

Uniaxial Pressure on Strongly Correlated Materials

Zieve lab, UC Davis

Owen Dix

Miles Frampton

Scooter Johnson

Adrian Swartz

Rena Zieve

Samples

Adam Dioguardi, UC Davis

Jason Cooley, LANL

Todd Sayles, UCSD

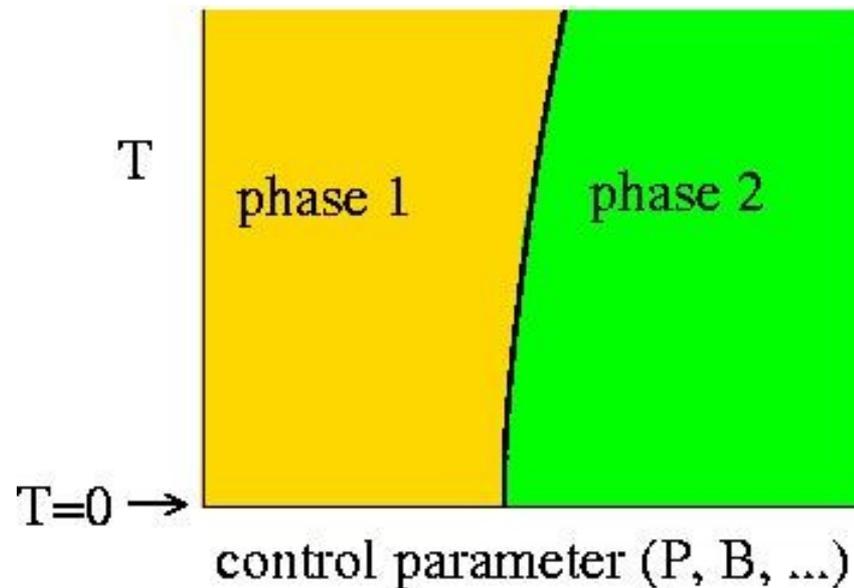
Brian Maple, UCSD

Funding from NSF, Division of Materials Research

Quantum Critical Points

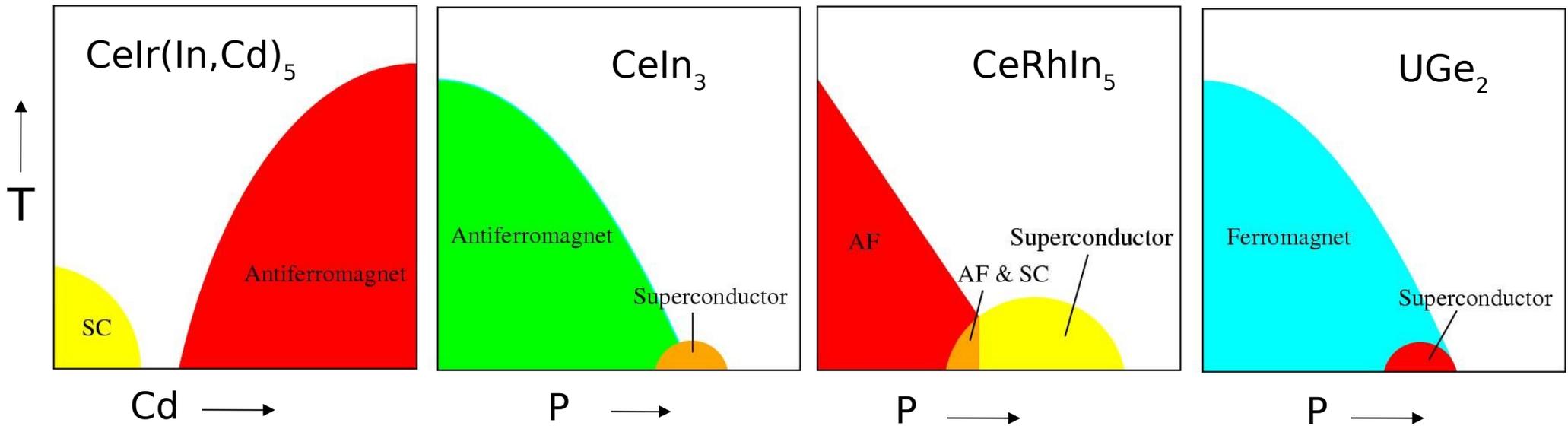
Phase Transitions at $T=0$

driven by control parameter: alloying, pressure, or magnetic field



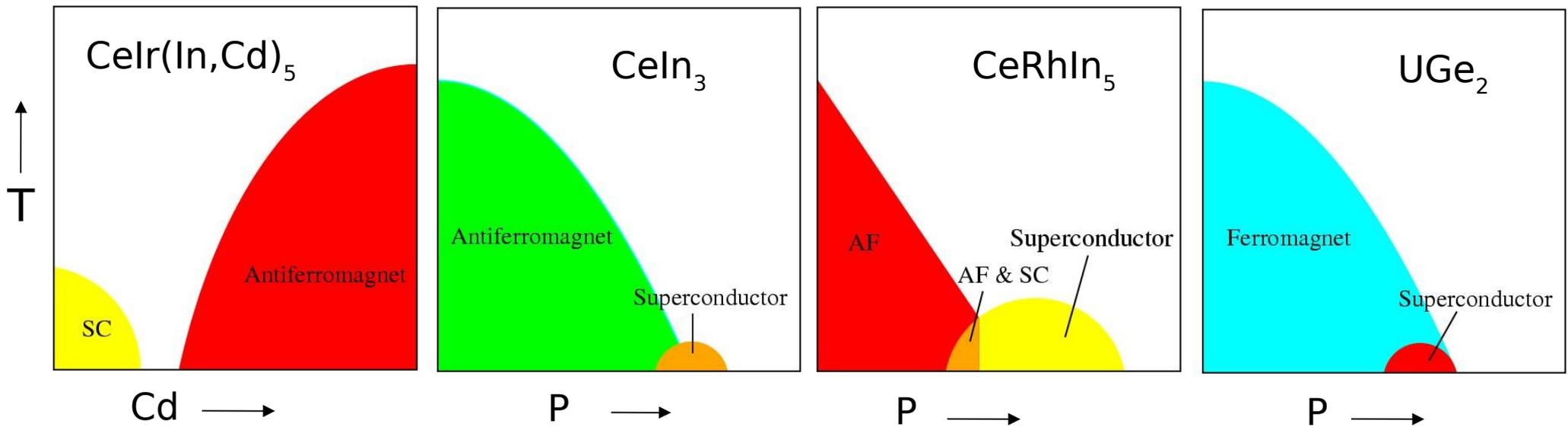
Can lead to a variety of unusual correlated behaviors

Phase diagrams follow certain trends...



- superconducting dome near disappearance of magnetism
- non-Fermi liquid regime in “normal metal”: unusual temperature dependence of resistivity, susceptibility, heat capacity, etc.
- unconventional superconductivity (nodes in energy gap, and resulting power-law temperature dependences)
- low dimensionality favors superconductivity

...but also have key differences



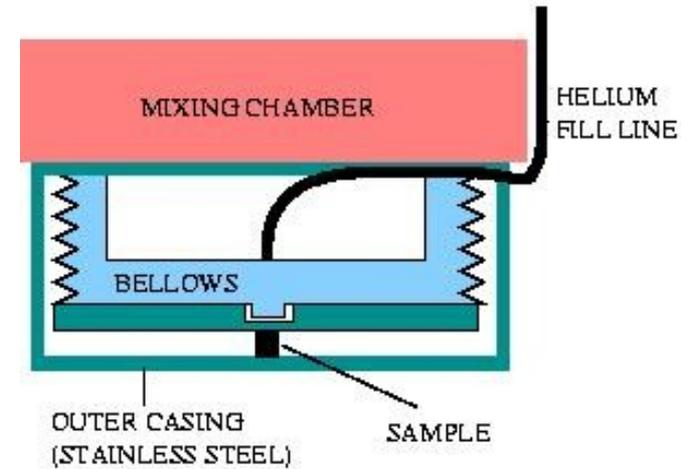
- different flavors of magnetism
- exact superconducting dome location varies; superconductivity may or may not coexist with magnetism
- widely different transition temperatures: <1K to >100K
- node structures differ among materials
- extra phase transitions in some compounds (structural; multiple magnetic or superconducting phases; or unknown phases)

Controlled studies needed!

- Changing samples alters
 - lattice constants or even crystal structure
 - electron concentration within sample
 - magnetic properties
 - sample purity considerations
- Methods for tuning an individual sample:
 - magnetic field
 - pressure (hydrostatic)
 - pressure (uniaxial)

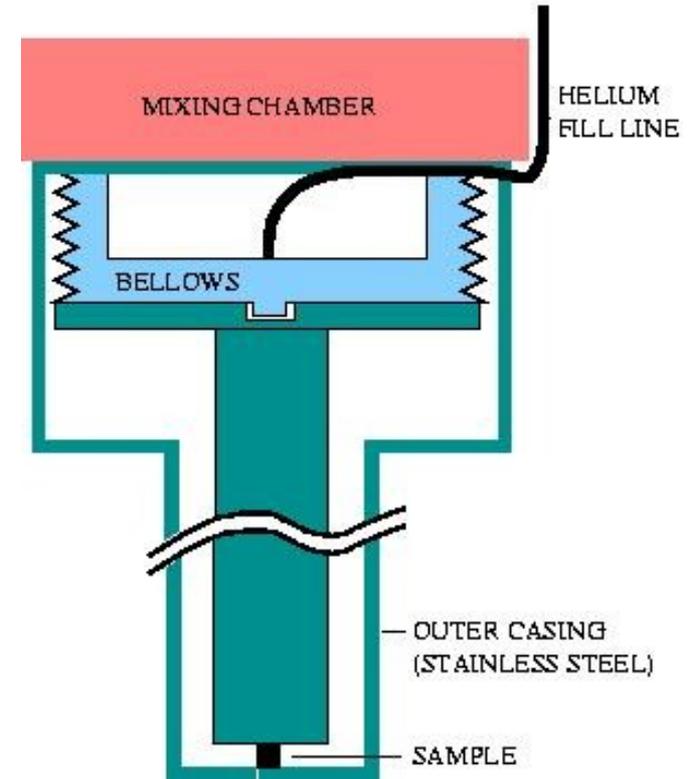
Pressure Cell Setup

- helium-activated bellows
- sample free to expand laterally



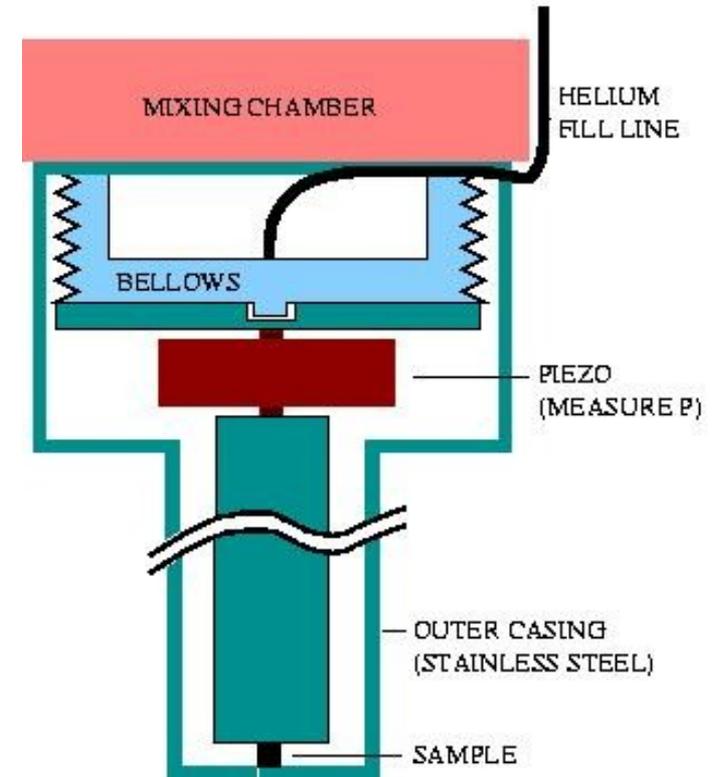
Pressure Cell Setup

- helium-activated bellows
- sample free to expand laterally
- centered in 8/10 Tesla magnet



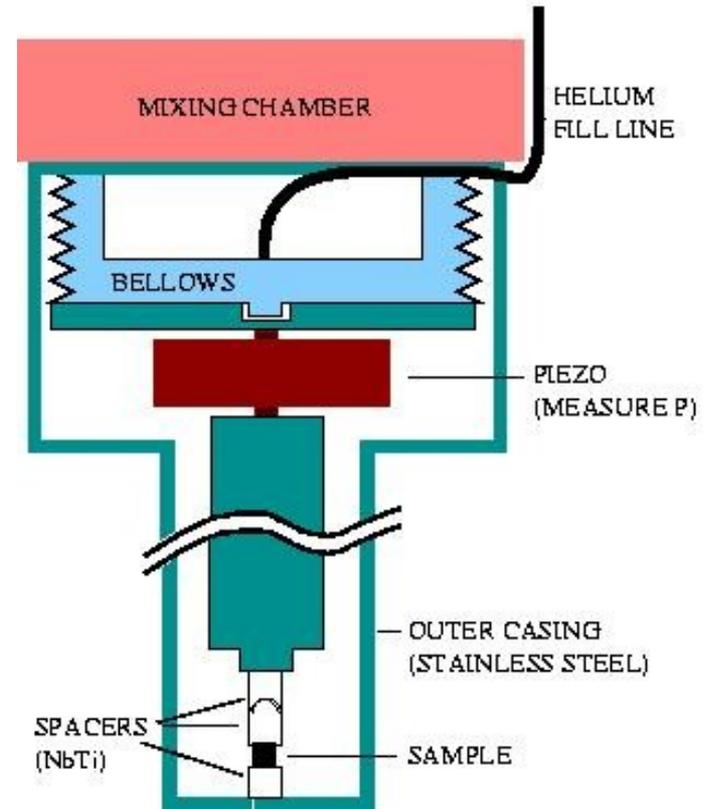
Pressure Cell Setup

- helium-activated bellows
- sample free to expand laterally
- centered in 8/10 Tesla magnet
- measure pressure with piezo



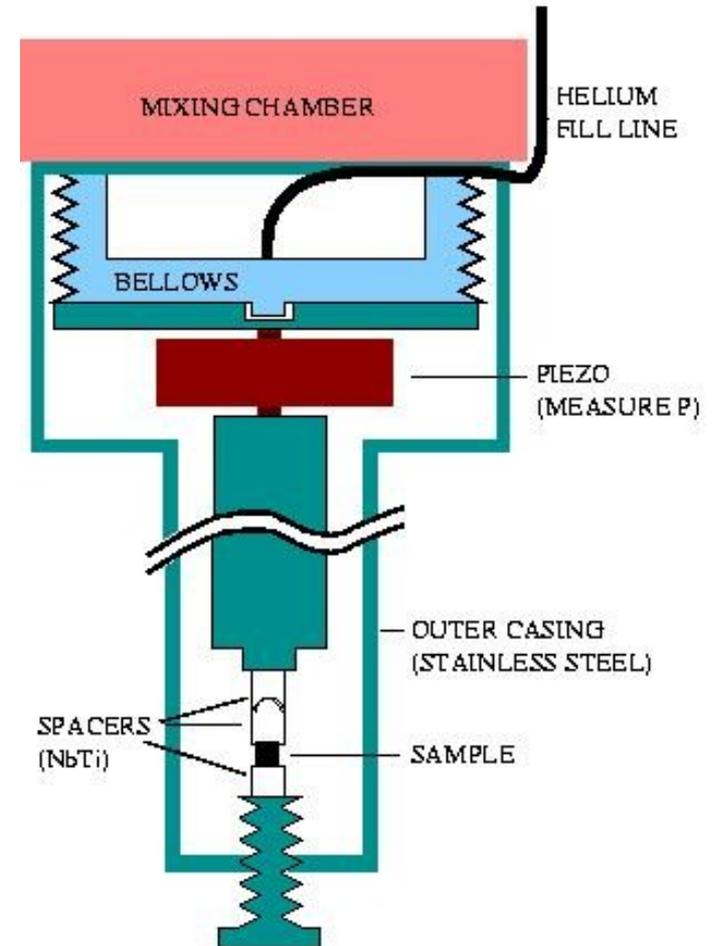
Pressure Cell Setup

- helium-activated bellows
- sample free to expand laterally
- centered in 8/10 Tesla magnet
- measure pressure with piezo
- spacers to adjust alignment



Pressure Cell Setup

- helium-activated bellows
- sample free to expand laterally
- centered in 8/10 Tesla magnet
- measure pressure with piezo
- spacers to adjust alignment
- adjust column length to cool under slight pressure
- cover *everything* with copper foil



Measurement Parameters

Sample size: at least 10 mg, after polishing, for heat capacity. Smaller samples can be used for χ measurements.

P_{\max} : about 10 kbar, depending on sample size. Helium solidifies at 25 bar, but area ratio between bellows and sample is several hundred. (Hydrostatic P goes much higher.)

ΔP : better than 0.1 kbar. (Much smaller than hydrostatic P.) We are limited by measurement sensitivity, not pressure step sizes.

Temperature range: have used setup from 100 mK to 200 K.

Temperature for pressure changes: as low as 200 mK.

Magnetic field: up to 10 T, for measurements below 4 K only.

Measure: resistivity, susceptibility, heat capacity

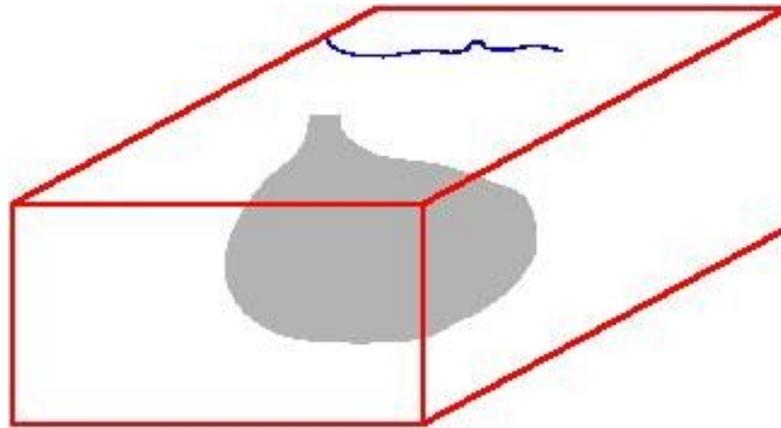
Sample Considerations

Polishing

two flat, parallel surfaces, for pressure application
all other surfaces perpendicular to these, for constant cross-section

No cracks

No occlusions

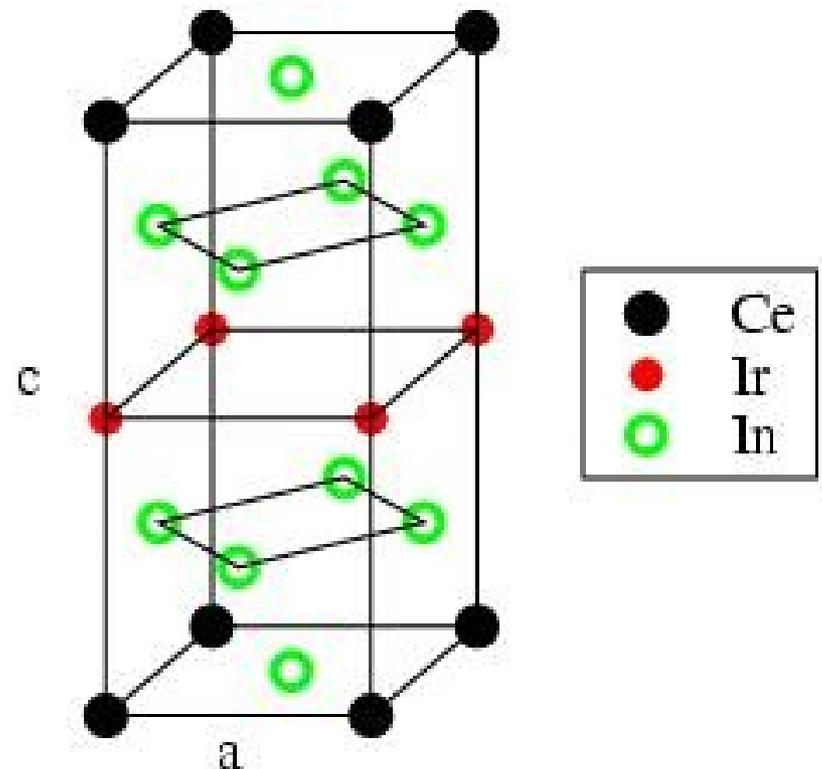


Need significant final size, for decent signal-to-noise

Often want single crystal samples

Ce(Ir,Rh,Co)In₅

- several types of behavior in a single crystal structure
- relatively easy to make; clean
- layered structure, similar to high-T_c superconductors
- special behavior:
 - ◆ disagreement in T_c
from ρ , C
 - ◆ exotic vortex phase in
CeCoIn₅ at high fields



Lattice Constants and Superconductivity

Trends in T_c :

CeCoIn₅ 2.3 K

CeIrIn₅ 0.4 K

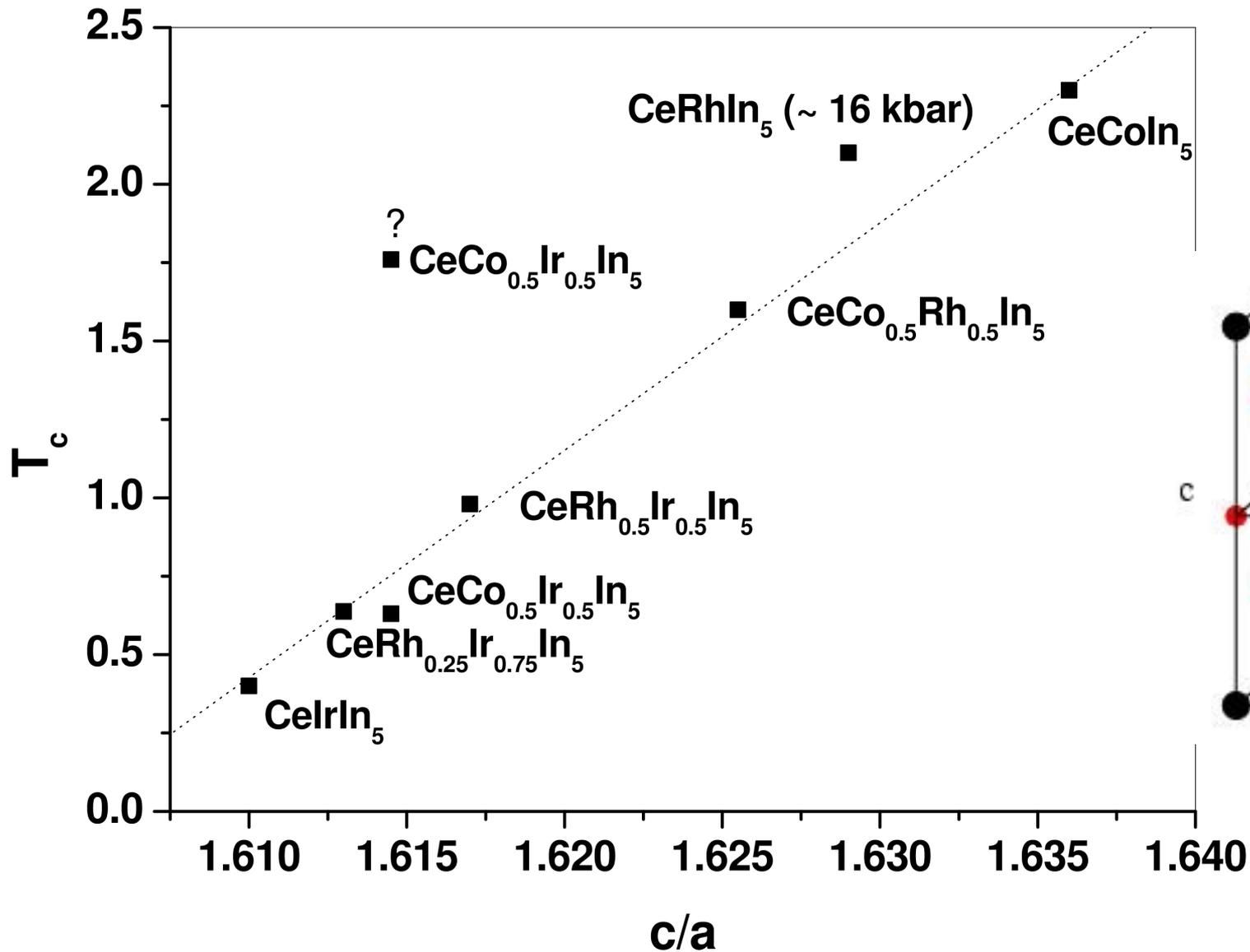
CeRhIn₅ low mK or with P

Doesn't match ion size, or individual lattice parameters.

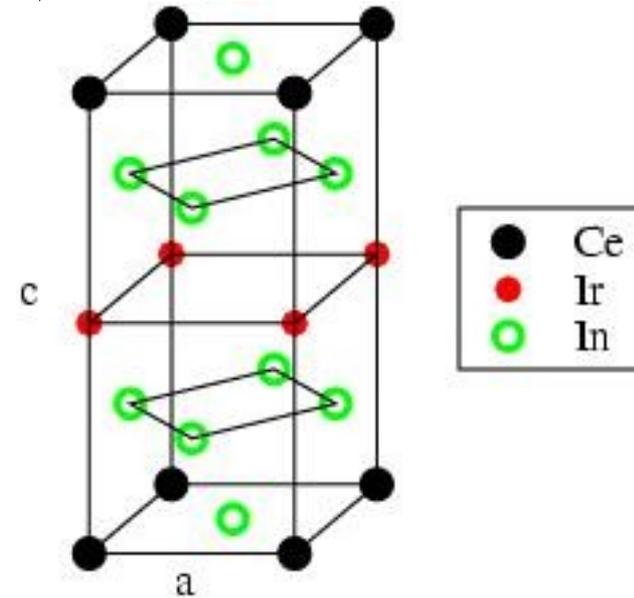
- **T_c linear in c/a for pure compounds and alloys**
- uniaxial P: vary c/a without sample-to-sample variations
- predictions from thermal expansion:

$$\frac{\partial T_c}{\partial P_a} = 54 \text{ mK/kbar}$$

$$\frac{\partial T_c}{\partial P_c} = -89 \text{ mK/kbar}$$



CeMIn₅
(M=Co, Ir, Rh)



[after Pagliuso et al., *Physica B* **131-132**, 129 (2001)]

To change c/a, use uniaxial pressure (NOT hydrostatic).

Lattice Constants and Superconductivity

Trends in T_c :

CeCoIn₅ 2.3 K

CeIrIn₅ 0.4 K

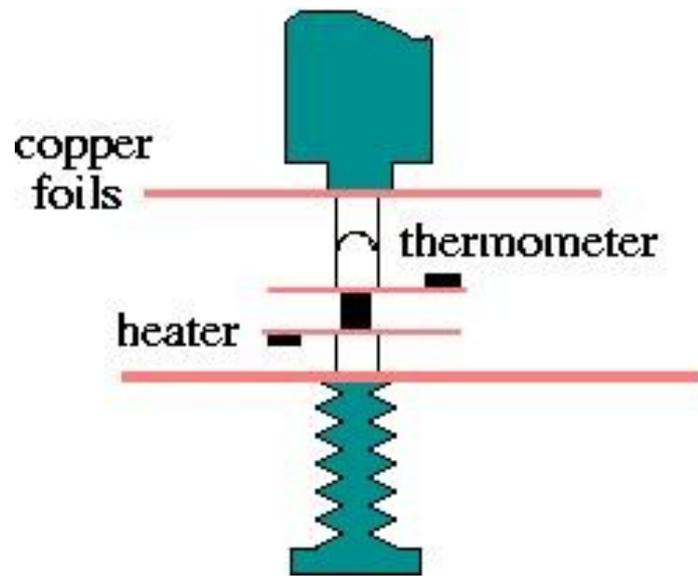
CeRhIn₅ low mK or with P

Doesn't match ion size, or individual lattice parameters.

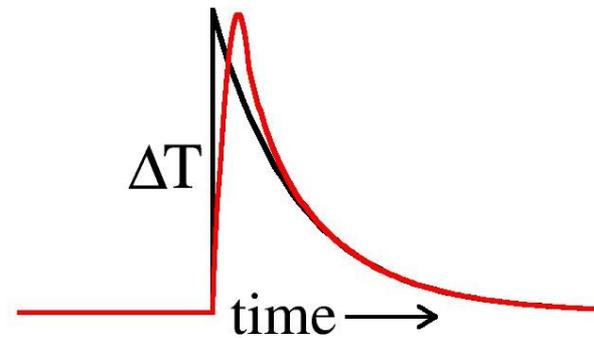
- **T_c linear in c/a for pure compounds and alloys**
- uniaxial P: vary c/a without sample-to-sample variations
- predictions from thermal expansion:

$$\frac{\partial T_c}{\partial P_a} = 54 \text{ mK/kbar}$$

$$\frac{\partial T_c}{\partial P_c} = -89 \text{ mK/kbar}$$



Thermometer - RuO₂ film
Heater - 50:50 Au:Cr alloy

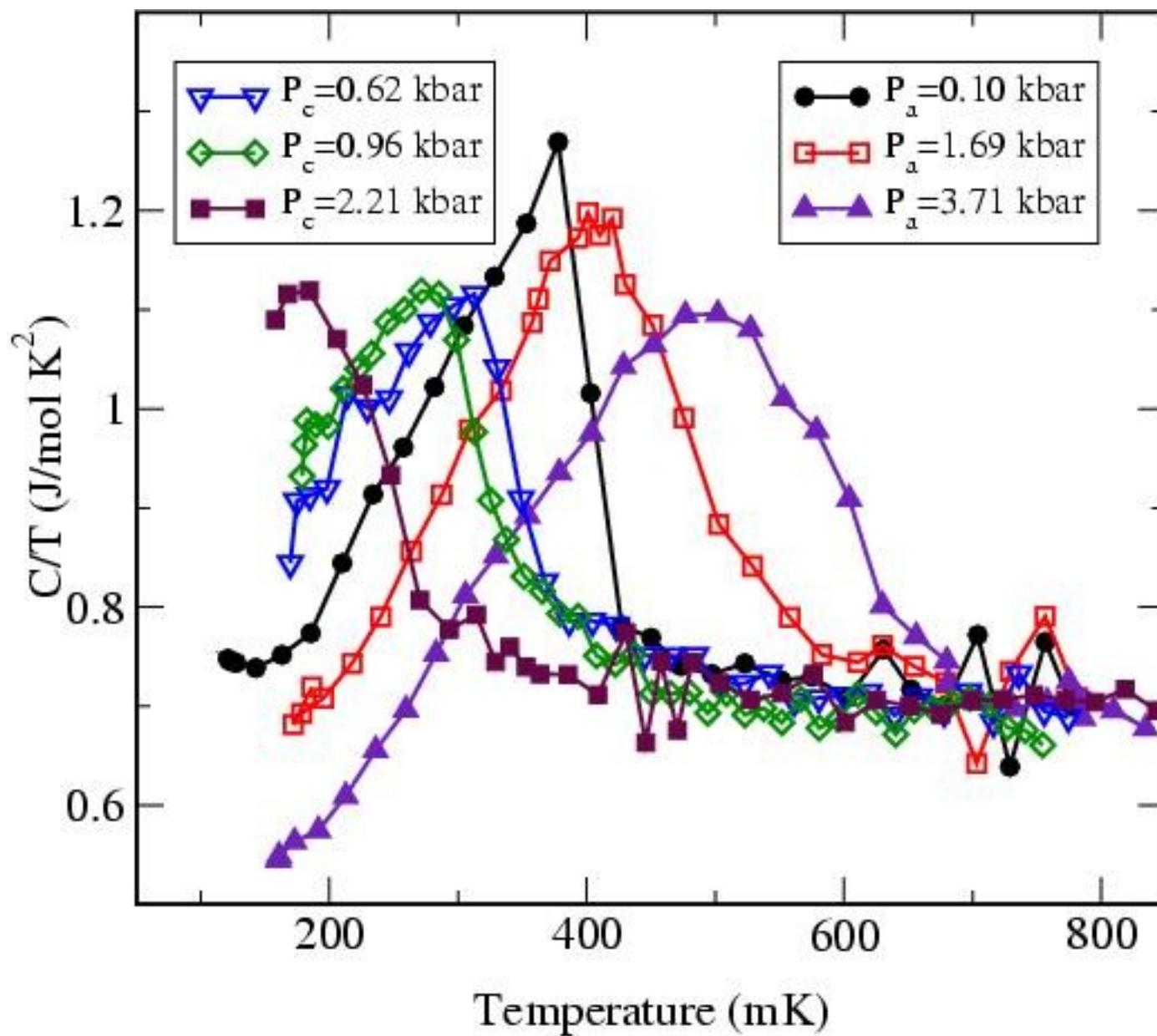


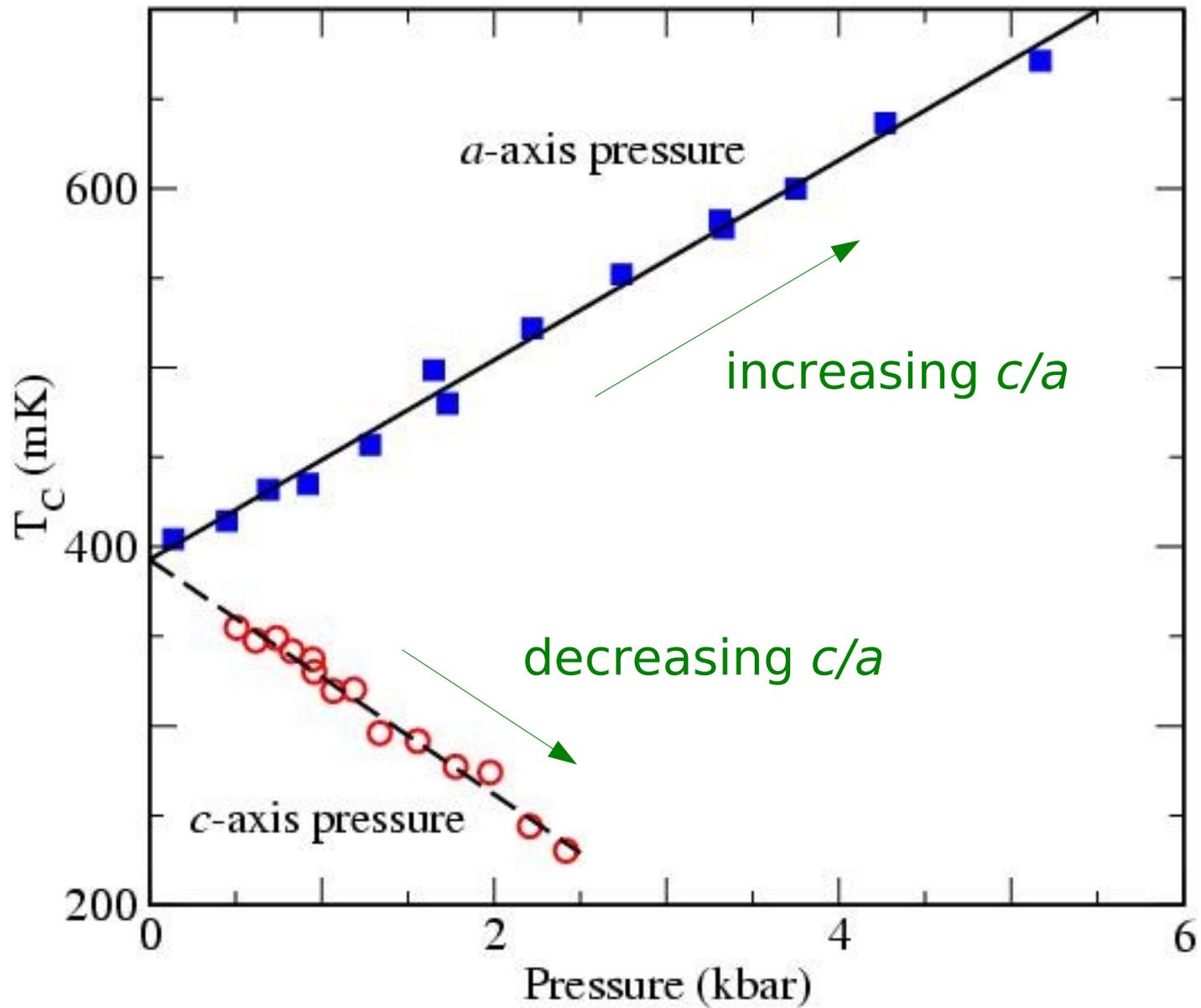
Use short heat pulse followed by exponential decay.

Time constant proportional to sample size.

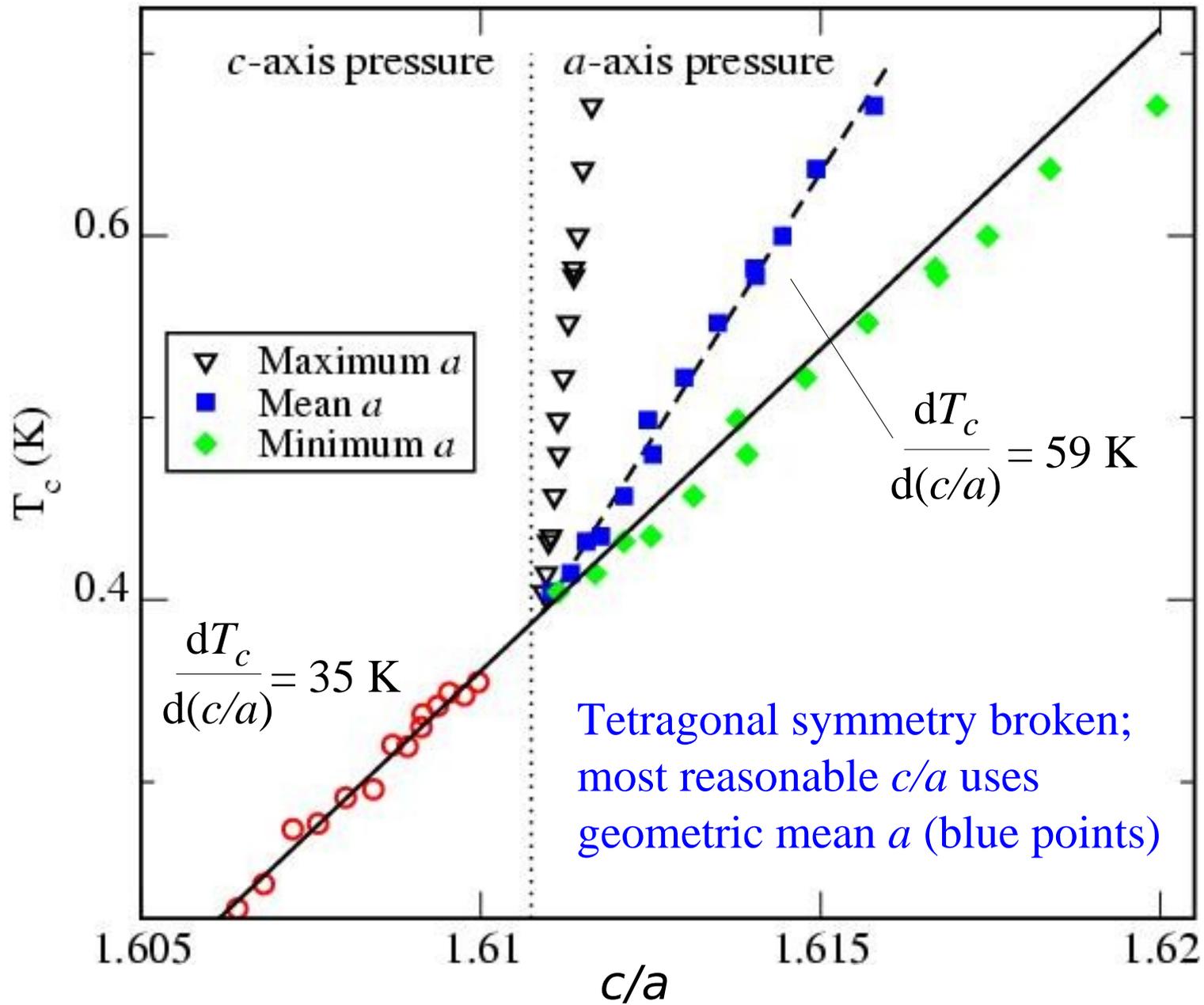
Place sample between heater and thermometer to reduce initial spike, but still have large background.

CeIrIn₅





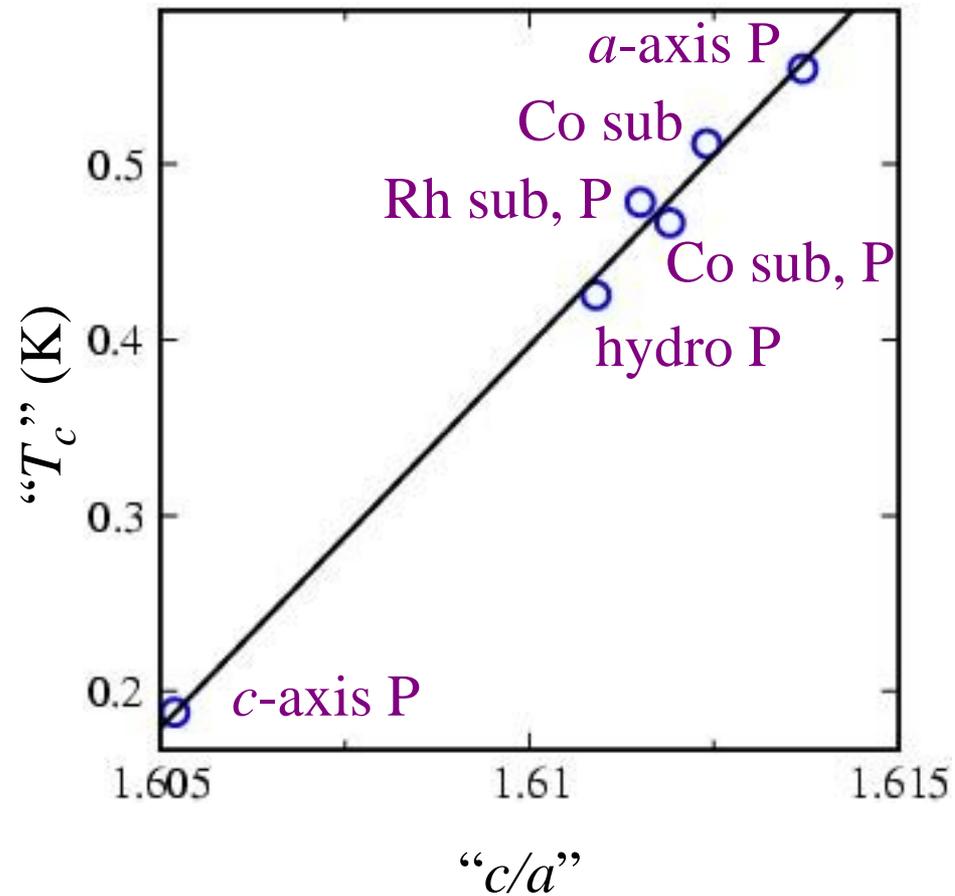
Linear change of T_C with pressure for both directions



Kink at zero pressure may indicate influence of hybridization

Calculate c/a and T_c
if control parameters
were scaled so all
achieved the same
hybridization

Uniaxial work
important: gives
extreme points



With hybridization variations removed, the resulting linear relationship shows the influence of dimensionality:

$$\frac{\partial T_c}{\partial (c/a)} = 44 \text{ K}$$

Further thermal expansion predictions

CeIrIn₅

$$\frac{\partial T_c}{\partial P_a} = 54 \text{ mK/kbar}$$

$$\frac{\partial T_c}{\partial P_c} = -89 \text{ mK/kbar}$$

CeCoIn₅

$$\frac{\partial T_c}{\partial P_a} = 29 \text{ mK/kbar}$$

$$\frac{\partial T_c}{\partial P_c} = -7.5 \text{ mK/kbar}$$

Why so small??

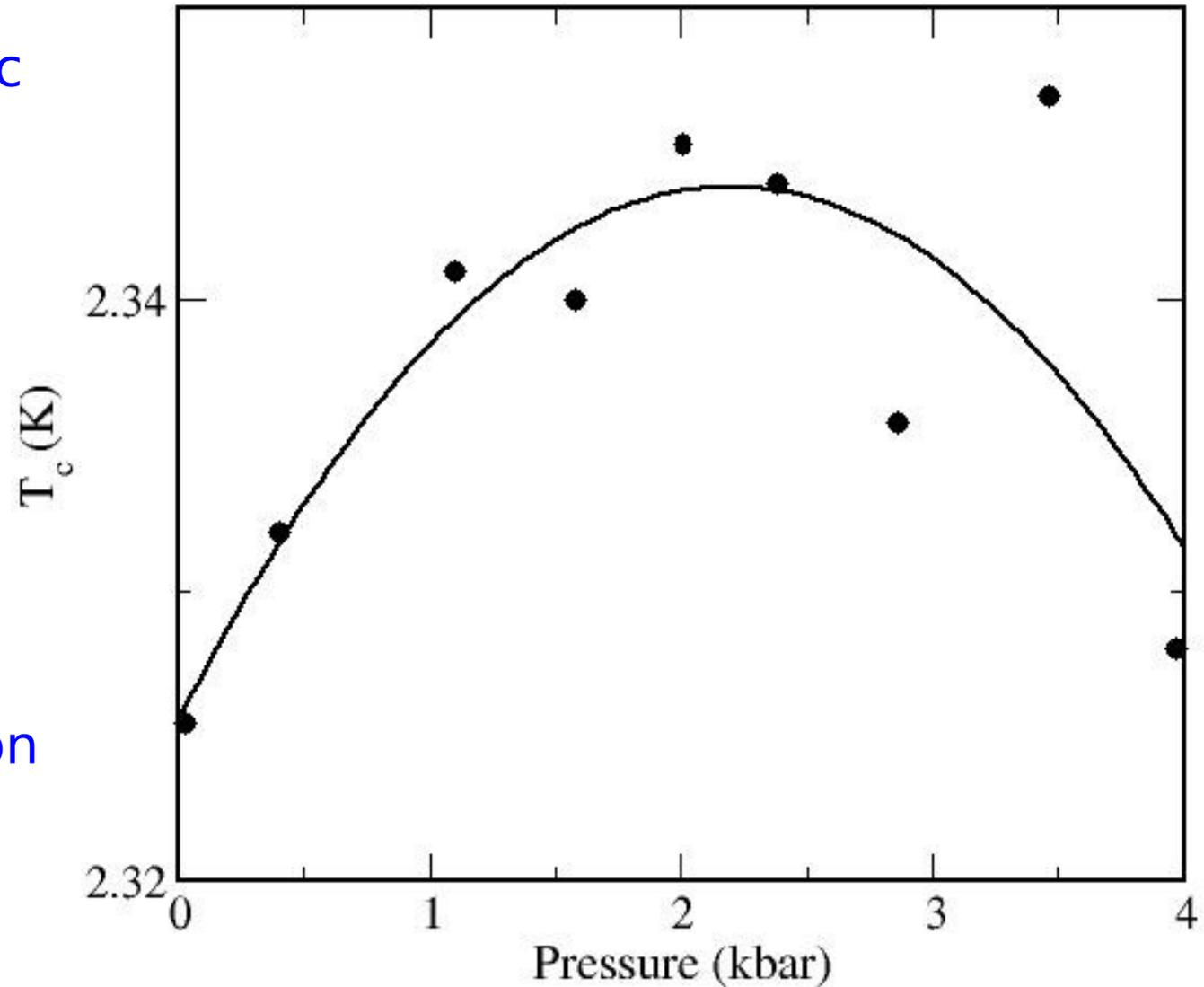
CeCoIn₅

Measure magnetic susceptibility

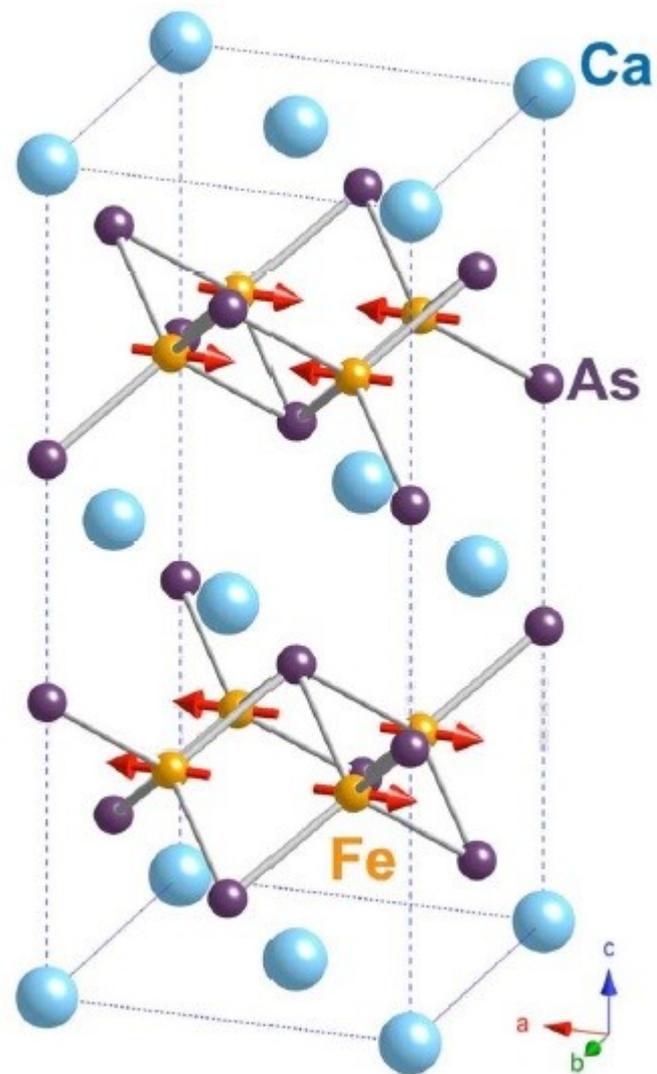
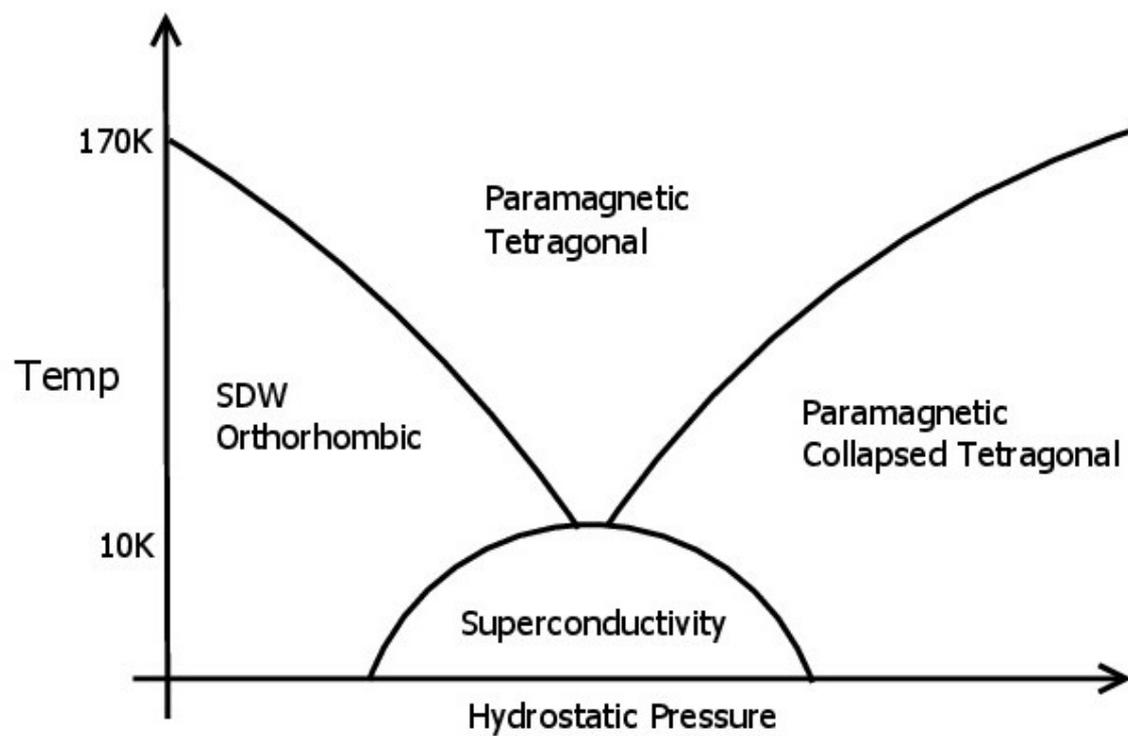
Very little P-dep
(20 mK rather than 200 mK)

Not linear!
Sample "tuned"
unusually well

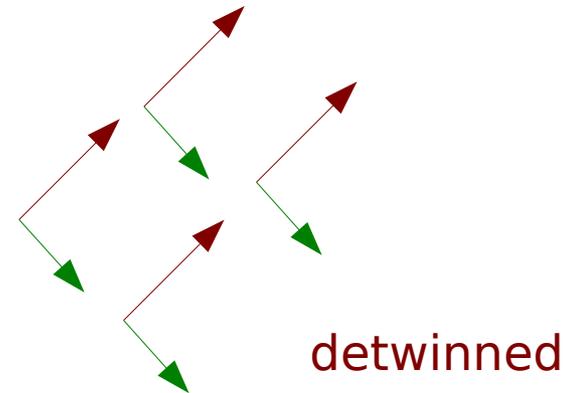
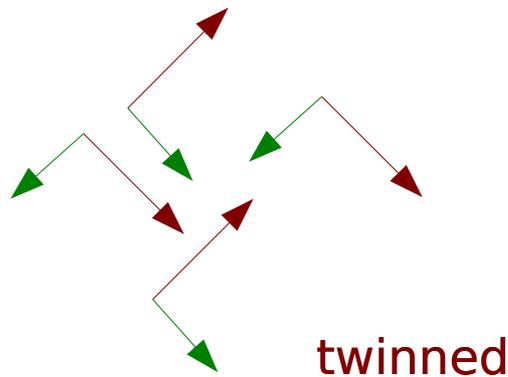
Thermal expansion
only gives info on
P=0 slope.



CaFe₂As₂



CaFe₂As₂

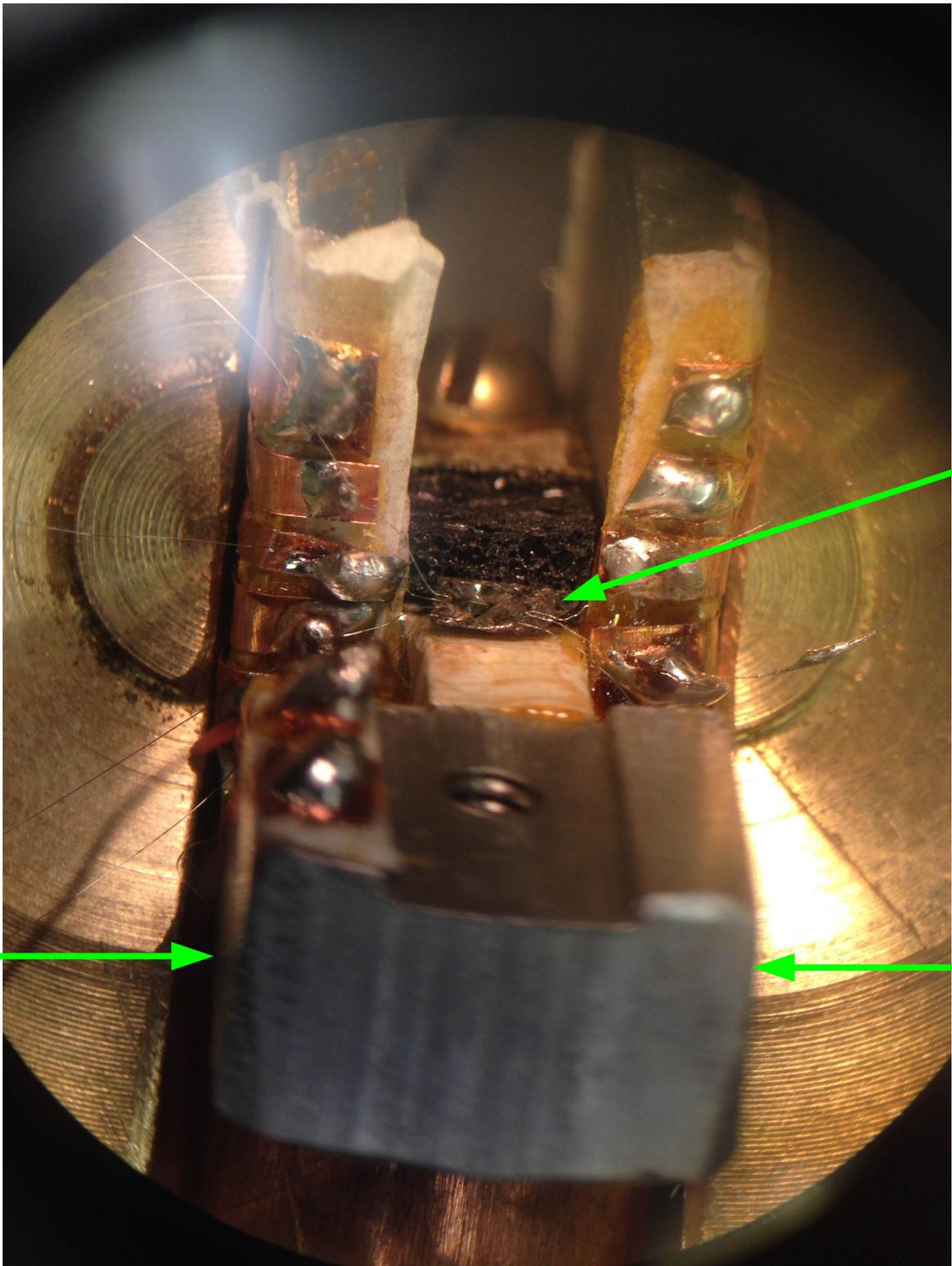


In orthorhombic phase, detwinned crystals have different $\rho(T)$ along the a and b axes.

In-plane pressure can accentuate the a - b difference.

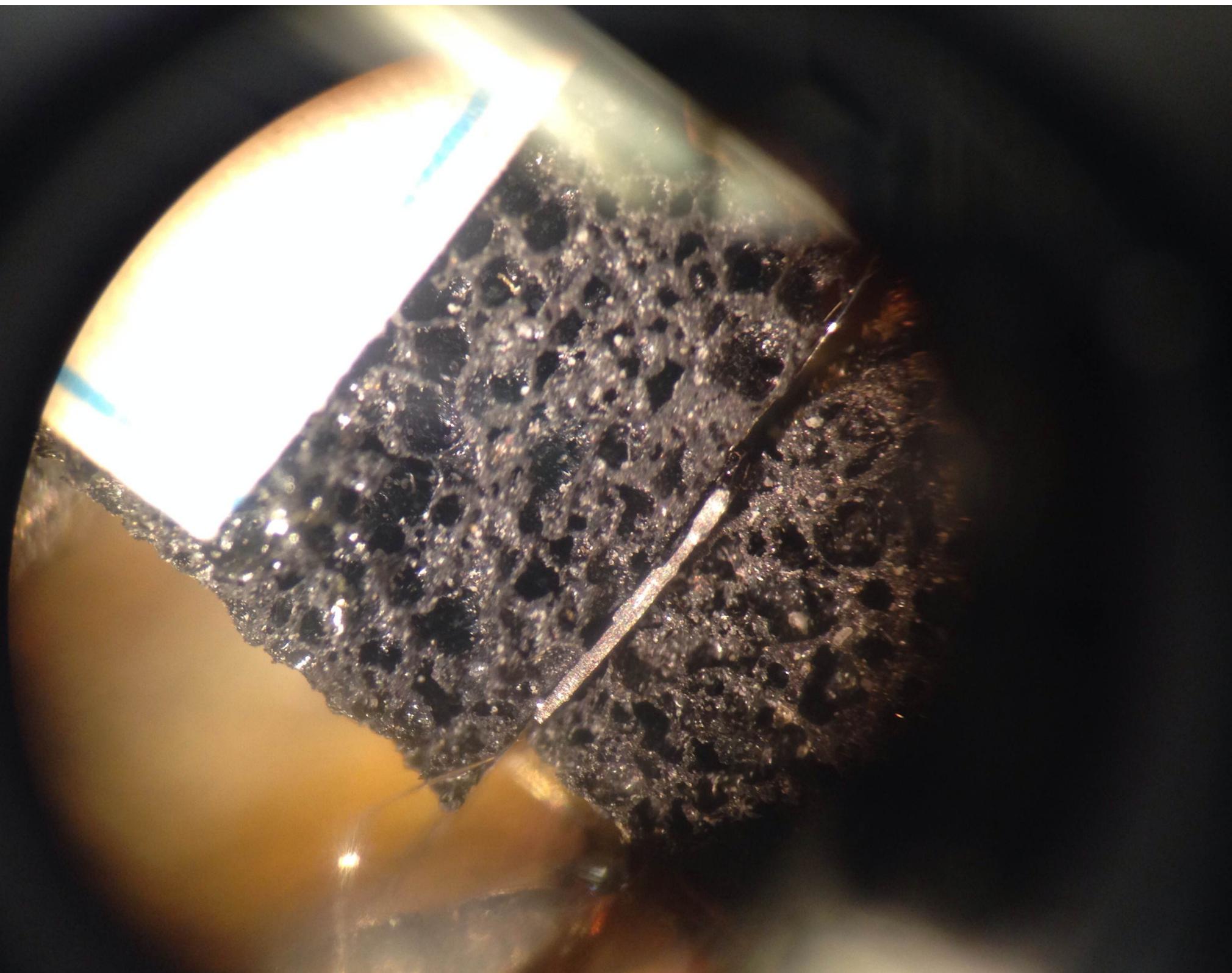
A technical challenge: platelet samples! (Unfortunately common among materials with layered crystal structures.)

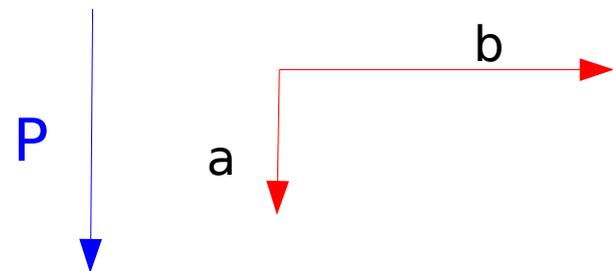
Use foam to keep alignment.



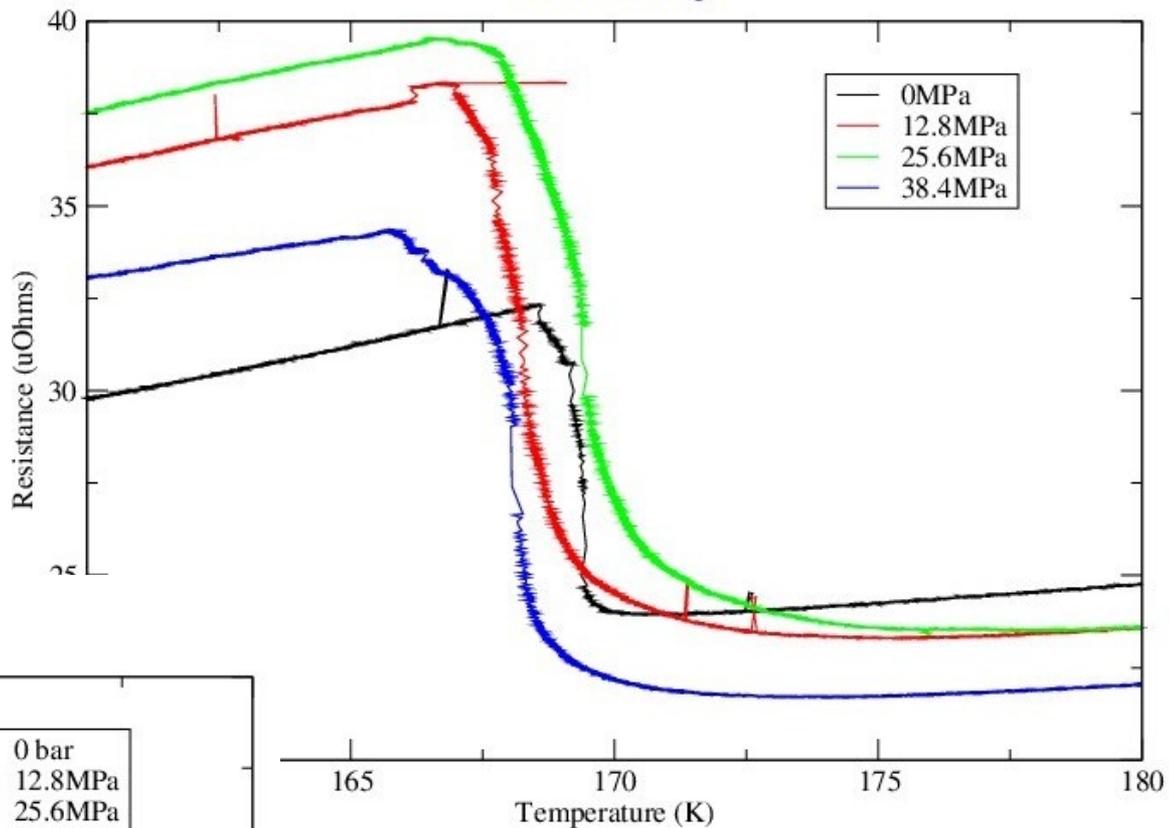
sample

$\frac{1}{2}$ inch

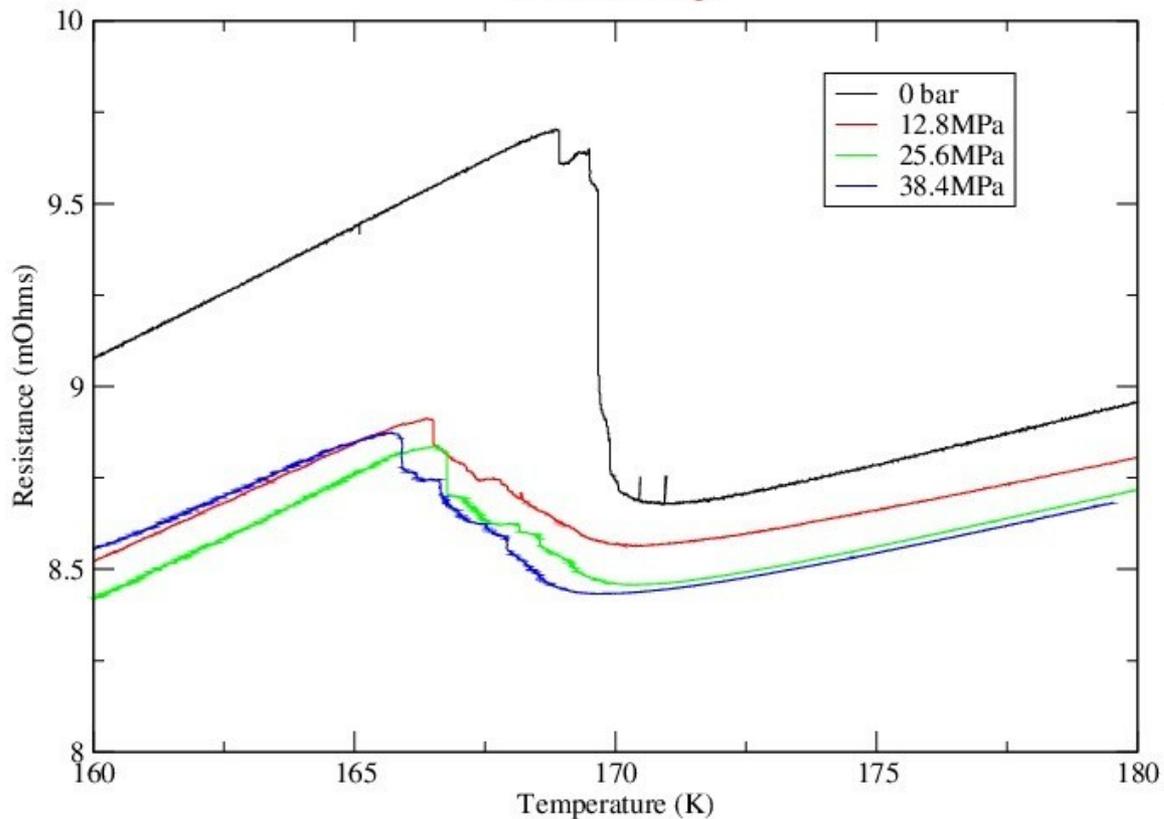




Resistance vs. Temperature
a-axis (warming)

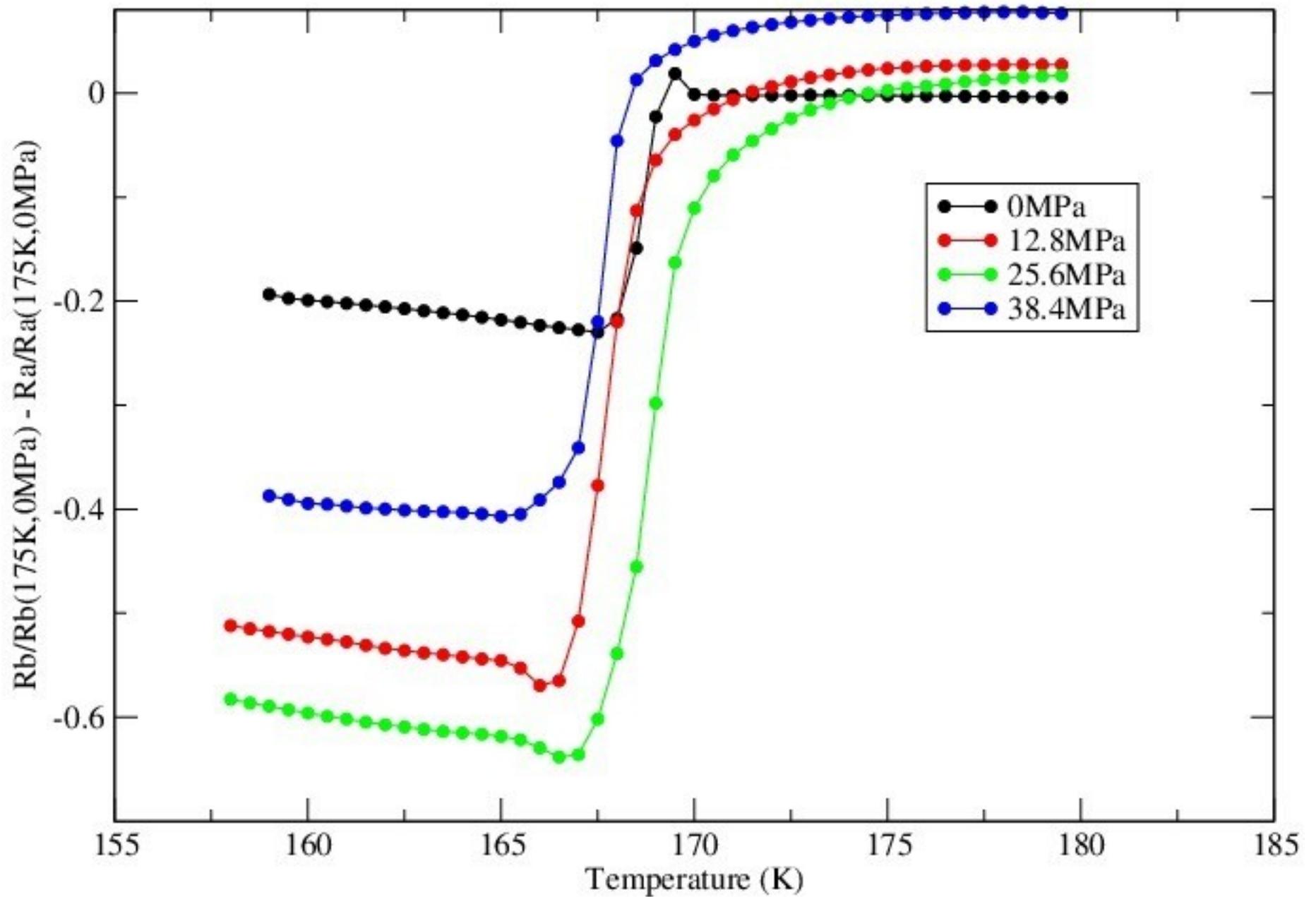


Resistance vs. Temperature
b-axis (warming)



Anisotropy vs. Resistance

Warming



Conclusions

- Uniaxial pressure provides a unique probe of strongly correlated systems.
- Our setup runs from below 100 mK to over 200 K, and to pressures of 1 GPa. At low temperature a magnetic field can also be applied along the pressure axis.
- We observe how dimensionality favors superconductivity in 115 materials and measure anisotropy from in-plane symmetry breaking in iron pnictides.