

## Temperature-dependent magnetic anisotropy in the layered magnetic semiconductors CrI<sub>3</sub> and CrBr<sub>3</sub>

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**Abstract:** Chromium trihalides are layered and exfoliable semiconductors and exhibit unusual magnetic properties with a surprising temperature dependence of the magnetization. By analyzing the evolution of the magnetocrystalline anisotropy with temperature in chromium iodide, CrI<sub>3</sub>, we find it strongly changes from  $K_u = 300 \pm 50$  kJ/m<sup>3</sup> at 5 K to  $K_u = 43 \pm 7$  kJ/m<sup>3</sup> at 60 K, close to the Curie temperature. We draw a direct comparison to CrBr<sub>3</sub>, which serves as a reference, and where we find results consistent with literature. In particular, we show that the anisotropy change in the iodide compound is more than 3 times larger than in the bromide. We analyze this temperature dependence using a classical model showing that the anisotropy constant scales with the magnetization at any given temperature below the Curie temperature indicating that the temperature dependence can be explained by a dominant uniaxial anisotropy where this scaling results from local spin clusters having thermally induced magnetization directions that deviate from the overall magnetization.

## I. Introduction

The materials class of layered chromium (Cr) trihalides  $\text{CrX}_3$ , with X being a halogen ion of fluorine (F), chlorine (Cl), bromine (Br) or iodine (I) experiences a revived attention recently with numerous publications on their magnetic properties [1-7]. Due to their layered structure with weak van der Waals bonds between the layers, they belong to those materials where exfoliation is possible and the prospect of magnetic monolayer systems is a source of the renewed interest. Only recently, Huang et al. demonstrated that ferromagnetism is persisting down to a monolayer of chromium iodide, which marks a milestone in the research of two-dimensional magnetic materials [5]. In all  $\text{CrX}_3$  compounds, the magnetic properties are closely tied to the details of the crystalline structure. Bulk  $\text{CrX}_3$  crystals all exhibit a phase transition from monoclinic ( $\text{AlCl}_3$ -type, space group  $C2/m$ ) at elevated temperatures to trigonal ( $\text{BiI}_3$ -type, space group  $R\bar{3}$ ) at cryogenic temperatures [3,6,7]. For both cases, the layers extend in the  $ab$ -plane and are stacked along the  $c$ -axis. Hansen [8], Tsubokawa [9] and Dillon Jr. [10,11] have pioneered the work on magnetism in these compounds finding them to be one of the rare instances where superexchange leads to ferromagnetism instead of antiferromagnetism. With their unusual magnetic properties and the possibility of existing in monolayer form, chromium trihalides are promising materials for ultra-thin spintronics [12]. While there are a number of publications dealing with optical properties and resonance studies of these materials [13-15], much work has focused on the chemical synthesis. Now that the fabrication of high-quality  $\text{CrX}_3$  crystals has been established, a careful analysis of the magnetic properties can be attempted in order to address the remaining open questions [3,6], where one of these questions is the stabilization of ferroic order in layered two dimensional systems such as  $\text{CrX}_3$  [5]. The magnetocrystalline anisotropy is for the moment the most promising ingredient that can allow one to overcome the limitations set by the Mermin-Wagner theorem [16], which prohibits magnetic ordering in two or less dimensions and here, we focus in particular on the magnetic anisotropy of  $\text{CrI}_3$  and  $\text{CrBr}_3$ . Since the magnetic

properties of bulk samples serve as a basis for the understanding of magnetic phenomena in reduced dimensions, e.g. in few- or monolayer samples of  $\text{CrX}_3$ , a profound understanding of these compounds is evidently needed, especially considering that magnetic monolayers of  $\text{CrI}_3$  have now become accessible [5]. The bromide compound has been studied in some detail with Tsubokawa (to our knowledge) having been the first to quantify the uniaxial anisotropy constant by torque magnetometry [9] and Dillon Jr. has provided the temperature dependence, using ferromagnetic resonance FMR [10,17]. In comparison,  $\text{CrI}_3$  has not been studied thoroughly. For instance, to the best of our knowledge, Ref. [11] is the only experimental determination of the magnetocrystalline anisotropy constant in  $\text{CrI}_3$ . Moreover, even though the compound exhibits a surprising temperature dependence of the magnetization (see discussion below), so far only the saturation field has been discussed without probing the value of the anisotropy constant, which is the key property that governs the anisotropic magnetic behavior [3,8]. In order to facilitate the understanding of magnetism in  $\text{CrX}_3$  compounds, we demonstrate in this paper how the temperature dependence of the anisotropy in  $\text{CrI}_3$  can be ascertained, showing a strong decrease as a function of increasing temperature. We directly compare the results for  $\text{CrI}_3$  to those measured in  $\text{CrBr}_3$ , where we confirm the value and temperature dependence given in the literature and show that the same is much stronger in the iodide. To understand this temperature dependence of the anisotropy, we demonstrate that for both,  $\text{CrI}_3$  and  $\text{CrBr}_3$ , it is consistent with a scaling law. The critical exponent found then allows us using a classical model to explain the magnetization as a function of temperature as resulting from a dominant uniaxial anisotropy, with local spin-cluster directions deviating from the overall magnetization direction due to thermal activation.

## II. Methods

### A. Sample preparation

$\text{CrBr}_3$  and  $\text{CrI}_3$  were prepared directly from the elements in an evacuated quartz tube similar to reports from literature [9]. Great care was taken to ensure the spatial separation of the reactants within the ampoule prior to heating. The halogen was kept slightly above room temperature, allowing the vapors to react with the metal at the hot zone at 1023 K. In the presence of a small excess of halogen, crystals grew at the cool end of the ampoule in the natural temperature gradient due to a chemical transport reaction.

### B. Magnetometry

After preparing and loading a  $\text{CrI}_3$  crystal or a  $\text{CrBr}_3$  crystal into a SQUID magnetometer (QuantumDesign MPMS XL), we measure the projection of the magnetic moment along the (vertical)  $z$ -direction at variable temperatures. Additionally weighing the crystals using a laboratory balance (Kern ABT 220-5DM) allows for the calculation of mass and resulting magnetization  $M$ , which is the measured magnetic moment divided by the volume. The small size of the crystals implies a low mass (typical mass is in the low milligram regime), resulting in the largest contribution to the absolute measurement error of the magnetization. However, relative changes exhibit comparatively low errors allowing us to deduce robust conclusions for instance about the temperature dependence. Furthermore, we do not consider effects of demagnetization because the impact of possible effective magnetic surface charges is negligible due to the bulk nature of our samples and therefore shape anisotropy plays no role.

### C. Ab-initio calculations

Density functional theory is employed by means of the full-potential linearized augmented plane wave method of the WIEN2k code [18]. The generalized gradient approximation of Perdew, Burke, and Ernzerhof is used for the exchange-correlation potential. Brillouin zone integrations are performed on a  $8 \times 8 \times 8$  mesh. The atomic sphere sizes of Cr and I are set to 2.5 Bohr radii and the plane-wave cutoff parameters to  $R_{Kmax} = 8$  and  $G_{max} = 12$ . We use

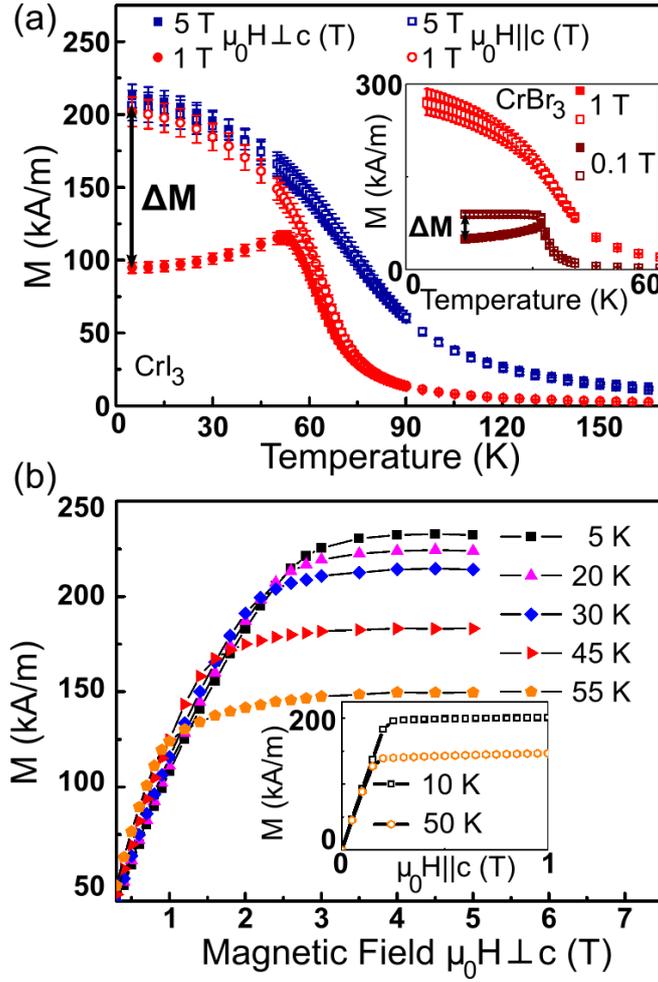
the experimental lattice constants ( $a = 6.867 \text{ \AA}$ ,  $c = 19.807 \text{ \AA}$ ) from Ref. [3] and relax the atomic coordinates until the atomic forces have declined below  $0.001 \text{ eV/\AA}$ . This procedure allows us to obtain the key structural information, namely the crystal structure that governs all electronic and magnetic properties.

### III. Results

In **Fig. 1 (a)** we present the magnetization  $M$  of  $\text{CrI}_3$  as a function of temperature for two different measurement configurations ( $H||c$  and  $H \perp c$ ) and at different field magnitudes of  $\mu_0 H = 1 \text{ T}$  and  $5 \text{ T}$ . Below the Curie temperature  $T_C$  (between  $61 \text{ K}$  [3,6] and  $68 \text{ K}$  [11] for  $\text{CrI}_3$ ) we observe with further decreasing temperature an increase of  $M$  which saturates eventually at liquid helium temperatures. When we apply a field of  $5 \text{ T}$ ,  $M$  saturates at a value of  $M_{S,5T} = 206.1 \pm 10.3 \text{ kA/m}$ , irrespective of whether  $H||c$  or  $H \perp c$ . This saturation magnetization corresponds to a magnetic moment of  $3.00 \pm 0.15 \mu_B/F.U.$ , where  $\mu_B$  denotes the Bohr magneton and  $F.U.$  stands for formula unit, in perfect agreement with the expected saturation moment in  $\text{CrI}_3$  [3,11]. However, for a smaller field of  $1 \text{ T}$ ,  $M$  saturates at the same value only in the  $H||c$  configuration, while in the  $H \perp c$  configuration,  $M$  remains at a significantly smaller saturation value of  $M_{S,1T}^{perp} = M_{S,1T}^{parallel} - \Delta M = 94.7 \pm 3.5 \text{ kA/m}$ . Generally, such a difference  $\Delta M$  can be attributed to the effect of the magnetocrystalline anisotropy and here, the  $c$ -axis is found to be the easy axis. If one looks at the temperature dependence of the magnetization in the plane, a surprising decrease of the magnetization below  $50 \text{ K}$  is observed, which cannot be explained by constant magnetocrystalline anisotropy and requires a more complex analysis as detailed below. In  $\text{CrBr}_3$ , the  $c$ -axis is also the easy axis, while the magnitude of the anisotropy is clearly lower (see inset of **Fig. 1 (a)**). As shown in the inset, for the bromide  $1 \text{ T}$  is enough to saturate  $M$  in both configurations at  $M_{S,1T} = 271 \pm 21 \text{ kA/m}$  (equivalent to  $3.09 \pm 0.24 \mu_B/F.U.$ ). Here, a discrepancy between  $M_S$  in the different configurations as explained above already for  $\text{CrI}_3$  occurs only at lower fields, for

example at 100 mT. Although 100 mT is also not sufficient to fully saturate the sample even in the configuration  $H||c$ , the behavior is qualitatively similar to the iodide compound. While  $M(T)$  monotonously increases with decreasing temperature for  $H||c$ , a decrease of  $M$  with decreasing temperature is found below 30 K for 0.1 T applied in the plane perpendicular to the  $c$ -axis.

In order to explain the unusual magnetization behavior discussed above which cannot be reproduced with the conventional assumption of a temperature independent anisotropy, we ascertain the impact of the magnetocrystalline anisotropy now in detail by analyzing magnetic hysteresis loops (**Fig. 1 (b)**) at different temperatures. In the inset of **Fig. 1 (b)**, we observe for  $H||c$  a sharp rise of the magnetic moment with increasing field, followed abruptly by a state of saturation when a certain magnetic field  $H_{sat}$  is reached. Increasing the temperature (while staying below  $T_C$ ) does not change this behavior, except that the saturation occurs at a lower magnetization. In **Fig. 1 (b)**, we show only the positive branches of the hysteresis loops, as within the measurement accuracy (800 A/m), it is neither possible to observe a remanent magnetization nor a coercive field  $H_C$ , thus setting an upper limit of  $H_C \leq 800$  A/m.



**Figure 1:** (a) Temperature dependence of the magnetization in  $\text{CrI}_3$ . (solid markers:  $H \perp c$ , open markers:  $H \parallel c$ ). For small fields ( $\leq 1$  T), the magnetocrystalline anisotropy leads to a strong difference of the effective saturation value for the different configurations and, below 50 K, to a decrease of  $M$  with decreasing temperature for  $H \perp c$  (see main text). The inset shows the same measurement for  $\text{CrBr}_3$ . (b) Positive branch of magnetic hysteresis loops for  $\text{CrI}_3$  between 5 K and 55 K in the  $H \perp c$  configuration. The saturation field  $H_{sat}$  is extracted from these curves and used to calculate the magnetocrystalline anisotropy. The inset shows for comparison the  $H \parallel c$  configuration at 10 K (black) and 50 K (pink). Error bars are omitted for the sake of clarity.

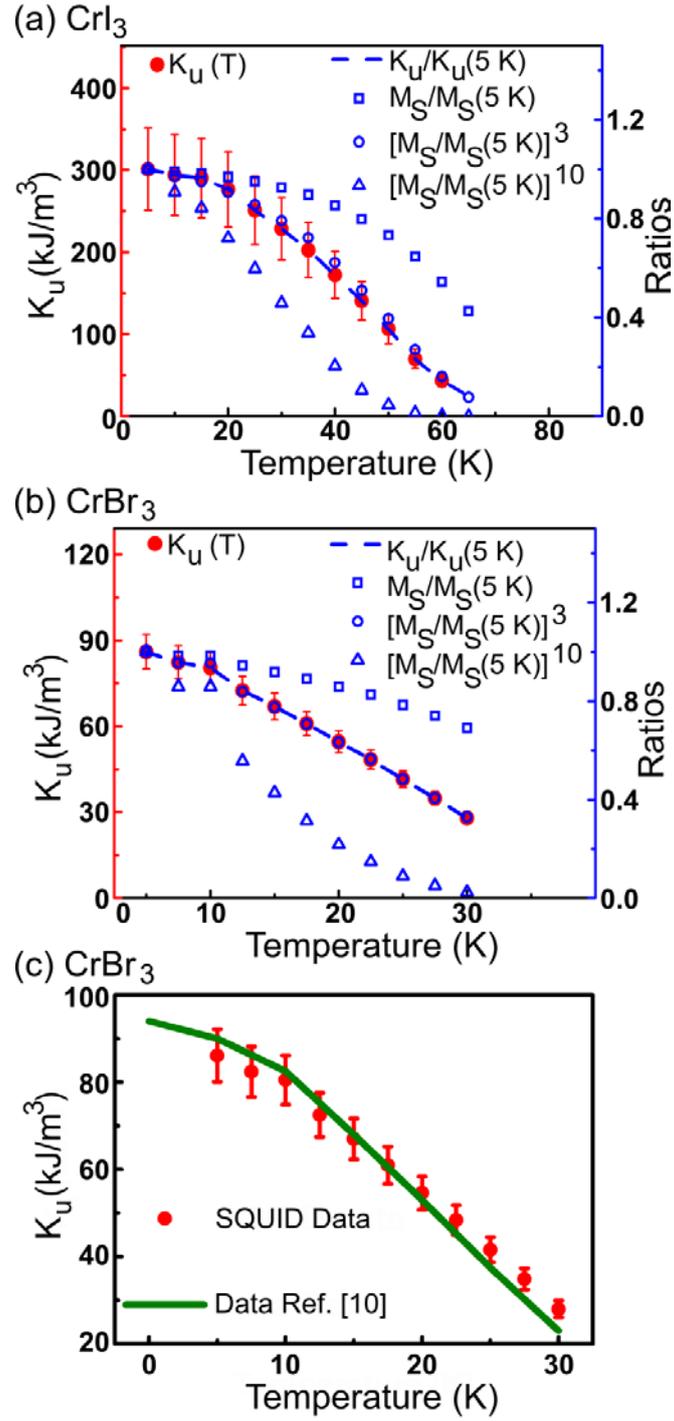
For  $\text{CrI}_3$  the hysteresis loops start to open slightly at 5 K, however, remanence and coercivity are still small at this temperature. Hence,  $\text{CrI}_3$  and  $\text{CrBr}_3$  are magnetic extraordinarily soft materials, where the magnetization follows the applied magnetic field direction virtually instantaneously during reversal and obviously not even pinning centers for domain walls in

the crystal structure lead to a remanent net magnetization at zero field. This observation is consistent with previous reports [3] and we suggest that one possible reason lies in the small interlayer exchange coupling of 1.8 K in CrI<sub>3</sub> and 0.5 K in CrBr<sub>3</sub> [19]. For CrI<sub>3</sub> this energy scale is 2.7 times lower than the lowest measurement temperature used here, leading to a vanishing interlayer coupling and thus to an immediate randomization of the magnetization between all layers, even though the magnetic coupling within individual layers is strong. In order to observe a remanent net magnetic moment and the resulting coercive field for the stacked system, the low-field regime should be revisited, ideally using remanence-free Helmholtz coils instead of superconducting ones and measured at millikelvin temperatures, which is however beyond the scope of this work and not possible with our current setup. Nevertheless, future investigations including thickness dependent measurements could shed light on this currently not completely understood issue.

We repeat the hysteresis loops in the  $H \perp c$  configuration, which show distinct differences as compared to the parallel configuration. First, the slope of  $M(H)$  in the low-field regime, i.e. the magnetic susceptibility  $\chi = dM/dH$ , is smaller. Furthermore, the curves measured at different temperatures clearly intersect, which is a consequence of a reduction in  $H_{sat}$  with increasing temperature. The magnetic field  $H_{sat}$  is a direct measure of the magnetocrystalline anisotropy, which contributes to the total micromagnetic energy density with a term  $E_A = K_u \sin^2(\theta - \varphi)$ . In this formula,  $K_u$  denotes the uniaxial anisotropy constant,  $\theta$  the direction of the preferred magnetization and  $\varphi$  the direction in which the magnetization points and when  $\theta - \varphi = 90^\circ$  (which is the case for  $H \perp c$ ) the effect of the magnetocrystalline anisotropy becomes maximal. If we consider the sample being fully magnetized above  $H_{sat}$ , the single domain state becomes a valid approximation, which can be described by the Stoner-Wohlfarth model [20]. Within this model, the anisotropy constant can be derived via

$$\frac{2K_u}{M_S} = \mu_0 H_{sat}, \quad (1)$$

with  $\mu_0$  denoting the vacuum permeability. This relation allows us to deduce  $K_u$  as a function of temperature below  $T_C$  for both  $\text{CrI}_3$  and  $\text{CrBr}_3$  (**Fig. 2**). However, one needs to obtain the saturation magnetization  $M_S$  and the saturation field  $H_{sat}$  in order to calculate  $K_u$  from **Eq. (2)**. Experimentally, we extract both quantities from the magnetization curves taken in the  $H \perp c$  configuration as follows: First, we consider a straight line parallel to the magnetic field axis fitted to the magnetization data above a magnetic field of 3 T, where the system is well above the saturation threshold. Second, we determine  $dM/dH$  in the low-field regime of the same curve. The intersection point of these two lines yields the saturation field as its abscissa and the saturation magnetization as its ordinate. In **Fig. 2 (c)**, we plot our measurement of the temperature dependence of  $\text{CrBr}_3$ 's anisotropy together with the temperature dependence extracted from Ref. [10]. For the whole temperature range, we find an excellent agreement, corroborating the validity of **Eq. (1)**, where the low- $T$  anisotropy amounts to  $K_u = 86 \pm 6 \text{ kJ/m}^3$ . For  $\text{CrI}_3$ , this is the first determination of a  $T$  dependence in  $K_u$ . As in  $\text{CrBr}_3$ , the anisotropy decreases towards  $T_C$ . At 5 K, the experimental value is  $K_u = 301 \pm 50 \text{ kJ/m}^3$ .



**Figure 2:** (a) and (b) Temperature dependence of  $K_u$  for  $\text{CrI}_3$  and  $\text{CrBr}_3$ . Left axes: For  $\text{CrI}_3$  as well as for  $\text{CrBr}_3$  a decrease of  $K_u$  is clearly visible as  $T$  is increased towards  $T_C$ . Right axes: The ratio  $K_u(T)/K_u(5\text{ K})$  plotted in comparison to the ratio  $[M_S(T)/M_S(5\text{ K})]^{n(n+1)/2}$ , where  $n = 1, 2$ , and  $4$ . For both compounds, the anisotropy scales to the third power of  $M_S(T)/M_S(5\text{ K})$ , in line with theoretical expectations for a predominant uniaxial anisotropy (leading to  $n = 2$ ). Error bars of the ratios are omitted for the sake of clarity. In (c), we show

the temperature dependence of  $K_u$  for  $\text{CrBr}_3$  in comparison with the temperature dependence for the anisotropy extracted from Ref. [10] where we find excellent agreement.

For  $\text{CrI}_3$ , where no published in-depth reports are available as a reference, we compare the experimental value of the anisotropy with numerical calculations based on density functional theory, which have been executed by the full-potential linearized augmented plane wave method of the WIEN2k code [18]. To understand the origin of the magnetic properties, we calculate as one of the key features of the system, the crystallographic structure. Based on the experimentally determined lattice constants ( $a = 6.867 \text{ \AA}$ ,  $c = 19.807 \text{ \AA}$ ) [3], we first model the crystal structure and check the stability. The relaxed atomic coordinates reproduce the experimental values with very good accuracy as shown in Table I.

	Experiment [3]	Theory
Cr	1/3, 2/3, 0.33299	1/3, 2/3, 0.33286
I	0.31667, 0.33453, 0.41230	0.32313, 0.33467, 0.41327

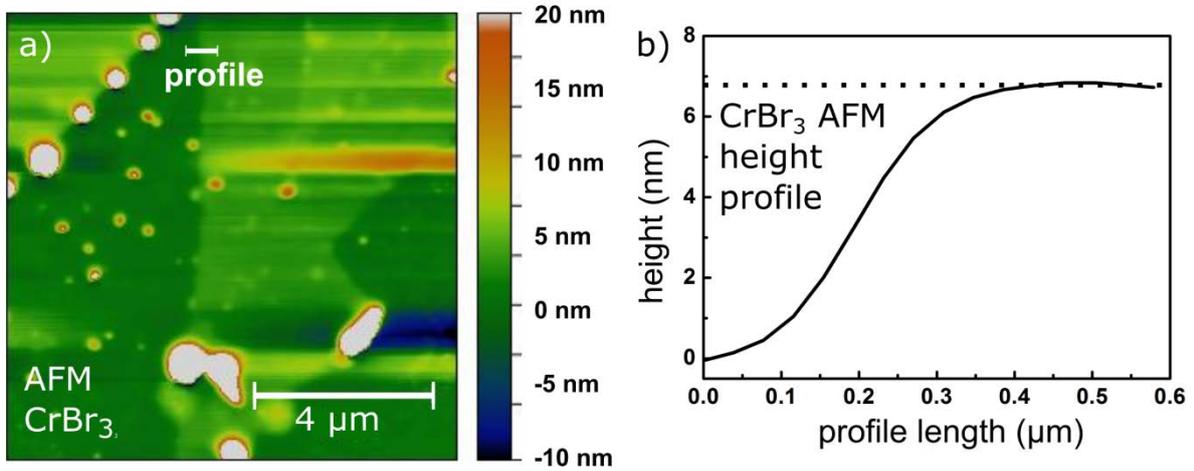
Table I. Experimental and theoretical atomic coordinates of Cr and I in  $\text{CrI}_3$ .

As the next step, we evaluate the total energies for magnetization along the  $a$ - and  $c$ -axes. The resulting energy difference of 0.5 meV divided by the unit cell volume yields the theoretically expected anisotropy constant. We find that the experimental value coincides with this calculated expectation of  $K_u^{theo} = 290 \text{ kJ/m}^3$  showing the validity of this approach. To analyze the temperature dependence of our anisotropy data, we consider that the magnetic anisotropy is a product of the interplay between the spontaneous magnetization and the crystal lattice. Often, the temperature dependence of the anisotropy is found to be stronger than that of the spontaneous magnetization. In a simple classical theory, anisotropy and magnetization are linked via the relation

$$\langle K^{(n)} \rangle \propto M_S^{n(n+1)/2}, \quad (2)$$

where  $\langle K^{(n)} \rangle$  is the anisotropy expectation value for the  $n^{\text{th}}$  power angular function [21,22]. In case of uniaxial anisotropy  $n = 2$  and in case of cubic anisotropy  $n = 4$ , leading to an exponent of 3 and 10, respectively. To check whether this model can describe our temperature dependence, we also show in **Fig. 2 (a)** and **(b)** the experimentally determined temperature dependences plotted as  $K_u/K_u(5\text{ K})$ ,  $M_S/M_S(5\text{ K})$ ,  $(M_S/M_S(5\text{ K}))^3$ , and  $(M_S/M_S(5\text{ K}))^{10}$ . The comparison clearly shows that the scaling  $\frac{K_u(T)}{K_{u,0}} = \left(\frac{M_S(T)}{M_{S,0}}\right)^3$  is the most appropriate fit. This is in line with the predominant uniaxial anisotropy found by our magnetometry measurements and thus explains the temperature dependence of the uniaxial anisotropy in both systems, CrI<sub>3</sub> and CrBr<sub>3</sub>. In the theory developed by Zener and Carr [21,22], the observed decrease of  $K_u$  with increasing temperature arises from solely from a large number of local spin clusters, fluctuating randomly around the macroscopic magnetization vector, activated by a non-zero thermal energy, while the anisotropy constants themselves are explicitly independent of temperature. These insights into the nature of the magnetocrystalline anisotropy are key for the understanding of ferromagnetism in CrX<sub>3</sub> compounds with a small number of layers. While in pure two-dimensional systems no magnetic order is expected [16], mechanical corrugations and magnetic anisotropy are possible pathways to establish magnetism in quasi-two-dimensional single CrX<sub>3</sub> layers nevertheless. While ferromagnetism has been shown to exist in a monolayer of CrI<sub>3</sub> [5], the magnetic properties of few- and monolayers of CrBr<sub>3</sub> and CrCl<sub>3</sub> are yet elusive. Like CrI<sub>3</sub>, both of these other two compounds can be subjected to mechanical exfoliation. For CrCl<sub>3</sub>, the exfoliation down to a double layer has been shown recently [7] and for CrBr<sub>3</sub> we conducted a set of cleavage experiments. For the procedure of mechanical exfoliation, we place the as-produced crystal on a piece of adhesive tape and exfoliate it at least 5 times (depending on the thickness of the initial crystal). In the last step, the tape with the crystals is pressed onto a heated substrate (120 °C) for 30 seconds. The resulting flakes with varying thicknesses on the

substrate surface can be located using an optical microscope. Flakes with low optical contrast are expected to be thin and we further investigate these flakes using atomic force microscopy (**Fig. 3 (a)**). We readily find stable thin crystals, where the one shown in **Fig. 3** has a thickness of approximately 6.7 nm corresponding to roughly 10 layers (**Fig. 3 (b)**). With these experiments, we demonstrate that also for the bromide the way towards exploration of magnetism in the two-dimensional limit is accessible and it is now equipped with a solid understanding of the magnetocrystalline anisotropy.



**Figure 3:** Atomic force microscopy image of a CrBr<sub>3</sub> flake on a GaAs substrate (a). The line profile (b) reveals a thickness of approximately 6.7 nm, which corresponds to roughly 10 layers.

#### IV. Conclusion

In summary, we carried out an in-depth analysis of the magnetocrystalline anisotropy in CrI<sub>3</sub> and in CrBr<sub>3</sub> revealing that the complex magnetization dependence on temperature results from temperature dependence of the magnetocrystalline anisotropy. While for CrBr<sub>3</sub>, our data's agreement with previous experiments is excellent, we find a more than three times stronger temperature evolution of  $K_u$  in CrI<sub>3</sub>. The experimentally obtained value at low temperature,  $K_u = 301 \pm 50 \text{ kJ/m}^3$ , is within the error identical to a theoretically derived

value,  $K_u^{theo} = 290 \text{ kJ/m}^3$ , for 0 K and agrees with a value determined by FMR. We find that the temperature dependence can be explained based on a classical model leading to a scaling of the temperature dependence with a critical exponent that results from a dominant uniaxial anisotropy combined with thermal activation effects that locally lead to magnetization deviating from the ensemble value.

Even with this understanding of the bulk compounds, our results show that the phenomenology of magnetism in  $\text{CrX}_3$  compounds still requires further exploration. For example, despite the strong ferromagnetic coupling within layers of  $\text{CrX}_3$ , global magnetic remanence is regularly absent. We explain this absence with the low interlayer coupling, calling for magnetometry in tailored set-ups, e.g. capable of reaching mK temperatures. To understand the bulk magnetic properties of chromium trihalide compounds is furthermore a key first step to using these compounds as single layers in spintronic applications, in particular as the anisotropy has a major influence for the stabilization of magnetic ordering in reduced dimensions.

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