

Twisted bilayer graphene reveals its flat bands under spin pumpingSonia Haddad^{1,2,3,*}, Takeo Kato², Jihang Zhu,³ and Lassaad Mandhour¹¹Laboratoire de Physique de la Matière Condensée, Faculté des Sciences de Tunis, Université Tunis El Manar, Campus Universitaire 1060 Tunis, Tunisia²Institute for Solid State Physics, University of Tokyo, Kashiwa, Chiba 277-8581, Japan³Max Planck Institute for the Physics of Complex Systems, Nöthnitzer Strasse 38, Dresden 01187, Germany

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The salient property of the electronic band structure of twisted bilayer graphene (TBG), at the so-called magic angle (MA), is the emergence of flat bands around the charge neutrality point. These bands are associated with the observed superconducting phases and the correlated insulating states. Scanning tunneling microscopy combined with angle resolved photoemission spectroscopy are usually used to visualize the flatness of the band structure of TBG at the MA. Here, we theoretically argue that spin pumping (SP) provides a direct probe of the flat bands of TBG and an accurate determination of the MA. We consider a junction separating a ferromagnetic insulator and a heterostructure of TBG adjacent to a monolayer of a transition metal dichalcogenide. We show that the Gilbert damping of the ferromagnetic resonance experiment, through this junction, depends on the twist angle of TBG, and exhibits a sharp drop at the MA. We discuss the experimental realization of our results which open the way to a twist switchable spintronics in twisted van der Waals heterostructures.

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Introduction. Stacking two graphene layers with a relative twist angle θ results in a moiré superstructure which is found to host, in the vicinity of the so-called magic angle (MA) $\theta_M \sim 1.1^\circ$, unconventional superconductivity and strongly correlated insulating states [1–3]. There is a general consensus that such strong electronic correlations originate from the moiré flat bands emerging at the MA around the charge neutrality point [4–11]. The tantalizing signature of the flat bands have been experimentally demonstrated by probing the corresponding peaks of the density of states using transport [1–3,12,13], electronic compressibility measurements [14,15], scanning tunneling microscopy (STM), and spectroscopy (STS) [16–23]. The direct evidence of these flat bands has been reported by angle resolved photoemission spectroscopy (ARPES) measurements combined to different imaging techniques [24–26]. However, spectroscopic measurements on magic-angle TBG raise many technical challenges related to the need of an accurate control of the twist angle, and the necessity to have nonencapsulated samples which can degrade in air [25].

Here we propose a noninvasive method to probe the flat bands of TBG and accurately determine the MA. This method is based on spin pumping (SP) induced by ferromagnetic resonance (FMR) [27–30], where the increase in the FMR linewidth, given by the Gilbert damping (GD) coefficient, provides insight into the spin excitations of the nonmagnetic (NM) material adjacent to the ferromagnet [31–33]. SP is expected to be efficient if the NM has high spin-orbit coupling (SOC) strength [34]. In our work, we consider spin injection from a ferromagnetic insulator (FI) into a TBG aligned on a

monolayer of transition metal dichalcogenides (TMD) which are considered as good substrate candidates to induce relatively strong SOC in graphene and TBG [35–58].

We theoretically study a planar junction of a FI and a TBG adjacent to WSe₂ (TBG/WSe₂) as depicted in Fig. 1. We consider the case where a microwave of a frequency Ω is applied to this junction, and focus on the twist angle dependence of the FMR linewidth [59].

Continuum model. In TBG with a twist angle θ , the Hamiltonian $h_l(\mathbf{k})$ of a graphene layer l ($l = 1, 2$), rotated at an angle θ_l , is $h_l(\mathbf{k}) = e^{i\frac{\theta_l}{2}\sigma_z} h_l^{(0)}(\mathbf{k}) e^{-i\frac{\theta_l}{2}\sigma_z}$, where $\theta_2 = -\theta_1 = \frac{\theta}{2}$, and $h_l^{(0)}(\mathbf{k})$ is the unrotated monolayer Hamiltonian. In the continuum limit, $h_1^{(0)}(\mathbf{k})$ reduces to $h_1^{(0)}(\mathbf{k}) = -\hbar v_F \mathbf{k} \cdot \sigma^*$, where v_F is the Fermi velocity, $\sigma^* = (\xi\sigma_x, \sigma_y)$ and σ_i ($i = x, y, z$) are the sublattice-Pauli matrices, and ξ is the valley index. We assume that the SOC is only induced in the graphene layer adjacent to the TMD layer, since the SOC arises from overlaps between atomic orbitals [50]. This assumption is consistent with recent studies on bilayer graphene and TBG aligned on TMD layers [50,56,60,61]. Layer (2), in contact with the WSe₂ monolayer, is then described by the Hamiltonian $h_2^{(0)}(\mathbf{k}) = h_1^{(0)}(\mathbf{k}) + h_{\text{SOC}} + \frac{m}{2}\sigma_z$ [56], where h_{SOC} is given by

$$h_{\text{SOC}} = \frac{\lambda_I}{2} \xi s_z + \frac{\lambda_R}{2} (\xi \sigma_x s_y - \sigma_y s_x) + \frac{\lambda_{\text{KM}}}{2} \xi \sigma_z s_z, \quad (1)$$

s_i ($i = x, y, z$) are the spin-Pauli matrices; λ_I , λ_R , and λ_{KM} correspond, respectively, to the Ising, Rashba, and Kane-Mele SOC parameters [56]. The variation ranges of these parameters are $\lambda_I \sim 1 - 5$ meV, $\lambda_R \sim 1 - 15$ meV, while λ_{KM} is expected to be small [37,44,50,51,62–64]. The last term in $h_2^{(0)}(\mathbf{k})$ is due to the inversion symmetry breaking induced by the TMD layer. Hereafter, we neglect this term regarding

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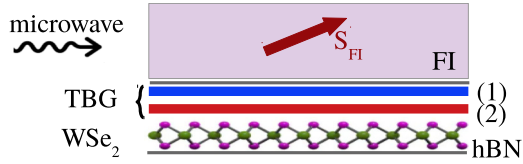


FIG. 1. Schematic representation of the junction between a ferromagnetic insulator (FI) and a heterostructure of TBG adjacent to a monolayer of WSe₂. The labels (1) and (2) denote the graphene layers of TBG represented by the red and the blue lines. The red arrow indicates the spin orientation of the FI characterized by an average spin $\langle \mathbf{S}_{FI} \rangle = (S_0, 0, 0)$, written in the coordinate frame of the FI magnetization. The gray lines represent the boron-nitride (hBN) layers encapsulating the TBG/WSe₂ heterostructure.

the small value of m compared to the SOC parameters [56]. As in the case of TBG [65], the low-energy Hamiltonian of TBG/WSe₂ reduces, at the valley ξ , to

$$H_{\xi, \text{SOC}}(\mathbf{k}) = \begin{pmatrix} h_1(\mathbf{k}) & T_1 & T_2 & T_3 \\ T_1^\dagger & h_{2,1}(\mathbf{k}) & 0 & 0 \\ T_2^\dagger & 0 & h_{2,2}(\mathbf{k}) & 0 \\ T_3^\dagger & 0 & 0 & h_{2,3}(\mathbf{k}) \end{pmatrix}. \quad (2)$$

$H_{\xi, \text{SOC}}(\mathbf{k})$ is written in the basis $\Psi = (\psi_0(\mathbf{k}), \psi_1(\mathbf{k}), \psi_2(\mathbf{k}), \psi_3(\mathbf{k}))$ constructed on the four-component spin-sublattice spinor $\psi_j(\mathbf{k})$ and $\psi_j(\mathbf{k})$, ($j = 1, 2, 3$) corresponding, respectively, to layer (1) and layer (2) (see Secs. I and II of the Supplemental Material [66] and Refs. [8,56,65,67–70]). The momentum \mathbf{k} is measured relatively to the Dirac point $\mathbf{K}_{1\xi}$ of layer (1). In Eq. (2), T_j are the spin-independent interlayer coupling matrices, $h_{2,j}(\mathbf{k}) = h_2(\mathbf{k} + \mathbf{q}_{j\xi})$, ($j = 1, 2, 3$) where $\mathbf{q}_{j\xi}$ are the vectors connecting $\mathbf{K}_{1\xi}$ to its three neighboring Dirac points $\mathbf{K}_{2\xi}$ of layer (2) in the moiré Brillouin zone (mBZ) [65], and are given by $\mathbf{q}_{1\xi} = \mathbf{K}_{1\xi} - \mathbf{K}_{2\xi}$, $\mathbf{q}_{2\xi} = \mathbf{q}_{1\xi} + \xi \mathbf{G}_1^M$, $\mathbf{q}_{3\xi} = \mathbf{q}_{1\xi} + \xi(\mathbf{G}_1^M + \mathbf{G}_2^M)$, where $(\mathbf{G}_1^M, \mathbf{G}_2^M)$ is the mBZ basis (see Sec. I of the Supplemental Material [66]). In the unrelaxed TBG, and choosing sublattice A as the origin of the unit cell in each layer, the T_j matrices take the form [8] $T_1 = w(\mathbb{I}_\sigma + \sigma_x)$, $T_2 = w(\mathbb{I}_\sigma - \frac{1}{2}\sigma_x + \xi \frac{\sqrt{3}}{2}\sigma_y)$, and $T_3 = w(\mathbb{I}_\sigma - \frac{1}{2}\sigma_x - \xi \frac{\sqrt{3}}{2}\sigma_y)$ [66], where $w \sim 110$ meV [71] is the interlayer tunneling amplitude and \mathbb{I}_σ is the identity matrix acting on the sublattice indices.

Using the perturbative approach of Ref. [65], we derive, from Eq. (2), the effective low-energy Hamiltonian $H_{\xi, \text{SOC}}^{(1)}$ of TBG/WSe₂ (see Sec. II of the Supplemental Material [66]). To the leading order in \mathbf{k} , $H_{\xi, \text{SOC}}^{(1)}(\mathbf{k})$ reads as [66]

$$H_{\xi, \text{SOC}}^{(1)}(\mathbf{k}) = \frac{\langle \Psi | H_{\xi, \text{SOC}} | \Psi \rangle}{\langle \Psi | \Psi \rangle} = \psi_0^\dagger [h_{\text{eff}}(\mathbf{k}) + h_{\text{eff}}^{\text{SOC}}] \psi_0, \quad (3)$$

$$h_{\text{eff}}(\mathbf{k}) = -\frac{\hbar v_F}{\langle \Psi | \Psi \rangle} \left\{ k_x \left[(1 - 3\alpha^2) \xi \sigma_x \mathbb{I}_s - \frac{3\alpha^2}{\hbar v_F q_0} (\xi \lambda_I \sigma_y s_z + \lambda_R (\xi \sigma_y s_y - \sigma_x s_x)) \right] \right. \\ \left. + k_y \left[(1 - 3\alpha^2) \sigma_y \mathbb{I}_s - \frac{3\alpha^2}{\hbar v_F q_0} (-\lambda_I \sigma_x s_z + \lambda_R (\sigma_x s_y + \xi \sigma_y s_x)) \right] \right\}, \quad (4)$$

$$h_{\text{eff}}^{\text{SOC}} = \frac{3\alpha^2}{\langle \Psi | \Psi \rangle} \left[\xi \lambda_I s_z \mathbb{I}_\sigma + \frac{\lambda_R}{2} (s_x \sigma_y - \xi s_y \sigma_x) \right], \quad (5)$$

where $\langle \Psi | \Psi \rangle \sim 1 + 6\alpha^2$, $\alpha = \frac{w}{\hbar v_F q_0}$, $q_0 = |\mathbf{q}_{j\xi}| = \frac{4\pi}{3a}\theta$, a is the graphene lattice constant, and σ_i , ($i = x, y, z$) act now on the band indices $\sigma = \pm$ of the eigenenergies of $H_{\xi, \text{SOC}}^{(1)}$, denoted $E_{\sigma, \pm}$, and given to the leading orders in \mathbf{k} and $\frac{\lambda_{I,R}}{\hbar v_F q_0}$ by

$$E(\mathbf{k})_{\sigma, \pm} = \frac{\sigma}{\langle \Psi | \Psi \rangle} \sqrt{f_1(\mathbf{k}) \pm 6\alpha^2 \sqrt{f_2(\mathbf{k})}}, \quad (6)$$

$$f_1(\mathbf{k}) = (\hbar v_F)^2 (1 - 3\alpha^2)^2 |\mathbf{k}|^2 + \frac{9}{2} \alpha^4 (2\lambda_I^2 + \lambda_R^2)$$

$$f_2(\mathbf{k}) = (\hbar v_F)^2 (1 - 3\alpha^2)^2 |\mathbf{k}|^2 (\lambda_I^2 + \frac{1}{4} \lambda_R^2) + \frac{9}{16} \alpha^4 \lambda_R^4.$$

Equation (5) shows that the SOC parameters λ_I and λ_R are renormalized by the moiré structure of TBG to

$$\tilde{\lambda}_I \sim \frac{6\alpha^2}{1 + 6\alpha^2} \lambda_I, \quad \tilde{\lambda}_R \sim \frac{3\alpha^2}{1 + 6\alpha^2} \lambda_R, \quad (7)$$

which increase by decreasing the twist angle. The expression of $H_{\xi, \text{SOC}}^{(1)}$ [Eq. (3)] can be taken as a starting point to unveil the

role of SOC in the emergence of the stable superconducting phase observed, at $\theta \sim 0.8^\circ$, in TBG adjacent to WSe₂ [56]. To probe the validity of the effective Hamiltonian $H_{\xi, \text{SOC}}^{(1)}$ [Eq. (3)], we compared the corresponding eigenenergies with the numerical band structure obtained within the continuum model and taking into account 148 bands per valley and spin projection (see Sec. II of the Supplemental Material [66]). The results show that $H_{\xi, \text{SOC}}^{(1)}$ describes correctly the band structure of TBG/WSe₂ down to a twist angle $\theta \sim 0.7^\circ$. At smaller angles, the effective Fermi velocities of $H_{\xi, \text{SOC}}^{(1)}$ are overestimated. Such a discrepancy is expected since the lattice relaxation effect is important at small angles [56]. It is worth noting that, for the sake of simplicity, we did not consider a relaxed TBG since we are interested in the SP around the MA.

Gilbert damping. In the absence of a junction, the magnon Green function of the FI is defined as [72–78] $G_0(\mathbf{q}_m, i\omega_n) = \frac{2S_0/\hbar}{i\omega_n - \omega_{\mathbf{q}_m} - \alpha_G |\omega_n|}$, where $\omega_n = 2\pi n/\hbar\beta$ are the Matsubara frequencies for bosons, S_0 is the amplitude of the average spin per site, and α_G is the GD strength. The term $-\alpha_G |\omega_n|$ describes the spin relaxation within the FI. In FMR experiments,

the microwave excitation induces a uniform spin precession, which limits the magnon self-energy to the processes with $\mathbf{q}_m = 0$ [79].

In the presence of the interfacial coupling, a correction, $\delta\alpha_G(\omega)$, to the GD term is induced by the adjacent heterostructure TBG/WSe₂. $\delta\alpha_G(\omega)$ can be expressed in terms of the self-energy $\Sigma_0^R(\omega) \equiv \Sigma_{\mathbf{q}_m=0}(i\omega_n \rightarrow \omega + i\delta)$, resulting from the interfacial exchange interactions, as [79]

$$\delta\alpha_G(\omega) \equiv -\frac{2S_0}{\hbar\omega} \text{Im} \Sigma_0^R(\omega). \quad (8)$$

For simplicity, we neglect the real part of $\Sigma_0^R(\omega)$ which simply shifts the FMR line and did not affect the linewidth, in which we are interested. The self-energy, in Eq. (8), includes the contributions of all the interfacial spin transfer processes and can be written as $\Sigma_0(i\omega_n) = \sum_{\mathbf{q}} \Sigma_0(\mathbf{q}, i\omega_n)$. Each process, described by the self-energy $\Sigma_0(\mathbf{q}, i\omega_n)$, is characterized by a momentum transfer \mathbf{q} and a matrix element $T_{\mathbf{q},\mathbf{q}_m=0} \equiv T_{\mathbf{q},0}$.

In the second order perturbation, with respect to the interfacial exchange interaction $T_{\mathbf{q},0}$, the self-energy $\Sigma_0(\mathbf{q}, i\omega_n)$ is written as [78]

$$\begin{aligned} \Sigma_0(\mathbf{q}, i\omega_n) &= \frac{|T_{\mathbf{q},0}|^2}{4\beta} \sum_{\mathbf{k}, i\omega_m} \text{Tr}[\sigma_s^{x',-} \hat{g}(\mathbf{k}, \omega_m) \\ &\times \sigma_s^{x',+} \hat{g}(\mathbf{k} + \mathbf{q}, i\omega_m + i\omega_n)]. \end{aligned} \quad (9)$$

$\sigma_s^{x',\pm}$ are the electronic spin ladder operators written in the coordinate system (x', y', z') of the FI magnetization characterized by an average spin $\langle \mathbf{S}_{FI} \rangle = (S_0, 0, 0)$. $\hat{g}(\mathbf{k}, i\omega_n)$ is the electronic Matsubara Green function given by $\hat{g}(\mathbf{k}, i\omega_n) = [i\omega_n \mathbb{I} - H_{\text{SOC}}^{(1)}(\mathbf{k})]^{-1}$, where $\omega_n = (2n+1)\pi/\hbar\beta$ are the fermionic Matsubara frequencies. In the basis of the spin-band four-component spinor $\Psi = (\psi_{+, \uparrow}, \psi_{+, \downarrow}, \psi_{-, \uparrow}, \psi_{-, \downarrow})$ [56]; $\hat{g}(\mathbf{k}, i\omega_n)$ reads as $\hat{g}(\mathbf{k}, i\omega_n) = \hat{g}_0(\mathbf{k}, i\omega_n) \mathbb{I}_s + \hat{\mathbf{g}}(\mathbf{k}, i\omega_n) \cdot \mathbf{s}$, where $\mathbf{s} = (s_x, s_y, s_z)$ are the spin-Pauli matrices; $\hat{\mathbf{g}} = (\hat{g}_x, \hat{g}_y, \hat{g}_z, \hat{g}_0)$, and \hat{g}_i ($i = x, y, z$) are expressed, to the leading order in the SOC, as a function of the band-Pauli matrices σ_i (see Sec. III of the Supplemental Material [66]).

Since the ferromagnetic peak, given by $\text{Im}G_0^R$ is sharp enough, namely $\alpha_G + \delta\alpha_G \ll 1$, one can replace the resonance frequency $\omega_{\mathbf{q}_m=0}$ by the FMR frequency Ω . The GD correction can then be expressed as [66,78]

$$\delta\alpha_G(\Omega) = -\frac{2S_0}{\hbar\Omega} \text{Im} \Sigma_0^R(\Omega). \quad (10)$$

In general, the interfacial spin transfer includes clean and dirty processes. The former (latter) take place with conserved (non-conserved) electron momentum, which turns out to take $\mathbf{q} = \mathbf{0}$ ($\mathbf{q} \neq \mathbf{0}$) in Eq. (9) [79]. We first consider a clean interface, for which an analytical expression of the GD correction [Eq. (10)] can be derived (see Sec. IV of the Supplemental Material [66] and reference [78,79]). The case of a dirty junction is discussed in the next section.

Carrying out the summation over ω_m in Eq. (9), we obtain the analytical expression of the interfacial self-energy (see Sec. IV of the Supplemental Material [66]). The sum over the electronic states $\mathbf{k} = (k, \varphi_{\mathbf{k}})$ runs over the states included within a cutoff, $k_c \sim q_0/2$, on the momentum amplitude k , where the low-energy Hamiltonian [Eq. (3)] is expected to hold (see Sec. IV of the Supplemental Material [66]).

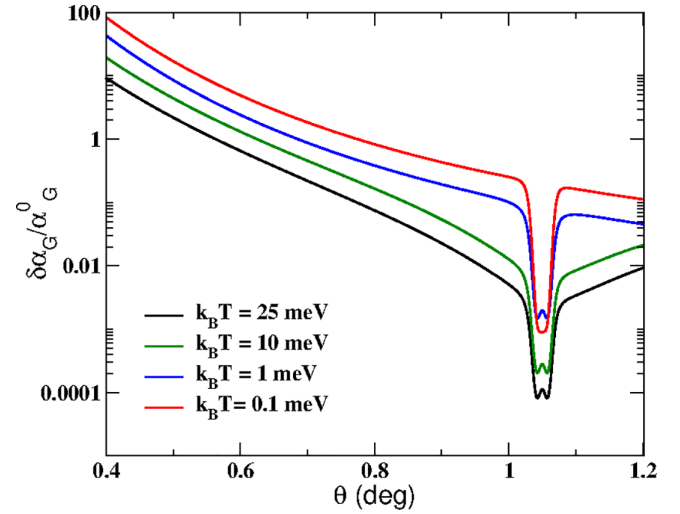


FIG. 2. Normalized GD, $\delta\alpha_G/\alpha_G^0$ Eq. (11), as a function of the twist angle θ at different temperature ranges. Calculations are done for $\lambda_l = 3$ meV, $\lambda_r = 4$ meV, $\mu = 0$, and for a FMR energy $\hbar\Omega = 0.06$ meV.

In the Following, we discuss the behavior of the normalized GD coefficient

$$\delta\alpha_G/\alpha_G^0 = \left(\frac{\lambda}{\hbar\Omega} \right)^2 \tilde{\Sigma}(\mathbf{q} = \mathbf{0}, \Omega), \quad (11)$$

where $\tilde{\Sigma}$ is a dimensionless function depending on the twist angle θ , temperature T , the chemical potential μ , and the orientation of the FI magnetization; $\alpha_G^0 = 2S_0(\frac{|T_0|}{\lambda})^2$ and $\lambda = \frac{\lambda_l + \lambda_r}{2}$ is the average SOC (for details, see Sec. IV of the Supplemental Material [66] and Ref. [80]).

Discussion. In Fig. 2, we plot $\delta\alpha_G/\alpha_G^0$ [Eq. (11)] as a function of the twist angle θ , for the undoped TBG, at different temperatures and for a fixed FMR energy $\hbar\Omega = 0.06$ meV which corresponds to the yttrium iron garnet. The SOC parameters are $\lambda_l = 3$ meV and $\lambda_r = 4$ meV, as in Ref. [56].

Figure 2 shows that regardless of the temperature range, $\delta\alpha_G$ increases by decreasing θ but drops sharply at the MA, where it exhibits a relatively small peak which is smeared out at low temperature.

Putting aside its drop at the MA, the enhancement of $\delta\alpha_G$ by decreasing θ can be, in a first step, ascribed to the dependence of the self-energy [Eq. (9)] on the effective SOC, given by Eq. (7), which increase by decreasing θ . However, to understand the behavior of $\delta\alpha_G$ at the MA one needs to go back to the band structure, $E_{\sigma,\pm}(\mathbf{k})$ [Eq. (6)], of the continuum Hamiltonian of TBG/WSe₂, which is depicted in Fig. 3 at different twist angles. The arrows indicate the out-of-plane electronic spin projection $\langle s_z \rangle$ which we have numerically calculated for different twist angles in Sec. II of the Supplemental Material [66].

Away from the MA, the band dispersion gets larger as θ decreases and, in particular, the separation between bands with opposite $\langle s_z \rangle$, involved in the SP process, increases. This behavior is due to the angle dependence of the effective Fermi velocity v^* of TBG/WSe₂, which reduces in the first order in the SOC to that of TBG, namely (see Secs. I and II of the

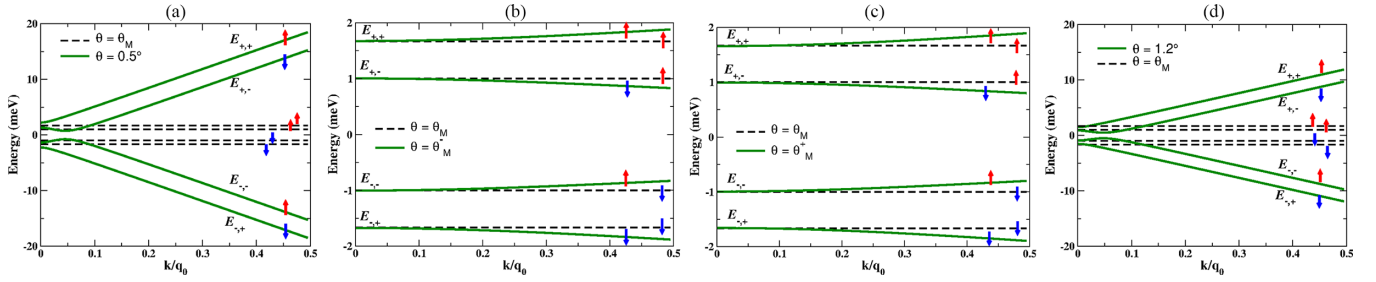


FIG. 3. Band structure of TBG/WSe₂ in the continuum limit [Eq. (6)] at $\theta = 0.5^\circ$ (a), $\theta = \theta_M^- = 1.043^\circ$ (b), $\theta = \theta_M^+ = 1.058^\circ$ (c), and $\theta = 1.2^\circ$ (d). The dashed lines represent the bands at the MA ($\theta_M = 1.05^\circ$). The red (blue) arrows correspond to the out-of-plane electronic spin projection $\langle s_z \rangle = +1$ ($\langle s_z \rangle = -1$) [56]. Calculations are done for $\lambda_I = 3$ meV and $\lambda_R = 4$ meV.

Supplemental Material [66])

$$v^* \sim v_F \frac{1 - 3\alpha^2}{1 + 6\alpha^2}. \quad (12)$$

The expression of the GD [Eq. (11)] includes transitions between bands with opposite $\langle s_z \rangle$ (see Sec. IV of the Supplemental Material [66]). These transitions depend on the statistical weight $\Delta f(E) = f(E_{\langle s_z \rangle}) - f(E_{-\langle s_z \rangle})$, where $f(x)$ is the Fermi-Dirac function and $E_{\langle s_z \rangle}$ is the energy band with a spin orientation $\langle s_z \rangle$.

In Fig. 4, we plot a pictorial representation of the band structure of the continuum model [Eq. (6)] and the Fermi-Dirac distribution $f(E)$ at a given temperature T . The band dispersion gets larger as θ moves away from the MA (Fig. 3) and the separation between the bands with opposite $\langle S_z \rangle$ increases. As a consequence, the corresponding statistical weight $\Delta f(E)$ is enhanced compared to the case around the MA. This behavior explains the drop of the GD at the MA.

Around the MA (θ_M^+ and θ_M^-), the statistical weight $\Delta f(E)$ is reduced compared to that at the MA since the bands $E_{+,-}$ and $E_{-,-}$ get closer (Fig. 3). This behavior gives rise to the small peak at the MA (Fig. 2), which disappears at low temperature ($k_B T < \lambda$), where bands around the MA have

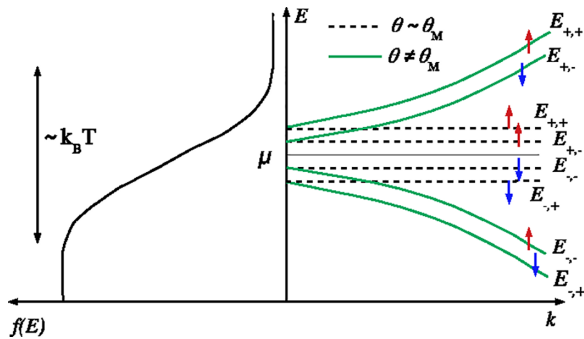


FIG. 4. Schematic representation of the band structure $E_{\sigma,\pm}$ [Eq. (6)] and the Fermi-Dirac distribution $f(E)$. The bands in dashed and green lines correspond, respectively, to the MA and to a twist angle θ far from the MA. The red (blue) arrows represent the projection of the out-of-plane spin projection $\langle S_z \rangle = +1$ ($\langle S_z \rangle = -1$). Around the MA, the bands are almost flat and the statistical weights $\Delta f(E)$, corresponding to the transitions between $E_{-,+} \rightarrow E_{+,+}$ and $E_{-,-} \rightarrow E_{+,-}$, are small compared to the case of a twist angle away from the MA, where the band dispersion is larger.

the same statistical weight $\Delta f(E) = 1$ (see Sec. IV of the Supplemental Material [66]). In this case, the GD is basically dependent on the effective Fermi velocity v^* [Eq. (12)] which vanishes at exactly the MA. Such dependence is responsible for the cancellation of several terms contributing to the self-energy [Eq. (9)], as they are proportional to v^* [Eq. (12)] (see Sec. IV of the Supplemental Material [66]). Let us now turn to the case of a dirty interface where the spin transfer should now also include the nonconserved momentum processes. The corresponding self-energy [Eq. (9)] can also be expressed in terms of the thermal weight $\Delta f(E)$ governing the interband transitions (see Sec. IV of the Supplemental Material [66]). Regarding the flatness of the bands, the dirty processes at the MA acquire, as in the clean limit, small thermal weights compared to the twist angles away from the MA, where the bands are dispersive. In the dirty limit, the Gilbert damping correction is then expected to drop at the MA as found in the case of a clean interface.

It comes out that the twist angle dependence of $\delta\alpha_G$ is a direct probe of the emergence of the flat bands in TBG. On the other hand, the temperature dependence of the fine structure around the MA provides an accurate measurement of the MA, with a precision below 0.005° (see Fig. S.4 of the Supplemental Material [66]). It also gives an estimation of the SOC induced in TBG adjacent to a monolayer of TMD.

It is worth stressing that in our model we did not take into account the electron-electron interactions which significantly distort the electronic band structure of TBG [80–83]. Near the MA, the dominant electron-electron interaction is found to be the Coulomb interaction with an amplitude estimated to be 10–15 meV [82], which is larger than the width of the flat bands ~ 2 –5 meV and the SOC considered in the present work. How are the results of Fig. 2 modified in the presence of Coulomb interaction? Treating this interaction within the Hartree-Fock approximation revealed that the Hartree term considerably widens the bands while the exchange term leads, basically, to broken-symmetry phases. At the charge neutrality, the Hartree term vanishes and the exchange potential, which concerns bands with identical spins, opens a gap of 4 meV [80,82,83], which is of the order of the SOC amplitudes. As a consequence, the statistical weight $\Delta f(E)$ of the bands with opposite $\langle S_z \rangle$ is expected to increase, but keeping larger values at small angles compared to the MA. Moreover, the bandwidth, around the MA, is found to relatively increase under the exchange term [80,82,83], but remains smaller than 3 meV, which preserve the flatness of the bands. It comes out

that our results hold in undoped TBG under Coulomb interaction, and can be used to extract the value of the MA at which the Gilbert damping correction drops. Away from the neutrality, the bands are substantially distorted by the Coulomb interaction [80,82,83] and our results should be taken with a grain of salt since they account for filling ν factors away from $-0.5 < \nu < 0.5$, where the bandwidth, at the MA, is less than 4 meV.

Besides interactions, strain is found to be a key parameter in the emergence of flat bands in TBG [68,70]. The effect of strain can be included in our model by deriving the strain induced correction to the Hamiltonian given by Eq. (3), taking into account the strain dependence of the vectors \mathbf{q}_j connecting the Dirac points [70]. The twist angle, at which $\delta\alpha_G$ drops, can then provide a way to measure the strain in TBG.

Experimental realization. Our proposed setup consists of an interface between a FI and a fully hBN encapsulated TBG/WSe₂ heterostructure (Fig. 1). The hBN layer acts as a tunnel barrier which prevents the diffusion of the FI atoms into the graphene layer [84]. On the other hand, the encapsulation provides a clean interface and prevents the graphene degradation [84] which is a challenging issue in the STM and ARPES experiments [24–26], carried out on nonencapsulated TBG samples. It should be stressed that the hBN encapsulated TBG/WSe₂ heterostructure has been already realized in Refs. [56,57]. Furthermore, the spin transport through a clean interface between a FI and 2D material has been experimentally achieved [84,85]. The 2D materials were fully encapsulated by hBN [84] or covered by a thin layer of an oxide insulator (as MgO) [85] to avoid the interdiffusion with the FI. Our proposed technique to measure the MA can then be implemented experimentally with

a clean interface and at room temperature. Moreover, an *in situ* manipulation of the twist angle can be realized as in Refs. [86–89].

Conclusion. To conclude, we have proposed an experiment to probe the flat bands of TBG and to measure its MA accurately. The experiment is based on a spin pumping measurement through a junction separating a FI and a TBG adjacent to a monolayer of WSe₂. We first derived the continuum model of TBG with SOC, which constitutes a first step to develop an analytical understanding of the emergence of a stable superconducting state at small twist angles observed in TBG in proximity to WSe₂ [56]. We then determined analytically the Gilbert damping correction $\delta\alpha_G$ induced by the presence of the TBG/WSe₂ heterostructure. Our results show that the twist angle dependence of $\delta\alpha_G$ exhibits a drop at the MA with a temperature-dependent fine structure. This feature provides an accurate determination of the MA and an estimation of the SOC induced in TBG by its proximity to the TMD layer. Our proposed set-up can be readily implemented regarding the state-of-the-art of the experimental realizations of SP in 2D materials and TBG-based heterostructure. Our work opens the gate to a twist tunable spintronics in twisted layered heterostructures.

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