Magnetic circular dichroism in the $dd$ excitation in the van der Waals magnet CrI$_3$
probed by resonant inelastic x-ray scattering

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We report on a combined experimental and theoretical study on CrI$_3$ single crystals by employing the polarization dependence of resonant inelastic x-ray scattering (RIXS). Our investigations reveal multiple Cr 3$d$ orbital splitting ($dd$ excitations) as well as magnetic dichroism (MD) in the RIXS spectra. The $dd$ excitation energies are similar on the two sides of the ferromagnetic transition temperature, $T_c \sim 61$ K, although MD in RIXS is predominant at 0.4 T magnetic field below $T_c$. This demonstrates that the ferromagnetic superexchange interaction that is responsible for the interatomic exchange field is vanishingly small compared with the local exchange field that comes from exchange and correlation interaction among the interacting Cr 3$d$ orbitals. The recorded RIXS spectra reported here reveal clearly resolved Cr 3$d$ intraorbital $dd$ excitations that represent transitions between electronic levels that are heavily influenced by dynamic correlations and multiconfiguration effects. Our calculations taking into account the Cr 3$d$ hybridization with the ligand valence states and the full multiplet structure due to intra-atomic and crystal field interactions in $O_h$ and $D_{3d}$ symmetry clearly reproduced the dichroic trend in experimental RIXS spectra.

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I. INTRODUCTION

Over the last few years, two-dimensional (2D) van der Waals materials have spurred enormous interest due to their unique magnetic properties and potential in the development of multifunctional electronics and spintronics devices [1–5]. Among various 2D systems, the family of CrX$_3$ (X = Cl, Br, and I) has recently been at the center of widespread research in developing high-quality samples, from monolayer to bulk single crystals. Although ferromagnetism in semiconducting bulk CrI$_3$ has been known since the mid-twentieth century [6], recent experiments reveal layer-dependent magnetic phases [7,8]: from ferromagnetism in the monolayer to antiferromagnetism in the bilayer and back to ferromagnetism in the trilayer and bulk. This has been postulated to arise from competing interlayer and intralayer antiferromagnetic and ferromagnetic interactions, respectively [8]. According to Mermin-Wagner theory [9], in the absence of intrinsic anisotropy, long-range magnetic order is strongly suppressed in a 2D isotropic Heisenberg system due to spin fluctuation at a finite temperature. However, any deviation from a pure Heisenberg interaction, such as the influence of the magnetocrystalline anisotropy (MCA), would be able to stabilize a magnetically ordered state, which is most likely what happens in CrI$_3$ [10]. In CrI$_3$, the MCA energy is $\Delta E_{\text{MCA}} \sim 0.5$ meV [11], a large value whose origin has been a topic of enormous interest in the past couple of years with continued debate. X-ray magnetic circular dichroism (XMCD) studies at Cr L$_{2,3}$ and I M$_{4,5}$ edges revealed strongly suppressed single-ion anisotropies of Cr 3$d$ and I 5$p$ states, whose energies are too small to explain the origin of $\Delta E_{\text{MCA}}$.  

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Despite this, there is a considerable spin-orbit coupling (SOC) strength (~0.63 eV) in the I 5p state and strong Cr 3d−I 5p hybridization [12,13]. The I 5p state SOC strength primarily contributes to the MCA (in contrast to the much smaller SOC of Cr 3d ∼ 0.05 eV), either through the Cr 3d−I 5p−Cr 3d superexchange hopping or via substantial admixture of band states [14]. Superexchange in this material is stabilized by the inherent CrI 3 crystal structure constituting of trivalent Cr ions in an octahedral environment of I ions and the in-plane Cr-I-Cr bond angle ∼ 95°. In this configuration, according to the Goodenough-Kanamori-Anderson (GKA) rule, the Cr-Cr coupling is expected to be primarily ferromagnetic [15]. Therefore the origin of long-range ferromagnetism could be attributed to the high SOC of I 5p states mediating the anisotropic Cr 3d−I 5p−Cr 3d superexchange hopping through 95° in-plane Cr-I-Cr bond angle [16]. Recent theoretical results suggest that the magnetic interaction is not purely ferromagnetic; instead there is a competition between the ferromagnetic interaction and the antiferromagnetic interaction, which involve different Cr 3d orbitals [10,11,17,18]. Calculations also suggest a sizable contribution of the orbital-resolved components of Cr 3d states from nearest-neighbor interactions [19,20]. In monolayer CrI 3, recent electronic structure calculations reveal substantial exchange splitting of the spin-polarized Cr 3d t 2g and e g states due to ferromagnetic Cr-I-Cr superexchange interaction [20]. However, the electronic structure of this (and similar) materials is complex, with expected competition from kinematic (band formation) effects and onsite Coulomb repulsion (e.g., as parametrized by the Hubbard U). It is expected that the Cr 3d interorbital interactions mediated by the I 5p orbital hold the key to understanding most electronic, magnetic, and transport properties of CrI 3 [21]. Therefore probing the Cr 3d orbital is indispensable for an in-depth understanding of the magnetic properties of CrI 3.

To directly access the Cr 3d interorbital interactions, we carried out resonant inelastic x-ray scattering (RIXS) studies at the Cr L 2,3 edge of bulk CrI 3 single-crystal samples with circularly polarized x rays. Soft x-ray RIXS, corresponding to the direct transition from Cr 2p core levels to the Cr 3d orbitals, allows us to study orbital (dd) excitations. In addition, magnetic circular dichroism (MCD) in RIXS enabled an in-depth understanding of magnetic interactions from the dd excitation which is also sensitive to the symmetry of the d orbitals. The experimental investigations are found to be well corroborated by electronic structure calculations.

II. EXPERIMENTAL AND THEORETICAL METHODS

Bulk CrI 3 single crystals were grown by chemical vapor transport, by reacting chromium powder (99.5%; Sigma-Aldrich) and anhydrous iodine beads (99.999%; same supplier) in a 1:3 ratio inside a glovebox with an argon atmosphere. Figures 1(a) and 1(b) show a scanning electron microscopy image of a CrI 3 single-crystal sample from the top and side, respectively, along with a schematic crystal structure shown at the bottom of each panel. The easily cleavable quality of the sample is clearly visible as multiple layers are observed particularly in Fig. 1(b), where the platelike crystal is bent to get a side view of the layers. Magnetization studies have been performed using a Quantum Design MPMS3 magnetometer with magnetic fields μ0H up to 7 T applied parallel and perpendicular to the crystallographic c axis, respectively. The calculations of the RIXS spectra of CrI 3 in this paper were performed in the framework of the Anderson impurity model (AIM) using the QUANTY software package (for detailed information, see Supplemental Material [22]). The spectra of Cr 3+ were calculated considering the Cr 3d hybridization with the ligand valence states and the full multiplet structure of more than 3 T is required to overcome the MCA energy of ~0.55 meV, in agreement with a previous report [11]. Figure 2(a) shows M vs temperature [M(T)] plots at an external magnetic field of 0.1 T applied parallel to the c axis and the ab plane. The out-of-plane (easy axis) spins will not completely polarize along the ab plane at a nominal field of 0.1 T due to MCA [see Fig. 2(a)]; a magnetic field of more than 3 T is required to overcome the MCA energy and polarsize all the spins along the ab plane [23]. However, the projections of the canted c-axis spins on the ab plane, at μ0H ∥ ab and 0.1 T, could be antiferromagnetic [AFM; bottom right inset in Fig. 2(b)], which explains the M(T)
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FIG. 2. (a) $M(\mu_0 H)$ measured at 2 K, and (b) $M(T)$ plots with applied dc magnetic field parallel to the $c$ axis and to the $ab$ plane. The top insets in (b) show the $dM/dT$ plots to extract $T_C$ and $T_S$. A line shape for $\mu_0 H \parallel ab$. Also, these in-plane spins are not collinear at a nominal magnetic field (0.1 T), which explains why at the lowest measured temperature (2 K) there is still a small but finite $M$. To probe the in-plane ferromagnetic (FM) interactions at a nominal in-plane magnetic field (above the $H \parallel c$ saturation field), we have carried out x-ray magnetic circular dichroism (XMCD) and RIXS studies at the Cr $L_{2,3}$ edge. Cr $L_{2,3}$-edge RIXS measurements were carried out at the SEXTANTS beamline [24,25] of the SOLEIL Synchrotron using the novel MAGELEC sample environment [26]. Left and right circular (CL and CR) polarized x rays of energy across the Cr $L_{2,3}$ edge ($E_{\text{in}}$) were focused at $20^\circ$ grazing to the sample surface ($ab$ plane). A schematic representation of the experimental geometry is presented in Fig. 3(a). A magnetic field, $H = 0.5$ T, was applied parallel to the incoming x-ray beam. Therefore the components of $H$ along the $c$ axis ($H_c$) and $ab$ plane ($H_{ab}$) are 0.17 and 0.33 T, respectively. Measurements were carried out at 20 and 100 K, i.e., below and above the Curie temperature.

FIG. 3. (a) Schematic illustration of experimental geometry. The sample was illuminated with circularly polarized x-ray photons of energy $E_{\text{in}}$ at $20^\circ$ grazing to the sample surface. The scattered rays ($E_{\text{out}}$) were detected by the spectrometer fixed at $85^\circ$ scattering angle. A magnetic field, $H = 0.5$ T, was applied parallel to the x rays, and $H_c$ and $H_{ab}$ are the components of $H$ parallel to the $c$ axis and $ab$ plane, respectively. (b) Top: Cr $L_{2,3}$ XAS and XMCD spectra measured at 20 K and $+H$ magnetic field. Bottom: Calculated XAS and XMCD spectra. The inset shows the area under the energy loss region between 0.6 and 5 eV of the RIXS spectra.

Figure 3(b) shows the Cr $L_{2,3}$-edge x-ray absorption spectra (XAS) and corresponding XMCD of a CrI$_3$ single-crystal sample at 20 K [ferromagnetic phase of rhombohedral (R3) crystal structure]. XAS were recorded in the total electron yield mode with CL and CR polarized x rays. XMCD is the difference between the CR and CL polarized XAS as shown at the top of Fig. 3(b). The bottom of Fig. 3(b) shows the calculated XAS and XMCD spectra. Calculations involve using the multiplet of the ground state configuration together with corresponding quantum numbers (including S, L, and J) using QUANTY [27,28] (see Supplemental Material for details of the calculations) in the framework of the Anderson impurity model (with spins aligned parallel to the magnetic field, along the $c$ axis). In the QUANTY calculations, parameters such as the crystal field splitting and the spin-orbit coupling are treated as adjustable parameters. The calculated XAS and XMCD
spectra, as shown at the bottom of Fig. 3(b), agree fairly well with the experimental data.

Figure 4(a) summarizes the RIXS energy loss spectra at various excitation energies (for CR and CL polarized x rays) mentioned alongside each spectrum. In a conventional RIXS process, a monochromatic beam of energy $E_{\text{in}}$ is incident onto the sample, and one measures the energy distribution of the emitted radiation ($E_{\text{out}}$) converted into an energy loss ($E_{\text{in}} - E_{\text{out}}$) scale to determine any low-energy excitations of the system. The energy loss at 0 eV corresponds to the elastically scattered photons (where $E_{\text{in}} = E_{\text{out}}$), while other features correspond to $dd$ excitations, charge transfer (CT), and Cr $L_\alpha$ fluorescence, in the order of increasing energy loss. For coherent $dd$ and CT excitations [29,30], the energy losses are independent of incident energy and are also known as Raman-like losses. We identify the $dd$ excitation features (1.45 ± 0.02, 1.75 ± 0.02, and 2.50 ± 0.05 eV) between 1 and 3 eV energy loss as shown in Fig. 4(a), and for better clarity we replot the RIXS spectrum at the resonance $L_3$ energy (574.3 eV) in Fig. 4(b). Above 575 eV excitation energy [Fig. 4(a)], the broad feature (between 2 and 5 eV) resembles the emission ($L_\alpha$) from Cr 3$d_{5/2}$ to 2$p_{3/2}$ in addition to the buried $dd$ and CT features. The broad feature in the RIXS spectra excited near the absorption threshold (near 572 eV) signifies the metal-ligand CT (3.50 ± 0.10 eV), which gets smeared out in intensity relative to the $dd$ excitation features, near the resonant energy. The difference in intensities between the CL and CR RIXS features at 20 K, in particular at $dd$ excitations, is clearly visible at resonant excitation and also at other excitation energies in Fig. 4(a). A clear contrast in the integrated intensities can also be observed in addition to the XMCD contrast in Fig. 2. Upon reversing the direction of the magnetic field (to $-H$), the CL and CR RIXS features in the $dd$ excitation also switch their relative intensity depending on the light helicity, and at zero magnetic field their intensities are similar [Fig. 4(b)]. This behavior is direct evidence of magnetic dichroism (MD) in RIXS and is not an experimental artifact. Above $T_C$, at 100 K, we multiply the intensity by 10 to make it comparable to that at 20 K. Such a reduction in intensity above $T_C$ is expected primarily due to the disordered orientation in the spin state, when the conductivity and crystal structure of CrI$_3$ remain similar at the two temperatures [31].

As discussed earlier in the present measurement geometry, the MD in RIXS is primarily due to the Cr spins parallel to the c axis. This is due to the fact that even at $H = 0.17$ T, the Cr spins parallel to the c axis are fully ferromagnetically saturated, in contrast to unsaturated in-plane Cr spins at $H_{\text{sat}} = 0.33$ T [see Fig. 2(a)]. However, at the nominal magnetic field, the in-plane spins can be antiferromagnetically oriented as already observed in Fig. 2(b). A similar situation is present in α-Fe$_2$O$_3$ [32], where the spin-flip excitations involving the $e_g$ states were interpreted as the origin of anisotropic SOC, which was responsible for the Dzyaloshinskii-Moriya (DM) interaction. Although structurally Fe$_2$O$_3$ and CrI$_3$ are entirely different systems, one cannot neglect the fact that similar MDs in $dd$ excitation are observed in these two systems, in addition to the coexistence of in-plane FM and AFM interactions. Therefore it may be worthwhile to consider the possibility that MD in $dd$ excitation in CrI$_3$ could be due to spin-flip excitation. DM interaction may also be present in CrI$_3$, which is believed to be the origin of the spin gap at the Dirac point of spin-wave (SW) excitation spectra, observed in inelastic neutron scattering experiments [33]. Several calculations [20,33,34] showed that in the absence of DM interaction the origin of the spin gap cannot be explained, and the system would eventually be similar to a spin analog of graphene (spin-gap-less at the Dirac point). There is, however, a continued debate on the strength of DM interaction. A realistic estimation of DM interaction strength, by Kvashnin et al. [20], as compared with the Heisenberg exchange interaction, underestimates the spin-gap value, although it fairly accurately predicts the splitting in the Cr 3$d$ orbital and corresponding spin-flip $dd$ excitations (1.45, 1.75, and 2.50 eV). Kitaev interaction [35] and electron correlation [36] may also contribute to the origin of the spin gap in SW excitation. Recently, we found possible evidence of a complex magnetic spin-liquid-like state at high pressures and very low temperatures in a CrI$_3$ single crystal [37]. However, further investigations are required to quantify the roles of DM and Kitaev interactions, associated with the SOC strength, in accurately estimating the magnitude of the spin-gap and spin-flip $dd$ excitations.

Importantly, we do not observe any significant peak shift or splitting between the RIXS spectra at 20 and 100 K. This demonstrates that the ferromagnetic superexchange interaction that is responsible for the interatomic exchange field is vanishingly small compared with the local exchange field that comes from intra-atomic exchange and correlation interaction among the interacting Cr 3$d$ orbitals of a given site. The latter interaction is driven by local exchange interactions of the order of eV, which is clearly much larger than estimates of interatomic exchange which is on the meV level. Figures 5(a) and 5(b) plot the experimental RIXS map near the Cr $L_3$ region for CL and CR polarization, respectively, at 20 K. RIXS spectra at the Cr $L_3$ resonance excitation for CL and CR are shown in Fig. 5(c) [replot of Fig. 4(b) for $+H$ at 20 K]. For comparison, we plot the corresponding calculated RIXS
FIG. 5. Experimental RIXS maps measured at 20 K with (a) CL and (b) CR polarized x rays incident grazing to the sample surface and a magnetic field along the +H direction. (c) Dichroic RIXS spectra at resonant excitation along the +H direction. Calculated RIXS maps with (d) CL and (e) CR polarized x rays. (f) Line spectra at 574.3 eV excitation energy obtained from (d) and (e).

map in Figs. 5(d) and 5(e) and the line spectra at resonance $L_3$ excitation in Fig. 5(f). In the calculations, we did not consider the self-absorption effects and the possibility of reflections from the sample surface, which might lead to discrepancies between the experimental and theoretical dichroic elastic peak signal. We also note that the present experimental resolution (around 180 meV at the Cr $L_3$ edge) prohibits the occurrence of well-resolved peaks particularly above 2.5 eV, although the integrated spectral features match well with the calculations. The dichroism and the energy loss peak positions in the calculated spectra, considering a SOC in the system, fairly agree with those of the experiment. A dichroic signal can only be recovered in these types of calculations if spin-orbit effects are included (see Fig. S1 of the Supplemental Material). Therefore SOC plays a significant role and may also be responsible for the DM interaction when considering the projected in-plane FM spins in the in-plane AFM matrix at a nominal in-plane magnetic field. However, further calculations are necessary to explicitly understand and quantify the effects of DM and Kitaev interactions, which are based on SOC, on the orbital excitations in CrI$_3$ and the related family of 2D materials.

IV. CONCLUSION

To summarize, we find that our experimental data are in good agreement with the spectra calculated using a theoretical method that allows for multiconfiguration effects, when spin-orbit correlations are included (see Supplemental Material for more details [22]). For the spectroscopic features we note that a theory on the multiconfigurational level reproduces experimental observations with high precision. The investigation presented here demonstrates that the electronic structure of bulk CrI$_3$ (and most likely of its 2D sister compounds) is complex in the sense that dynamical electron correlations are important, meaning that the electronic structure signals multiconfiguration effects. This is signaled from both the ground state (magnetic moment) and spectroscopic, excited state properties. The recorded RIXS spectra here reveal clearly resolved Cr 3d intraorbital $dd$ excitations that represent transitions between electronic levels heavily influenced by multiconfiguration effects. The MD observed in the $dd$ excitations could reflect spin flip and, in conjunction with SOC, could support DM interaction as well, imparting tremendous interest in this and similar 2D materials for future investigations.

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