

RKKY interaction between extended magnetic defect lines in grapheneP. D. Gorman,^{1,*} J. M. Duffy,¹ S. R. Power,² and M. S. Ferreira^{1,3}¹*School of Physics, Trinity College Dublin, Dublin 2, Ireland*²*Center for Nanostructured Graphene (CNG), DTU Nanotech, Department of Micro- and Nanotechnology, Technical University of Denmark, DK-2800 Kongens Lyngby, Denmark*³*CRANN, Trinity College Dublin, Dublin 2, Ireland*

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Of fundamental interest in the field of spintronics is the mechanism of indirect exchange coupling between magnetic impurities embedded in metallic hosts. A range of physical features, such as magnetotransport and overall magnetic moment formation, are predicated upon this magnetic coupling, often referred to as the Ruderman-Kittel-Kasuya-Yosida (RKKY) interaction. Recent theoretical studies on the RKKY in graphene have been motivated by possible spintronic applications of magnetically doped graphene systems. In this paper a combination of analytic and numerical techniques are used to examine the effects of defect dimensionality on such an interaction. We show, in a mathematically transparent manner, that moving from single magnetic impurities to extended lines of impurities effectively reduces the dimensionality of the system and increases the range of the interaction. This has important consequences for the spintronic application of magnetically-doped systems, and we illustrate this with a simple magnetoresistance device.

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I. INTRODUCTION

Interest in graphene has been spreading within the scientific community due to its potential for applications in myriad fields such as photonics, sensor technology, and spintronics [1–3]. Graphene’s weak spin-orbit and hyperfine interactions, which are the sources of spin relaxation and decoherence in other materials, make spintronic applications particularly attractive.

Of particular interest in the field of spintronics is the mechanism of the interaction, mediated by the conduction electrons of the host material, between localized magnetic moments embedded in nanoscale systems. This indirect exchange coupling (IEC) manifests as an energy difference between different alignments of the localized moments and is usually calculated within the Ruderman-Kittel-Kasuya-Yosida (RKKY) approximation [4–7]. This interaction has been extensively studied in graphene, nanoribbons, and nanotubes for a wide variety of impurities. The behavior of this interaction has been found to depend on the host [8–11], impurity type [12–17], and impurity configuration [18–21]. Since a range of effects are predicated on exchange interactions, methods of modifying the interaction via strain, edges, magnetic fields, doping, and lattice defects have also been studied [22–26].

An important aspect of the RKKY interaction is the rate at which it decays as a function of the separation D between magnetic impurities. In undoped graphene a decay rate of D^{-3} for substitutional, top-adsorbed, and bridge-adsorbed impurities is found, while a much faster decay rate of D^{-7} is found for center-adsorbed impurities [13,14,20,23]. This decay rate is faster than the D^{-2} decay expected for conventional two-dimensional materials and arises from the vanishing density of states at the Fermi energy in graphene [27]. This fast decay rate results in the interaction being very short ranged, and any method of amplifying the coupling to extend its range

could prove useful for both the experimental detection of the RKKY interaction and future spintronic applications.

The sign of the coupling, which determines the ferromagnetic (FM) or antiferromagnetic (AFM) alignment of the moments, should oscillate as a function of their separation, but in graphene this is masked by the coincidence of the Fermi surface and the corners of the Brillouin zone when the Fermi level coincides with the Dirac point. This causes the sign of the coupling, within the RKKY interaction, to depend only on whether the two moments occupy the same or opposite sublattices. When graphene is doped or gated, the Fermi surface and Brillouin zone no longer coincide in this manner [27], so sign-changing oscillations are recovered and the interaction is found to decay as D^{-2} .

There is currently a large and growing interest in nanopatterning graphene. Atomically precise tailoring of 2D materials, the addition of absorbants or vacancies, allows for a complex manipulation of the electronic properties of the material. Recent studies have examined how larger structures of impurities or vacancies can be used to modify the electronic properties of graphene allotropes. One such study looks at how the controlled growth of a line of defects in graphene can be used for gate-tunable valley filtering [28], while another study looks at how transition-metal nanowires result in long-range magnetic order and magnetic anisotropy in graphyne [28,29].

The possibility and ease of depositing a line of magnetic impurities in graphene has increased with the recent work of Duesberg *et al.* [30,31]. In these works the electronic structure of graphene is altered via folding—sometimes called graphene origami. This folding is achieved by depositing the graphene over a corrugated surface which, in conjunction with an applied magnetic field, has recently been shown to open a band gap in graphene [32]. The ridges created during this process act as traps for magnetic impurities and open up the possibility of depositing impurities in straight parallel lines. Alternatively, increased reactivity near grain boundaries [33] or simple kinks [34] may also lead to the formation of impurity lines. A sublattice preference of the impurities along zigzag

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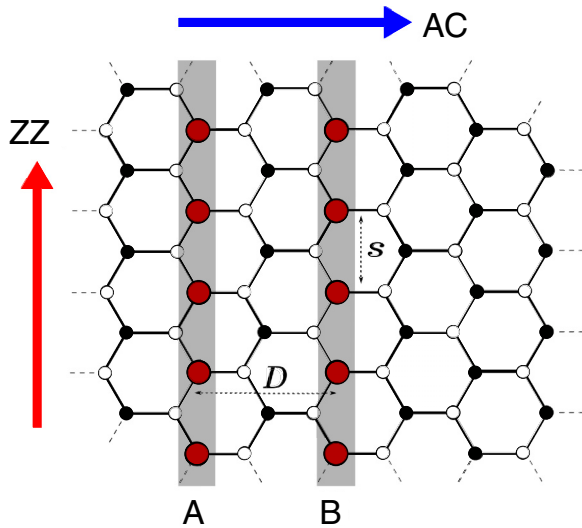


FIG. 1. (Color online) Schematic representation of the graphene lattice showing the armchair (AC) and zigzag (ZZ) directions. The filled and hollow symbols (\bullet, \circ) represent sites on different sublattices. The red circles represent two parallel lines of magnetic impurities, A and B , separated by a distance $D = 1$ (in units of $3a$, where $a = 2.64 \text{ \AA}$) in the armchair direction, with a dimensionless quantity s (here $s = 1$) indicating the zigzag separation between moments.

lines, as shown in Fig. 1, is expected due to similar behavior predicted for zigzag edged nanoribbons [35]. Although the RKKY interaction in graphene has been intensively studied, one aspect that has yet to be examined is how lines of impurities, which change the dimensionality of the interaction, affect the coupling.

With the motivation of understanding the magnetic interaction strength, and providing some theoretical foundation, we will examine the role of dimensionality in the RKKY interaction. In this paper we will apply analytical and numerical techniques to a system containing two parallel lines of magnetic impurities (Fig. 1), and present a method for calculating the coupling between lines of impurities embedded or adsorbed in a host material. Some analogues can be made between our setup and 1D systems such as carbon nanotubes and nanoribbons [11], as well as systems with two ferromagnetic monolayers embedded in a nonmagnetic metal [36,37], where the Ruderman-Kittel theory can be used to derive a D^{-2} decay rate for the coupling between the ferromagnetic layers [38]. We start by introducing the general formalism used to calculate the magnetic coupling, which is written entirely in terms of the real-space single-particle Green's functions (GFs) of the host graphene sheet. We show in a mathematically transparent manner how the standard coupling equation may be modified to calculate the RKKY interaction between two infinite lines of impurities. We find that the interaction of finite lines of impurities quickly tends towards the interaction of infinite lines, and that the reduced dimensionality of the system leads to smaller decay rates and longer ranged interactions. These predictions are confirmed using numerical calculations. Finally, we examine the magnetoresistance response of a simple device based on the setup discussed in this paper and which may motivate further

studies of graphene spintronics based on magnetic defect lines. In this paper we only consider a graphene host, but the method is easily generalized to other two-dimensional materials.

II. METHODS

The indirect exchange coupling J is defined as the change in energy between impurities in FM and AFM alignments. This can be calculated conveniently using the Lloyd formula [39], which allows the total change in energy ΔE between FM and AFM alignments to be determined without requiring that the total energy of either system be calculated. This is given as

$$J = \Delta E(E_F) = \frac{1}{\pi} \text{Im} \int dE f(E) \ln(\det(\hat{I} - \hat{G}\hat{V}_s)), \quad (1)$$

where $f(E)$ is the Fermi function, \hat{G} represents the GF of the system containing impurities, and \hat{V}_s is the spin perturbation matrix which rotates specific impurities by an angle θ . A nearest-neighbor tight-binding Hamiltonian is used to describe the electronic properties of the system and calculate the GFs. The impurities are introduced to the pristine system via Dyson's equation

$$\hat{G} = (\hat{I} - \hat{g}\hat{V}_I)^{-1}\hat{g}, \quad (2)$$

where the characteristics of the impurities are contained within the perturbation matrix \hat{V}_I , and \hat{g} is the pristine graphene GF calculated from a nearest-neighbor tight-binding approximation Hamiltonian.

The calculation of the coupling between two parallel lines of impurities A and B of length N , which we will denote J_{AB}^N , is relatively simple. However, the necessary introduction of a $2N \times 2N$ perturbation matrix \hat{V}_I creates a computational problem: The matrix inversion present in the coupling calculation becomes cumbersome as N becomes large. What follows is a method which allows for the rapid calculation, as well as yielding an analytic form, of the coupling between two infinite parallel lines of impurities. The finite-line coupling J_{AB}^N and infinite-line coupling \mathcal{J}_{AB} are then compared in the results section.

The coupling calculation can be greatly simplified by keeping the GF partially in reciprocal space when introducing the perturbation. We will introduce our perturbation as a line of impurities along the y (zigzag) direction, though a similar procedure may be used for impurities in the x (armchair) direction (Fig. 1). When dealing with lines of impurities in the y direction, we make the projection $\langle x, k_y, \alpha | \hat{g} | x', k_y, \beta \rangle = g_{xx'}^{\alpha\beta}(k_y)$, where \hat{g} is the GF of the pristine system, and α and β are sublattice labels. This can be written as

$$g_{xx'}^{\alpha\beta}(k_y) = \frac{1}{N_x} \sum_{k_x} \frac{N^{\alpha\beta}(E, k_y) e^{ik_x(x'-x)}}{E^2 - |f(k_x, k_y)|^2}, \quad (3)$$

where $N^{\alpha\beta}$ is a sublattice dependent term

$$N^{\alpha\beta} = \begin{cases} E & \alpha = \beta \\ f(k_x, k_y) & \alpha \neq \beta \end{cases}, \quad (4)$$

and $f(k_x, k_y)$ is the dispersion relation

$$f(k_x, k_y) = t + 2t \cos(k_x) e^{ik_y}. \quad (5)$$

For neatness we will neglect the superscripts α and β from the following equations, which should be taken to be same sublattice ($\alpha = \beta$), unless otherwise stated. Since we are considering two infinite parallel lines that extend in the zigzag direction, the separation between these lines must necessarily be in the armchair direction. We no longer refer to the sites as x and x' , but as A and B , which represent each line of impurities, respectively; the new separation is now just $D = x' - x$. The sum over k_x may be converted into an integral over the k_x direction in the Brillouin zone, which can then be performed by contour integration. This results in an analytic form for the GF,

$$g_{AB}(k_y) = \frac{i}{4t^2} \frac{N(E, k_y) e^{iq(ak_y)D}}{\cos\left(\frac{ak_y}{2}\right) \sin(q(ak_y))}, \quad (6)$$

with

$$q(k_y) = \pm \cos^{-1} \left[\frac{E^2 - t^2 - 4t^2 \cos^2\left(\frac{ak_y}{2}\right)}{4t^2 \cos\left(\frac{ak_y}{2}\right)} \right], \quad (7)$$

where t is the hopping integral and a is the graphene lattice parameter.

Considering Dyson's equation, we may greatly reduce our analytical workload by writing our GFs as Bloch matrices $\hat{g}_{AA} = \sum_{k_y} |A, k_y\rangle g_{AA}(k_y) \langle A, k_y|$, so that

$$\hat{I} - \hat{g}\hat{V} = \hat{I} - \begin{pmatrix} \hat{g}_{AA} & \hat{g}_{AB} & \cdots \\ \hat{g}_{BA} & \hat{g}_{BB} & \\ \vdots & & \ddots \end{pmatrix} \begin{pmatrix} \tau \hat{I} & 0 & \cdots \\ 0 & \tau \hat{I} & \\ \vdots & & \mathbf{0} \end{pmatrix}, \quad (8)$$

where our perturbation, with change in onsite energy τ , has been trivially transformed to the same basis by a Fourier transformation

$$\begin{aligned} \hat{V}_I &= \sum_n \tau |A, ns\rangle \langle A, ns| + \tau |B, ns\rangle \langle B, ns|, \\ &= \frac{1}{s} \sum_{k_y} \tau |A, k_y\rangle \langle A, k_y| + \tau |B, k_y\rangle \langle B, k_y|, \end{aligned} \quad (9)$$

where s is a dimensionless quantity describing the separation of impurities along the zigzag direction as shown in Fig. 1, and n is summed over all sites. Since all but the first two columns are zero, we can find the desired elements by just considering the 2×2 matrix multiplication. Similar identities facilitate the inversion, and our new GF \hat{G} can be calculated in a relatively transparent manner

The spin perturbation is introduced as

$$\hat{V}_s = \sum_{k_y} |B, k_y\rangle \hat{V}_\theta \langle B, k_y|, \quad (10)$$

where \hat{V}_θ is a 2D spin perturbation matrix which introduces the exchange splitting V_{ex} between the bands and leads to a greater occupation of up-spin electrons and hence to the formation of a magnetic moment, and rotates the spins on line B through an angle of θ , and is defined as

$$\hat{V}_\theta = -V_{\text{ex}} \begin{pmatrix} \cos \theta - 1 & \sin \theta \\ \sin \theta & 1 - \cos \theta \end{pmatrix}. \quad (11)$$

Calculating $\hat{I} - \hat{g}\hat{V}$ for the spin perturbation is more involved, but the basic method is the same. For $\theta = \pi$ we can now write

$$\begin{aligned} \det(\hat{I} - \hat{G}\hat{V}_s) &= \det((\hat{I} - 2V_{\text{ex}}\hat{g}_{BB}^\uparrow)(\hat{I} + 2V_{\text{ex}}\hat{g}_{BB}^\downarrow)), \\ &= \prod_{k_y} [(1 - 2V_{\text{ex}}\hat{g}_{BB}^\uparrow(k_y))(1 + 2V_{\text{ex}}\hat{g}_{BB}^\downarrow(k_y))], \end{aligned} \quad (12)$$

where all of the terms are diagonal Bloch matrices in k_y space, which allows us to write the determinant as a product.

To see the distance dependence in the coupling, we use Dyson's equation to write the coupling in terms of the spin dependent Green's function between each line of impurities, $g_{AB}^{\uparrow/\downarrow}$. This is done by calculating the effect at site B of raising the energy of the down-spin channel on sites A and B

$$g_{BB}^\downarrow = g_{BB}^\uparrow + 2V_{\text{ex}}\hat{g}_{BB}^\uparrow\hat{g}_{BB}^\downarrow + 2V_{\text{ex}}\hat{g}_{BA}^\uparrow\hat{g}_{AB}^\downarrow. \quad (13)$$

By rearranging this, expanding out Eq. (12), and using the properties of logarithms, we can now write the coupling per atom pair AB , \mathcal{J}_{AB} , in a convenient form

$$\begin{aligned} \mathcal{J}_{AB} &= -\frac{s}{N_y\pi} \text{Im} \int_{-\infty}^{E_F} dE \\ &\times \sum_{k_y} \ln(1 + 4V_{\text{ex}}^2\hat{g}_{BA}^\uparrow(E, k_y)\hat{g}_{AB}^\downarrow(E, k_y)). \end{aligned} \quad (14)$$

The sum may be converted to an integral over the Brillouin zone in the k_y direction. We may also write $k_z = \frac{a}{2}k_y$, exploiting the symmetries of the GF, and use the analyticity of the integrand in the upper half plane to change the integration to the imaginary axis. This gives us a form of the coupling that is suitable for evaluation using numerical methods,

$$\begin{aligned} \mathcal{J}_{AB} &= \frac{2}{\pi^2} \int_\eta^\infty dy \int_0^{\frac{\pi}{2}} dk_z \\ &\times \ln(|1 + 4V_{\text{ex}}^2\hat{g}_{BA}^\uparrow(E_F + iy, k_z)\hat{g}_{AB}^\downarrow(E_F + iy, k_z)|). \end{aligned} \quad (15)$$

To obtain a useful analytic form of the coupling we make several approximations which are similar in form to the RKKY approximation. We first expand the log in Eq. (14) to first order by assuming a small exchange coupling V_{ex} . We then make the approximation $g_{AB}^\downarrow = g_{BA}^\uparrow = g_{AB}$ so that the coupling, which we now write as \mathcal{J}_{AB} to distinguish it from the numerical form above, is given by

$$\mathcal{J}_{AB} \approx -\frac{4V_{\text{ex}}^2}{\pi^2} \text{Im} \int_{-\infty}^{E_F} dE \int_{-\frac{\pi}{2}}^{\frac{\pi}{2}} dk_z [g_{AB}(E, k_z)]^2. \quad (16)$$

At large separations the integrand oscillates quickly and only select points contribute to the integral over k_z . This allows us to use the stationary phase approximation (SPA) to write the integral in a fully analytic form. The SPA has previously been used [27] to write the pristine Green's functions of graphene in an analytic form. This allows one to calculate the coupling between two impurities. Here we use the SPA to calculate the integral over Green's functions that already contain impurities,

$$\int_{-\frac{\pi}{2}}^{\frac{\pi}{2}} dk_z [g_{AB}(E, k_z)]^2 = \frac{\mathcal{A}(E)e^{i2Q(E)D}}{\sqrt{D}}, \quad (17)$$

where $\mathcal{A}(E)$ captures the specific sublattice dependencies, and noncontributing terms have been ignored. The symmetry in the zigzag direction allows us to use the SPA in this manner.

We can now write our coupling equation as

$$\mathcal{J}_{AB} = -\frac{4V_{\text{ex}}^2}{\pi^2} \text{Im} \int_{-\infty}^{E_F} dE \frac{\mathcal{A}(E)e^{i2Q(E)D}}{\sqrt{D}}. \quad (18)$$

The integration procedure is now identical to that for single impurities in graphene [27]. The functions $\mathcal{A}(E)$ and $Q(E)$ are expanded around E_F and the integral can be reduced to a sum over Matsubara frequencies, which in the low temperature limit gives

$$\mathcal{J}_{AB} = V_{\text{ex}}^2 \text{Im} \sum_{\ell} \frac{\mathcal{A}^{(\ell)} e^{i2Q(E)|D|} (-1)^{\ell}}{(2iQ^{(1)})^{\ell+1} D^{\ell+\frac{3}{2}}}. \quad (19)$$

The decay rate of the coupling is now determined by the first nonvanishing $\mathcal{A}^{(\ell)}$, where this notation is defined as the ℓ^{th} derivative of \mathcal{A} with respect to energy evaluated at the Fermi energy.

III. RESULTS

Unless otherwise specified the results that follow refer to the case $s = 1$, i.e., that the impurities are separated by the minimum distance in the zigzag direction. The stationary phase approximation (SPA) used in the derivation of Eq. (18) relies on the assumption of large separations between the lines of impurities, so it is important to check the accuracy of this approximation at reasonable separations. The SPA coupling \mathcal{J}_{AB} was compared against the full numerical calculation of the coupling \mathcal{J}_{AB} for a variety of Fermi energies and exchange splittings. A representative sample of these checks is shown in Fig. 2, where the agreement is seen to be very good even for reasonably small separations and increases in accuracy as separation increases. The approximation is therefore concluded to be sufficiently accurate for our uses here.

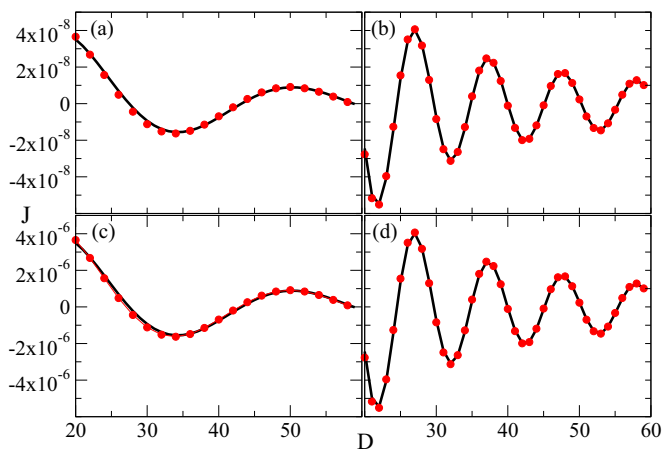


FIG. 2. (Color online) A comparison of the coupling, $\mathcal{J}_{AB}^{\alpha\alpha}$ (lines), and the SPA of coupling, $\mathcal{J}_{AB}^{\alpha\alpha}$ (dots), for different exchange splittings V_{ex} and at different Fermi energies E_F . (a) $V_{\text{ex}} = 0.01$, $E_F = 0.1$. (b) $V_{\text{ex}} = 0.01$, $E_F = 0.3$. (c) $V_{\text{ex}} = 0.1$, $E_F = 0.1$. (d) $V_{\text{ex}} = 0.1$, $E_F = 0.3$.

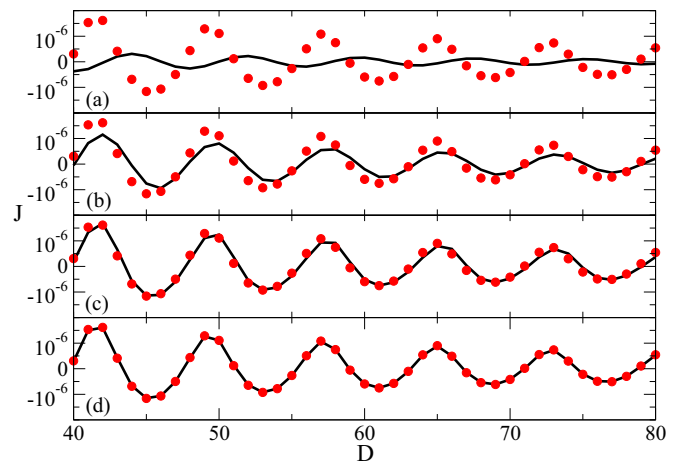


FIG. 3. (Color online) A plot of the numerical evaluation of the finite-line coupling \mathcal{J}_{AB}^N against separation D for a pair of finite lines of length (a) $N = 1$, (b) $N = 10$, (c) $N = 50$, (d) $N = 500$. In each case the numerical evaluation of the infinite-line coupling \mathcal{J}_{AB} (Eq. 15) is represented by the red dots.

Figure 3 plots the coupling between several pairs of line segments, of length N , against an infinite pair of lines. As we increase the length of the line segments the magnitude and phase of the coupling are seen to approach that of the infinite line segment (red dots). At a length of $N = 500$ [Fig. 3(d)], the coupling can be seen to converge to that of an infinite line of impurities. For shorter lines, $N \sim 50$ [Fig. 3(c)] the coupling closely resembles that between the infinite lines, with only slight discrepancies in magnitude and phase. These discrepancies increase as the line is shortened to $N \sim 10$ [Fig. 3(b)], however even the coupling between line segments as short as $N = 10$ are seen to behave much closer to the infinite case than to the case of individual impurities [cf. Fig. 3(a)]. Thus Eq. (19) is a useful tool for quickly calculating the coupling between finite lines of impurities, since it is clear that as the length N of the lines increases the interaction converges to that of the infinite case.

An important feature of the coupling is its overall rate of decay with separation $D^{-\alpha}$, which is determined from the analytic form of the coupling in Eq. (19). Here the coupling is represented as a sum of terms over ℓ , with decay rates that increase as $D^{-\ell-\frac{3}{2}}$. The decay rate is therefore determined by the first nonzero term in the series, as subsequent terms decay much more rapidly. It is the quantity $\mathcal{A}^{(\ell)}(E)$ that determines whether or not the whole term will vanish. For the case of lines of impurities occupying the same sublattice ($\alpha = \beta$), where the interaction is ferromagnetic, we have

$$(\mathcal{A}^{\alpha\alpha}(E))^{(0)} = -\sqrt{\frac{\pm i\pi}{(E^2 + 3t^2)}} \frac{E^2}{2(E^2 t^2 - E^4)^{\frac{3}{4}}}. \quad (20)$$

It is clear from examination that $\mathcal{A}^{(0)}$ is nonzero whenever E is nonzero, and thus away from the Fermi energy we would expect a decay rate that goes as $\sim D^{-\frac{3}{2}}$.

In Fig. 4 we compare the coupling with the expected value of its decay rate at several Fermi energies. We find that for values of the Fermi energy not equal to zero ($E_F \neq 0$)

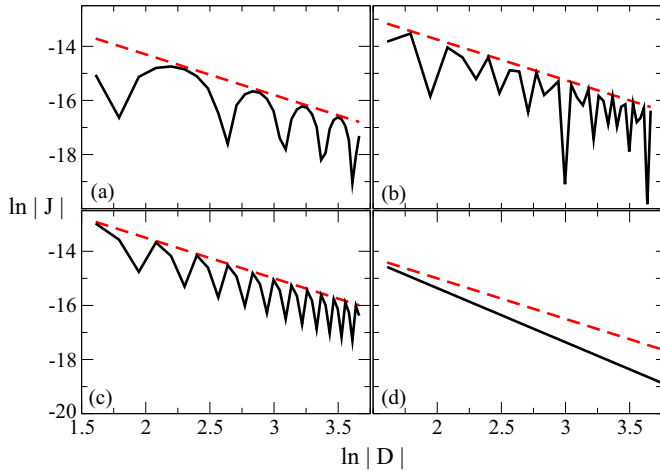


FIG. 4. (Color online) A log-log plot of the numerical evaluation of the coupling, J , between two lines of impurities as a function of their separation, D , for (a) $E_F = 0.1$, (b) $E_F = 0.3$, (c) $E_F = 0.5$, (d) $E_F = 0$. In each case the log of the coupling is represented by the solid black line and the dashed red line represents a decay rate of 1.5.

that our theoretical predictions hold. At precisely $E_F = 0$, corresponding to an undoped graphene sheet, the analytical approximation no longer holds and a faster decay rate of $\sim D^{-2}$ is found. This difference of behavior at the Dirac point is a hallmark of such interactions in graphene and is also seen for the single impurity case. However, both of the linear defect results correspond to an increased interaction range compared to single site impurities, where we would expect to find decay rates of D^{-2} and D^{-3} for doped and undoped graphene, respectively [27].

The form of $\mathcal{A}^{\alpha\beta}(E)$ was also examined for the case of opposite sublattice occupation ($\alpha \neq \beta$), where the interaction is antiferromagnetic, as well as the case where one line is offset by half a lattice spacing. These cases produced similar results to the initial case examined above in both coupling decay rate and the convergence of finite lines to the infinite case. For the cases where the impurities are spaced in the zigzag direction by a larger length $s > 1$, we again find that the coupling between lines quickly converges to that of the infinite case as N is increased.

IV. MAGNETORESISTANCE

RKKY interactions in multilayered systems play a key role in the giant magnetoresistance (GMR) effect. Up- and down-spin electrons have different transmission probabilities through a magnetic layer of a certain orientation. The most energetically favorable configuration of layer orientations is determined by RKKY interactions and will have a certain total transmission. Aligning all the layers with an applied magnetic field can lead to markedly different transmission probabilities and a resulting change in the resistance of the system [40,41]. Previous studies have considered the possibility of magnetoresistance devices based on graphene systems [42–44], but it is worth examining whether a magnetoresistance signal emerges using devices based on the system we have considered in this paper. We consider two lines of

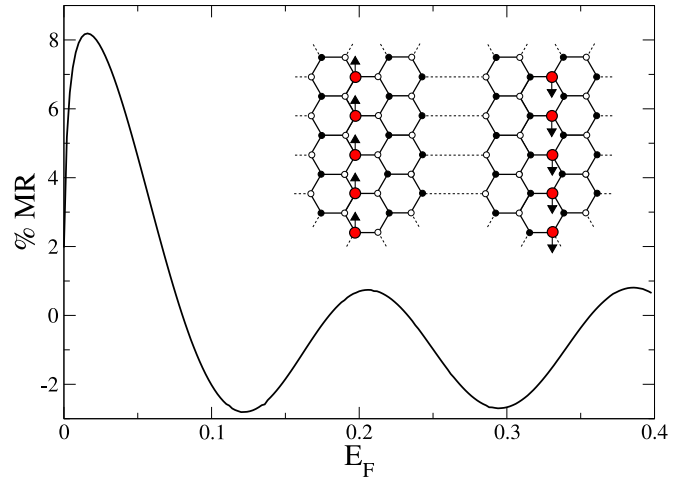


FIG. 5. (Color online) Magnetoresistance percentage as a function of Fermi energy for a simple device based on magnetic lines in a graphene sheet. The inset shows the initial AFM orientation of two lines of moments on opposite sublattices with $D = 9$.

magnetic impurities, one on each sublattice, as shown in the inset of Fig. 5. The total conductance $\Gamma_{\text{AFM/FM}}$ through such a system is calculated as the sum of the conductances of up-spin and down-spin electrons. These are calculated using the standard Landauer-Buttiker approach with recursively calculated Green's functions and spin-dependent potentials at each of the magnetic impurity sites. We assume that these lines initially have an antiferromagnetic relative orientation [45], and the magnetoresistance is calculated from the relative change in the resistance when a small magnetic field is applied to force a ferromagnetic alignment

$$MR = \frac{\Gamma_{\text{FM}}^{-1} - \Gamma_{\text{AFM}}^{-1}}{\Gamma_{\text{AFM}}^{-1}}. \quad (21)$$

The magnetoresistance, expressed as a percentage, is plotted in the main panel of Fig. 5. The overall magnitude of the magnetoresistance will vary with V_{ex} and thus depend on the exact impurity species and may be considerably smaller than our calculated value with an arbitrarily chosen $V_{\text{ex}} = 0.2|t|$. However, the qualitative features observed here are robust over a wide range of impurity parametrizations. Furthermore, it should be noted that this calculation only considers two impurity lines. A more realistic device could contain many more lines, and this would increase the magnetoresistance significantly by increasing the discrepancy in potential landscapes experienced by electrons with or without an applied magnetic field. Nonetheless, this simple example demonstrates the potential for magnetic impurity lines to play a part in graphene-based spintronics and motivates further studies of such systems and the RKKY interactions on which magnetoresistance effects are predicated.

V. CONCLUSIONS

In this paper we have derived an analytic expression for the RKKY interaction in graphene between two lines of magnetic impurities separated along the armchair direction. We have shown that the coupling between lines of impurities

quickly converges to the coupling of the infinite case as their length increases. The analytic form is therefore a useful way to approximate the interaction between line segments. We also use this analytic form to predict the rate of decay of this interaction, away from $E_F = 0$, which is slower than the rate of decay for two impurity interactions. Furthermore we have shown that it is the reduced dimensionality of the system that increases the range of the RKKY interaction. This increased range may ease the detection of the RKKY which is notoriously hard to examine experimentally. We also show that these extended line defects may have interesting magnetoresistance properties which motivates further study. Since a whole range of physical features, such as magnetotransport and overall magnetic moment formation, are predicated upon the

magnetic coupling, it is hoped that this work may lead to interesting spintronic applications.

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