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Superconductivity and Low Temperature Physics I



**Lecture Notes
Winter Semester 2022/2023**

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Chapter 2

Thermodynamic Properties of Superconductors

2. Thermodynamic Properties of Superconductors

2.1 Basic Aspects of Thermodynamics

2.1.1 Thermodynamic Potentials

2.2 Type-I Superconductor in an External Field

2.2.1 Free Enthalpy

2.2.2 Entropy

2.2.3 Specific Heat

2.3 Type-II Superconductor in an External Field

2.3.1 Free Enthalpy

2. Thermodynamic Properties of Superconductors

- **already in 1924:**
 - W.H. Keesom tries to describe the superconducting state using basic laws of thermodynamics
 - Meißner effect not known that time: unclear whether superconducting state is a *thermodynamic phase*
- **after 1933:**
 - after discovery of Meißner effect it is evident that *superconducting state is real thermodynamic phase*
 - this fact is used in development of GLAG theory
- **questions:**
 - what is the suitable thermodynamic potential to describe superconductors ?
 - what can we learn on superconductors from their basic thermodynamic properties (e.g. specific heat) ?

2.1 Thermodynamic Potentials

- **thermodynamics:**
 - describes systems with large particle number by small quantity of variables: T, V, N, \dots
- **extensive variables:** V, S, N, \dots
 - depend on system size (amount of substance)
- **intensive variables:** T, p, n, \dots
 - do not depend on system size
- **thermodynamic potentials:**
 - used for the description of equilibrium states
 - equilibrium state: determined by extremal value of the potential
- **example:**
 - S, V and N are the natural variables of a systems, all other fixed by external constraints
 - **internal energy** $U(S, V, N)$ yields full information on the system

$$dU = TdS - pdV + \mu dN$$
 - **adiabatic-isochore processes** are characterized by minimum of U
- **other potentials:**
 - **Helmholtz free energy** $F(T, V, N)$
 - **enthalpy** $H(S, p, N)$
 - **free enthalpy or Gibb energy** $G(T, p, N)$

2.1 Thermodynamic Potentials

- **question:**
what is the suitable potential to represent the system under consideration ?
→ find the set of independent variables
- **discussion of magnetic and electronic systems:**
→ additional variables such as *polarization P* and *magnetization M*
- **discussion of systems with $N = const.$**

A: Internal Energy

- differential of the internal energy

$$dU = \delta Q + \delta W_{\text{mech}} + \delta W_{\text{em}} = TdS - pdV + \mathbf{B} \cdot d\mathbf{m}$$

infinitesimal heat flow into the system infinitesimal mechanical work done on the system infinitesimal electromagnetic work done on the system

$$\delta W_{\text{em}} = \sum_i \mathbf{F}_{Z_i} \cdot d\mathbf{Z}_i$$

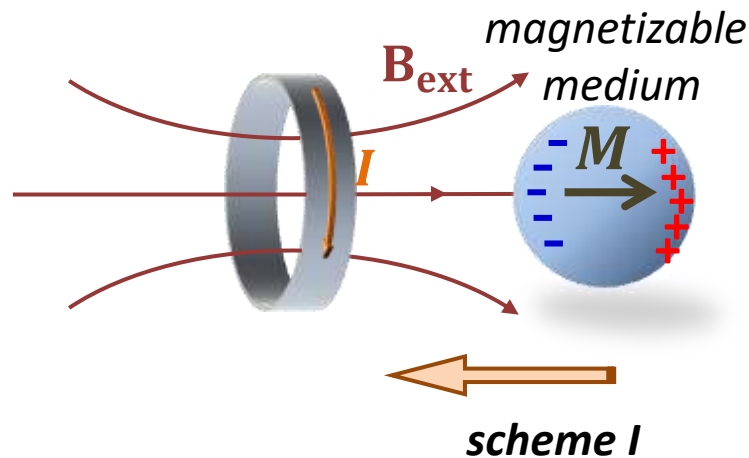
generalized force state variable

\mathbf{m} = magnetic moment
 $\mathbf{M} = \mathbf{m}/V$ = magnetization

$\mathbf{m} \cdot \mathbf{B} = m B$ or $-m B$, since magnetization is mostly parallel or anti-parallel to B

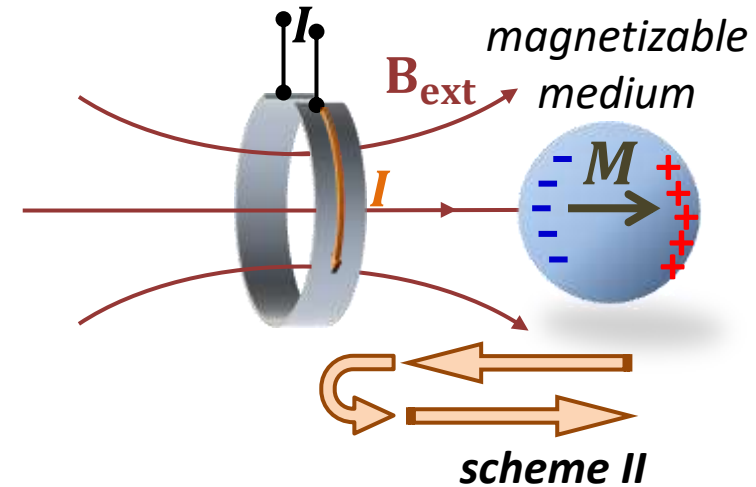
2.1 Thermodynamic Potentials

- **important question:**
 - how to calculate the em work ?
 - do we have to use mdB or Bdm in the expression of dU ?
- **answer:** depends on experimental situation



closed metallic ring with infinite conductivity:
 → flux density in ring stays constant

*interaction energy of dipole with field
 has to be taken into account*

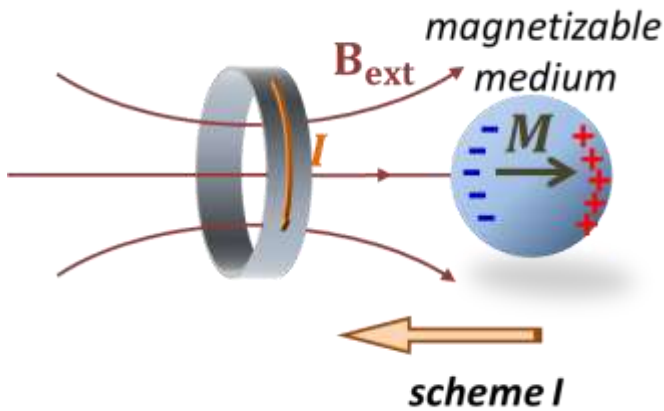


open metallic ring with current source attached:
 → current through ring (magnetic field) stays constant

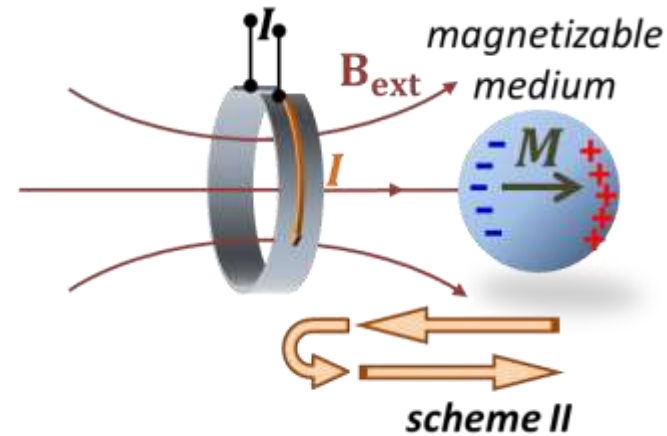
no interaction energy of dipole and field

2.1 Thermodynamic Potentials

- (1) bring magnetic moment \mathbf{m} from ∞ to position r_1 in field \mathbf{B}_{ext}



- (1) bring unmagnetized sample from ∞ to r_1 in field \mathbf{B}_{ext}
 (2) freeze-in magnetic moment \mathbf{m}_1
 (3) bring magnetic moment \mathbf{m}_1 from r_1 to ∞



$$\mathbf{F}_{\text{mag}} = -\mu_0 \mathbf{m} \cdot \nabla \mathbf{H}_{\text{ext}}$$

$$W_I = \int_{\infty}^{r_1} \mathbf{F}_{\text{mag}} \cdot d\mathbf{r} = - \int_0^{H_1} \mu_0 \mathbf{m} \cdot d\mathbf{H}_{\text{ext}} = - \int_0^{B_1} \mathbf{m} \cdot d\mathbf{B}_{\text{ext}}$$

$$W_{II}^{(1)} = \int_{\infty}^{r_1} \mathbf{F}_{\text{mag}} \cdot d\mathbf{r} = - \int_0^{H_1} \mu_0 \mathbf{m} \cdot d\mathbf{H}_{\text{ext}} = - \int_0^{B_1} \mu_0 \mathbf{m} \cdot d\mathbf{B}_{\text{ext}}$$

$$W_{II}^{(3)} = - \int_{H_1}^0 \mu_0 \mathbf{m}_1 \cdot d\mathbf{H}_{\text{ext}} = -\mu_0 \mathbf{m}_1 \cdot \int_{H_1}^0 d\mathbf{H}_{\text{ext}} = \mu_0 \mathbf{m}_1 \cdot \mathbf{H}_1 = \mathbf{m}_1 \cdot \mathbf{B}_1$$

$$\delta W_I = -\mathbf{m} \cdot d\mathbf{B}_{\text{ext}}$$

$$\delta W_{II}^{(1)} = -d(\mathbf{m}_1 \cdot \mathbf{B}_{\text{ext}}) + \mathbf{B}_{\text{ext}} \cdot d\mathbf{m}$$

$$\delta W_I = -d(\mathbf{m} \cdot \mathbf{B}_{\text{ext}}) + \mathbf{B}_{\text{ext}} \cdot d\mathbf{m}$$

$$\delta W_{II}^{(3)} = +d(\mathbf{m}_1 \cdot \mathbf{B}_{\text{ext}})$$

change of the interaction energy

magnetization work

$$\delta W_I = -\mathbf{m} \cdot d\mathbf{B}_{\text{ext}}$$

$$\delta W_{II} = \mathbf{B}_{\text{ext}} \cdot d\mathbf{m}$$

2.1 Thermodynamic Potentials

A: Internal Energy U

$$dU_I = TdS - pdV - \mathbf{m} \cdot d\mathbf{B}_{\text{ext}}$$

$$dU_{II} = TdS - pdV + \mathbf{B}_{\text{ext}} \cdot d\mathbf{m}$$

two different expressions \rightarrow depend on definition of the considered systems

Scheme I: interaction energy $-d(\mathbf{m}_1 \cdot \mathbf{B}_1)$ is included into the considered systems

Scheme II: interaction energy $-d(\mathbf{m}_1 \cdot \mathbf{B}_1)$ is not included

\rightarrow it is assigned to external circuit which performs work on the system and is not part of the considered system

2.1 Thermodynamic Potentials

B: Helmholtz Free Energy F

- definition: $F = U - TS$

- differential: $dF = dU - TdS - SdT$

$$dU_I = TdS - pdV - \mathbf{m} \cdot d\mathbf{B}_{\text{ext}}$$

$$dU_{II} = TdS - pdV + \mathbf{B}_{\text{ext}} \cdot d\mathbf{m}$$

$$dF_I = -SdT - pdV - \mathbf{m} \cdot d\mathbf{B}_{\text{ext}}$$

$$dF_{II} = -SdT - pdV + \mathbf{B}_{\text{ext}} \cdot d\mathbf{m}$$

F_I is potential for the natural variables T, V, B_{ext} , F_{II} for T, V, m

→ isothermal-isochore processes at $B_{\text{ext}} = \text{const.}$ are characterized by minimum of F_I

→ isothermal-isochore processes at $m = \text{const.}$ are characterized by minimum of F_{II}

→ problem: ***in experiments it is difficult to keep V and m constant***

→ ***use free enthalpy or Gibbs energy***

2.1 Thermodynamic Potentials

C: Free Enthalpy G

- definition: $G = U - TS + pV - \mathbf{m} \cdot \mathbf{B}_{\text{ext}}$
- differential: $dG = dU - TdS - SdT + pdV + Vdp - d(\mathbf{m} \cdot d\mathbf{B}_{\text{ext}})$

$$dU_{\text{I}} = TdS - pdV - \mathbf{m} \cdot d\mathbf{B}_{\text{ext}}$$

$$dU_{\text{II}} = TdS - pdV + \mathbf{B}_{\text{ext}} \cdot d\mathbf{m}$$

$$dU_{\text{I}} = TdS - pdV - d(\mathbf{m} \cdot d\mathbf{B}_{\text{ext}}) + \mathbf{B}_{\text{ext}} \cdot d\mathbf{m}$$

interaction energy already included in dU_{I}
and thus has to be omitted in expression
for G to avoid double counting

interaction energy not included in dU_{II} , if
it is included into expression for G , we
obtain $dG_{\text{I}} = dG_{\text{II}}$

$$dG_{\text{I}} = -SdT + Vdp - \mathbf{m} \cdot d\mathbf{B}_{\text{ext}}$$

$$dG_{\text{II}} = -SdT - pdV - \mathbf{m} \cdot d\mathbf{B}_{\text{ext}}$$

G is potential for the natural variables $T, p, B_{\text{ext}} = \mu_0 H_{\text{ext}}$

- isothermal-isobaric processes at $B_{\text{ext}} = \text{const.}$ are characterized by minimum of G
- this is appropriate for most experimental situations

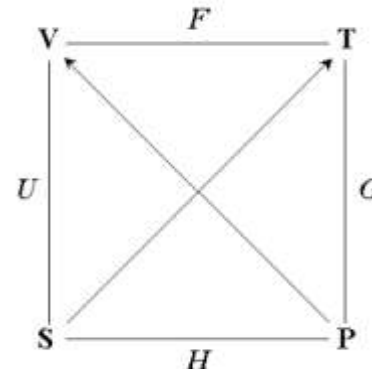
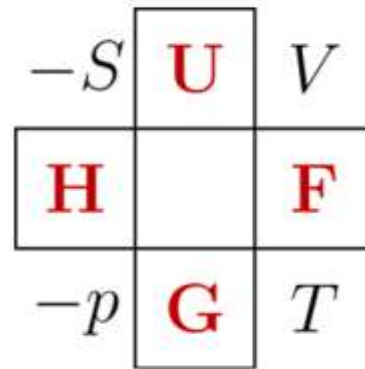
2.1 Thermodynamic Potentials

- if the suitable thermodynamic potential is known
 → *derivation of various thermodynamic quantities is obtained by partial differentiation*

$$-\left(\frac{\partial G}{\partial T}\right)_{p, B_{\text{ext}}} = S$$

$$\left(\frac{\partial G}{\partial p}\right)_{T, B_{\text{ext}}} = V$$

$$-\left(\frac{\partial G}{\partial B_{\text{ext}}}\right)_{p, T} = m$$



- specific heat** $C \equiv \Delta Q / \Delta T$

for $p, B_{\text{ext}} = \text{const.}$

$$\rightarrow dU = \delta Q_{\text{rev}} = TdS$$



$$C_p = T \left(\frac{\partial S}{\partial T}\right)_{p, B_{\text{ext}}} = -T \left(\frac{\partial^2 G}{\partial T^2}\right)_{p, B_{\text{ext}}}$$

2.1 Thermodynamic Potentials

superconductor in external magnetic field: free energy F

- normal state: $F_n = V_S f_n + \frac{1}{2} \mu_0 H_{\text{ext}}^2 V$ ← total volume
 - superconducting state: $F_S = V_S f_S + \frac{1}{2} \mu_0 H_{\text{ext}}^2 V_a$ ← volume outside SC
- ➔ $F_n - F_S = V_S (f_n - f_S) + \frac{1}{2} \mu_0 H_{\text{ext}}^2 V_S$ ← volume of SC

with $f_n - f_S = \frac{1}{2} \mu_0 H_{\text{cth}}^2$ we obtain for $H_{\text{ext}} = H_{\text{cth}}$

$$F_n - F_S \Big|_{H_{\text{ext}}=H_{\text{cth}}} = \mu_0 H_{\text{cth}}^2 V_S = \frac{B_{\text{cth}}^2}{\mu_0} V_S$$

→ problem: free energies of normal and superconducting state do not coincide at phase boundary

→ origin of discrepancy: interaction energy of magnetic moment of SC with H_{ext}

with $\mathbf{M} = -\mathbf{H}_{\text{ext}}$ and $\mathbf{m} = V_S \mathbf{M}$ we obtain

$$F_n - F_S \Big|_{H_{\text{ext}}=H_{\text{cth}}} = -\mu_0 \mathbf{m}_{\text{cth}} \cdot \mathbf{H}_{\text{cth}} = -\mathbf{m}_{\text{cth}} \cdot \mathbf{B}_{\text{cth}}$$

2.1 Thermodynamic Potentials

- where is the energy coming from ?

→ it is provided by the current source maintaining $B_{\text{ext}} = \mu_0 H_{\text{ext}} = \text{const.}$ by doing work against the back electromotive force induced as the flux is entering the sample

- at $H_{\text{ext}} = H_{\text{cth}}$, superconductivity collapses

→ flux density $B_{\text{ext}} = \mu_0 H_{\text{ext}} = B_{\text{cth}}$ enters the volume V_s of the superconductor

- work done by current source $W = \int_a^b U I_{\text{coil}} dt = \int_a^b -N\dot{\Phi} I_{\text{coil}} dt = \int_{a'}^{b'} -N I_{\text{coil}} d\Phi$

$$W = N I_{\text{coil}} \int_0^{B_{\text{cth}}} A dB_{\text{ext}} = N I_{\text{coil}} A B_{\text{cth}}$$

with $B_{\text{ext}} = \mu_0 I_{\text{coil}} N / L$ and $V_s = A L$

$$W = N \frac{B_{\text{cth}} L}{\mu_0 N} A B_{\text{cth}} = V_s \frac{B_{\text{cth}}^2}{\mu_0}$$

2.1 Thermodynamic Potentials

superconductor in external field: free enthalpy G

with $G = U - TS + pV - \mathbf{m} \cdot \mathbf{B}_{\text{ext}} = F + pV - \mathbf{m} \cdot \mathbf{B}_{\text{ext}}$ and $\mathbf{m} = V_s \mathbf{M} = -V_s \mathbf{H}_{\text{ext}}$ we obtain

- normal state:
($\mathbf{m} = 0$) $G_n = V_s f_n + \frac{1}{2} \mu_0 H_{\text{ext}}^2 V_s + \frac{1}{2} \mu_0 H_{\text{ext}}^2 V_a$ (we neglect volume changes)
- superconducting state:
($\mathbf{m} = -V_s \mathbf{H}_{\text{ext}}$) $G_s = V_s f_s + \frac{1}{2} \mu_0 H_{\text{ext}}^2 V_a + \mu_0 H_{\text{ext}}^2 V_s$



$$G_n - G_s \Big|_{H_{\text{ext}}=H_{\text{cth}}} = V_s (f_n - f_s) - \frac{1}{2} \mu_0 H_{\text{cth}}^2 V_s$$

with $f_n - f_s = \frac{1}{2} \mu_0 H_{\text{cth}}^2$ we obtain for $H_{\text{ext}} = H_{\text{cth}}$

$$G_n - G_s \Big|_{H_{\text{ext}}=H_{\text{cth}}} = 0$$

Summary of Lecture No. 2 (1)

- **discussion of basic properties of superconductors**
 - perfect conductivity – perfect diamagnetism – type-I and type-II superconductivity – fluxoid quantization
- **superconducting materials and transition temperatures**
 - discussion of different families of superconducting materials and their transition temperature
- **thermodynamic properties of superconductors**
 - revision of key aspects of thermodynamics
 - thermodynamic potentials: inner energy, free energy, free enthalpy
 - free energy is suitable potential for describing situations at constant volume
 - free enthalpy is suitable potential for describing situations at constant pressure
- **free energy and free enthalpy of a superconductor in an applied magnetic field**
 - discussion of pitfalls in deriving the free energy



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Superconductivity and Low Temperature Physics I



Lecture No. 3

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2. Thermodynamic Properties of Superconductors

2.1 Basic Aspects of Thermodynamics

2.1.1 Thermodynamic Potentials



2.2 Type-I Superconductor in an External Field

2.2.1 Free Enthalpy

2.2.2 Entropy

2.2.3 Specific Heat

2.3 Type-II Superconductor in an External Field

2.3.1 Free Enthalpy

2.2 Type-I Superconductor in B_{ext}

- perfect diamagnetism $\rightarrow \mathbf{M} = \frac{\mathbf{m}}{V} = -\mathbf{H}_{\text{ext}} = -\frac{B_{\text{ext}}}{\mu_0}$
- we assume $p, T = \text{const.}$

$$\rightarrow dG = -SdT + Vdp - \mathbf{m} \cdot d\mathbf{B}_{\text{ext}} = -\mathbf{m} \cdot d\mathbf{B}_{\text{ext}}$$

$$dG_S = \frac{V}{\mu_0} B_{\text{ext}} dB_{\text{ext}} \quad d\mathcal{G}_S = dG_S/V$$

\rightarrow integration yields

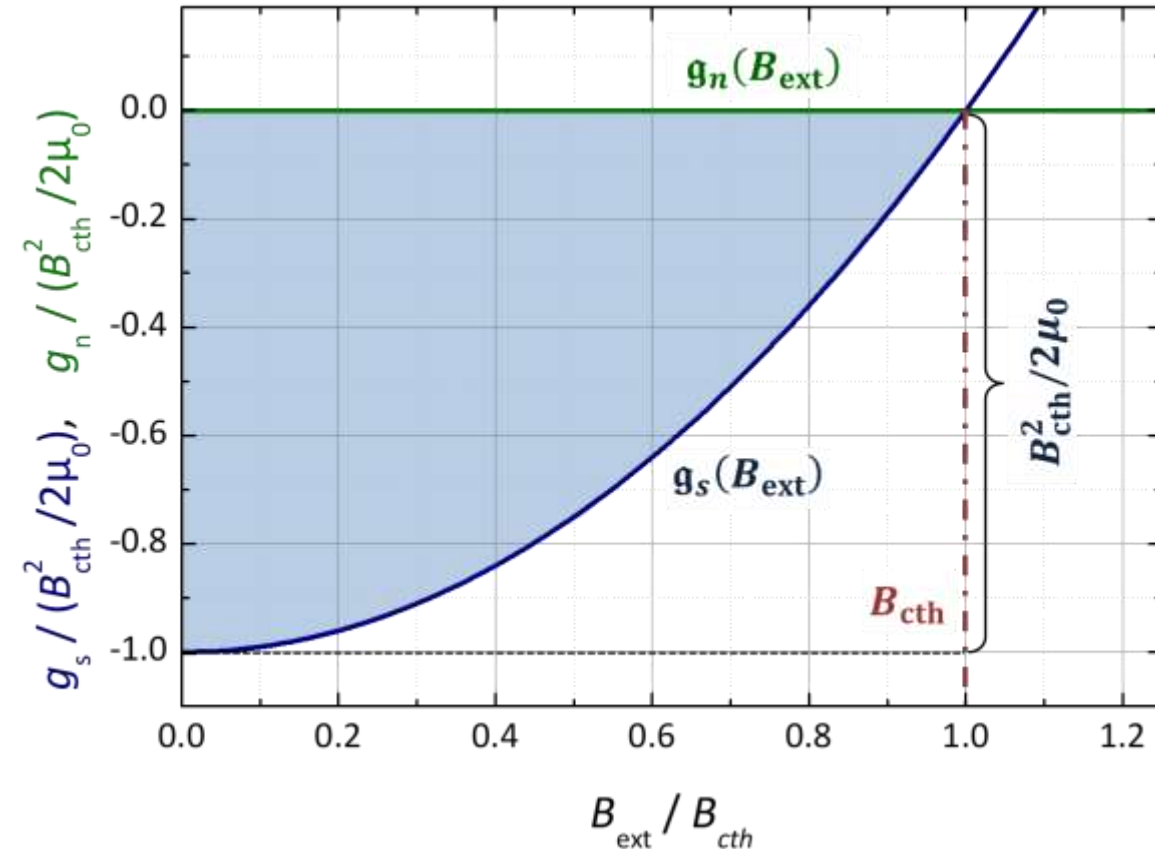
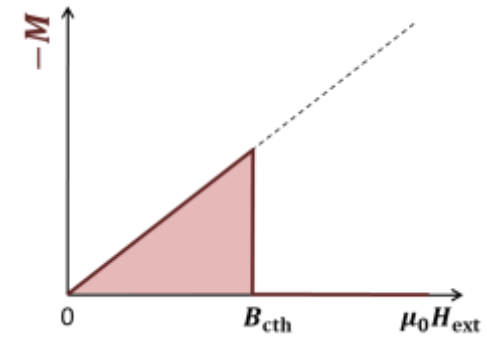
$$\mathcal{G}_S(B_{\text{ext}}, T) - \mathcal{G}_S(0, T) = \frac{1}{\mu_0} \int_0^{B_{\text{ext}}} B' dB' = \frac{B_{\text{ext}}^2}{2\mu_0}$$

@ $B_{\text{ext}} = B_{\text{cth}}$: $\mathcal{G}_S(B_{\text{cth}}, T) = \mathcal{G}_n(B_{\text{cth}}, T) \simeq \mathcal{G}_n(0, T)$

$$\Delta\mathcal{G}(T) = \mathcal{G}_n(0, T) - \mathcal{G}_S(0, T)$$

$$= \mathcal{G}_S(B_{\text{cth}}, T) - \mathcal{G}_S(0, T) = \frac{B_{\text{cth}}^2(T)}{2\mu_0}$$

\rightarrow $\Delta\mathcal{G}(T) = \frac{B_{\text{cth}}^2(T)}{2\mu_0}$ **condensation energy**



2.2 Type-I Superconductor in B_{ext}

temperature dependence of the free enthalpy densities g_n and g_s

$$g_s(T) = g_n(T) - \Delta g(T), \quad \text{with } \Delta g(T) = \frac{B_{\text{cth}}^2(T)}{2\mu_0}$$

$$\rightarrow g_s(T) = g_n(T) - \frac{B_{\text{cth}}^2(T)}{2\mu_0}$$

– temperature dependence of B_{cth} :

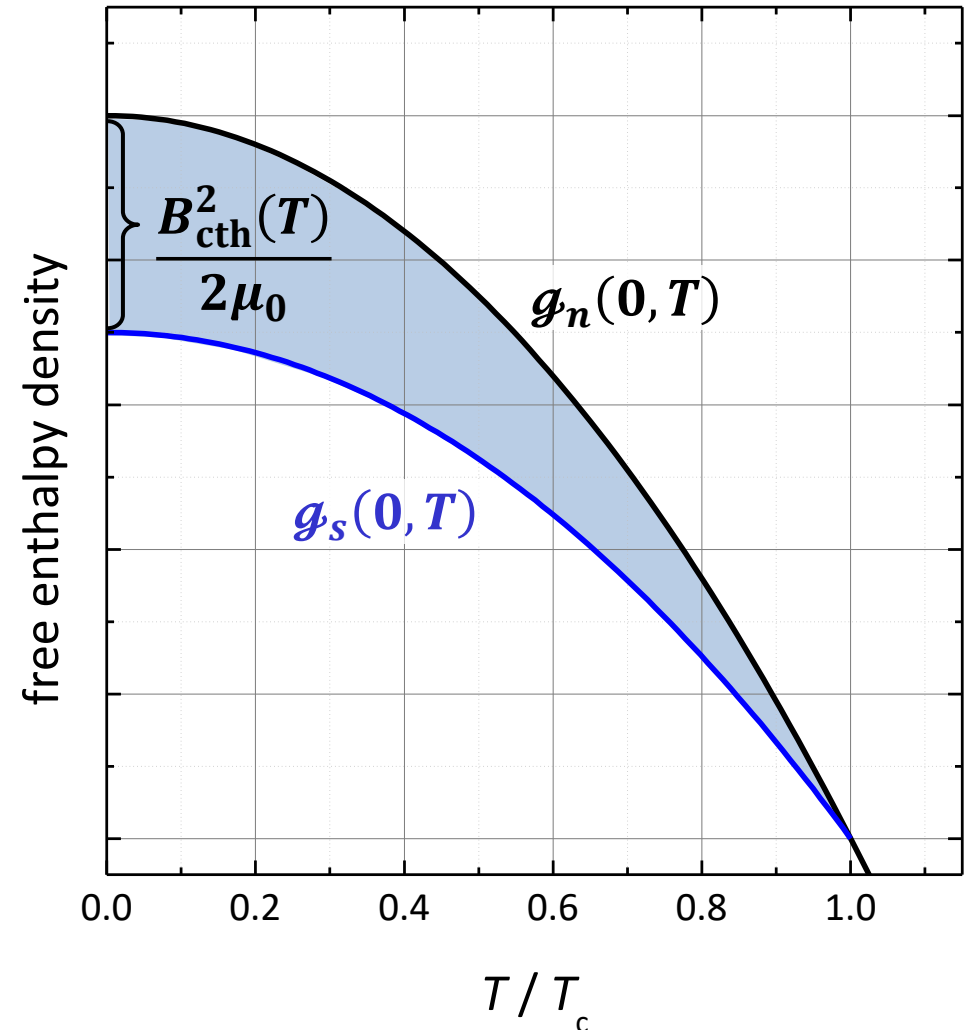
$$B_{\text{cth}}(T) = B_{\text{cth}}(0) \left[1 - \left(\frac{T}{T_c} \right)^2 \right]$$

(empirical relation, calculation within BCS theory)

– entropy density of normal metal (free electron gas):

$$s_n(T) \propto T, \quad dg_n = -s_n dT \quad (@ B_{\text{ext}} = 0)$$

$$g_n(T) = - \int_0^T s_n(T') dT' \propto -T^2$$



\rightarrow determination of entropy density $s_s(T)$ and specific heat $c_p(T)$ by calculating the temperature derivative of $g_s(T)$

2.2 Type-I Superconductor in B_{ext}

temperature dependence of the entropy density $\mathcal{s}_s = S_s/V$

- with $-\left(\frac{\partial G}{\partial T}\right)_{p, B_{\text{ext}}} = S$ and $\mathcal{s}_s = S_s/V$, $\mathcal{s}_n = S_n/V$

$$\mathcal{s}_{s,n}(T) = -\left(\frac{\partial \mathcal{g}_{s,n}}{\partial T}\right)_{p, B_{\text{ext}}}$$

$$\Rightarrow \Delta \mathcal{s}(T) = \mathcal{s}_n(T) - \mathcal{s}_s(T) = -\left(\frac{\partial \Delta \mathcal{g}(T)}{\partial T}\right)_{p, B_{\text{ext}}}$$

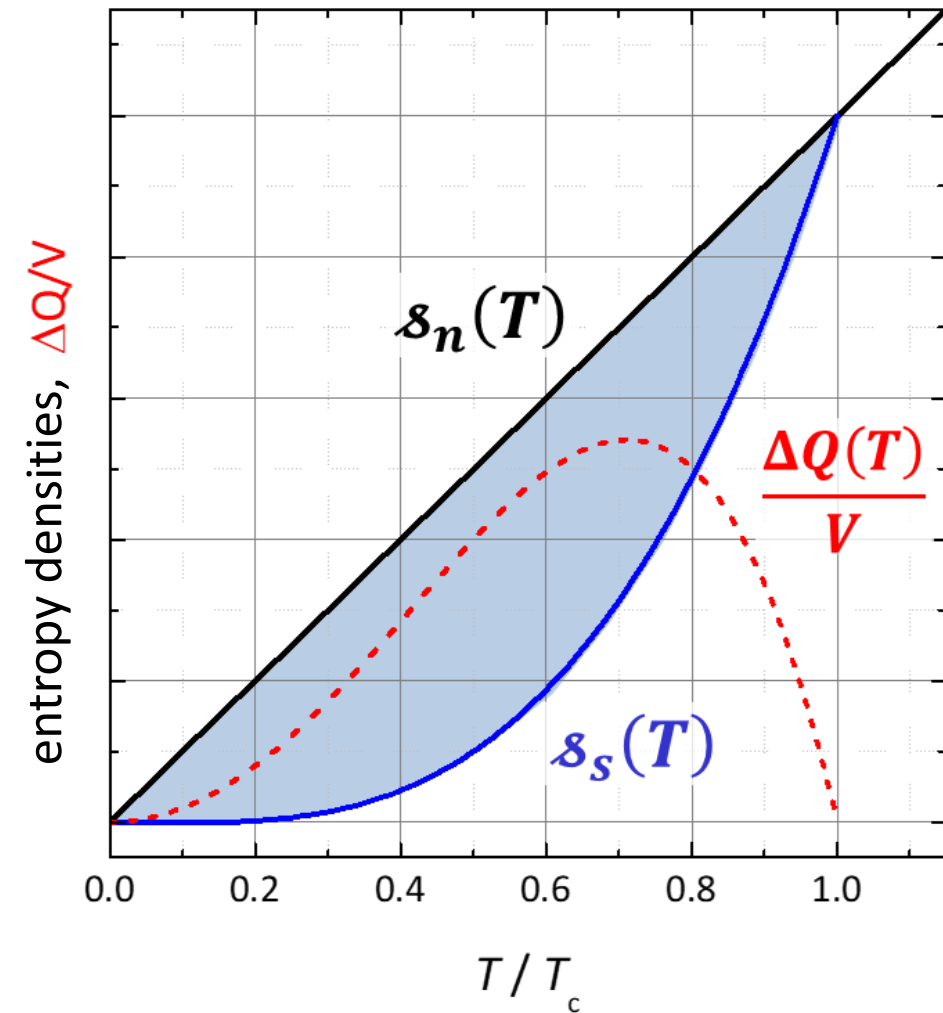
- with $\Delta \mathcal{g}(T) = \frac{B_{\text{cth}}^2(T)}{2\mu_0}$ we obtain

$$\Delta \mathcal{s}(T) = -\frac{B_{\text{cth}}}{\mu_0} \frac{\partial B_{\text{cth}}}{\partial T} \quad \text{with } B_{\text{cth}}(T) = B_{\text{cth}}(0) \left[1 - \left(\frac{T}{T_c}\right)^2\right]$$

- how does $\mathcal{s}_n(T)$ look like?

we use $c_p = T (\partial \mathcal{s}_n / \partial T)_{B_{\text{ext}}, p}$ and $c_p = \gamma T$ (free electron gas)

$\Rightarrow \mathcal{s}_n$ proportional T



2.2 Type-I Superconductor in B_{ext}

discussion of the temperature dependence of the entropy difference Δs

$$\Delta s(T) = -\frac{B_{\text{cth}}}{\mu_0} \frac{\partial B_{\text{cth}}}{\partial T} \quad \text{mit } B_{\text{cth}}(T) = B_{\text{cth}}(0) \left[1 - \left(\frac{T}{T_c} \right)^2 \right]$$

i. $T \rightarrow T_c$:

$B_{\text{cth}} \rightarrow 0$ and therefore $\Delta s \rightarrow 0$ and $\frac{\Delta Q}{V} = T_c \Delta s \rightarrow 0$
(no latent heat, 2nd order phase transition)

ii. $T \rightarrow 0$:

$\frac{\partial B_{\text{cth}}}{\partial T} \rightarrow 0$ and therefore $\Delta s \rightarrow 0$
(3. law of thermodynamics, Nernst theorem)

iii. $0 < T < T_c$:

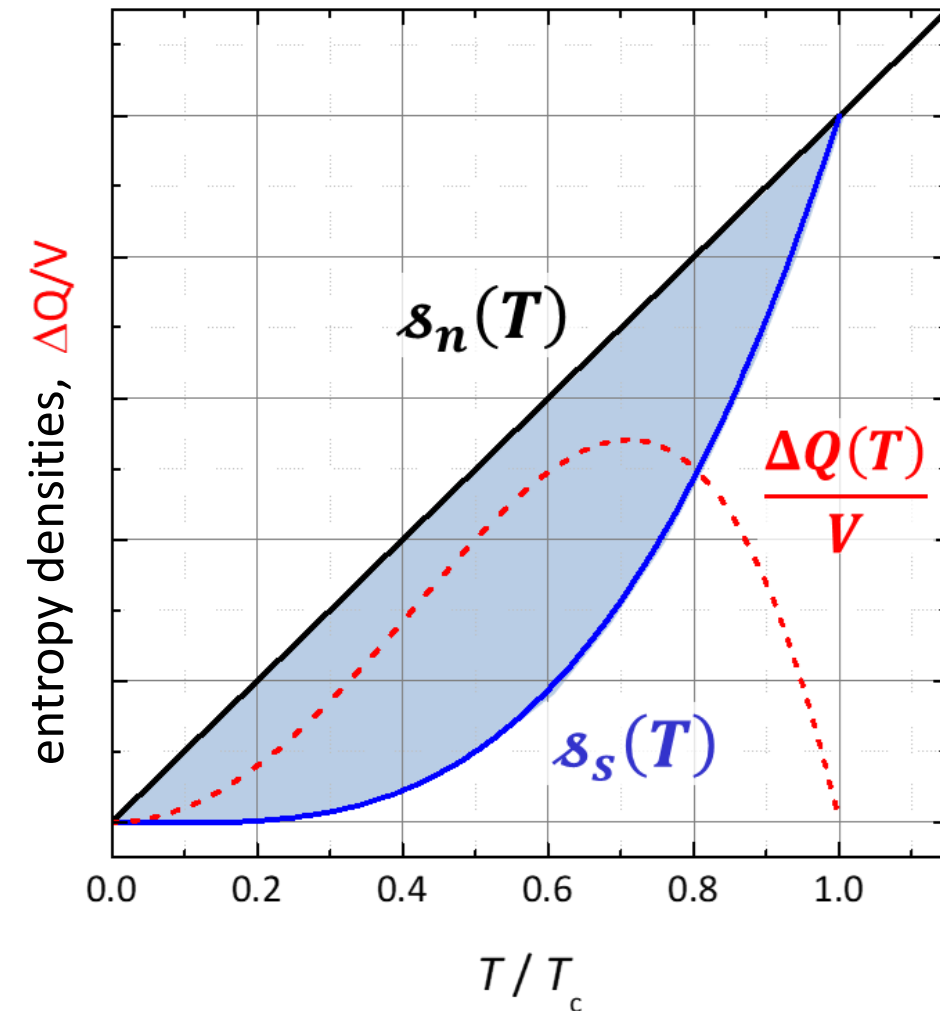
$\frac{\partial B_{\text{cth}}}{\partial T} < 0$, $B_{\text{cth}} > 0$ and therefore $\Delta s > 0$

→ entropy density larger in N-phase than in S-phase

→ S-phase is phase with larger order
(correlation of electrons to Cooper pairs)

→ since $\Delta s > 0$ also $\frac{\Delta Q}{V} = T_c(B_{\text{ext}}) \Delta s > 0$

(finite latent heat, 1st order phase transition)



2.2 Type-I Superconductor in B_{ext}

temperature dependence of specific heat

- we use the general relations

$$C_p = T \left(\frac{\partial S}{\partial T} \right)_{p, B_{\text{ext}}} = -T \left(\frac{\partial^2 G}{\partial T^2} \right)_{p, B_{\text{ext}}}$$

$$\text{and } \Delta \mathcal{G} = \mathcal{G}_n(T) - \mathcal{G}_s(T) = \frac{B_{\text{cth}}^2(T)}{2\mu_0}$$

- jump of specific heat at $T = T_c$

$$\Delta c_{T=T_c} = -\frac{T_c}{\mu_0} \left(\frac{\partial B_{\text{cth}}}{\partial T} \right)_{T=T_c}^2$$

- agrees well with experiment
- $c_s = c_n$ at temperature, where $\Delta s = s_n - s_s = \text{max}$

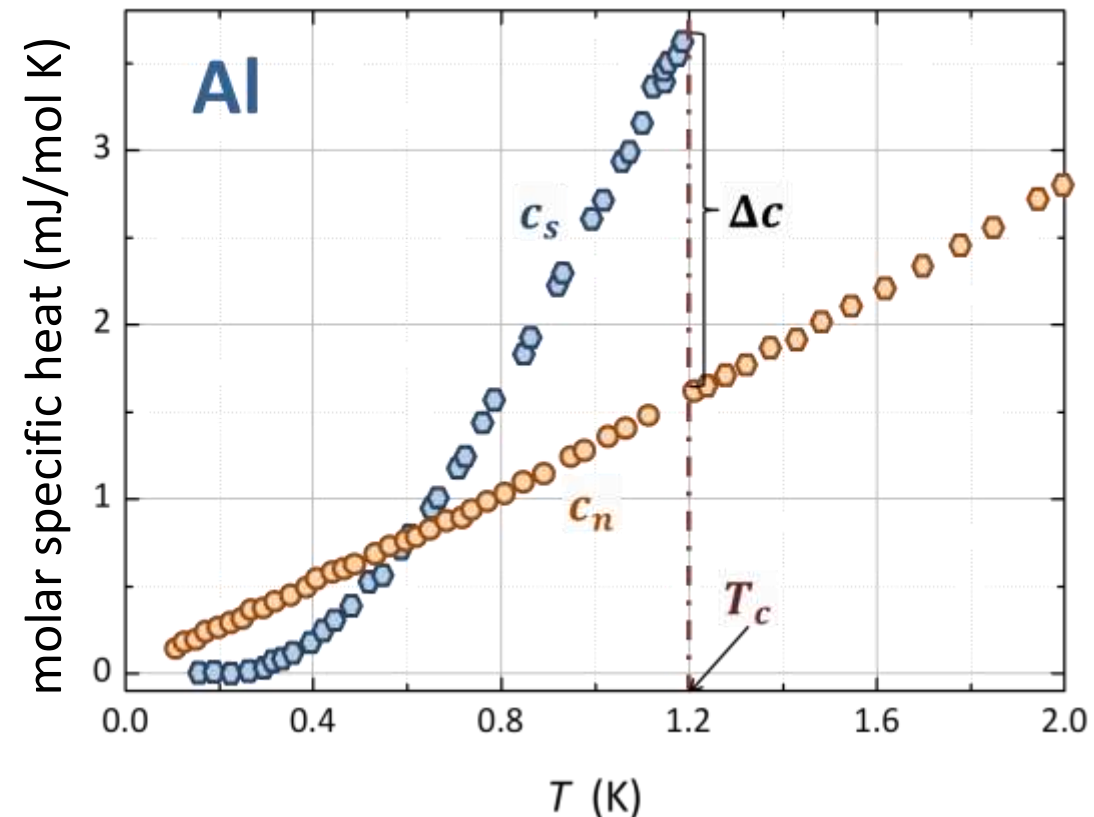
$$\text{for } B_{\text{cth}}(T) = B_{\text{cth}}(0) \left[1 - \left(\frac{T}{T_c} \right)^2 \right]:$$

$$\Delta c_{T=T_c} = -\frac{8}{T_c} \frac{B_{\text{cth}}^2(0)}{2\mu_0}$$

Rutgers formula

$$\Delta c(T) = c_n(T) - c_s(T) = -\frac{T}{\mu_0} \left[B_{\text{cth}} \frac{\partial^2 B_{\text{cth}}}{\partial T^2} + \left(\frac{\partial B_{\text{cth}}}{\partial T} \right)^2 \right]$$

< 0 for all T decreases with T
 ➔ Δc changes sign



2.2 Type-I Superconductor in B_{ext}

determination of the Sommerfeld coefficient γ :

- for $T \ll T_c$, we can neglect c_s compared to c_n
- we use $c_n(T) = \gamma \cdot T$ (γ = Sommerfeld coefficient)

$$\Rightarrow \gamma = \frac{\Delta c}{T} = -\frac{1}{\mu_0} \left[B_{\text{cth}} \frac{\partial^2 B_{\text{cth}}}{\partial T^2} + \left(\frac{\partial B_{\text{cth}}}{\partial T} \right)^2 \right] \approx -\frac{1}{\mu_0} B_{\text{cth}} \frac{\partial^2 B_{\text{cth}}}{\partial T^2}$$

$B_{\text{cth}} \frac{\partial^2 B_{\text{cth}}}{\partial T^2} \gg \left(\frac{\partial B_{\text{cth}}}{\partial T} \right)^2$

- with $B_{\text{cth}}(T) = B_{\text{cth}}(0) \left[1 - \left(\frac{T}{T_c} \right)^2 \right]$, we obtain $\frac{\partial^2 B_{\text{cth}}}{\partial T^2} = -2B_{\text{cth}}(0)/T_c^2$ and hence:

$$\gamma = \frac{4}{T_c^2} \frac{B_{\text{cth}}^2(0)}{2\mu_0}$$

→ by measuring T_c and $B_{\text{cth}}(0)$, we can determine γ and, in turn, the density of states at the Fermi level, $D(E_F)$

$$\gamma = \frac{\pi^2}{3} k_B^2 \frac{D(E_F)}{V}$$

(free electron gas model)

2.2 Type-I Superconductor in B_{ext}

- with $c_n(T_c) = \gamma T_c$ we obtain

$$\frac{\Delta c_{T=T_c}}{c_n} = -\frac{8}{\gamma T_c^2} \frac{B_{\text{cth}}^2(0)}{2\mu_0}$$

- BCS theory predicts $g_n - g_s(0) = \frac{B_{\text{cth}}^2(0)}{2\mu_0} = \frac{1}{4} D(E_F) \Delta^2(0)/V$, then with $\gamma = \frac{\pi^2}{3} k_B^2 \frac{D(E_F)}{V}$ we obtain

$$\frac{\Delta c_{T=T_c}}{c_n} = -\frac{8}{\frac{\pi^2}{3} k_B^2 \frac{D(E_F)}{V} T_c^2} \frac{\frac{1}{4} D(E_F) \Delta^2(0)}{V}$$

$$\frac{\Delta c_{T=T_c}}{c_n} = -\frac{6}{\pi^2} \left(\frac{\Delta(0)}{k_B T_c} \right)^2$$

note that the factor $\frac{6}{\pi^2} \simeq 0.6079 \dots$ comes from $B_{\text{cth}}(T) = B_{\text{cth}}(0) \left[1 - \left(\frac{T}{T_c} \right)^2 \right]$ (phenomenological approximation), BCS theory yields instead **0.4583 ...**

$\Delta c_{T=T_c}/c_n$	Al	Nb	Pb
direct measurement	1.4	1.9	2.7
derived from measured $B_{\text{cth}}(0), T_c, \gamma$	1.6	1.9	2.4
derived from measured $\Delta(0), T_c$ with factor $6/\pi^2$	1.7	2.2	2.9
derived from measured $\Delta(0), T_c$ with factor 0.4583	1.3	1.7	2.2

larger numbers for Pb caused by stronger electron-phonon coupling (discussed later)

2.2 Type-I Superconductor in B_{ext}

calculation of $c_s(T)$ at low temperatures

$$c_n(T) - c_s(T) = -\frac{T}{\mu_0} \left[B_{\text{cth}} \frac{\partial^2 B_{\text{cth}}}{\partial T^2} + \left(\frac{\partial B_{\text{cth}}}{\partial T} \right)^2 \right]$$

$$B_{\text{cth}}(T) = B_{\text{cth}}(0) \left[1 - \left(\frac{T}{T_c} \right)^2 \right]$$

$$\frac{\partial B_{\text{cth}}(T)}{\partial T} = -B_{\text{cth}}(0) \frac{2T}{T_c^2}, \quad \frac{\partial^2 B_{\text{cth}}(T)}{\partial T^2} = -B_{\text{cth}}(0) \frac{2}{T_c^2}$$

$$c_n(T) - c_s(T) = -\frac{T}{\mu_0} \left[B_{\text{cth}}(0) \left[1 - \left(\frac{T}{T_c} \right)^2 \right] \left(-B_{\text{cth}}(0) \frac{2}{T_c^2} \right) + \left(-B_{\text{cth}}(0) \frac{2T}{T_c^2} \right)^2 \right]$$

$$c_n(T) - c_s(T) = -\frac{4}{T_c} \frac{B_{\text{cth}}^2(0)}{2\mu_0} \left[- \left[1 - \left(\frac{T}{T_c} \right)^2 \right] \left(\frac{T}{T_c} \right) + 2 \left(\frac{T}{T_c} \right)^3 \right] = -\frac{4}{T_c} \frac{B_{\text{cth}}^2(0)}{2\mu_0} \left[3 \left(\frac{T}{T_c} \right)^3 - \left(\frac{T}{T_c} \right) \right]$$

- with $\gamma = \frac{4}{T_c^2} \frac{B_{\text{cth}}^2(0)}{2\mu_0}$ we obtain $c_n(T) = \gamma T = \frac{4T}{T_c^2} \frac{B_{\text{cth}}^2(0)}{2\mu_0}$

$$c_s(T) = \frac{4}{T_c} \frac{B_{\text{cth}}^2(0)}{2\mu_0} \left[3 \left(\frac{T}{T_c} \right)^3 - \left(\frac{T}{T_c} \right) \right] + \frac{4T}{T_c^2} \frac{B_{\text{cth}}^2(0)}{2\mu_0} \quad \Rightarrow \quad c_s(T) = \frac{12}{T_c} \frac{B_{\text{cth}}^2(0)}{2\mu_0} \left(\frac{T}{T_c} \right)^3$$

- obviously, the phenomenological description predicts $c_s(T) \propto T^3$, therefore experimentalists initially tried to fit their data to a power law dependence
- later BCS predicts the correct exponential temperature dependence

2.2 Type-I Superconductor in B_{ext}

volume change at S/N phase transition:

- we use the general relation

$$V = \left(\frac{\partial G}{\partial p} \right)_{T, B_{\text{ext}}}$$

$$\text{and } \Delta \mathcal{G} = \mathcal{G}_n(T) - \mathcal{G}_s(T) = \frac{B_{\text{cth}}^2(T)}{2\mu_0}$$

$$\Rightarrow \left(\frac{V_n - V_s}{V_n} \right)_{T, B_{\text{cth}}(T)} = \frac{B_{\text{cth}}(T)}{\mu_0} \left(\frac{\partial B_{\text{cth}}(T)}{\partial p} \right)_{T, B_{\text{cth}}(T)}$$

→ volume change approaches zero for $T \rightarrow T_c$ since $B_{\text{cth}}(T) \rightarrow 0$

→ $T < T_c$: usually $V_s > V_n$ as $\frac{\partial B_{\text{cth}}(T)}{\partial p} < 0 \Rightarrow$ **superconductor has a larger volume**

typical relative volume change is very small: $\frac{V_n - V_s}{V_n} \approx 10^{-7} - 10^{-8}$

2.3 Type-II Superconductor in B_{ext}

thermodynamic properties of type-II superconductors

- $B_{\text{ext}} < B_{c1}$ (**Meißner-phase**): same behavior of type-I and type-II superconductors
 - $B_{c1} < B_{\text{ext}} < B_{c2}$ (**Shubnikov-phase**): different (more complicated) behavior of type-II superconductors
- ➔ functional form of $g_s(T, B_{\text{ext}})$ depends on details

