ and a large $\chi$; these are manifestations of a small AF exchange interaction $J$.

The most striking feature of Fig. 3 is the strong suppression of the AF exchange around $\nu = 2/3$. The width of the feature in the filling factor agrees well with the width of the Wigner–Mott insulator at $\nu = 2/3$ (Fig. 1c, d). A similar effect is also likely at around $\nu = 1/3$, but because the local moments were far apart, the exchange energy scale was too small to be fully resolved in this experiment. Suppression of the AF exchange at other fractional fillings between 1/3 and 2/3 was not observed. As the magnetism in the triangular-lattice Hubbard models at generic fillings that involve itinerant carriers is too complex to be resolved in one study, we focused on the most notable observation at fillings around 2/3 below.

To elucidate the role of the Wigner–Mott insulator on the magnetic response at 2/3 filling, we compared the behaviour of devices 1 and 2. In device 2 (Fig. 1a), the bottom gate electrode effectively screened out the long-range Coulomb repulsion and quenched the Wigner–Mott insulator. This is consistent with recent reports and is supported by the absence of any enhancement in the exciton RC at $\nu = 2/3$ (Extended Data Fig. 6). Also no longer observable in device 2 were the pronounced peak in $\chi$ and the dip in $B_s$ at around 2/3 filling at 1.7 K (Fig. 3a, c). In addition, we were able to perform CW measurements on the temperature-dependent $\chi$ (Extended Data Fig. 7). The presence of a strong correlation probably contributes to the applicability of the CW analysis for interacting local moments here. The extracted CW temperature (Fig. 3b) was again negative with a larger magnitude for the entire filling range ($\theta = -7$ K at $\nu = 2/3$), which reflects stronger AF

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**Fig. 2 | Magneto-optical measurements (device 1).** a, MCD spectrum as a function of magnetic field at $\nu = 2/3$ and $T = 1.7$ K. The spectrum is integrated over a narrow window around the fundamental moiré exciton resonance (within the two vertical lines) to represent the sample magnetization. B, Magnetic-field dependence of integrated MCD at representative temperatures. The MCD saturates above -1 T at 1.7 K. The slope at zero magnetic field is extracted to represent $\chi$. c, Filling dependence of $\chi$ at varying temperatures. A sharp peak emerges at 2/3 filling as the temperature decreases. d, CW analysis of the temperature dependence of the inverse susceptibility (symbols). The error bars are one-sigma uncertainties of the zero-field slope in b. CW temperatures that correspond to the best fits (solid lines) are included. Data for different filling factors are vertically displaced for clarity. The horizontal dashed lines mark where 1/$\chi$ is zero. a.u., arbitrary units.
interactions between the local moments. The dip in the filling dependence of $\theta$ at around $\nu = 2/3$ was no longer observable. All these results indicate that the suppressed AF exchange interaction at a 2/3 filling in device 1 is associated with the Wigner–Mott insulator.

We considered the exchange mechanisms between local moments. The Anderson superexchange, which involves virtual hopping of particles between two neighbouring sites, was AF. When the long-range interactions were negligible, the superexchange was inversely proportional to the on-site $U(\sim \frac{1}{U})$. This mechanism contributed to the observed net AF exchange and its enhancement on screening of the Coulomb repulsions. However, the superexchange mechanism alone cannot explain the observed suppression of the net AF exchange near 2/3 filling (Fig. 3b). Other contributions, such as the interaction-assisted hopping and direct exchange, have to be considered.3–5 These non-local contributions to the exchange interaction are relevant in semiconductor moiré materials because of the harmonic trapping potentials and the extended electron Wannier functions1,2,10,13. In particular, the direct exchange is FM and proportional to Coulomb interactions; it can be tuned by screening to compete with the AF superexchange to induce frustrated magnetism.

In the Wigner–Mott insulator at $\nu = 2/3$, holes form a crystal with a honeycomb lattice30 (Fig. 1f). As the system remains insulating when immediately doped away from a 2/3 filling32, the spin physics near $\nu = 2/3$ can be captured by a (doped) spin model on the honeycomb lattice (Methods). The spin model is a good approximation in the flat band limit, as in angle-aligned WSe$_2$/WS$_2$ (refs. 6, 7). For simplicity, we considered a model with both nearest and second neighbour Heisenberg interactions. We did not make assumptions of the spin of the Heisenberg interactions to begin with, but we showed (Methods) that the experimental data favours an AF nearest-neighbour coupling $J_1$ and a FM second-neighbour coupling $J_2$ (both $J_1$ and $J_2$ are taken to be positive in our convention). A recent microscopic analysis3 showed that such a regime is possible. The choice of a FM $J_2$ is consistent with the observed increase in $|\theta|$ when the long-range interaction was screened (Fig. 3b).

Exactly at $\nu = 2/3$, there were three nearest neighbours and six second neighbours per hole. The CW temperature, which is a weighted sum of the exchange interactions $2\nu J_2$, is hence given by $\theta = -\nu J_1 - 2J_2/\nu$. The exchange is always AF as long as $\nu J_1 > 2J_2$.

For $\nu > 2/3$, the extra holes naturally resided at the centre moiré sites of the honeycomb to minimize the Coulomb repulsion (Fig. 1g). They increased the AF nearest-neighbour links, but did not increase the FM second-neighbour links in the dilute limit. This simple picture implies that filling above 2/3 drove the system ‘more AF’. For $\nu < 2/3$, vacancies were introduced into the honeycomb lattice (Fig. 1h). Introducing a vacancy removes twice as many FM links as AF links. This is identical to the original lattice and hence cannot explain the stronger AF exchange observed in the experiment. However, if we further assume that a fraction of the vacancies ($0 < \nu < 1$) reside on a pair of nearest-neighbour sites, to introduce a pair of vacancies removes 12FM links but only 5 AF links (Fig. 1i), and hence drives the system more AF (detailed calculations in Methods). Note that we did not attempt to prove that vacancies were, indeed, introduced in pairs here, but a recent theoretical study on TMD moiré materials suggested that doping in pairs could cost less Coulomb repulsion energy under the right conditions33. Nevertheless, the spin model with only $J_1$ and $J_2$ adopted here is probably too simple to account for all the experimental data near $\nu > 2/3$.

In conclusion, we reveal the frustrated magnetic interactions in semiconductor moiré bilayers in a strong interaction limit through a strongly suppressed AF exchange in the Wigner–Mott insulator at $\nu = 2/3$. We did not observe any clear enhancement or suppression of the net exchange interaction at other commensurate fractional fillings between 1/3 and 2/3, for which charge-ordered states have been reported34,35. In general, the magnetic interaction in an extended Hubbard system at generic fillings is a complicated problem. Future studies are required to better understand the mechanism of magnetic exchange in the presence of charge order, nematicity36 and itinerant electrons.

Online content
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Dual-gate devices of angle-aligned WSe\(_2\)/WS\(_2\) heterobilayers were fabricated using a dry-transfer method reported in the literature\(^6\). In short, atomically thin flakes of each constituent were exfoliated from bulk crystals onto SiO\(_2\)–Si substrates, and identified by their RC under an optical microscope. The thickness was determined more accurately by atomic force microscopy. A polymer stamp was used to pick up all of the flakes sequentially. The finished stack was released onto SiO\(_2\)–Si substrates with patterned Au electrodes. Second harmonic generation was employed to determine the crystal axes of the WSe\(_2\) and WS\(_2\) monolayers before stacking\(^6\). The main text reports results from a 60°-aligned sample. The results were reproduced in a second 60°-aligned sample (Extended Data Fig. 8). Similar results were also observed in a 0°-aligned sample (Extended Data Fig. 8). For the second type of device, an extra few-layer TaSe\(_2\) flake was introduced into the device (Fig. 1a). It was separated from the WSe\(_2\)/WS\(_2\) heterobilayer by a bilayer h-BN spacer. The TaSe\(_2\) flakes were exfoliated inside a nitrogen-filled glove box to avoid sample degradation. We followed the procedure described in Tang et al.\(^4\) to calibrate the doping density and filling factor of the moiré bilayer. Specifically, we measured the h-BN gate dielectric thickness by atomic force microscopy and used the known dielectric constant of h-BN (-3) to calculate the hole doping density using a parallel plate capacitor model.

**Magnetooptical measurements**

Details of the MCD measurements were reported in Tang et al.\(^6\) and Li et al.\(^5\). In short, the devices were mounted in a closed-cycle cryostat with a superconducting magnet (attoDRY 2100). The magnetic field was applied perpendicular to the sample plane. White light from a tungsten halogen lamp was collimated and focused onto the devices (with the power less than 1 nW). The reflected light was detected by a liquid nitrogen-cooled charge-coupled device attached to a grating spectrometer. The polarization of the white light was controlled by a combination of a polarizer and a broadband quarter-wave plate. The WSe\(_2\)/WS\(_2\) heterobilayers were grounded during the optical measurements. In device 1, both the top and bottom gate voltages were controlled independently by two Keithley source meters. In device 2, the TaSe\(_2\) flake was grounded, and only the top gate voltage was applied to tune the filling factor. In contrast to that of Wang et al.\(^5\), the excitation intensity was kept substantially lower to minimize the perturbation on the correlated states. We also analysed the MCD spectrum rather than a single-wavelength MCD signal so that a sign change in the MCD signal as well as a spectral shift in the exciton resonance with doping can be properly accounted for.

The MCD spectrum is defined as
\[
\text{MCD} = \frac{I_+ - I_-}{I_+ + I_-},
\]
where \(I_+\) and \(I_-\) are, respectively, the spectrum of the reflected left- and right-handed incident light from the sample. The integrated MCD signal is computed as
\[
\int_\varepsilon \frac{\text{MCD}}{\varepsilon} d\varepsilon,
\]
where \(\varepsilon\) is the photon energy and the limits of integration are given by the dashed lines in Fig. 2a. We focus on the MCD response in this study rather than the exciton Zeeman splitting as in an earlier study\(^5\). Although the two methods are generally in good agreement with each other, the MCD gives a better signal-to-noise ratio. It directly reflects the optical Hall response and the spin-valley polarization of the system, and is therefore more straightforward to interpret\(^6\). In particular, the MCD is easier to analyse than the exciton Zeeman splitting in the presence of multiple exciton peaks at \(v = 2/3\) which partially overlap in energy (Extended Data Fig. 1). We focused on hole doping, for which a strong MCD is observed (Extended Data Fig. 9). To isolate the local moment response, we also subtracted a small, constant slope at high magnetic fields from the magnetic-field-dependent MCD; this slope arises from the bare exciton Zeeman effect, which is nearly temperature independent.

**Determination of the saturation magnetic field**

We used the Brillouin function,
\[
\frac{M}{M_c} = \tanh \left( \frac{B}{B_c} \right),
\]
to describe the magnetic-field dependence of the integrated MCD signal or the sample magnetization \(M\). Here \(B\) denotes the applied magnetic field and \(M_c\) and \(B_c\) are the saturation magnetization and magnetic field, respectively. An example is shown in Extended Data Fig. 10 for \(v = 0.96\) and \(T = 1.7\) K. The filling dependence of \(B_c\) is shown in Fig. 3c.

**Spin model**

At 2/3 filling, the charges form a Wigner–Mott crystal with the shape of a honeycomb lattice\(^6\), which is 2/3 of the lattice sites of the original triangular moiré lattice (Fig. 1). We considered filling factors of around 2/3. Although in our experiment the moiré lattice was filled with holes for the entire doping range, we refer to fillings slightly below and above 2/3 as ‘electron doping’ and ‘hole doping’, respectively. One key observation we made is that, when immediately doped away from a 2/3 filling, the system remained insulating at low temperatures, based on capacitance measurements\(^6\). This implies that the doped charges in the flat band system are always immobile, probably due to the presence of long-wavelength moiré disorder and/or a strong correlation. The spin physics of the system near a 2/3 filling was therefore captured by a (doped) spin model on the honeycomb lattice with each immobile hole carrying a spin-1/2 degree of freedom. The effect of moiré disorder was not treated explicitly in our model; it only enters in localizing the doped holes/electrons.

The spin model we adopted has AF nearest-neighbour (nn) coupling \(J_1\) and FM next-nearest-neighbour (nnn) coupling \(J_2\) (in our convention, both \(J_1\) and \(J_2\) are taken to be positive). The Hamiltonian of the spin model takes the following form:
\[
H = \sum_{\text{nn bonds}} J_1 \mathbf{S}_i \cdot \mathbf{S}_j - \sum_{\text{nn bonds}} J_2 \mathbf{S}_i \cdot \mathbf{S}_j + \sum_{\text{sites}} \mathbf{h} \cdot \mathbf{S}_i, \tag{1}
\]
where \(\mathbf{S}_i\) is the spin on site \(i\) and \(\mathbf{h}\) is the external magnetic field. As the main physics we focused on is observed at a finite temperature, we used a high-temperature expansion to analyse this model. Note that the qualitative results of our high-temperature expansion calculation were insensitive to whether the spin was treated as a classical three-component vector or a quantum spin-1/2 variable. The following calculation was performed using the latter treatment.

With hole doping away from 2/3 filling, namely, \(v > 2/3\), there were two main qualitative findings in the experiments: (1) The CW temperature \(\theta\) remained negative, but its magnitude increased under hole doping (Fig. 3b); this implies that, speaking overall, the system is AF, and it becomes even more AF under hole doping. (2) The spin susceptibility decreased with hole doping at a low temperature (Fig. 3a), but increased with hole doping at a high temperature (Extended Data Fig. 5).

Figure 1g depicts the set-up for hole doping away from 2/3 filling. The extra hole doped to the system naturally resides at the centre moiré sites of the honeycomb. A doped hole increases the nearest-neighbour AF links in the system (labelled as dashed lines in Figure 1g), but does not increase the FM next-nearest-neighbour links. This simple picture implies that hole doping drives the system ‘more AF’, and increases \(|\theta|\). At filling \(v = 2 (1 + p)/3\), the high temperature expansion leads to the following result for \(\theta\), expanded to the leading order of doping \(p > 0\):
\[
\theta = -(J_1 - 2J_2) + (3J_1 + 2J_2)p + O(p^2). \tag{2}
\]
As long as \(J_1 > 2J_2\), \(\theta\) is always negative, which signifies AF behaviour. Also, \(|\theta|\) increases with hole doping \(p\) at the leading order expansion of \(p\), which is consistent with the experimental data. The spin susceptibility \(\chi\) reads:
\[
\chi = \frac{n (1 + p)}{3(T - \theta)} \sim \frac{n (1 + p)}{3T} + \frac{n (-J_1 + 2J_2 - 4J_2p)}{3T^2} + O(p^2), \tag{3}
\]
where \(n\) is the number of holes of the honeycomb at \(v = 2/3\).
As an example, let us choose \(J_0 = 2.5J_1 = 2.5A\). We then have \(\theta = -0.5A\). One immediately notices that when \(T < 10A\), \(\chi\) decreases with doping \(p\), and when \(T > 10A\), \(\chi\) increases with \(p\). All these are consistent with the experimental observations.

With electron doping away from 2/3 filling, namely, \(\nu < 2/3\), there are also two main qualitative findings in the experiments: (1) The CW temperature \(T\) remains negative, and the magnitude of \(|\theta|\) increases under hole doping (Fig. 3b); this implies that the system still becomes ‘more AF’ under hole doping. (2) The spin susceptibility always decreases with doping within the temperature range in the experiment (Fig. 3a).

Electron doping was modelled as doping vacancies in our spin model. The theoretical facts are somewhat counterintuitive, as doping vacancies normally do not increase \(|\theta|\). However, in the following discussion, we show that with a simple extra assumption of the configurations of doped vacancies, our model can explain the two experimental facts listed above. The set-up for vacancies doping is depicted in Fig. 1h: we assume that the doped vacancies always reside on a pair of nearest-neighbour sites. Although we do not prove here that the vacancies do form pairs, we note that a similar pair-wise doping in TMD moiré superlattices does form pairs, we note that a similar pair-wise doping in TMD moiré materials in the flat band limit was discussed by Slagle and Fu11 as a way to minimize the Coulomb repulsion energy. At filling \(\nu = 2(1 - p)/3\), the high temperature expansion calculation gives the following results:

\[
\theta = -\left[\frac{2\rho}{3} - 2J_1\right] + O(p^2) \tag{4}
\]

As long as \(3J_2 > J_1 > 2J_0\), \(\theta\) is always negative, and \(|\theta|\) increases with \(p\), again consistent with the experimental data. The reason \(|\theta|\) increases with \(p\) is that, on the original lattice model, there are twice as many second neighbour links (FM interactions) as nearest-neighbour links (AF interactions). As the vacancies are ‘paired up’, doping a pair of vacancies removes 12 FM links, but removes only 5 AF links. Hence, doping vacancies can indeed drive the system more AF, and increase \(|\theta|\). The spin susceptibility \(\chi\) is calculated as:

\[
\chi = \frac{n(1-p)}{3(T-\theta)} \sim \frac{n(1-p)}{3T} + \frac{n\left[-J_1 + 2J_2 + \left(\frac{5}{3} - 4J_2\right)p\right]}{3T^2} + O(p^2) \tag{5}
\]

Again, as an example, let us choose \(J_0 = 2.5J_1 = 2.5A\). Then we obtain \(\theta = -0.5A\). We found that as long as \(T > A/6\), \(\chi\) always decreases with \(p\), which is consistent with experimental findings.

Data availability
All data that support the findings of this study are available from the corresponding author upon reasonable request. Source data are provided with this paper.

Code availability
All codes to analyse the reflectance spectrum are available from the corresponding author upon reasonable request.

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Author contributions
Y.T. performed the optical measurements and the analysis. Y.T. fabricated the devices with assistance from L.L. and Y.X. S.L. and J.H. grew the bulk TMD crystals. K.W. and T.T. grew the bulk h-BN crystals. K.S., C.-M.J. and C.X. conceived and developed the spin model. K.F.M., J.S. and Y.T. designed the scientific objectives and oversaw the project. K.F.M., J.S., K.S., C.-M.J. and C.X. co-wrote the manuscript. All the authors discussed the results and commented on the manuscript.

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Extended Data Fig. 1 | Helicity-resolved reflectance contrast spectra at varying magnetic fields. a–c, Results for filling factor 0.88 (a), 0.67 (b) and 0.44 (c). The black and red lines denote the left (LCP) and right circularly polarized (RCP) excitations, respectively. The spectra are vertically displaced for clarity.
Extended Data Fig. 2 | Analysis of MCD spectra. a, MCD spectrum as a function of magnetic field at $\nu = 2/3$ and 1.7 K. The vertical dashed lines denote the three energy windows (M1, M2 and M3) for the MCD analysis. b, Magnetic-field dependent MCD integrated in window M1, M2 and M3 and normalized to 1 upon saturation at large fields. The solid lines are fits to the Brillouin function. The saturation field $B_s$ obtained from the fit (shown in the key) is insensitive to the choice of the spectral window.
Extended Data Fig. 3 | Magnetic response of device 1 at varying temperatures for $\nu = 0.33$ (a), 0.57 (b), 0.8 (c) and 0.96 (d). The color of the symbols varying from darkest to lightest denotes the temperature from 1.7 to 29.9 K.
Extended Data Fig. 4 | Magnetic response of device 1 at 1.7 K. a, Integrated MCD as a function of magnetic field and filling factor. b, Linecuts of a at selected filling factors. The curves are displaced vertically for clarity.
Extended Data Fig. 5 | The magnetic susceptibility as a function of fillings at high temperature (device 1). The dashed straight line is just an eye guidance to the linear filling dependence.
Extended Data Fig. 6 | Optical characterization of device 2 at 1.7 K. **a**, Filling-dependent reflectance contrast (RC) spectrum. Enhanced RC near the fundamental moiré exciton resonance of WSe₂ is observed only at integer filling factors. The absence of the RC enhancement at $\nu = 1/3$ and $2/3$ supports the absence of these Wigner-Mott insulator states. **b**, Linecut of a at 1.673 eV.
Extended Data Fig. 7 | Magneto-optical measurements (device 2). a, MCD spectrum as a function of magnetic field at \( \nu = 2/3 \) and \( T = 1.7 \) K. The spectrum is integrated over a narrow window around the fundamental moiré exciton resonance (within the two vertical lines) to represent the sample magnetization.

b, Filling dependence of the magnetic susceptibility, \( \chi \), at varying temperatures. The susceptibility is extracted from the linear slope of the magnetic-field dependence of integrated MCD at zero field. The solid lines are guides to the eye.

c, Curie-Weiss analysis (solid lines) of the inverse susceptibility (symbols) as a function of temperature. The error bars are propagated from the uncertainty of \( \chi \). The CW temperatures corresponding to the best fits (solid lines) are included. Data for different filling factors are vertically displaced for clarity. The horizontal dashed lines mark where \( 1/\chi \) is zero. Linear dependences are observed at temperatures smaller than the CW temperatures; no sign of magnetic ordering is observed (that is no kink in the temperature dependence). The absence of magnetic ordering at temperatures smaller than the mean-field CW temperature signifies the importance of magnetic frustrations in the system.
Extended Data Fig. 8 | Results from additional devices. a, Comparison of the filling factor dependent saturation magnetic field of a 0- and 60-degree-aligned sample at 3.5 K. b–d, Results from another 60-degree-aligned sample, including the filling factor dependence of the magnetic susceptibility at 1.7 K (b), the saturation magnetic field at 1.7 K (c), and the Curie-Weiss temperature (d). The dashed lines denote 2/3 filling, where suppression of AF interactions is observed in all samples. A weaker suppression is observed in the 0-degree-aligned sample, which could be caused by the different moiré structure. The error bars correspond to the uncertainties of the Brillouin-function fit (a and c), the linear fit to the field dependence of the MCD near zero field (b), and the CW analysis (d).
Extended Data Fig. 9 | Helicity-resolved reflectance contrast spectrum as a function of doping for both electron and hole doping. a, b, d, e Gate voltage dependent left- and right-handed reflectance contrast spectrum (LCP and RCP) for the WSe$_2$ (a, b) and WS$_2$ (d, e) layer. The magnetic field is at 2 T. Positive and negative fillings correspond to hole and electron doping, respectively. The dashed lines label the integer filling factors. c, f Extracted gate voltage dependent MCD spectrum for the WSe$_2$ (c) and WS$_2$ (f) layer. Strong MCD response is observed only in the WSe$_2$ exciton resonance and only for the hole doping case. The results can be understood based on the type II band alignment in WSe$_2$/WS$_2$ bilayers; electron (hole) doping fills the WS$_2$ (WSe$_2$) conduction (valence) band. Strong MCD in the WSe$_2$ layer is expected for hole doping because states near the Fermi level are directly involved in the optical transitions. On the other hand, states near the Fermi level are not involved in the optical transitions for electron doping in W-based TMDs. The MCD response is weak in all cases for electron doping.
Extended Data Fig. 10 | Determination of the saturation magnetic field (device 1). The red curve is a Brillouin-function fit to the integrated MCD as a function of magnetic field (at 0.96 filling and 1.7 K). The corresponding saturation magnetic field is 0.63 ± 0.01 T.