Suppression of ferromagnetism in CeSi_{1.81} under temperature and pressure

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We have studied the pressure dependence of the magnetization of single crystalline $CeSi_{1.81}$. At ambient pressure, ferromagnetism develops below $T_c = 9.5$ K. Below \sim 5 K an additional shoulder in low-field hysteresis loops and a metamagnetic crossover around 4 T suggest the appearance of an additional magnetic modulation to the ferromagnetic state. The suppression of the magnetic order in $\text{CeSi}_{1.81}$ as function of temperature at ambient pressure and as function of pressure at low temperature are in remarkable qualitative agreement. The continuous suppression of the ordered moment at $p_c \approx 13.1$ kbar suggests the existence of a ferromagnetic quantum critical point in this material.

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I. INTRODUCTION

Quantum phase transitions have recently attracted great scientific interest. In contrast to phase transitions at finite temperatures where thermal order-parameter fluctuations are dominant, quantum phase transitions, e.g., phase transitions at *T*= 0 controlled by a non-thermal parameter like pressure or magnetic field, are driven by quantum fluctuations. Magnetic quantum phase transitions involving the formation of a spin density wave out of an itinerant electron system have been studied theoretically for quite a long time.¹⁻³ However, experiments on intermetallic Ce and Yb compounds undergoing an antiferromagnetic quantum phase transition defy a simple explanation, notably the systems $CeCu_{6-x}Au_x,^{4,5}$ and $YbRh_2(Si_{1-x}Ge_x)_2$,^{6,7} which have been studied extensively. The ferromagnetic to paramagnetic quantum phase transition of itinerant electron systems is considered to be the simplest quantum phase transition of the conduction electrons in metals. To date all pure materials in which ferromagnetic quantum phase transitions have been studied, e.g., UGe_{2} , ZrZn_2 , ZrZn_2 and $Ni₃Al$ (Ref. 10) display a behavior that can be well explained in terms of first-order transition. On the theoretical side, several mechanisms have been identified that preempt ferromagnetic quantum criticality by a first-order transition.¹¹

The search for ferromagnetic quantum phase transitions focused so far on transition-metal compounds like $MnSi$, 12 $ZrZn_2$,⁹ and Ni₃Al.¹⁰ Among *f*-electron materials, in particular the ferromagnetic superconductors UGe_2 ,⁸ URhGe,¹³ and $UIr,$ ¹⁴ have attracted interest. In Ce-based ferromagnets, the zero-temperature phase transition from a ferromagnetic to an antiferromagnetic state has been studied in $CeRu₂Ge₂$.¹⁵ For CePt,¹⁶ the ferromagnetic quantum phase transition has been studied. However, no measurements of the magnetization being the order parameter were presented, leaving many questions unanswered.

In CeSi_x, ferromagnetism, antiferromagnetism and paramagnetism have been reported to exist within a wide homogeneity range. Brauer and Haag¹⁷ reported that stoichiometric CeSi₂ crystallizes in the α -ThSi₂ structure. The wide homogeneity range $1.78 < x < 2$ that makes $CeSi_x$ of interest in our study was first noticed by Ruggiero and Olcese.¹⁸

Yashima *et al.*¹⁹ inferred from the specific heat $C(T)$ and the magnetic susceptibility $\chi(T)$ of CeSi₂ and CeGe₂ that CeSi₂ is a nonmagnetic intermediate-valent system with a metallic Fermi-liquid ground state down to 100 mK. In contrast, CeGe₂ displays a magnetic transition at T_C =7 K, where the susceptibility suggested ferromagnetic order. Electron diffraction gave evidence for a superlattice structure in $CeGe₂$, while no such feature was observed in CeSi₂.

Further studies^{20,21} of CeSi_x for $x=1.7, 1.8, 1.85, 1.9, 2$ suggested an even wider homogeneity range. At the same time, a ferromagnetic ground state was observed below $T_c \approx 10 \text{ K}$ for $x=1.8$ and a nonmagnetic intermediatevalence state for $1.85 \le x \le 2$. The spontaneous ferromagnetic moment of $\mu_S \approx 0.3 \mu_B$ (Ce atom⁻¹) is strongly reduced by comparison to the free Ce^{3+} moment of μ_{free} $= 2.54 \mu_B$ (Ce atom⁻¹), which is attributed to Kondo screening in a lattice. Hence, CeSi_x has been referred to as a ferromagnetic dense Kondo system.

The magnetic-nonmagnetic boundary as function of Si content was investigated in various studies. Measurements of $C(T)$ and $\chi(T)$ (Ref. 22) indicated a divergent Sommerfeld constant $\gamma = C_V/T$ in the Si concentration range $1.8 < x < 1.85$. Sato *et al.*²³ reported measurements of the electrical resistivity and the paramagnetic susceptibility for the *a* and *c* axis of a $CeSi_{1.86}$ single crystal, i.e., near the ferromagnetic instability. The lattice parameters of the crystal grown by a floating zone technique were $a = 4.182$ A and *c*= 13.85 Å. Strongly anisotropic behavior is observed in $\rho(T)$ and $\chi(T)$. At low temperatures both quantities show a *T*² dependence that indicates Fermi-liquid behavior. Based on the thermal expansion and magnetic susceptibility of two CeSi_x single crystals $(x=1.70 \text{ and } x=1.86)$ Sato *et al.*²⁴ suggested a Γ_7 crystal-field ground state. Measurements of the magnetization established that the *c* axis is the magnetically hard axis. For the easy axis the spontaneous magnetization at *T*=4.2 K is given by $\mu_s \approx 0.44 \mu_B$ (Ce atom⁻¹). At magnetic fields H_{c1} =30 kOe and H_{c2} =47 kOe anomalous behavior was observed, where measurements of the magneto-caloric effect for magnetic field along the *a* axis was interpreted as providing a weak antiferromagnetic modulation. Later Sato *et al.*²⁵ also reported measurements of the specific heat

 $C_V(T)$, electrical resistivity $\rho(T)$ and magnetic susceptibility $\chi(T)$ of two CeSi_x single crystals grown by an inductive floating zone method with the same Si concentrations. To determine the Si content, the specific heat was compared with results for polycrystalline samples. The analysis of $\chi(T)$ conjectured the ground state to be a Γ_7 -like doublet, as also suspected earlier;²⁴ $\rho(T)$ was found to exhibit a broad maximum indicating the onset of coherence.

Shaheen and Schilling²⁶ reported a transition from the tetragonal α -ThSi₂ structure to an orthorhombic α -GdSi₂ phase for low Si concentrations $x \le 1.84$. For $x \le 1.84$ ferromagnetic order was observed with evidence for further magnetic components. At ambient and high pressure the spontaneous magnetic moment was found to be strongly reduced as compared to the free Ce^{3+} moment. This reduction was attributed to strong Kondo screening. Precision measurements of $\rho(T)$ on polycrystalline CeSi_x samples in the range 1.6 $\lt x \lt 1.9$ were also made by Lee *et al.*²⁷

In neutron scattering experiments on a single crystal with $x=1.8$ ($T_C=13.4$ K), a spin-wave-like response was observed for $T < T_c$ over the entire Brillouin zone with abnormally large linewidths both above and below T_c ²⁸ This suggests that the magnetic excitations are unstable at low *T* because of the Kondo effect.

Further measurements of $\rho(T)$ and $\chi(T)$ on single crystals indicated strong anisotropies and confirmed the (ab) -plane as easy magnetic plane.²⁹ For $x=1.71$ magnetic order was observed below T_C = 12.5 K, while no magnetic order could be determined in a sample with $x = 1.86$. In the ordered system a trend toward Kondo behavior is suggested because the magnetic moment $\mu_s \approx 0.47 \mu_B$ (Ce atom⁻¹) is strongly reduced compared to the free Ce moment $\mu_{\text{free}} = 2.54 \ \mu_B(\text{Ce atom})^{-1}$.

The influence of annealing $CeSi_x$ in the range of $1.7 < x$ 1.84 was reported by Shaheen and Mendoza.³⁰ The ac susceptibility $\chi_{ac}(T)$ in the range 10–15 K shows, depending on annealing conditions, either a single peak or multiple peaks, which are attributed to multiple magnetic transitions. This suggests that the magnetism of this system is not that of a simple ferromagnet, but has been ascribed to two different types of magnetic (ferromagnetic and antiferromagnetic) order. However, it is also suggested that a nonuniform distribution of Si-vacancies can lead to this complex behavior.

Although $CeSi_x$ has been considered as a system with broad homogeneity range, recently Souptel *et al.*³¹ successfully grew comparatively large single crystals of high perfection within the narrow region $1.81 \le x \le 1.82$ only, using float zoning with an image furnace. For the optically floatzoned samples the structural phase transition is found at $x=1.85$. This is exactly the Si concentration of the paramagnetic to ferromagnetic phase transition, thus clarifying the relation between structure and magnetic order.

In light of these new results, we have revisited the question of the ferromagnetic to paramagnetic transition in CeSi*^x* in a detailed study of the magnetic properties of $CeSi_{1.81}$ as function of pressure in order to compare the suppression of ferromagnetism as function of temperature with the suppression as function of pressure. The magnetic anisotropy is associated in an obvious manner to the structural anisotropy. Both anisotropies are rather strong, in particular the magnetic anisotropy field is very large. We measured the magnetization only along the *a* axis of the single crystal and hence cannot draw a conclusion about the anisotropy within the easy plane. But we feel safe to assume that the applied pressures do not tilt the easy axis away from the (ab) -plane since it is unlikely that the underlying orbital orientation of the Ce atoms, which drives the magnetic anisotropy, will change for the very weak changes of lattice constants imposed by the pressures applied in our study. A direction within the easy magnetic plane was therefore chosen to obtain information of the pressure dependence of magnetic properties.

II. EXPERIMENTAL TECHNIQUES

Single crystals of $CeSi_{1.81}$ were grown at IFW Dresden with an image furnace as described previously.³¹ Cylindrical feed rods were prepared from high-purity Ce $(>\!\!>\!\!99.9\%)$ and Si (>99.99%) on a water-cooled copper hearth. The Si content of the resulting single crystals was determined from the unit-cell volume as measured by powder *x*-ray diffraction and compared with standards of known concentration. The crystal structure for $x=1.82$ was carefully studied by fourcircle *x*-ray diffraction, and an incommensurate superstructure was suggested to occur.³⁷ Neutron scattering experiments on the same samples do not confirm these findings, suggesting that the superstructure is a surface property.³⁸ Closer inspection by Laue x-ray diffraction of the single crystal studied in the present investigation revealed a tiny amount of twinning (<2) ^o on parts of the sample. Samples were cut with a diamond wire saw from the ingot. For our high-pressure magnetization study we used a rectangular sample cut along the *a* axis. The lattice parameters for *x* $= 1.81$ are $a = 4.173$ Å, $b = 4.181$ Å, and $c = 13.846$ Å.

The magnetization was measured with a vibrating sample magnetometer (VSM) at temperatures down to 1.5 K under magnetic field *B* up to 12 T at the University of Karlsruhe. The magnetization as function of pressure *p* up to 17 kbar was measured with a bespoke nonmagnetic Cu:Be clamp cell. The empty pressure cell was measured separately and the signal corrected accordingly. The pressure transmitting medium was a methanol:ethanol mixture of 4:1 volume ratio. The pressure was determined from the superconducting transition of Sn, after carefully demagnetizing the superconducting solenoid of the VSM.

III. RESULTS

The presentation of the results of our high-pressure magnetization study proceeds as follows. We first focus on the effect of temperature at ambient pressure. Here the magnetic order "melts" with increasing temperature. Next we present the suppression of magnetic order under high pressure. This addresses the "melting" of magnetic order as function of a nonthermal control parameter. Finally, we present the temperature dependence at high pressure, i.e., changes of thermally melting magnetic order at various pressures.

A. Ambient pressure

Shown in Fig. 1 are typical hysteresis loops in the range -0.5 T $\leq B \leq 0.5$ T for three characteristic temperatures

FIG. 1. Field dependence of the magnetization at typical temperatures $T = 1.6, 5.1, 10$ K. The magnetic field *B* as applied $\vert \vert \langle 100 \rangle$ was varied between -0.5 T \rightarrow 0.5 T \rightarrow −0.5 T. The widest hysteresis loop for $T=1.6$ K shows a small shoulder which vanishes at 5 K.

 $T=1.6, 5.1,$ and 10 K, i.e., well below T_C , near T_C and above T_c , respectively. The shape of the magnetic field dependence down to $T^* \approx 5$ K ($\approx T_C/2$) is characteristic of a purely ferromagnetic state with a small ordered moment. Above a certain coercive field the sample is in a single domain state. Below *T** a shoulder emerges as an additional feature that suggests an antiferromagnetic component on top of the otherwise ferromagnetic signal. We have extracted the ordered magnetic moment μ_s from magnetization loops by extrapolating $B \rightarrow 0$ from field values that are clearly above the coercive field. The presence of nonlinearities in the Arrott plots does not allow to extract T_C unambiguously. However, where possible an analysis with Arrott plots leads to the same results as a direct extrapolation of $M(B)$ for $B \to 0$. The ordered moment as function of temperature is shown in Fig. $2(a)$. For $T = 1.7$ K, the lowest temperature studied, \approx 0.18 μ_B (Ce atom⁻¹). With increasing temperature $\mu_S(T)$ passes over a shallow maximum at 4 K which is close to T^* =5 K; $\mu_S(T)$ vanishes continuously at T_C =9.5 K characteristic of a second-order phase transition.

The shallow maximum of $\mu_S(T)$ and the emergence of the additional shoulder in the magnetization loops signal the presence of an additional magnetic component on top of the otherwise ferromagnetic signal below *T** . However, the coercive field *H** as function of temperature does not exhibit any pronounced features as shown in Fig. 2(b). While the weak structure around 6 K is within the error bar, an influence of the onset of the change of the hysteresis loops (Fig. 1) cannot be excluded.

The Curie temperature T_c is normally defined directly from the order parameter, i.e., $\mu_s \rightarrow 0$. Alternatively, it may be defined from $\chi^{-1}(T_C) \equiv 0$. The dc susceptibility $\chi \approx M/B$, has been determined at $B=0.5$ T and $B=1$ T, while the differential susceptibility $\chi_d(T) = dM/dB$ has been extracted from low-field cycles above T_C for $B > 0$, corresponding to a measurement for $B \rightarrow 0$ (see Fig. 1 for an example). Shown in Fig. 3 is a comparison of the inverse dc susceptibility $\chi^{-1}(T)$ and the differential inverse susceptibility $\chi_d^{-1}(T)$ (full dots) for temperatures below *T*=25 K. $\chi_d^{-1}(T)$ drops continuously to zero for T_C =9.5 K in excellent agreement with the

FIG. 2. (a) Ordered magnetic moment μ_S as function of temperature *T* for *B* $\vert \vert$ $\langle 100 \rangle$. The data points were obtained by extrapolation of the hysteresis loops to $B=0$. The line connecting the points serves as guide to the eye. (b) Temperature dependence of the coercive field $H^*(T)$ at ambient pressure as extracted from magnetic hysteresis loops between −0.5 T→0.5 T→−0.5 T.

temperature below which μ_S emerges, whereas the inverse dc susceptibility reaches a finite constant value corresponding to a finite polarization in the external magnetic field as can be directly seen in Fig. 1.

Far above T_c the temperature dependence of χ is Curie-Weiss-type with an effective moment μ_{eff} with an effective moment μ_{eff}

FIG. 3. Temperature dependence of the inverse dc susceptibility $\chi^{-1}(T)$ for low fields (B=0.5, 1 T) applied along the *a* axis of the $CeSi_{1.81}$ single crystal. In addition, the inverse differential susceptibility $\chi_d^{-1}(T)$ (circles) extrapolated from hysteresis loops above T_C is displayed. Near T_c an additional downward curvature appears in both data sets. The inset shows the inverse dc susceptibility $\chi^{-1}(T)$ at high temperatures together with an extrapolation to $\chi^{-1}(T) = 0$.

FIG. 4. Magnetic field dependence of the magnetization $M(B)$ up to 12 T at different temperatures. The magnetic field was directed along the *a* axis $(B \parallel \langle 100 \rangle)$ of the CeSi_{1.81} single crystal. The magnetization decreases monotonically with increasing temperatures.

 $= 2.05 \mu_B$ (Ce atom⁻¹) as shown in the inset of Fig. 3. The slight reduction of the effective moment as compared with the free-ion value, $\mu_{\text{free}} = 2.54 \mu_B(\text{Ce atom}^{-1})$, may be related to the presence of crystal electric fields and the magnetic anisotropy. Alternatively, a phenomenological interpretation in terms of a spin fluctuation model³² may also be possible. However, the negative curvature of $\chi^{-1}(T)$ below ~100 K speaks against this possibility. In comparison to the effective moment, the ordered moment μ_s is reduced by over an order of magnitude. In intermetallic Ce compounds this is normally taken as a signature characteristic of the Kondo effect.

Above \sim 100 K the extrapolated Curie-Weiss temperature is given by $\Theta \approx -20$ K. This suggests effective antiferromagnetic interactions between the magnetic moments at high temperatures and may signal that the system is in fact a ferrimagnet (or "canted ferromagnet") as already inferred from the magnetization and saturation moment. For $T < 100$ K the inverse susceptibility curves downwards and differs from the Curie-Weiss dependence observed at high temperatures. The downward curvature below \sim 20 K may signal the effect of low-lying crystal electric fields. It may be a fortuitous coincidence that the inverse susceptibility above 20 K, when extrapolated to zero, vanishes around 5 K, i.e., the temperature *T** below which the magnetization loops display the additional shoulder.

We next turn to the magnetic properties at ambient pressure under high magnetic fields. Shown in Fig. 4 is the easyplane magnetization of single-crystal $CeSi_{1.81}$ as function of magnetic field up to 12 T for temperatures in the range $2.3 < T < 20$ K. Below $T \le 10$ K the magnetization displays an initial jump to a finite value for $B \rightarrow 0$ consistent with ferromagnetic order. With increasing field up to 12 T the magnetization continues to grow without apparent bound. At the lowest temperatures studied it reaches $\mu \approx 0.65 \mu_{\rm B}$ (Ce atom⁻¹), which is a small fraction of the effective Curie-Weiss moment and the free-ion value. While the reduced high-field magnetization may be the consequence of low-lying crystal electric fields and/or the Kondo effect, it is also taken as the typical signature of weak itinerant magnetism in transition metal compounds. As an un-

FIG. 5. Hysteresis loops between -0.5 T \rightarrow 0.5 T \rightarrow −0.5 T for different pressures at *T*= 2.3 K. The field was orientated along the *a* axis of the single crystal as for $p=0$. The small spikes and humps are remains of the Sn gauge. Applied pressures lie in the range $0 < p < 14.4$ kbar. The high-field magnetization decreases monotonically as the pressure is increased from 7.6 to 14.4 kbar. For 4.2 and 5.6 kbar M increases slightly [see also Fig. $6(b)$].

usual feature the magnetization displays an additional increase around $B \approx 4$ T. An inflection point of the magnetization as function of magnetic field is now commonly referred to as an indication of metamagnetism. Such a metamagnetic transition has been observed in several heavyfermion systems without magnetic oder, e.g., $CeRu₂Si₂³³$ or $CeCu₆.³⁴$ A possible interpretation for our data, as for the case of nonmagnetic $CeRu₂Si₂$ or $CeCu₆$, would be the weakening of the Kondo effect in high magnetic fields. The new feature here is that this transition exists "above" a ferromagnetic ground state. The inflection point vanishes for temperatures above $T^* \approx 5$ K. It will be a challenge to establish a possible link of the shoulder in the low-field hysteresis loops and the inflection point in the high-field magnetization. Indeed, the features in the hysteresis loop are strongly tied to the hump in $M(B)$ since both vanish at the same temperature or pressure (see below). A definite possibility is therefore a ferrimagnet arising from different Ce sublattices where a large temperature or pressure renders the two sites equivalent. Two different Ce lattice sites might arise from the nonstoichiometric Si concentration leading to differences in the atomic Ce environment.

B. High pressure

We now present the pressure dependence of the magnetization for magnetic field applied along the *a* axis, i.e. the (zero-pressure) easy magnetic axis. As pressure suppresses ferromagnetism we proceed in correspondence with the presentation of the data as function of temperature. Data were recorded for the same combination of temperature and field sweeps as studied at ambient pressure.

We begin by discussing typical magnetization loops in the range -0.5 T \rightarrow 0.5 T \rightarrow −0.5 T as measured for temperatures in the range $2.3 K < T < 15 K$ at various pressures. Data at *T*= 2.3 K for different pressures are shown in Fig. 5. At ambient pressure the hysteresis loop is characterized by the coercive field and the shoulder as already shown above.

FIG. 6. (a) Pressure dependence of the coercive field $H^*(p)$ extracted from hysteresis loops between −0.5 T→0.5 T→−0.5 T at $B(M=0)$ for different pressures at $T=2.3$ K. (b) Spontaneous magnetic moment μ_s as function of pressure at $T=2.3$ K and $T=1.7$ K. (c) Curie temperature T_C as function of pressure $T=2.3$ K. As for $p=0$ the field was directed along the *a* axis of the single crystal. Applied pressures lie in the range 4.2 kbar \leq *p* 14.4 kbar.

With increasing pressure the ordered magnetic moment and the size of the coercive field decrease monotonically and vanish for pressures above \sim 13 kbar [Fig. 6(a)]. Small sharp features in the $M(B)$ data and a tiny residual offset for $B\rightarrow 0$ at the highest pressures are attributed to spurious effects and show the level to which the signal of the empty pressure cell, notably the contribution of the superconducting piece of Sn, could be subtracted.

The evolution of the ordered magnetic moment at $T=2.3$ K as function of pressure, $\mu_S(p)$, is shown in Fig. 6(b). With increasing pressure, $\mu_S(p)$ increases slightly and displays a shallow maximum for $p^* \approx 5$ kbar before dropping off gradually and vanishing continuously above an extrapolated critical pressure $p_c = 13.1$ kbar. There is a remarkable qualitative analogy of the suppression of the ordered moment under temperature and pressure, respectively, that is not expected. The behavior suggests that temperature and pressure have the same qualitative effect on the magnetic properties. In particular, the similarity suggests that the critical point at

FIG. 7. Temperature dependence of the spontaneous moment μ_s for different pressures.

 $p=0$ as function of temperature $T\rightarrow T_c$ has a counterpart in a quantum critical point at $T=0$ for $p \rightarrow p_c$.

It is now interesting to establish if the suppression of the ordered magnetic moment as function of temperature is continuous at all pressures and whether the line $T_C(p)$ [Fig. 6(c)] is a line of second-order phase transitions. Shown in Fig. 7 is the spontaneous moment μ_S as function of temperature for various pressures in the range $0 < p < 12.9$ kbar. These data have been extrapolated directly from the hysteresis loops at various temperatures in the range 2.3 K \lt T \lt 15 K, where an extrapolation by means of Arrott plots led to the same results. For all pressures at which a temperature dependence could be extracted from our data the magnetization vanishes continuously at T_c . The shallow maximum of $\mu_S(T)$ seen at ambient pressure near $T^* \approx 5$ K disappears and is not present even at the lowest applied pressure of 4 kbar. When taken together this suggests that the line $T_c(p)$ in the temperaturepressure plane corresponds indeed to a line of second-order phase transitions.

An irregularity that we cannot account for are variations in the initial slope of $d\mu_S/dT$ just below T_C , e.g., at 10.4 and 12.9 kbar. In particular, the pressure dependence of the ordered moment shown in Fig. 7 does not represent the strict zero-temperature limit. The variations in initial slope may in principle lead to considerable qualitative differences of the pressure dependence of μ_s as compared to the temperature dependence shown in Fig. $2(a)$. To resolve this issue requires further measurements to lower temperatures, which are outside the technical possibilities of our apparatus.

The pressure dependence of the Curie temperature $T_C(p)$, shown in Fig. 6(c), has been determined from the data shown in Fig. 7 (filled squares) and slow temperature sweeps at a very low field of 0.01 T (open circles; raw data is not shown). These temperature sweeps only served to check for anomalies that mark the Curie point, but cannot be analyzed properly further. The solid line connecting the data points serves as guide to the eye. Down to 1.5 K, the lowest temperatures accessible with our apparatus, $T_c(p)$ is consistent with a quantum critical point at $p_c \approx 13.1$ kbar.

Characteristic features of the inverse susceptibility as function of temperature at ambient pressure (Fig. 3) are a Curie-Weiss dependence of large fluctuating moments at high temperatures and a strong downward curvature below

FIG. 8. Inverse dc susceptibility as function of temperature for different pressures (4.2 kbar $\leq p \leq 14.4$ kbar) at *B*=0.5 T. The fast increase near T_c does not change with p . The curves are shifted to lower temperatures with increasing pressure.

about 20 K. These features characterize the environment out of which the ferromagnetic ground state at low temperatures emerges. To interpret the suppression of the ordered moment as function of pressure, it appears therefore important to track the general features of the susceptibility as function of pressure as well.

Shown in Fig. 8 are dc susceptibility data recorded in a magnetic field of 0.1 T as function of temperature for various pressures. Data have only been recorded up to 30 K as the signal of the sample at higher *T* is too small in comparison to that of the empty pressure cell to be resolved. At high temperatures the slope of the susceptibility is essentially constant and the curves are shifted to the left (lower temperatures). Further, even for temperatures approaching T_C the curves are remarkably parallel. Deviations from being parallel for the lowest temperatures and highest pressures may be readily attributed to the polarizing effects of the applied magnetic field of 0.1 T.

Our susceptibility data are consistent with a smooth evolution of the size of the ordered magnetic moments and unchanged second-order behavior with increasing pressure. In particular, the effective Curie-Weiss moment and the important energy scale marked by the downturn below 20 K are essentially constant as function of pressure. Pressure appears to affect only the initial susceptibility, defined as $\chi(T)$ \rightarrow 0,*B* \rightarrow 0), which in present-day models of quantum phase transitions^{11,35} is used as the control parameter of the transition.

We finally turn to the pressure dependence of the highfield magnetization as measured at *T*= 2.3 K for fields up to 12 T. Typical data are shown in Fig. 9. With increasing pressure the curves are monotonically shifted to lower values and stay essentially parallel. Thus the susceptibility at the highest fields, i.e., the slope of $M(B)$, is roughly constant as function of pressure, signaling that the magnetization is highly unsaturated. The shoulder at $B^* \approx 4$ T which dominates $M(B)$ at *p*= 0 vanishes almost completely under a pressure of only 4.2 kbar and disappears altogether at higher pressures. The suppression of the metamagnetic transition with increasing pressure is in line with the interpretation given above. The increased pressure strengthens the Kondo effect and thereby

FIG. 9. Magnetization under pressure as function of the magnetic field at $T=2.3$ K. As for $p=0$ the field was directed along the *a* axis of the single crystal. Applied pressures lie in the range between $0 < p < 14.4$ kbar. The magnetization decreases monotonically with increasing pressure.

suppresses the metamagnetic transition. Despite a remarkable qualitative analogy with the variation of $M(B)$ at $p=0$ for increasing temperature, there is also an important difference. Because the transition at $p=0$ as function of temperature is second order, $M(B)$ at 7.5 K, i.e., just below T_C , is highly nonlinear (Fig. 4) and exhibits strong additional curvature over the entire magnetic field range up to 12 T, in particular below \sim 3 T, in comparison to the $M(B)$ curves at other temperatures. This nonlinearity is expected for secondorder behavior. In contrast, the curvature of $M(B)$ as function of pressure is much less affected by the proximity to p_c .

IV. DISCUSSION

The discontinuous suppression of ferromagnetism in pure ferromagnets^{8–10,12} has led to the suggestion that for pure compounds ferromagnetism is *always* suppressed discontinuously under pressure. The conclusion drawn from the work on $ZrZn₂$ is related to the peculiar structure of the density of states near the Fermi level, notably there being a pronounced maximum in the vicinity of E_F .^{9,36} The suppression of the magnetic order in weak transition-metal magnets as function of temperature at ambient pressure is essentially driven by a broadening of the density of states (DOS) due to thermal fluctuations. Although hydrostatic pressure also acts to broaden the DOS via increased *d*-electron overlap, this transition is discontinuous, i.e., temperature and pressure suppress the magnetic order in a different fashion.

In contrast, the suppression of the easy-axis magnetization of $CeSi_{1.81}$ under temperature and pressure shows a remarkable similarity. In particular, the continuous suppression of the ordered magnetic moment appears to suggest that $CeSi_{1.81}$ is the first example of a genuine ferromagnetic quantum critical point. However, four features in our magnetization data clearly establish that the magnetic state at ambient pressure and low temperatures is not that of a pure ferromagnet. These features are: (i) a broad maximum in the ordered magnetic moment at $T^* \approx 5$ K, (ii) the emergence of an additional shoulder in the low-field magnetization below $\sim T^*$,

(iii) the presence of metamagnetism at $B^* \approx 4$ T, and (iv) a broad maximum at a low pressure around $p^* \approx 5$ kbar. In this respect, the role of Si deficiencies has to be considered. Detailed four-circle *x*-ray diffraction studies of the crystallographic structure of samples from the same section of the ingot where the magnetization samples were cut has suggested an incommensurate superstructure of the Si vacancies.³⁷ As mentioned above, neutron scattering experiments do not confirm this superstructure and suggest that it may be a surface property.³⁸ Finally, as noted above we also find hints of twinning in Laue x-ray diffraction (<2) ^o on parts of the sample studied here.

Since all investigated ferromagnets exhibit a first-order transition when T_c approaches absolute zero, $\text{CeSi}_{1.81}$ appears at first sight as an exception. However, a possible firstorder transition in this material might be smeared due to the Si lattice disorder. The deviation from simple ferromagnetic behavior as described by the four features mentioned above may be due to an incommensurability of the Fermi surface introduced by the strain fields of an incommensurate structural modulation. It may alternatively be due to variations of the magnetic moments of the Ce atoms from a local-moment point of view, that are introduced by incommensurate variations of the crystalline environment surrounding the Ce sites, keeping in mind that the Kondo effect arises essentially from the coupling of Ce 4*f* moments to the conduction-electron "configuration" in the local environment around a given Ce atom. A possible metamagnetic transition induced by a magnetic field might also play a role. An additional effect may be a change of the local crystalline electric field. In either case, Fermi-surface effect or local-environment picture, the density of states near the Fermi level is expected to display a complex structure that is clearly outside present-day models of pure ferromagnets.

By the same token, the effect of pressure in $Cesi_{1.81}$ may be considerably more complex than just inducing a broadening of the conduction bands and associated suppression of the DOS near the Fermi level that leads to a reduction of T_c , as in weak transition-metal ferromagnets. Furthermore, the proximity of the nonmagnetic α -ThSi₂ phase has to be considered. However, the nonmagnetic α -ThSi₂ phase at large Si concentrations *x* has a larger volume than the magnetically ordered α -GdSi₂ phase found for smaller *x*. Therefore it is unlikely that pressure induces a structural change toward the nonmagnetic α -ThSi₂ phase. The incommensurate structural modulation also implies the presence of a modulation in the response to pressure and the buildup of inhomogeneous strains within the sample. These internal concomitant strains may lead to smearing of the ferromagnetism and a suppression that is not accompanied by nonlinearities in the magnetic-field dependence.

V. CONCLUSIONS

We have studied the easy-axis magnetization of $CeSi_{1.81}$ as function of temperature and pressure. The qualitative similarity of the temperature dependence at ambient pressure with the pressure dependence at low temperatures suggests that $CeSi_{1.81}$ may be a compound where a ferromagnetic quantum critical point may be induced by hydrostatic pressure. However, the presence of an additional magnetic modulation at low temperatures and ambient pressure and the observation of a possible incommensurate superstructure of the Si vacancies questions a straightforward interpretation in terms of quantum criticality in pure ferromagnets. Clearly, elastic neutron scattering, also under hydrostatic pressure, currently under way, 38 is needed to disentangle the possible antiferromagnetic component which may be operative in this system, and its relation to the structure properties.

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