



direct and exchange processes together

$$t^{(2)} = \sum_{\mathbf{k}'', \sigma} \frac{1}{E(\mathbf{k}) - E(\mathbf{k}'')} \left[(1 - f_{\mathbf{k}''}) \langle \mathbf{k}' \uparrow | \mathcal{H}_{sd} | \mathbf{k}'' \sigma \rangle \langle \mathbf{k}'' \sigma | \mathcal{H}_{sd} | \mathbf{k} \uparrow \rangle + f_{\mathbf{k}''} \langle \mathbf{k}'' \sigma | \mathcal{H}_{sd} | \mathbf{k} \uparrow \rangle \langle \mathbf{k}' \uparrow | \mathcal{H}_{sd} | \mathbf{k}'' \sigma \rangle \right]$$

- without spin flip \longrightarrow small temperature independent contribution (only terms with S_z contribute)
- with spin flip \longrightarrow Kondo effect

Kondo resistance

algebraic calculation leads to $t^{(2)} = J^2 S_z \sum_{\mathbf{k}''} \frac{2f_{\mathbf{k}''} - 1}{E(\mathbf{k}) - E(\mathbf{k}'')}$

in addition: $D(E) \approx D(E_F)$ $f_{\mathbf{k}''} \longrightarrow$ step function ($T = 0$)

$\delta E < |E_F \pm \mathcal{D}|$ $\sum \rightarrow \int$

$$t^{(1)} + t^{(2)} = -JS_z \left[1 - 2JD(E_F) \ln \frac{\mathcal{D}}{|E_F - E(\mathbf{k})|} \right]$$



scattering probability $w(\mathbf{k} \uparrow, \mathbf{k}' \uparrow) = t_K^2$

→ $w(\mathbf{k} \uparrow, \mathbf{k}' \uparrow) \propto J^2 S_z^2 \left[1 - 4 J D(E_F) \ln \frac{\mathcal{D}}{|E_F - E(\mathbf{k})|} + \dots \right]$

terms of the order of $O(J^4)$ are omitted, **integration** over **all vectors** and **energies** with $E(\mathbf{k}) \approx E_F \pm k_B T$

→ $\varrho(T) \propto \varrho_0 \left[1 - 4 J D(E_F) \ln \frac{\mathcal{D}}{k_B T} \right]$
 $J < 0 \longrightarrow$ increase with decreasing temperature

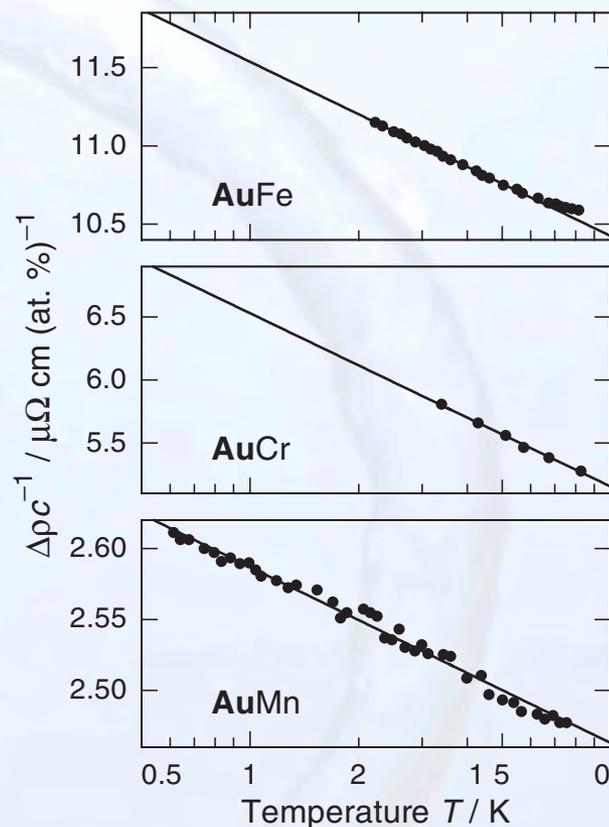
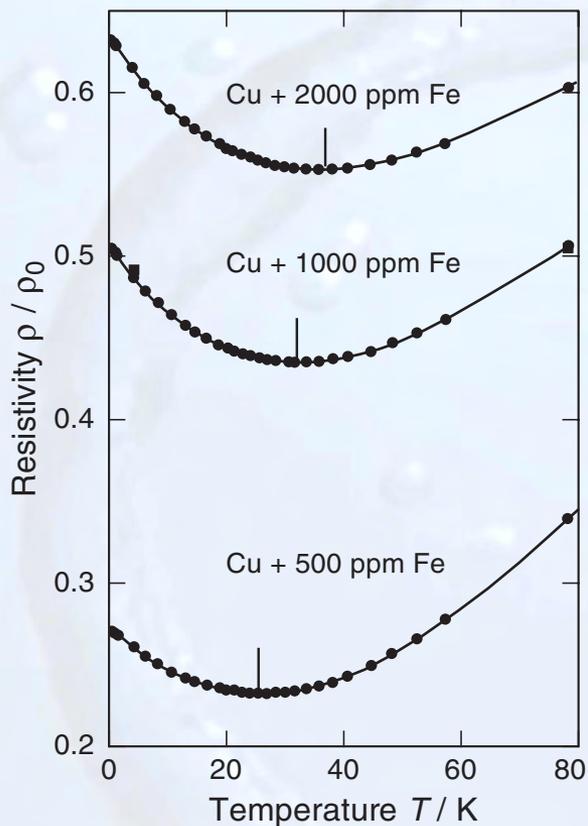
adding the lattice contribution $\varrho_{\text{ph}} = aT^5 \longrightarrow$ total resistance

→ $\varrho = aT^5 + c\varrho_0 + c\varrho_1 \ln \frac{\mathcal{D}}{k_B T}$
concentration of magnetic impurities

minimum expected at $T_{\text{min}} = \left(\frac{c\varrho_1}{5a} \right)^{1/5}$



experimental observations:



► **concentration**-dependent minimum

► **logarithmic temperature** dependence



dependence for $T \rightarrow 0$? \longrightarrow logarithmic divergence is nonphysical

$T > T_K$ weak coupling regime ($-\ln T$)

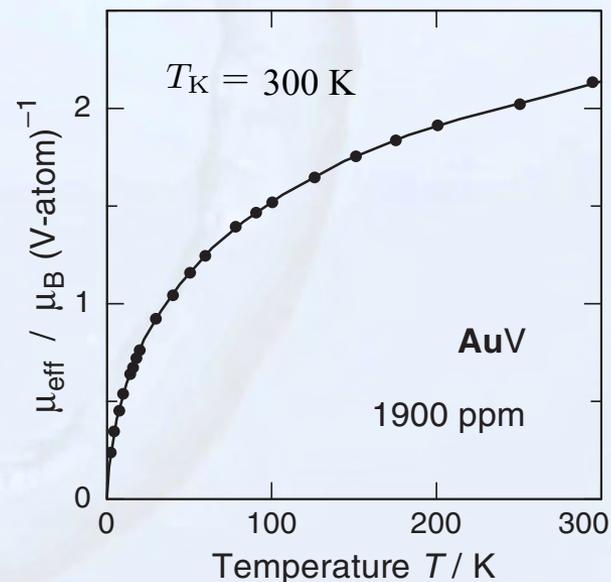
$T < T_K$ strong coupling regime

Kondo temperature: $T_K \approx T_F e^{-1/JD(E_F)}$

strong coupling regime

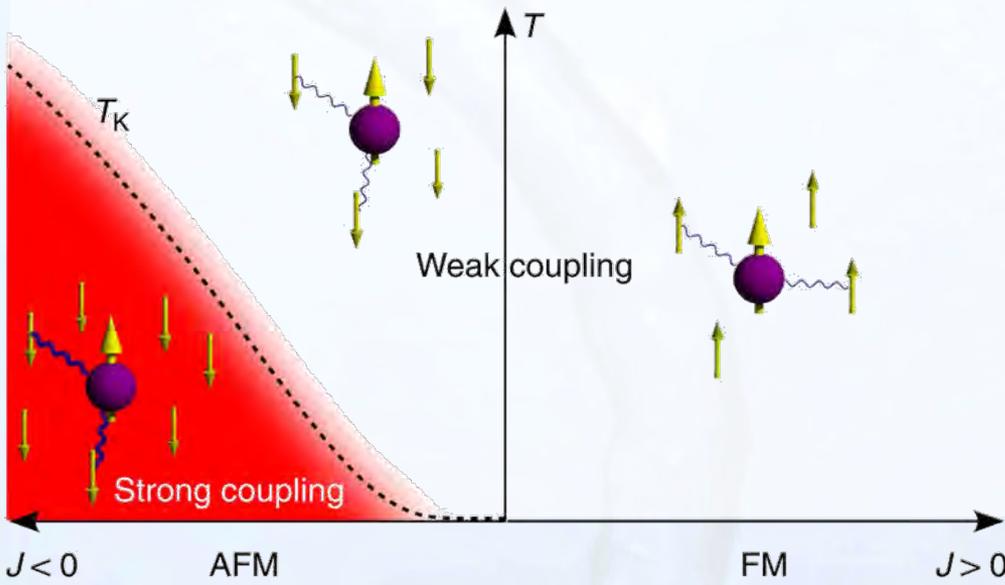
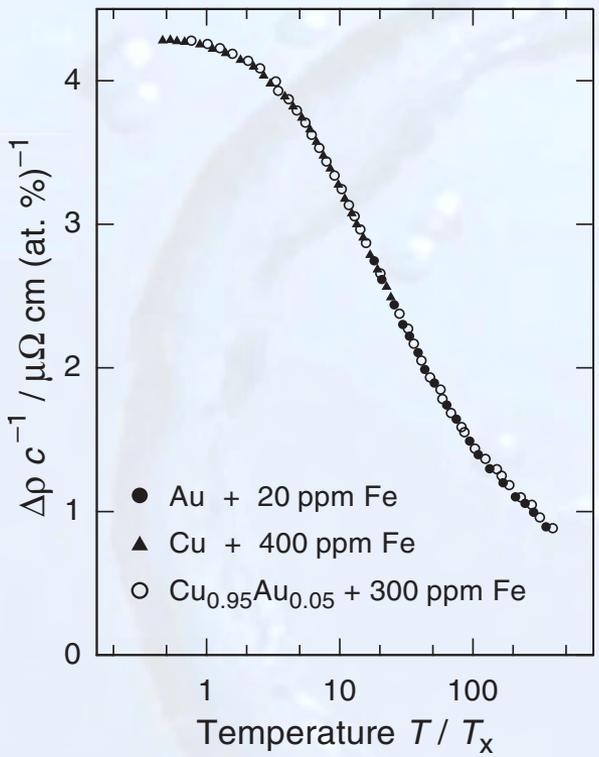
- ▶ strong screening through surrounding conduction electrons
 - \longrightarrow spin-compensated **singlet ground state**
 - \longrightarrow **coherent** Kondo state
- ▶ **transition** from a **magnetic** to a **non-magnetic** system
- ▶ energy necessary to form the spin-compensated cloud: $k_B T_K$
- ▶ **maximum** in **specific heat** of Kondo systems at T_K
- ▶ magnetic moment **disappears** below T_K

T_K $\begin{cases} \text{CuMn} & 1 \text{ mK} \\ \text{AlMn} & 1000 \text{ K} \end{cases}$





resistivity flattens towards low temperatures



- ▶ normalized temperature dependence
- ▶ Kondo resistance is scalable by $T/T_x \approx T/T_K$ and $\Delta\rho/c$



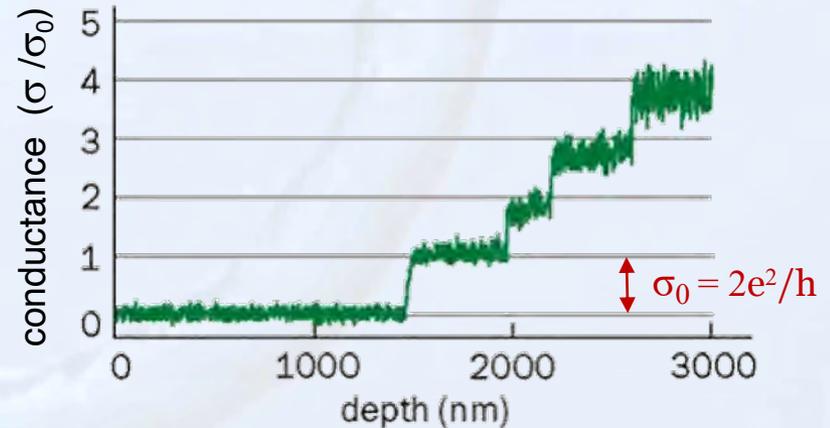
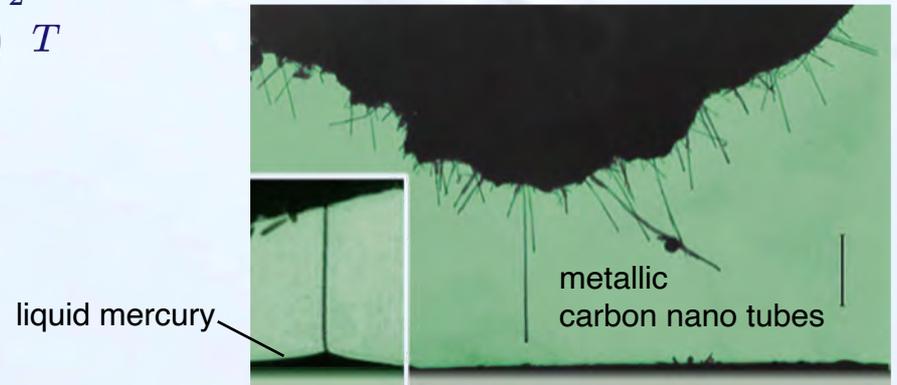
Starting point

quantized thermal conductivity: $\Lambda_0 = \frac{\pi^2}{3} \frac{k_B^2 T}{h}$

Wiedemann-Franz law: $\frac{\Lambda_{el}}{\sigma} = \frac{\pi^2}{3} \left(\frac{k_B}{e}\right)^2 T$

→ $\widetilde{\sigma}_0 = \frac{\Lambda_0}{(\pi^2/3)(k_B^2/e^2)T} = \frac{e^2}{h}$

2 spin directions: $\sigma_0 = 2\widetilde{\sigma}_0 = \frac{2e^2}{h}$





magnetic order of nuclear spins

first observation 1969 in CaF_2 (Chapellier, Goldman, Chan, Abragam)

^{19}F

→ **isolators**: dipole-dipole interaction $E \propto \frac{\mu_n^2}{r^3}$

→ **metals**: dipole-dipole and indirect exchange interaction $H = H_{\text{dip}} + H_{\text{RK}}$

↙ **weak** electron-nuclei coupling
 ↘ **strong** electron-nuclei coupling

$$H_{\text{RK}} \approx H_{\text{dip}}$$

$$H_{\text{RK}} > H_{\text{dip}}$$

↘ Ruderman Kittel

metals:

strength of coupling important for relaxation time τ

→ Korringa relation: $\tau = \frac{\kappa}{T_e}$



a) strong electron-nuclei coupling

nuclear ordering in **thermal equilibrium** ($T_e = T_n$): there are only a **few examples**

- ▶ solid ^3He antiferromagnetic ordering at 0.9 mK
- ▶ $\text{PrCu}_6, \text{PrNi}_5, \dots$ ferromagnetic ordering at 2.5 mK, 0.4 mK,

Van Vleck – paramagnets \longrightarrow magnetic field at the nuclei are **enhanced** by **hyperfine interaction** through **polarization of the electrons**

- ▶ AuIn_2 is the only **non-hyperfine-enhanced** compound showing nuclear ordering in **thermal equilibrium**

$I = 9/2, \mu = 5.5 \mu_n$, small Korringa constant $\kappa = 0.09 \text{ Ks}$ (pure In)

pure In: tetragonal \longrightarrow nuclear **quadruple interaction** \longrightarrow suppresses nuclear order

and $B_c = 28 \text{ mT}$ (superconductivity) $\longrightarrow B_{\text{ext}} > B_c$ to stay normal conducting

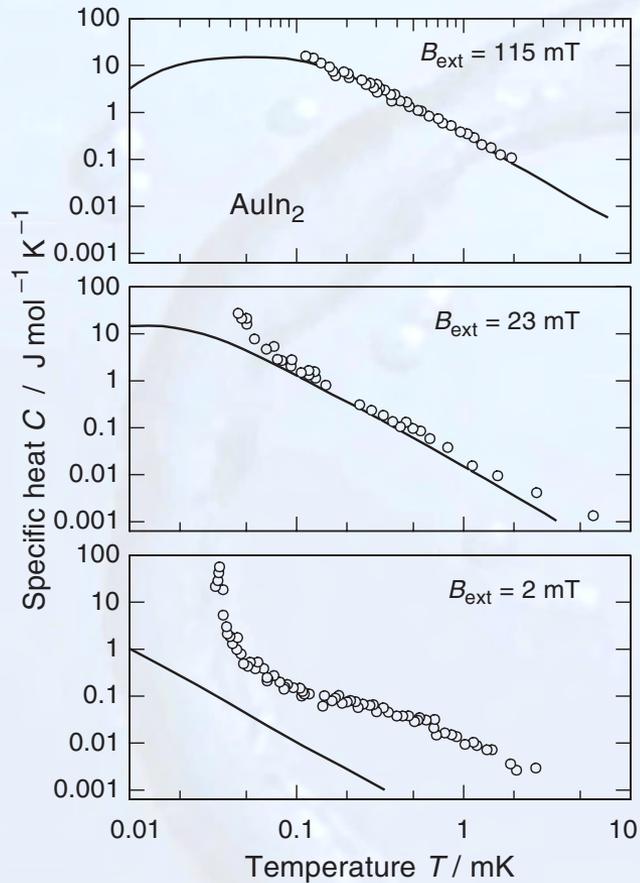
\longrightarrow demagnetization has to stop above B_c

\longrightarrow cannot demagnetized as deep

AuIn_2 : fcc lattice, $B_c = 1.45 \text{ mT}$, $\kappa = 0.11 \text{ Ks}$ (similar to pure In)

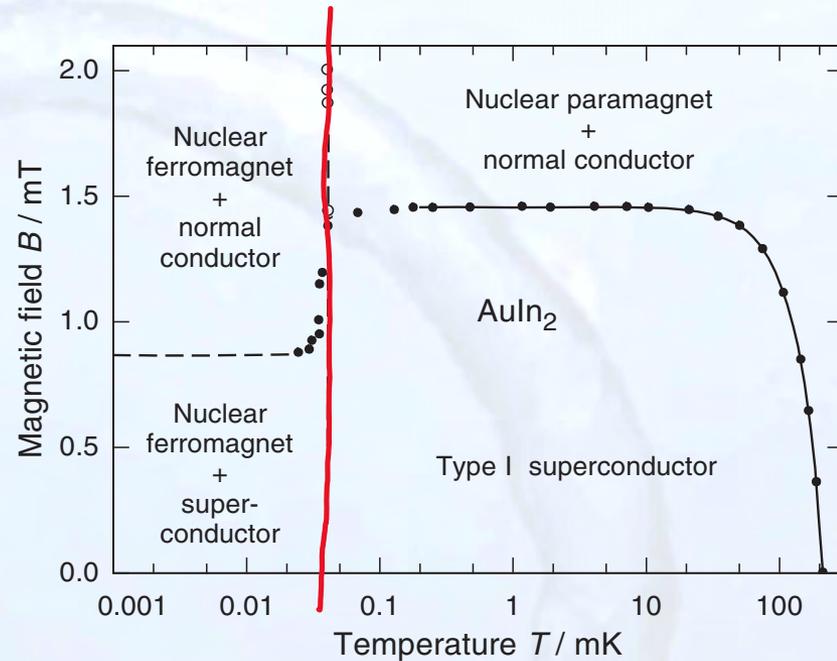
no quadruple interaction

$100 \mu\text{K} \longrightarrow \sim 1000 \text{ s}$ (doable)



- ▶ $B_{\text{ext}} = 2 \text{ mT} \longrightarrow$ phase transition at $35 \mu\text{K}$
- ▶ heat capacity is very large in absolute terms
- ▶ this phase transition is suppressed at higher fields

Phase diagram AuIn_2



- ▶ the phase transition at $35 \mu\text{K}$ is related to nuclear ferromagnetic ordering
- ▶ as a consequence, the critical field for superconductivity is partially suppressed



b) weak electron-nuclei coupling $H_{RK} \approx H_{dip}$

→ nuclear order expected in the **nK range**

not yet possible in thermal equilibrium

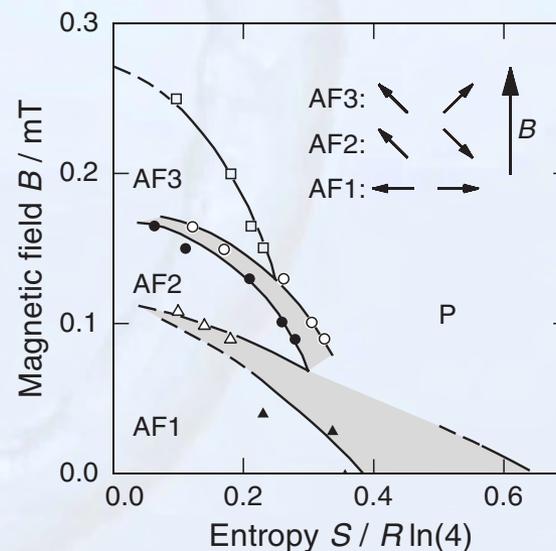
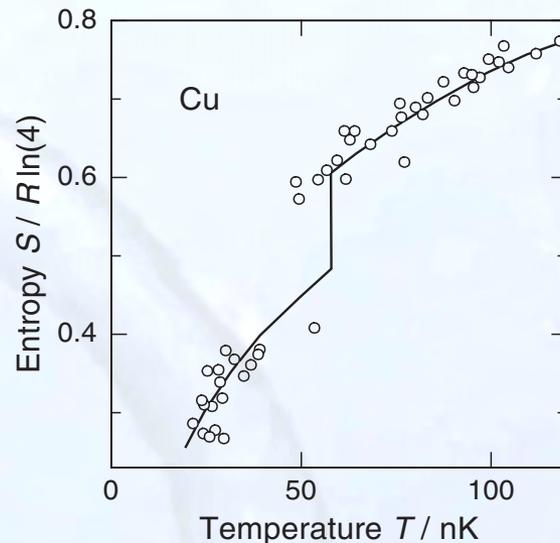
Example: Copper, $I = 3/2$, 69.1% ^{63}Cu , 30.9% ^{65}Cu

Korringa constant $\kappa = 1.2 \text{ Ks}$

→ 100 μK → ~ 3 hours

fast cooling: $T_n \neq T_e$

- ▶ **phase transition** at $T_n = 58 \text{ nK}$
- ▶ first order phase transition → reduction of entropy
→ antiferromagnetic phase
- ▶ in magnetic fields **3 different antiferromagnetic phases**





Negative temperatures

1938 Casimir, DuPré : concept

1951 Purcell, Pound : first realization LiH

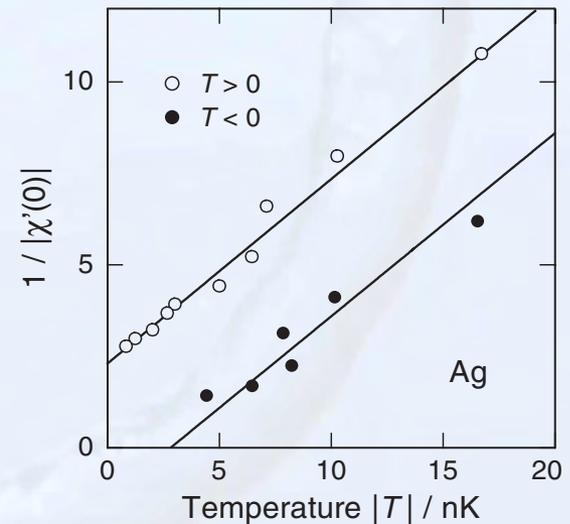
$$\tau_1 = 5 \text{ min}, T = -1 \text{ K}$$

prerequisites:

- ▶ system with **finite number of levels**
- ▶ thermal equilibrium within the system $\tau_1 \gg \tau_2$

example: silver: $T_e = 200 \mu\text{K}$ $\tau_1 = 5 \times 10^4 \text{ s}$
 $\tau_2 = 10 \text{ ms}$

susceptibility at **positive** and **negative** temperature



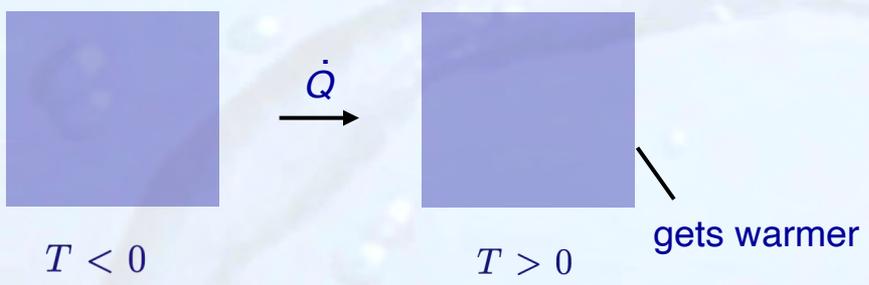


8. Magnetic Moments – Spins



Thermodynamic at negative temperatures

negative temperatures are warmer than positive temperatures



$$+0\text{ K} \rightarrow 300\text{ K} \rightarrow \pm\infty\text{ K} \rightarrow -300\text{ K} \rightarrow -0\text{ K}$$

system cannot be cooled further, it cannot release any energy

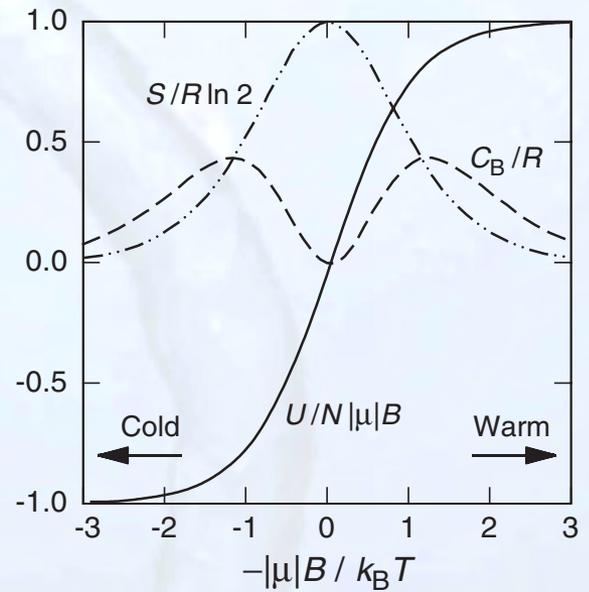
system cannot be warmed further, it cannot take up any energy

NMR: absorption \rightarrow stimulated emission (no laser)

TS term in free energy $F = U - TS$ changes sign \rightarrow spin system maximizes free energy!

systems that orders antiferromagnetically at positive temperatures
order ferromagnetically at negative temperatures and vice versa

internal energy, entropy, specific heat of two-level systems at positive and negative temperatures



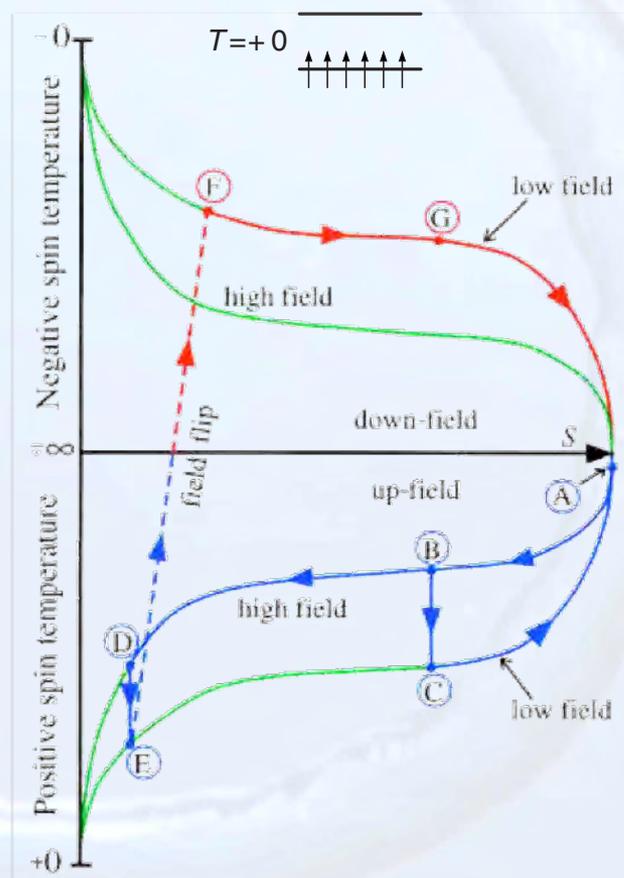
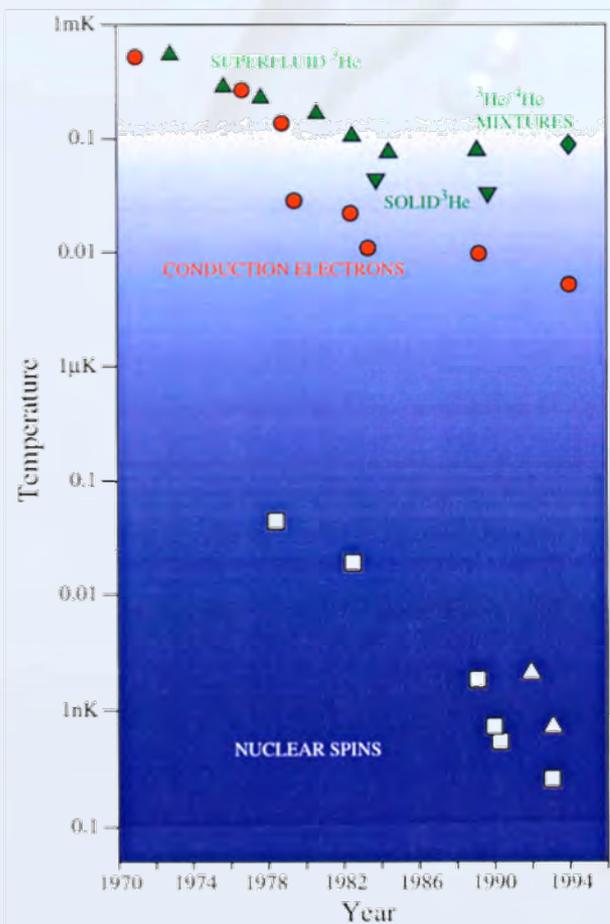
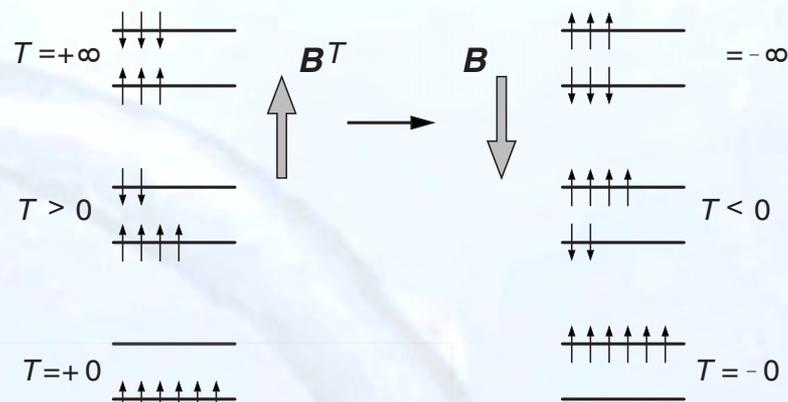


8. Magnetic Moments – Spins



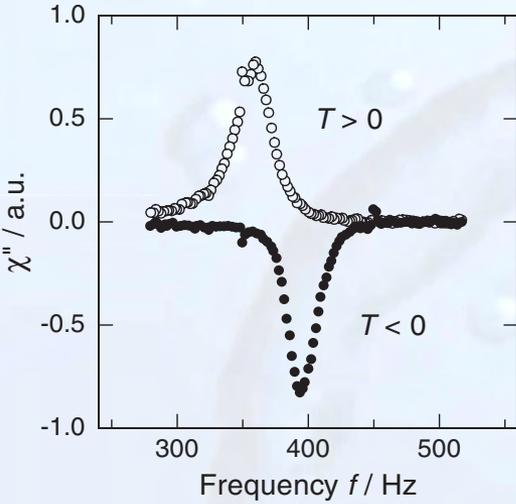
Production of negative temperatures

- ▶ precool system in magnetic field
- ▶ isolate system sufficiently
- ▶ adiabatic reversal of magnetic field





NMR experiments

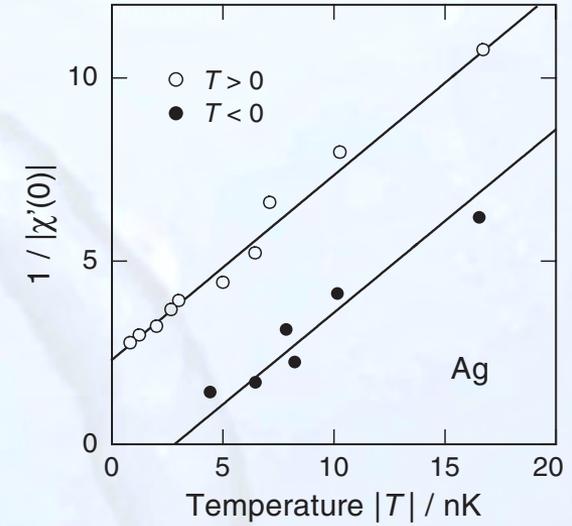


- ▶ normal NMR absorption $T > 0$
- ▶ stimulated emission $T < 0$

$T > 0$ anti-ferromagnetic order
 $T_N = 560 \pm 60$ pK

$T < 0$ ferromagnetic order
 $T_c = -1.9 \pm 0.4$ nK

Nuclear Susceptibility



Phase diagram of silver at positive and negative temperature

- antiferromagnetic phase region $T > 0$
- - - ferromagnetic phase region $T < 0$
- paramagnetic phase region

