Magnetostructural Transitions in a Frustrated Magnet at High Fields

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Ultrasound and magnetization studies of bond-frustrated $ZnCr_2S_4$ spinel are performed in static magnetic fields up to 18 T and in pulsed fields up to 62 T. At temperatures below the antiferromagnetic transition at $T_{N1} \approx 14$ K, the sound velocity as a function of the magnetic field reveals a sequence of steps followed by plateaus indicating a succession of crystallographic structures with constant stiffness. At the same time, the magnetization evolves continuously with a field up to full magnetic polarization without any plateaus in contrast to geometrically frustrated chromium oxide spinels. The observed high-field magnetostructural states are discussed within a H-T phase diagram taking into account the field and temperature evolution of three coexisting spin structures and subsequent lattice transformations induced by the magnetic field.

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Frustrated magnets with spinel structure AB_2X_4 manifest an intriguing behavior and unusual ground states, such as composite spins [1], spin dimerization [2,3], heavyfermion properties [4,5], spin-orbital liquid [6], and orbital glass [7,8], which originate from magnetic frustration but also from the intimate interplay of spin, charge, and orbital degrees of freedom and their coupling to the lattice. In the magnetic B-site Cr spinels with strong spin-phonon coupling, a novel type of structural transformation has been identified experimentally, the so-called spin Jahn-Teller effect [9–12]. In an octahedral crystal field the t_{2g} levels of the Cr^{3+} ions are half filled and the spin-orbit coupling is negligible. Therefore, the conventional Jahn-Teller scenario related to magnetic ions with an orbitally degenerate state is not applicable here, and the structural deformation is believed to be driven purely by spin ordering. The ground-state properties of frustrated magnets are characterized by a large degeneracy and are highly susceptible to external perturbations. An external magnetic field can change the balance between the competing interactions, and unusual phenomena, such as magnetization plateaus at half or fractional saturation, are observed [13,14].

In geometrically frustrated $A\text{Cr}_2X_4$ oxide (X=O) spinels, the Cr ions forming a pyrochlore lattice of corner-sharing tetrahedra are strongly coupled by direct antiferromagnetic (AFM) interactions of the order of 100–400 K. Emerging phenomena in geometrically frustrated oxides are dominated by local "tetrahedron" physics [15,16]. In sulfide (X=S) and selenide (X=Se) spinels, the direct AFM exchange is reduced due to the increasing distance between the magnetic ions and at the same time 90° ferromagnetic (FM) exchange becomes important.

Where FM and AFM exchanges are of comparable strength, the ground state again is strongly frustrated, a situation which has been named bond frustration [17].

In ZnCr₂S₄, the subject of the present study, competing FM and AFM interactions indeed are of equal strength resulting in a Curie-Weiss temperature close to zero [11]. Neutron-diffraction measurements [18–20] established two subsequent magnetic transitions in ZnCr₂S₄: the first one to an incommensurate helical AFM order at $T_{N1} \approx$ 14 K and the second one to a coexisting commensurate spin order at $T_{N2} \approx 7$ K. The helical state is characterized by a spin spiral with a propagation vector $k_1 \approx (0, 0, 0.787)$ along the crystallographic c direction and with the spins rotating in the a-b plane. This is also the ground-state structure of the bond-frustrated AFM ZnCr₂Se₄ with increased FM exchange as compared to ZnCr₂S₄ [12]. At temperatures below T_{N2} , the spiral phase coexists with two additional collinear spin structures with propagation vectors $k_2 \approx (0.5, 0.5, 0)$ and $k_3 \approx (1.0, 0.5, 0)$ [20]. These collinear ordering wave vectors resemble those of geometrically frustrated AFM ZnCr₂O₄, which exhibits composite spin structures of weakly interacting self-organized spin clusters [1,21]. The magnetic ground state of ZnCr₂S₄ can be regarded as a combination of spin orders known from geometrically frustrated ZnCr₂O₄ and bond-frustrated ZnCr₂Se₄. An external field favors the parallel spin alignment, and one can expect that the system passes through a sequence of exotic states when frustration is released via strong magnetoelastic coupling.

Several bulk properties of ZnCr₂S₄, such as specific heat and thermal expansion, exhibit significant anomalies at the magnetic transitions. Also, a pronounced splitting of the

phonon modes in the IR reflectivity spectra below the magnetic transitions was found, clearly indicative of broken symmetry [11]. These anomalies suggest structural transformations due to a strong spin-phonon coupling [17]. Recent high-resolution synchrotron x-ray powder diffraction measurements indeed revealed two subsequent structural transformations in ZnCr_2S_4 from a cubic $Fd\bar{3}m$ to a tetragonally distorted intermediate phase (space group $I4_1/amd$) below I_{N1} and a further transition into a low-temperature orthorhombic phase (space group Imma) below I_{N2} [20].

In this Letter, we report on ultrasound and magnetization studies on ZnCr₂S₄ single crystals performed in static (up to 18 T) and pulsed magnetic fields (up to 62 T). We explore the unique case of almost fully compensated AFM and FM exchange interactions in the spinel ZnCr₂S₄ with strong spin-lattice coupling and several coexisting magnetic structures with different spin arrangement. We expected that strong spin-phonon coupling leads to significant fingerprints in the temperature and magnetic-field dependence of sound waves. Ultrasound techniques are known to be highly sensitive probes for magnetoelastic interactions [22]. It has been demonstrated by Kino and Lüthi that the magnetic phase transition in ZnCr₂O₄, recognized now as the spin Jahn-Teller effect, is accompanied by pronounced anomalies in the temperature dependence of sound waves [23].

ZnCr₂S₄ single crystals were grown by chemical transport reactions. The magnetic susceptibility reveals a sharp peak at $T_{N1}=13.8~\rm K$ in good agreement with published data [11]. The measurements of the velocity and attenuation of longitudinal waves with wave vector **k** and polarization **u** parallel to the $\langle 001 \rangle$ axis (corresponding to the c_{11} acoustic mode for a cubic crystal) were performed for temperatures between 1.5 and 300 K and in static magnetic fields utilizing an experimental setup similar to that in Ref. [24] with a phase-sensitive detection technique based on a pulse-echo method. The measurements in pulsed fields with a rise time of 35 ms and a pulse duration of 150 ms were done in the range 1.5–20 K.

In Fig. 1, the relative changes of the sound velocity and attenuation for different static magnetic fields are presented as a function of temperature. In zero field, the sound velocity exhibits significant softening on decreasing the temperature below 60 K and a well-defined anomaly at $T_{N1} = 13.8$ K, in agreement with susceptibility and thermal-expansion data [11]. The sound attenuation raises sharply, approaching T_{N1} , and exhibits a well-defined peak. With an increasing magnetic field, the magnitude of the anomalies in the sound velocity and attenuation at T_{N1} are reduced and shifted to lower temperatures. We recall that at T_{N1} the sample transforms from the cubic paramagnetic state into the tetragonal helimagnetic phase [20]. The softening of the sound velocity probably results from strong spin fluctuations in the cooperative paramagnetic state

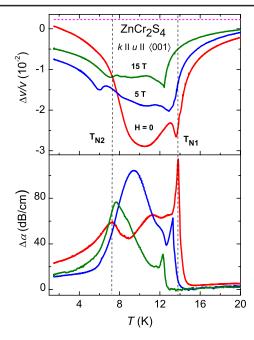


FIG. 1 (color online). Temperature dependencies of the relative change of the sound velocity $\Delta v/v_0$ (upper panel) and sound attenuation $\Delta \alpha$ (lower panel) for ZnCr₂S₄ measured in different static magnetic fields. The vertical dashed lines mark the magnetic phase transitions T_{N1} and T_{N2} in zero field. The horizontal dashed line shows the extrapolated undisturbed sound velocity estimated from a fit to the data above 60 K (see the text for details).

where the thermal energy is lower than the leading frustrated magnetic exchange. In contrast, at $T_{N2} = 7.3$ K, where $\rm ZnCr_2S_4$ transforms into the orthorhombic structure, only a smooth change of the sound velocity is found, and a broad peak in the attenuation appears. According to Ref. [18], the collinear structure evolves already at 12 K, which could explain the observation of two peaks in the attenuation below T_{N1} and of a weak anomaly in the sound velocity at T_{N2} . The anomaly in the attenuation at 11 K shifts to lower temperatures with increasing fields. Its magnitude first increases with the field up to 5 T but then decreases at higher fields.

Figure 2 documents the main results of the pulsed-field studies presenting the relative change of the sound velocity $\Delta v/v_0$, attenuation $\Delta \alpha$, and magnetization M as a function of the applied field for different temperatures. We notice rather significant changes of the sound velocity with field of the order of 3%–5% which prove the strong magnetoelastic coupling in ZnCr₂S₄. At 1.5 K, the sound velocity shows a nonmonotonic behavior with the field; $\Delta v/v_0$ first decreases and beyond 30 T increases again with increasing fields. Both the sound velocity and the attenuation exhibit four prominent anomalies at 7, 27, 38, and 44 T, suggesting changes in the spin state and structural phase transitions. The anomalies (labeled, respectively, from 4 to 1) are visible as minima or clear

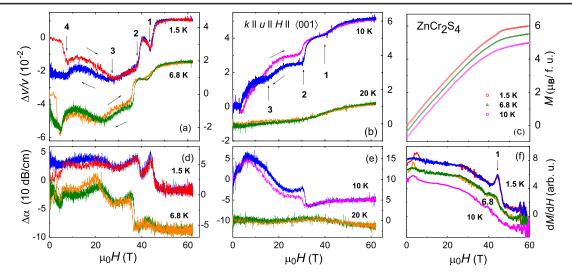


FIG. 2 (color online). Relative change of the sound velocity $\Delta v/v_0$ [(a) and (b)] and attenuation $\Delta \alpha$ [(d) and (e)] in ZnCr₂S₄ vs the magnetic field at 1.5 and 10 K (left scale) and at 6.8 and 20 K (right scale). The vertical arrows mark the magnetostructural anomalies labeled from 1 to 4. (c) Magnetization curves with the field aligned along the $\langle 001 \rangle$ axis and (f) derivatives of the magnetization dM/dH for 1.5, 6.8, and 10 K. For clarity, the curves are shifted along the vertical axis. Data for field sweep up and down are shown.

changes of slope in the sound velocity and as maxima in the attenuation. Anomalies 1–3 (at 1.5 K) in the sound velocity at fields above 27 T are free of hysteresis, whereas anomaly 4 shows a marked hysteresis on increasing and decreasing fields. Such hysteretic behavior indicates firstorder transformations induced by the magnetic field. With an increasing temperature, all anomalies in the sound velocity shift to lower fields. At 10 K, the sound velocity develops into distinct plateaus with steplike features at 16.5, 30, and 40 T. This signals abrupt changes of the lattice stiffness, probably induced by structural phase transitions, which are followed by plateaus with a given structure and, therefore, constant stiffness [Fig. 2(b)]. At the same time, between 7.5 and 40 T the changes of the magnetization M with the field are rather gradual [Fig. 2(c)] and occur with two different slopes below and above 27 T [Fig. 2(f)]. This indicates that the structural changes are not accompanied by significant changes in the spin structure. In Fig. 2(f), the field derivatives of the magnetization dM/dH for different temperatures are shown. The sharp maximum in dM/dH at 44 T (at 1.5 K) correlates well with anomaly 1 in $\Delta v/v_0$ and $\Delta \alpha$. Finally, at 47 T the full saturated polarization is achieved with the net ordered moment close to 6 μ_B . A well-defined change of slope in M appears just before reaching full polarization. With an increasing temperature, the maximum in dM/dH is shifted to lower fields and broadens considerably.

The magnetic-field dependence of the sound velocity and attenuation below 15 T is dominated by dynamic effects as revealed by measurements in static fields. A comparison of $\Delta v/v_0$ in pulsed and static fields shows a general agreement. However, below 7 T the variations of $\Delta v/v_0$ in static fields are much smaller than in pulsed fields which can be attributed to the relaxation dynamics

of domains reorientations that should be comparable to the high sweep rate in pulsed fields.

The observed temperature and field evolution of the anomalies in the sound characteristics detected in both static and pulsed fields is summarized in a tentative phase diagram in Fig. 3. We interpret it within the scenario which considers the interplay of different magnetic phases with structural transformations. In zero field at T = 1.5 K (phase V), two commensurate collinear $(k_2 + k_3)$ spin structures coexist with an incommensurate helical (k_1)

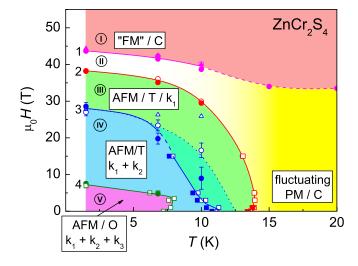


FIG. 3 (color online). *H-T* phase diagram of ZnCr₂S₄. Circles correspond to ultrasound data for pulsed fields, squares for static fields, up triangles to magnetization data for pulsed fields, and down triangles for static fields. Open symbols are for up sweeps in the field (temperature), closed symbols for down sweeps. Anomalies 1–4 and different magnetostructural phases I–V are described in the text.

spin structure and the crystal structure is orthorhombic [20]. At fields above 7.5 T (phase IV), the commensurate structure (k_3) becomes suppressed, but the second commensurate structure (k_2) survives and coexists with the helical spin structure (k_1) . At the same time, the suppression of the splitting of the lowest phonon by this field strength observed in the IR experiments [11,17] indicates the change of the lattice symmetry, from orthorhombic to tetragonal, which identifies the origin of anomaly 4 in the ultrasound data.

The transition from phase IV into phase III can be traced by the anomaly in the attenuation (Fig. 1, static data) and by anomaly 3 in the pulsed-field scans in Fig. 2. Above 7 K in the static fields this transition is clearly defined, whereas in the pulsed fields the boundary between phases IV and III is much broader probably due to relaxation effects. But we cannot exclude an intermediate phase in between phases III and IV for temperatures 7 K < T < 12 K. Phase III corresponds to a tetragonal phase with the Cr spins forming a spiral, such as in ZnCr₂Se₄.

The best defined features in the sound velocity and in the attenuation in both temperature and field dependencies are reflected by anomaly 2. Its steplike shape is only slightly affected by temperature, indicating a well-defined phase transformation. Since the magnetization shows only a gradual change in this temperature and field range, and taking into account neutron-diffraction and x-ray data [18–20], we associate anomaly 2 with the second structural transition from the tetragonal phase III to the cubic phase II. It is also important to note that the magnitude of the total change of $\Delta v/v_0$ by the magnetic field is close to that expected for the full recovery of the cubic state estimated from a fit to the experimental data in the true paramagnetic state (above 60 K) shown by the dashed line in Fig. 1, by using an anharmonic approximation according to Ref. [25]. We suppose that at zero temperature in fields close to 40 T the AFM spin spiral arrangement becomes fully suppressed and the system enters into a strongly polarized cubic paramagnetic state because no additional phase boundary to the paramagnetic phase above T_{N1} is evidenced in the ultrasound data. Anomaly 1 in the ultrasound and magnetization data probably has a purely magnetic origin. Finally, phase I corresponds to a state with full polarization induced by the magnetic field.

In conclusion, ultrasound and magnetization studies in magnetic fields up to 62 T of bond-frustrated $ZnCr_2S_4$ with strong magnetoelastic coupling revealed a sequence of magnetostructural states. We evidenced novel effects, namely, plateaus in the sound velocity on the way towards the recovery of the lattice symmetry in the polarized state which are ascribed to different crystallographic phases

with constant stiffness. In contrast to geometrically frustrated antiferromagnets $A\mathrm{Cr_2O_4}$ ($A=\mathrm{Zn}$, Cd, Hg), which reveal magnetization plateaus accompanied by lattice distortions [13,14,26–29], the magnetization of bond-frustrated $\mathrm{ZnCr_2S_4}$ evolves continuously without any anomalies up to full polarization. Our study provides a new insight into the physics of bond-frustrated spinels which is clearly distinct from that of the geometrically frustrated oxides. The origin of the observed intriguing effects, in particular, of anomalies 1–3, as well as of the coupling mechanism of different magnetic structures to lattice strain which generates crystal structures of different symmetry, is yet to be clarified and demands further experimental and theoretical studies.

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