サブゼミ 磁 性

テーマ「 近藤半導体 CeNiSn 」

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近藤格子化合物は、価数変動や重い電子的挙動などの特異な物性を示すため、発見以 来、盛んに研究されています。そのなかで Ce 化合物は磁気秩序の有無にかかわらず低温 では金属的に振る舞います。しかしながら、CeNiSn の電気抵抗は低温で半導体的に増加 を示します。CeNiSn は他にも比熱、磁化率、加圧下での抵抗等でも異常がみられるとい うような大変興味深い物質です。

今回のサブゼミではf電子系のこのような「近藤半導体 CeNiSn」についてお話していただく予定です。

内容的には講義のみとなりますが、この分野での現状を積極的に学習し、活発な質疑 応答での討論を期待します。

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KONDO SEMICONDUCTOR CeNiSn

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1. INTRODUCTION

Since the discovery of semiconducting behavior of SmB₆ in 1969,¹ the formation of a small energy gap in 4f-electron systems has been the subject of intensive studies. The presence of energy gap of several 10 K was found in valence-fluctuating (VF) compounds gold SmS, TmSe and YbB₁₂ with cubic structures.²⁻⁴ A simple picture of these systems is that the hybridization of the 4f electron states and the conduction band leads to the small gap at the Fermi level.⁵ However, the detailed mechanism of the gap formation remains unsettled. Recently, CeNiSn has been found to be the first example of a cerium compound showing the behavior of a small-gapped semiconductor.⁶ From the activation-type resistivity, the gap energy Eg was estimated to be 6 K. This compound crystallizes in an orthorhombic structure (Pn2₁a),⁷ which is closely related to the ε -TiNiSi type structure. Subsequently, similar gap formation has been found in Ce₃Pt₃Bi₄ and CeRhSb,^{8,9} where the values of Eg are 70 and 8 K, respectively. The latter crystallizes in the same type of structure as CeNiSn, whereas the former in the cubic Y₃Au₃Sb₄-type structure. These findings have renewed the interest in the problem of the insulating ground state of the Kondo lattice.^{10,11}

In this article, we review the experimental studies of the transport, magnetic and thermal properties of CeNiSn.¹²⁻¹⁷ The results obtained on single crystalline samples demonstrate that an anisotropic gap opens in the heavy-fermion bands as temperature is reduced below 6 K. Because of the small gap energy of several Kelvins, we expect strong effects of magnetic field and pressure on the gapped state. The strength of hybridization in this compound can be controlled by substituting Co, Cu and Pt for Ni in the nonmagnetic sublattice. The substituted samples serve as systems where one can examine the coherence effect on the gap formation. Furthermore, we compare the physical properties of CeNiSn with those of the

isostructural compound CePtSn, the latter of which is an antiferromagnetic Kondo compound with $T_N = 7.5 \text{ K.}^{18,19}$

2. SAMPLE PREPARATION AND CHARACTERIZATION

Polycrystalline samples were prepared from stoichiometric starting materials by arcmelting in a purified argon atmosphere. The samples were homogenized by annealing in quartz ampoules for 10 days at 1000°C. Single crystals were grown from Ames Laboratory Ce by a Czochralski technique in a triarc furnace or by a floating-zone method in an infrared mirror furnace. From metallographic examination and electron-probe microanalysis (EPMA) of the as-grown crystal, impurity phases of CeNi₂Sn₂ and cerium oxides were detected at the tail end and on the surface. As shown in Fig. 1, a line-shaped impurity phase of Ce₃Ni₄Sn₃ of 1~10 μ m width was detected even in the central part of the crystal. The volume fraction of this phase is less than about 1% of the sample, which is much smaller than that assumed by Kasuya²⁰ to explain the anomalous properties of CeNiSn based on a two-phase model. The impurity phase of Ce_{1.05}NiSn_{1.03} and Ce_{1.07}NiSn_{1.05}. However, any deviation from the 1-1-1 stoichiomerty larger than the resolution of about 0.3 at.%. was not detected by EPMA for the host phase in spite of the off-stoichiometric starting compositions.

The unit cell of CeNiSn consists of four formula units and hence contains an even number of valence electrons irrespective of the valence states of the Ce ions. We note here that in CeRhSb, which exhibits similar semiconducting behavior, the number of valence electrons is supposed to be same as in CeNiSn. In the orthorhombic structures of the two compounds, the Ce atoms form a zigzag chain along the a axis. Recently, band structure calculations on CeNiSn have been performed by Yanase and Harima²¹ using a selfconsistent LAPW method. According to their calculations, CeNiSn is a semimetal with a hole Fermi surface on the Δ axis and two electron Fermi surface centered at the X points. The overlap of the valence band with the conduction band is about 600 K. Since the 4f components are about 80 % both at the top of the valence band and the bottom of the conduction band, we expect large effect of strong correlation among 4f electrons on the transport and magnetic properties of this compound.





3. BASIC PROPERTIES OF CeNiSn

3.1 Magnetic Susceptibility

In Fig. 2, magnetic susceptibility $\chi(T)$ of a single crystalline sample of CeNiSn along the three principal axes is compared with that of CePtSn along the *a* axis labelled as $\chi_a(T)$.^{13,19} We note here that the relationship $\chi_a > \chi_b > \chi_c$ at low temperatures is the same for the two compounds. Hence, we may attribute the magnetic anisotropy in CeNiSn to the effect of the hybridization similar to that in CePtSn where Ce ions are almost trivalent. However, the temperature dependence is weaker in CeNiSn at low temperatures below 100 K, which is an indication of valence fluctuation in this compound. The VF character has been confirmed by inelastic neutron scattering experiments,²² by which no well-defined crystal-field (CF) excitations are observed. The maximum in $\chi_a(T)$ at 12 K for CeNiSn is not associated with a long-range magnetic order as will be discussed later. Instead, it can be attributed to the development of antiferromagnetic correlations among quasiparticles as observed in neutron scattering experiments.²³



Fig. 2. Magnetic susceptibility vs temperature for single-crystalline CeNiSn (H//a, H//b,H//c) and CePtSn (H//a).

3.2 Electrical Resistivity

Figure 3 represents the resistivity $\rho(T)$ of CeNiSn and CePtSn as a function of temperature.^{13,19} As temperature is decreased from 300 K, the $\rho(T)$ curves of CeNiSn initially show a quasi-logarithmic increase in all directions, whereas those of CePtSn are metallic. From a local maximum around 100 K for $\rho_a(T)$ and 60 K for $\rho_b(T)$, the Kondo temperature T_K of CeNiSn is estimated to be about 100 K for the degenerate J = 5/2 manifold. Another lnT dependence for $\rho_a(T)$ appears between 15 and 40 K, which is similar to that



Fig. 3. Electrical resistivity vs temperature for single-crystalline CeNiSn and CePtSn.

found in the magnetic part of the resistivity in CePtSn.¹⁹ The close similarity suggests the presence of another Kondo scale of about 30 K for the CF ground state in CeNiSn although the CF level scheme is not well defined.

A local maximum in $\rho_a(T)$ appears at the same temperature of 12 K at which $\chi_a(T)$ shows the peak. The decrease in both $\rho_a(T)$ and $\rho_b(T)$ below 12 K indicates that the system goes into the coherent scattering regime. With further decrease of temperature below 6 K, the resistivities turn to increase. An activation-type variation appears only in a small temperature range between 2.5 and 4.6 K. Nevertheless, the gap energy $E_g/k_B=T_g$ in the formula $\rho(T)$ = $\rho_0 \exp(E_g/2k_BT)$ was deduced as 1.0, 4.8 and 8.0 K for the *a*, *b* and *c* axes, respectively. Such a large difference in T_g implies an anisotropic gapping of the density of states on the Fermi surface. The value of T_g along the *c* axis is still one order of magnitude smaller than that reported for Ce₃Pt₃Bi₄. As shown in the inset of Fig. 3, $\rho_c(T)$ and $\rho_b(T)$ pass through a weak maximum at 0.4 and 0.6 K, respectively, and decrease with further decreasing temperature. A plausible explanation for this temperature dependence is that the gapping of the quasiparticle band is incomplete or there exist impurity bands in the gap.

3.3 Hall Effect

The temperature dependence of the Hall coefficient R_H for three configurations H//a (I//b), H//b (I//c) and H//c (I//b) is represented in Fig. 4 from Ref. 15. The three curves rise rapidly with decreasing temperature below 100 K and exhibit a positive peak around 9 K, followed by a precipitous drop to a negative value. The positive R_H peak of the size of 5×10^{-3} cm³/C is in common with those found in heavy-fermion compounds like CeAl₃ and CeRu₂Si₂.²⁴ Acording to the theory of Fert and Levy,²⁵ the strong temperature dependence of R_H with a positive peak arises from the contribution of the intrinsic skew scattering. Further, the extraordinary part of R_H is proportional to the product of $\chi(T)$ and magnetic resistivity $\rho_m(T)$. Between 100 and 10 K, R_H(H//a) approximately follows the product. Since the peak temperature of R_H in heavy-fermion systems is regarded as the



Fig. 4. Hall coefficient of CeNiSn as a function of temperature taken for three configuration; H//a (I//b), H//b (I//c) and H//c (I//b) (after ref. 15).

onset of coherence,²⁴ the above result indicates that the electronic state in CeNiSn gradually goes into a coherent scattering regime below 9 K. This temperature is somewhat lower than the maximal temperature in $\chi_a(T)$ and $\rho_a(T)$ at 12 K. The strong decrease in R_H below 5 K can be ascribed to the reduction of carrier density caused by the opening of the energy gap in the density of states. If we assume a single type of carriers, the concentration at 1.3 K is estimated to be 4.6×10^{-3} per formula unit from the data for H//c.

3.4 Specific Heat

The specific heat C of a single crystalline sample of CeNiSn is shown in Fig. 5 in a plot of C/T vs T² for T < 16 K. No appreciable anomaly exists near 12 K, and hence the peaking in both $\chi_a(T)$ and $\rho_a(T)$ at 12 K does not originate from a long-range magnetic order. The value of C/T decreases almost linearly with T² down to nearly 6 K and then suddenly diminishes. This temperature dependence is consistent with the opening of a gap below 6 K as inferred from the transport properties. Furthermore, the large value of C/T of 0.2 J/K²mol near 6 K indicates the development of a narrow band of heavy quasiparticles antecedent to the gap opening. Using the relation between γ and T_K derived for a single Kondo impurity, T_K = 0.68R/ γ (R is the gas constant),²⁶ T_K is estimated as 28 K. This temperature is within the lower temperature range where $\rho_a(T)$ increases as ln T, and thus this T_K can be regarded as the Kondo temperature for the CF ground state.

The magnetic contribution to the specific heat, C_m , was estimated by subtracting the data of LaNiSn from that of CeNiSn. As shown in the inset of Fig. 5, C_m/T reveals a pronounced maximum near 6.7 K. The magnetic entropy up to 20 K amounts to only half of Rln2. Between 1.5 and 5 K, C_m/T shows a linear variation, $C_m/T = \gamma + AT$, as opposed to the T² dependence of C_m/T in usual metallic systems. The observed temperature dependence is consistent with the renormalized density of states with a V-shaped gap near the Fermi level E_F, which was first proposed based on the results of NMR experiments.²⁷ The nuclear-spin lattice relaxation rate $1/T_1$ of ¹¹⁹Sn in CeNiSn was found to be proportional to T³ between 0.4 and 1.3 K. The proposed density of states is proportional to $|E-E_F|/E_g$, which yields a T³ dependence for $1/T_1$ and a T² dependence for C_m at low temperatures below $T_g = E_g/k_B$.



Fig. 5. Specific heat divided by temperature C/T vs T^2 for CeNiSn. The inset shows the magnetic contribution to the specific heat divided by temperature C_m/T vs T.

3.5 Magnetic Correlations Sudied by NMR and μ SR

Both NMR and μ SR techniques have been employed to study the magnetic correlations in CeNiSn at low temperatures. Kyogaku et al.²⁸ extended the temperature range of NMR experiments down to 80 mK. They observed a strong deviation of $1/T_1$ of ¹¹⁹Sn from the T³ behavior below 0.4 K. With further decrease of temperature below 0.13 K, $1/T_1$ decreases exponentially together with an increase in the line width. The results suggest that the pseudogap state becomes unstable below 0.4 K and a spin excitation gap is induced by the development of quasistatic magnetic correlations.

Krazer et al.²⁹ revealed from μ SR experiments that CeNiSn exhibits properties typical of a paramagnet moving towards magnetic order below 1 K. However, no transition into long-range order was observed down to 33 mK. The formation of extended spin correlation up to short-range order is deduced from the unusual dependence of muon spin relaxation rate and muon spin precession frequency on external field.

4. HIGH-MAGNETIC FIELD STUDIES

4.1 Magnetization

The field dependence of magnetization M(H) of single crystal CeNiSn at 1.3 K is represented in Fig. 6 from Ref. 16. The M(H) curve only along the *a* axis exhibits a weak metamagnetic-like transition near 13 T. It is more clearly seen in the derivative susceptibility dM/dH vs H. However, the increase in M associated with the transition is much smaller than that found in nonmagnetic heavy-fermion compounds like CeRu₂Si₂.³⁰ This fact suggests that the weak transition in CeNiSn is not due to the suppression of antiferromagnetic intersite interactions as found in CeRu₂Si₂ but due to the collapse of the pseudogap. Above 20 T, M_a(H) increases linearly with increasing field and attains 0.3 μ_B/Ce at 36 T. The size of the magnetization is only one fourth of that found in the isostructural, antiferromagnetic compound CePtSn, 1.2 μ_B/Ce .¹⁹ The small and linearly increasing moment in CeNiSn can be interpreted as a result of strong Kondo-type interaction persisting even after the pseudogap has collapsed. Assuming the effective moment of 0.3



Fig. 6. Magnetization of CeNiSn along the three principal axes at 1.3 K (after ref. 16).



Fig. 7. Magnetoresistance of CeNiSn at 1.4 K (after ref. 14).

 $\mu_{\rm B}/{\rm Ce}$, the magnetic energy at the transition field of 13 T corresponds to the thermal energy of 2.6 K, which is comparable to the gap energy estimated from the *a*-axis resistivity.

4.2 Magnetoresistance

Strong suppression of the energy gap in CeNiSn by application of magnetic field was first demonstrated by the magnetoresistance measurements on a polycrystalline sample.¹² The resistivity at 1.7 K was found to decrease from 730 to 200 $\mu\Omega$ cm as the field is raised to 24 T. The results obtained on single crystalline samples show very strong anisotropy.¹⁴ The normalized magnetoresistance $\Delta\rho(H)/\rho(0)$, where $\Delta\rho(H) = \rho(H)-\rho(0)$, at 1.4 K are presented in Figs. 7(a) and 7(b). At H = 15 T, the negative magnetoresistance for H//*a* attains -88%, which is much larger than for H//*b* and H//*c*. Thus, the energy gap is most sensitive to the magnetic field applied along the easy *a* axis. For H//a, the dependence of magnetoresistance on the current direction was further examined. In Fig. 7(b), the field dependence of $\Delta\rho(H)/\rho(0)$ for I//*c* is similar to that for I//*b*, whereas $\Delta\rho(H)/\rho(0)$ for I//*a* is almost constant up to 4 T and then gradually decreases with increasing field. These results suggest a strong anisotropic scattering mechanism under magnetic fields.

Shown in Fig. 8 is the temperature dependence of the resistivity along the three principal axes in fields of 0, 12 and 14 T. At H = 0 T, $\rho_a(T)$ exhibits a maximum near 11.4 K, which has been ascribed to the development of antiferromagnetic correlations. In a field of 14 T parallel to the *a* axis, this maximum is almost smeared out. The strong suppression of the upturn below 6 K is a result of the gap suppression by magnetic fields. A drastic effect occurs in the resistivity for the configuration I//b and H//a, which indicates metallic behavior and is in contrast to the semiconductor-like behavior for H//c. Furthermore, at temperatures below 4 K, it obeys a T² dependence with a coefficient of 1.0 $\mu\Omega$ cm/K² and a residual resistivity of 39 $\mu\Omega$ cm. The size of this coefficient is typical for moderately heavy fermion systems.





Fig. 8. Electrical resistivity vs temperature for CeNiSn in external fields 0, 12 and 14 T for electrical currents along the three principal axes (after ref. 14).

Fig. 9. Specific heat of CeNiSn plotted as C/T vs T in magnetic fields for (a) H//a and (b) H//c (after ref. 14).

4.3 Specific Heat in Magnetic Fields

The suppression of the energy gap by magnetic field was further studied by specific-heat measurements.¹⁴ Temperature variations of the specific heat of CeNiSn in magnetic fields parallel to the *a* and *c* axes are shown in Figs. 9(a) and 9(b), respectively. At H = 0 T, C/T is almost proportional to T between 0.3 and 0.8 K and the linear extrapolation to T= 0 K yields a γ value of 57 mJ/K²mol. This size of γ value seems to be too large to be ascribed to the contribution from impurity phases. Rather, it may be the contribution from the residual density of states at E_F in the pseudogap, as inferred from the saturation of the resistivity below 1 K. The origin of the upturn in C/T below 0.2 K is not clear yet.

When magnetic field is raised to 12 T, the value of C/T is strongly enhanced for H//a, whereas it is almost unchanged for H//c. For H//a, the field dependence of C/T was measured at 0.15, 0.42 and 0.76 K.¹⁴ The values of C/T at these temperatures stay constant for H < 4 T and then increase monotonically from about 70 to 125 mJ/K²mol. This large enhancement is consistent with the heavy-fermion behavior in $\rho_b(T)$ at 12 T in Fig. 8(b). These results support the idea that the density of states in the minimum of the V-shaped pseudogap is increased by application of magnetic field along the easy axis of

magnetization. In this anisotropic suppression of the pseudogap, a strong spin polarization of the renormalized band should play an important role.

5. HIGH-PRESSURE STUDIES

The effect of application of pressure on the gapped state in CeNiSn has been studied by Kurisu et al.^{31,32} from resistivity measurements. As a typical result, the *a*-axis resistivity is shown in Fig. 10. With increasing pressure, the upturn in $\rho_a(T)$ below 6 K is strongly suppressed, and the local maximum at 12 K shifts to higher temperatures. Above 12 kbar, the resistivity shows metallic behavior with a single maximum around 100 K. This maximum, which originates in the interplay of the Kondo effect and the CF effect, shifts also to higher temperatures. Generally, application of pressure on cerium-based Kondo-lattice compounds increases the degree of the 4f-ligand hybridization so that the renormalized band broadens.³³ As a result, the value of T_K is increased and hence the maximal temperature of the resistivity due to the Kondo effect is elevated. In fact, we find large shift of both the peaks in $\rho_a(T)$, which are originally at 12 and 65 K, respectively, to 30 and 180 K. The broadening of the renormalized band should also be responsible for the suppression of the gap at low temperatures. In other words, the gap formation in CeNiSn is



Fig. 10. Electrical resistivity vs $\ln T$ for singlecrystalline CeNiSn along the *a* axis under various applied hydrostatic pressures.

Fig. 11. Electrical resistivity vs temperature for CeNiSn at hydrostatic pressures 0, 18 and 20 kbar.



strongly suppressed by the increase of the degree of hybridization. We will see in the next section that the reduced hybridization by negative chemical pressure also suppresses the gap.

It is interesting to compare the effect of pressure on the resistivity with that of magnetic field. Comparing the resistivity curves in Figs. 8 and 11, we notice that the effect of application of 20 kbar is comparable with that of a magnetic field of 14 T applied along the *a* axis. However, a simple scaling is rather difficult because the negative magnetoresistance is significant only below 30 K, whereas the overall temperature dependence is changed by application of pressure.

6. EFFECT OF ALLOYING

Low-temperature properties of heavy-fermion compounds are generally very sensitive to a small amount of substitution.³⁴ We expect that the substitution of 3d element Co or Cu for the Ni atom in CeNiSn would change the number of conduction electrons. If the Ni atom is replaced by the 4d or 5d atom in the same column in the periodic table, i.e., Pd or Pt, then the unit cell volume would increase without changing the number of conduction electrons so much. Keeping this in mind, we have performed systematic studies of the substituted systems CeNi_{1-x}T_xSn with T= Co, Cu and Pt.

Figure 12 shows variations of the lattice parameters with x. The substitution of Co for Ni hardly changes the lattice parameters, whereas that of Cu leads to a linear increase in the three parameters. By the substitution of Pt, the *c* parameter increases significantly but the *a* parameter slightly decreases. For CeNi_{1-x}Cu_xSn, we previously reported that the energy gap disappears near x=0.1 and then a long-range antiferromagnetic order develops for $x \ge 0.13$.¹² Hereafter, we compare the magnetic and thermal properties of the Co and Pt substituted systems. Temperature dependences of the magnetic susceptibility $\chi(T)$ of



Fig. 12. Variations of lattice parameters of $\text{CeNi}_{1-x}T_xSn$ (T = Co, Cu and Pt).

-596 -



Fig. 13. Magnetic susceptibility vs temperature for field-aligned polycrystalline samples of (a) CeNi1-xCoxSn and (b) CeNi1-xPtxSn.

polycrystalline samples of $\text{CeNi}_{1-x}T_x$ Sn for T= Co and Pt are shown in Figs. 13 (a) and 13(b), respectively. The measurements were performed on powdered samples which were pre-aligned by application of a magnetic field of 14 T at 4.2 K.

The data of CeNiSn with a maximum around 12 K agree with those of $\chi(T)$ along the *a* axis of the single crystalline sample as was shown in Fig. 2. This agreement guarantees that the powders are aligned along the easy *a* axis of the magnetization. In Fig. 13(a), the values of $\chi(T)$ for T<100 K decrease as x is increased in CeNi_{1-x}Co_xSn. At higher Co concentration $x \ge 0.2$, a broad maximum appears around 100 K and the value of the paramagnetic Curie temperature estimated from the Curie-Weiss fitting exceeds -130 K. These features are the signals for the transition to the VF regime. The Pt substitution has an opposite effect on the system and leads to the Kondo regime. In Fig. 13(b), the low-temperature value of $\chi(T)$ increases with increasing x in CeNi_{1-x}Pt_xSn, resulting in the recovery of the Curie-Weiss behavior down to low temperatures. The paramagnetic Curie temperature about -70 K.

The trend toward either the VF or the Kondo regime is confirmed by the result of magnetization measurements at 4.2 K. As shown in Fig. 14(a), the slope of the magnetization curve decreases with increasing Co concentration. A downward curvature near 4 T for $x \ge 0.1$ may indicate the saturation of impurity contribution. All the magnetization curves of the Pt substituted samples are almost linear with field. The value of M(H) at H = 14.6 T increases linearly with increasing Pt concentration, whereas it decreases with Co concentration at almost the same rate of dM/dx = 0.42 μ_B .

As mentioned above, Co substitution for Ni in CeNiSn may decrease the number of conduction electrons. The Fermi level accordingly lowers toward the level of the unrenormalized 4f states, E_{f} . For a Cerium impurity in metal, the Kondo temperature T_{K} is proportional to exp[-1/JI N(E_{F})], where J is the effective exchange integral given by



Fig. 14. Magnetization vs magnetic field for powdered polycrystalline samples of (a) CeNi_{1-x}Co_xSn and (b) CeNi_{1-x}Pt_xSn.

 $J = |V_{k,f}|^2/(E_f - E_F)$ with the hybridization matrix element $V_{k,f}$ between the 4f state and the conduction electrons.³⁵ Then, T_K is expected to be strongly enhanced by Co substitution and the system is converted to the VF regime, as is experimentally observed. On the contrary, the negative chemical pressure induced by the Pt substitution may weakens the hybridization of the 4f states with the Sn-5p and Ni-3d states. In fact, recent photoemission study by Nohara et al.³⁶ have revealed that the hybridization weakens significantly on going from CeNiSn to CePtSn. The decrease of $V_{k,f}$ in the above expression for T_K results in the lowering of the value of T_K . This is consistent with the transition into the Kondo regime observed for the Pt substituted system .

The effect of the substitution on the gapped state in CeNiSn was studied by the measurements of resistivity and specific heat. However, the temperature dependence of resistivity of polycrystalline samples of CeNi_{1-x} T_x Sn (T= Co and Pt) was found to be very sample-dependent even for a fixed value of x. This is presumably caused by the preferred orientation of the sample, as is expected from the highly anisotropic behavior in the the resistivity of the single crystalline sample (see Fig. 3). This situation did not allow us to study how the gap energy changes as a function of x.

The low-temperature specific heat of $\text{CeNi}_{1-x}T_x \text{Sn}$ (T= Co and Pt) changes smoothly with x as shown in Fig. 15.^{37,38} For pure CeNiSn, the sudden decrease in C/T below 6 K originates from the gap formation in the density of states. With increasing Pt concentration, the value of C/T increases and an upturn in C/T appears for $x \ge 0.12$. This change implies that the density of quasiparticle states grows within the gap and eventually a heavy fermion band is formed. The Kondo temperature for the x = 0.2 sample is estimated to be 25 K from an analysis of C/T using the isolated Kondo impurity model.³⁸ The pronounced peak at 2.1 K for x = 0.33 indicates the development of a long-range magnetic order for higher Pt concentration. This trend is consistent with the increased magnetic moment with Pt concentration, as was shown in Fig.14. We recall here that the transition into an antiferromagnetically ordered state is induced by the Cu substitution at a smaller concent-



Fig. 15. Specific heat divided by temperature C/T vs T for $CeNi_{1-X}T_XSn$ (T= Co and Pt).

ration x = 0.13.¹² In this case, the increase in both the number of 3d electrons and the unitcell volume may promote the transition.

On the other hand, the substitution of only 3 at.% Co is enough to reduce the value of C/T substantially throughout the temperature range below 30 K. Thereby, the anomaly due to the gap formation below 6 K is also smeared out. This in turn suggests that a moderate mass-enhancement is a necessary condition for the gap formation. Comparing the results of the Co, Cu and Pt substituted systems, we find that any replacement of about 10% of the Ni sublattice in CeNiSn closes the energy gap. This fact implies that the loss of the periodicity in the Kondo-lattice is very destructive for the gap formation.

7. CONCLUDING REMARKS

In this article, we reviewed our experimental attempts to understand the semiconductorlike behavior in CeNiSn at low temperatures. The results of transport, thermal and magnetic measurements characterize this compound as a rare example located near the crossover between the Kondo and VF regimes. The highly anisotropic behavior reflects the strong hybridization of the 4f states with the Sn-5p and Ni-3d states in the orthorhombic structure. The Kondo temperature of this system is about 28 K. At low temperatures below 12 K, coherence of local spin fluctuations develops, as indicated by the peaking in both $p_a(T)$ and $\chi_a(T)$ at 12 K and in R_H(T) at 9 K. An anisotropic gap opens in the heavy-quasiparticle bands as temperature is reduced below 6 K. This semiconducting state with a charge excitation gap of several Kelvins is destructible by applying either hydrostatic pressure of 20 kbar or magnetic field of 13 T along the easy a axis. Substitutions of Co and Pt for a few percent of Ni in CeNiSn destroy the energy gap and shift the system toward the VF and the heavy-fermion regimes, respectively. The combined results indicate that the gap formation in CeNiSn is very sensitive to the degree of hybridization of the 4f and the conduction electron states. It is also emphasized that the development of coherence in the Kondo lattice is crucial for the gap formation.

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REFERENCES

- 1. A. Menth, E. Buehler and T.H. Geballe, Phys. Rev. Lett. 22:295(1969).
- 2. A. Jayaraman, V. Narayanamurti, E. Bucher and R.G. Maines, Phys. Rev. Lett. 25:1430(1970).
- 3. P. Haen, F. Holzberg, F. Lapierre, T. Penny and R. Tournier, in Valence Instabilities and Related Narrow Band Phenomena, ed. by R.D. Parks (Plenum, New York, 1977) p. 1495.
- 4. M. Kasaya, F. Iga, M. Takigawa and T. Kasuya, J. Magn. Mater. 47&48:429(1985).
- 5. N.F. Mott, Phil. Mag. 30:403(1973).
- 6 T. Takabatake, Y. Nakazawa and M. Ishikawa, Jpn. J. Appl. Phys. Suppl. 26-3:547(1987).
- 7. I. Higashi, K. Kobayashi, T. Takabatake and M. Kasaya, to be published in J. Alloys Compds. (1992).
- 8. M.F. Hundley, P.C. Canfield, J.D. Thompson, Z. Fisk and J.M. Lawrence, Phys. Rev. B42:6842 (1990).
- 9. S.K. Malik and D.T. Adroja: Phys. Rev. B43:6277(1991).
- 10. P.S. Riseborough, Phys. Rev. B45:13984(1992).
- 11. P. Schlottmann, Phys. Rev. B46:998(1992).
- T. Takabatake, Y. Nakazawa, M. Ishikawa, T. Sakakibara, K. Koga and I. Oguro, J. Magn. Magn. Mater. 76&77:87(1988).
- T. Takabatake, F. Teshima, H. Fujii, S. Nishigori, T. Suzuki, T. Fujita, Y. Yamaguchi, J. Sakurai and D. Jaccard, Phys. Rev. B41:9607(1990).
- 14. T. Takabatake, M. Nagasawa, H. Fujii, G. Kido, M. Nohara, S. Nishigori, T. Suzuki, T. Fujita, R. Helfrich, U. Ahlheim, K. Fraas, C. Geibel and F. Steglich, Phys. Rev. B45:5740(1992).
- T. Takabatake, M. Nagasawa, H. Fujii, M. Nohara, T. Suzuki, T. Fujita, G. Kido and T. Hiraoka, J. Magn. Magn. Mater. 108:155(1992).
- T. Takabatake, M. Nagasawa, H. Fujii, G. Kido, K. Sugiyama, K. Senda, K. Kido and M. Date, Physica B172:177(1992).
- 17. T. Takabatake and H. Fujii, to be published in Jpn. J. Appl. Phys. Ser.8 (1992).
- J. Sakurai, Y. Yamaguchi, S. Nishigori, T. Suzuki and T. Fujita, J. Magn. Magn. Mater. 90&91: 442(1990).
- 19. T. Takabatake, H. Iwasaki, G. Nakamoto, H. Fujii, H. Nakotte, F.R. de Boer and V. Sechovsky, to be published in Physica B182(1992).
- 20 T. Kasuya, J. Phys. Soc. Jpn. 61:1863(1992).
- 21. A. Yanase and H. Harima, Progr. Theor. Phys. Jpn. Suppl. 108:19(1992).
- 22. M. Kohgi, K. Ohoyama, T. Osakabe and M. Kasaya, J. Magn. Magn. Mater. 108:187(1992).
- 23. T.E. Mason, G. Aeppli, A.P. Ramirez, K.N. Clausen, C. Broholm, N. Stücheli, E. Bucher and T.T.M. Palstra, Phys. Rev. Lett. 69:490(1992).
- 24. F. Lapierre, P. Haen, R. Brigs, A. Hamezić, A. Fert and J.P. Kappler, J. Magn. Magn. Mater. 63&64: 338(1987).
- 25. A. Fert and P.M. Levy, Phys. Rev. B36:1907(1987).

26. N. Andrei, K. Furuya and J.K. Loewenstein, Rev. Mod. Phys. 55:331(1983).

- 27. M. Kyogaku, Y. Kitaoka, H. Nakamura, K. Asayama, T. Takabatake, F. Teshima and H. Fujii, J. Phys. Soc. Jpn. 59:1728(1990).
- 28. M. Kyogaku, Y. Kitaoka, K. Asayama, T. Takabatake and H. Fujii, J. Phys. Soc. Jpn. 61:43(1992).
- 29. A. Kratzer, G.M. Kalvius, T. Takabatake, G. Nakamoto, H. Fujii and S.R. Kreitzman, Europhys. Lett. 19:649(1992).
- J.M. Mignot, J. Flouquet, P. Haen, F. Lapierre, L. Puech and J. Voiron, J. Magn. Magn. Mater. 76&77:97(1988).
- 31. M. Kurisu, T. Takabatake and H. Fujiwara, Solid State Commun. 68:595(1988).
- 32. M. Kurisu, T. Takabatake and H. Fujii, in this issue.
- 33. J.D. Thompson, J. Magn. Magn. Mater. 63&64:358(1987).
- 34. N. Grewe and F. Steglich, in *Handbook on the Physics and Chemistry of Rare Earths*, ed. by K.A. Gschneidner, Jr. and L. Eyring (Elsevier, Amsterdam, 1991) Vol.14, p. 343.
- 35. B. Cornut and B. Coqblin, Phys. Rev. B5:4541(1972).
- 36. S. Nohara, H. Namatame, A. Fujimori and T. Takabatake, to be published in Phys. Rev. B (1992).
- T. Fujita, T. Suzuki, S. Nishigori, T. Takabatake, H. Fujii and J. Sakurai, J. Magn. Magn. Mater. 104-107:1415(1992).
- 38. S. Nishigori, H. Goshima, T. Suzuki, T. Fujita, G. Nakamoto, T. Takabatake, H. Fujii and J. Sakurai, to be published in Physica B (1992).