

Critical current anisotropy in clean superconducting pnictide thin films

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Outline

- Why pnictide thin films?
- Deposition methods
- Strain dependence of T_c
- Scaling of J_c
- Intrinsic Pinning

Acknowledgment



PLD of 11 and 122:

**K. Iida, F. Kurth, J. Engelmann, S. Trommler,
O. Rodriguez, S. Molatta**

Targets: M. Schulze, S. Wurmehl

Pulsed field:

A. Kauffmann, J. Freudenberg, N. Kozlova

TEM: T. Thersleff, E. Reich

Discussions:

S. Haindl, L. Schultz, B. Holzapfel

High-field: C. Tarantini, J. Jaroszynski

MBE of 1111: S. Ueda, M. Naito

TEM: A. Ichinose, I. Tsukada

Discussion: M. Weigand, B. Maiorov

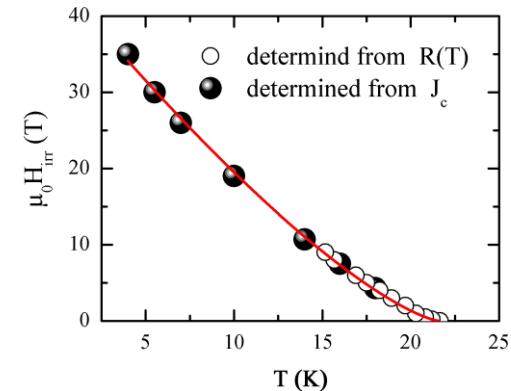
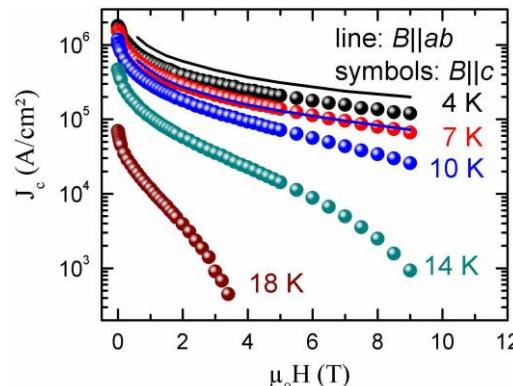
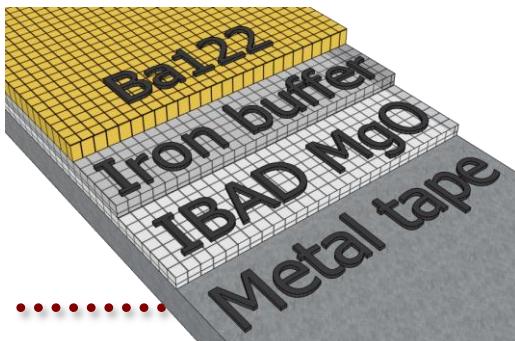
Why pnictide thin films?

Research:

- **Geometry:** allows measurements on large areas (spectroscopy) or thin filaments (transport critical currents)
- **Dimensionality:** allows investigation of 2D-3D crossover ($d \sim \lambda, \xi, I_0 \dots$)
- **Kinetics:** allows to grow metastable compounds, heterostructures and phases not available as single crystals

Applications (low G_i, g , high T_c, H_{irr}):

- **Electronics:** Projects on SQUIDs, bolometers, ...
- **Wires and tapes:** Coated conductors possible (prototypes already demonstrated)



Fe-based sc thin film efforts around the world

Madison:
122 on oxide
buffers, GBs

Texas
A&M:
11

Mexico:
CSD, 1111
on oxides

Brookhaven:
11 on oxides,
RABiTS and
IBAD

Genova:
11 on
oxides

Tokyo IoT:
122 on
oxides, IBAD,
GBs

Tokyo, Kyushu,
and CRIEPI:
11 on oxides

Darmstadt:
112 on
oxides

Beijing:
MOCVD,
11 on GaAs

Sunkyunkwan
(Korea):
Sr122 K:Ba122

Tokyo:
ED, 122 on
RABiTS

Bogota:
Sputtering,
122 on oxides

IFW Dresden:
11, 122, 1111.
Fe buffer, GBs,
IBAD

Nagoya:
P:Ba122 and
1111

ISTEC-SRL:
PLD, P:Ba-
122 MBE, Ca-
122

Tokyo A&T:
K:Ba-122, FeSe,
1111,
flouride substrates

PLD

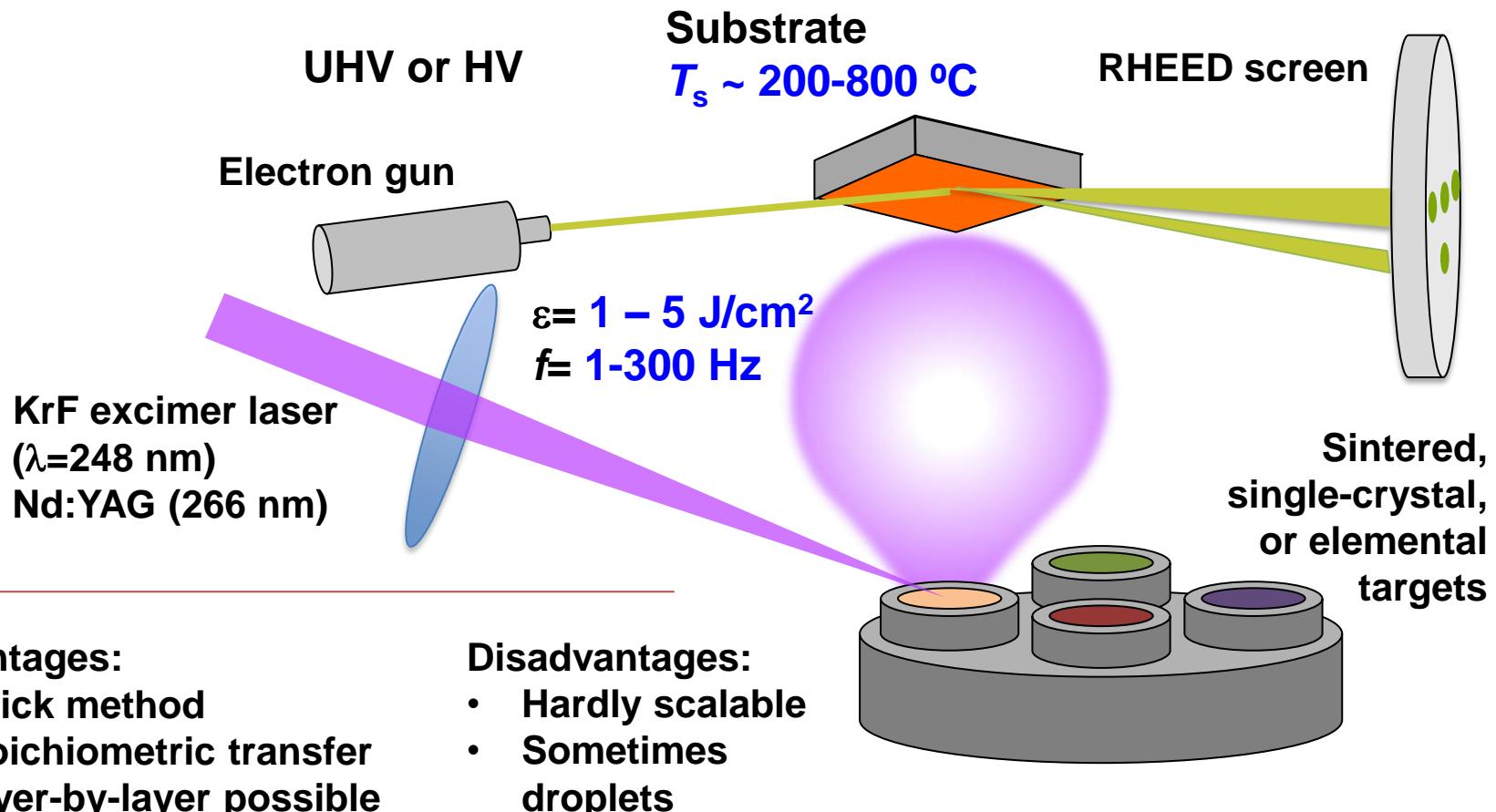
MBE

New methods

So far no films of 111 and 32522

Pulsed Laser Deposition (PLD)

Focussing or imaging of pulsed laser beam on target → stoichiometric plasma of target material → deposition on substrate



Advantages:

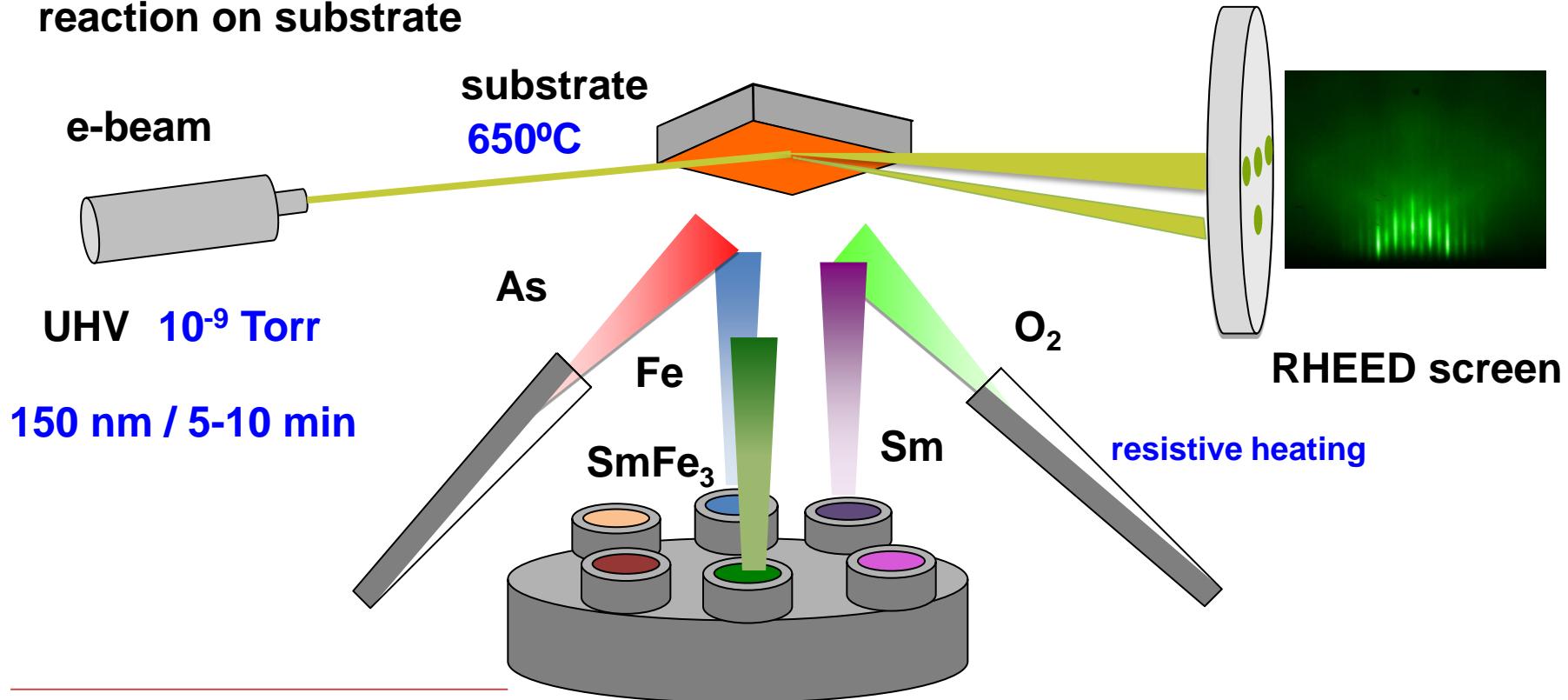
- Quick method
- Stoichiometric transfer
- Layer-by-layer possible

Disadvantages:

- Hardly scalable
- Sometimes droplets

Molecular Beam Epitaxy (MBE)

Individual sources (e.g. Knudsen cells) → atomar or „molecular“ beams → reaction on substrate



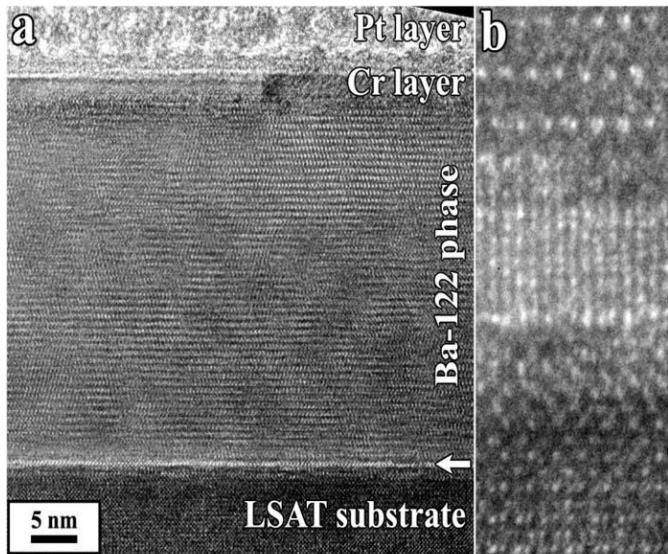
Advantages:

- Slow deposition rates possible
- Clean films
- Good epitaxy

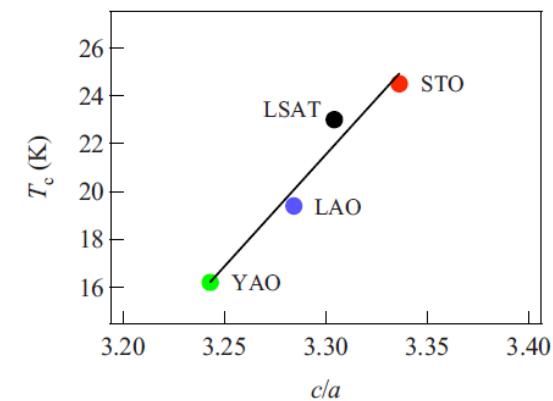
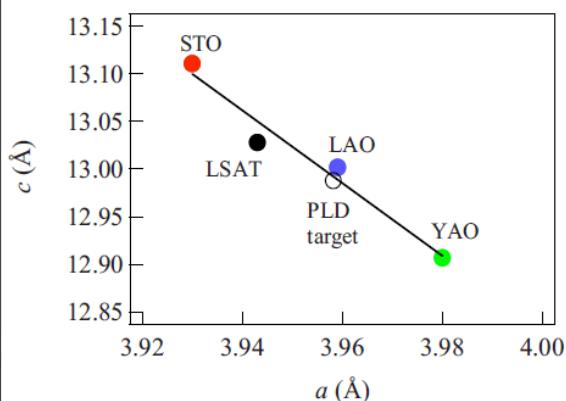
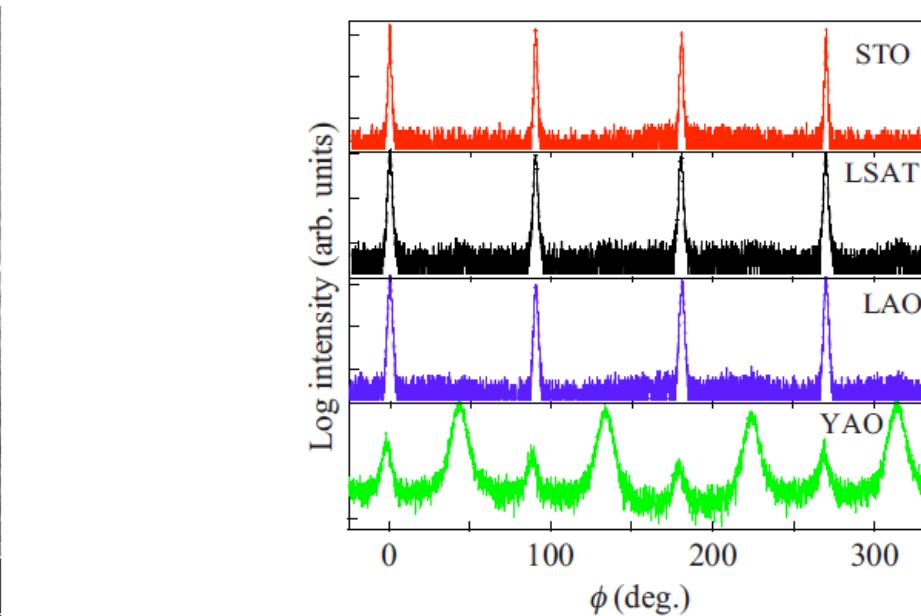
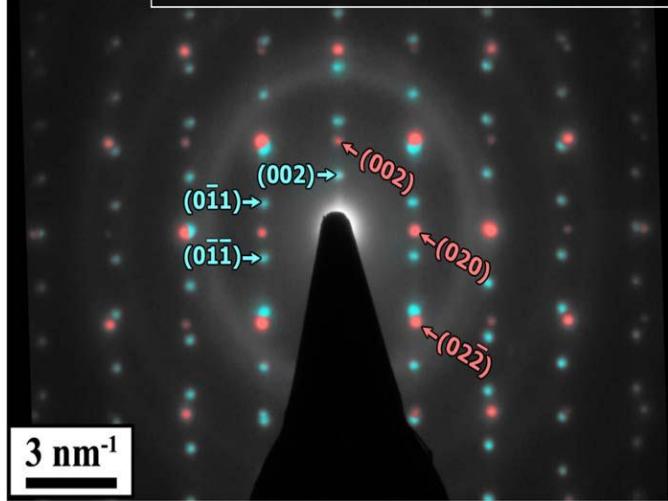
Disadvantages:

- Rather expensive
- UHV needed (low rates!)

Strain dependence of T_c - Oxides



c Ba-122 [1 0 0] || LSAT [1 0 0]

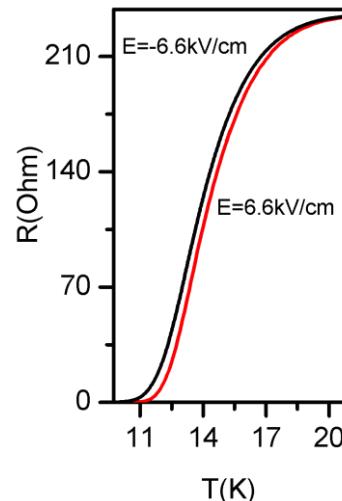
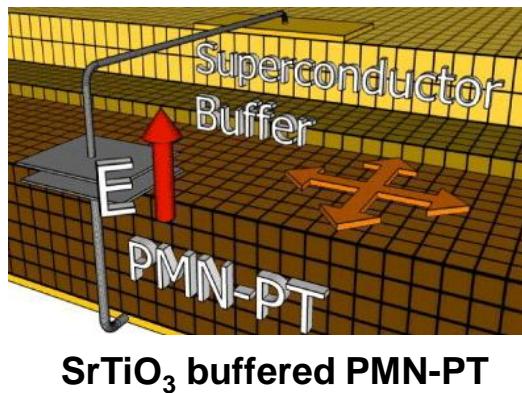
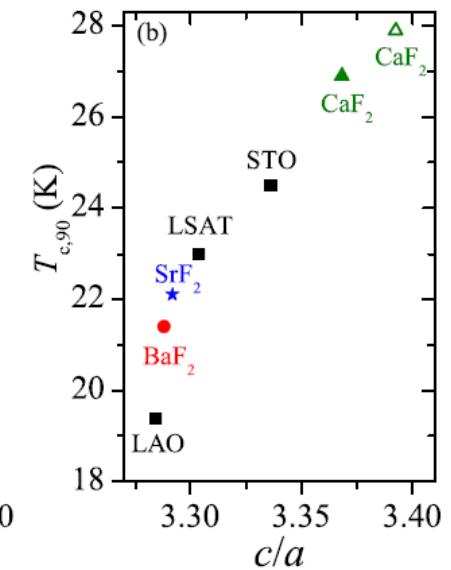
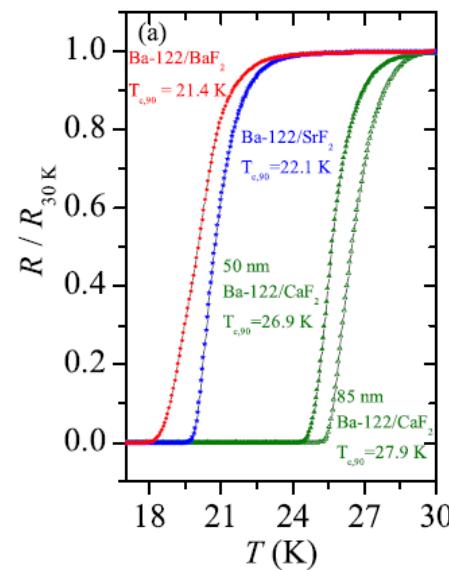
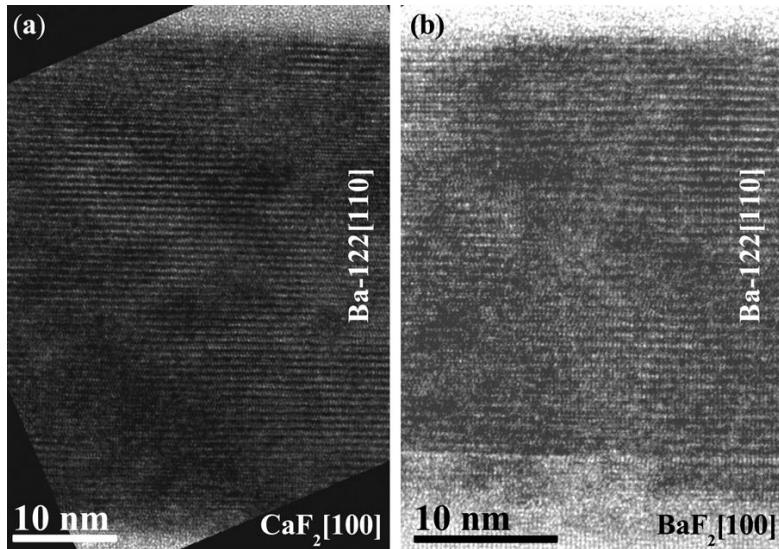


K. Iida et al., APL 95, 192501 (2009)

K. Iida et al., PRB 81, 100507(R) (2010)

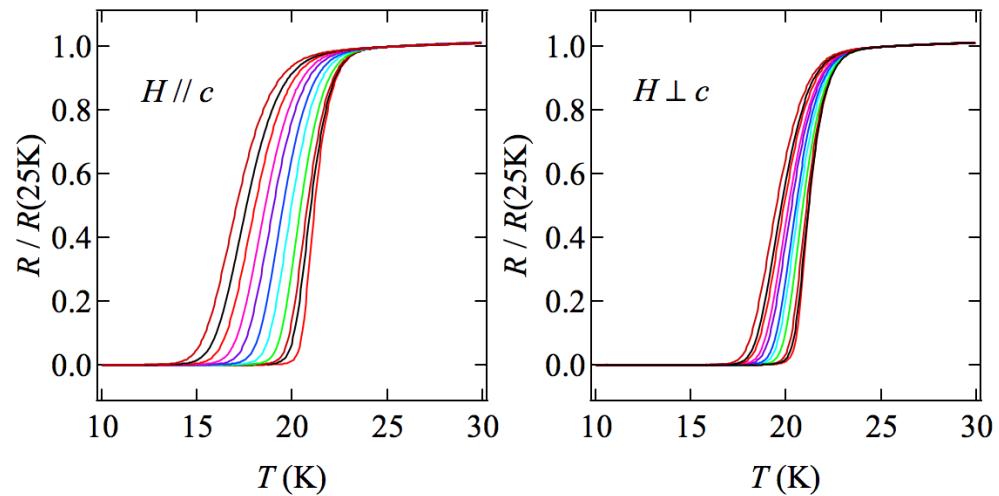
T. Thersleff et al., APL 97, 022506 (2010)

Strain dependence of T_c – Flourides and Piezo

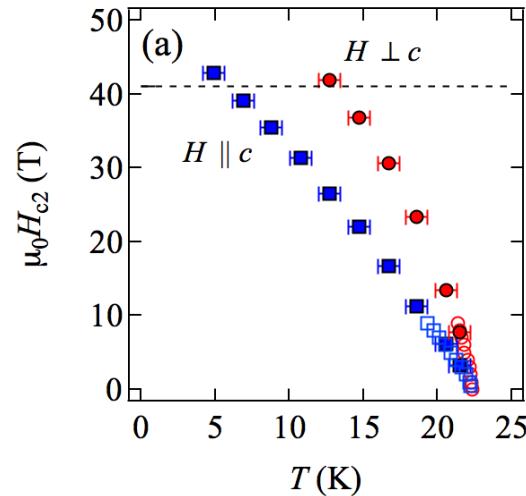
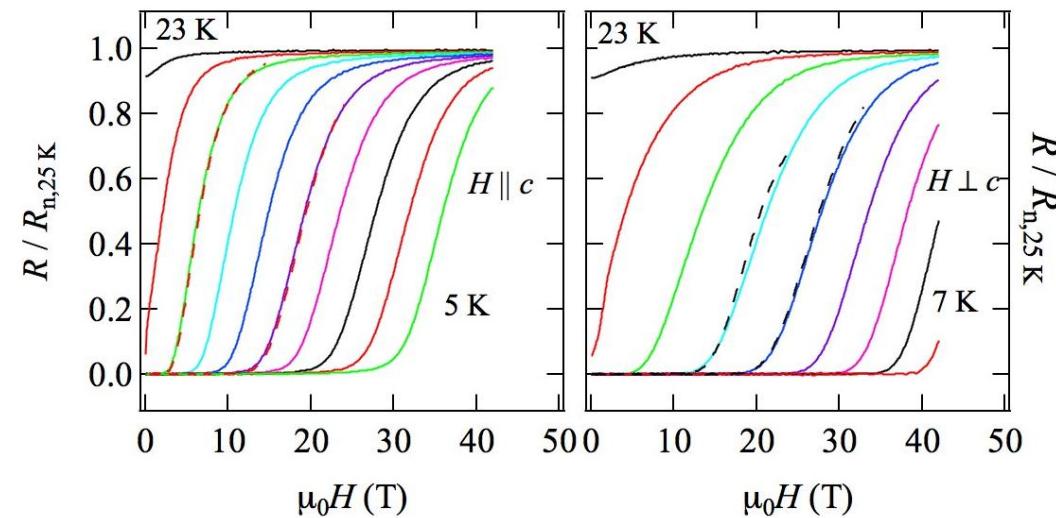


$$\frac{\Delta T_c}{\epsilon_{ab}} = 17 \frac{K}{\%}$$

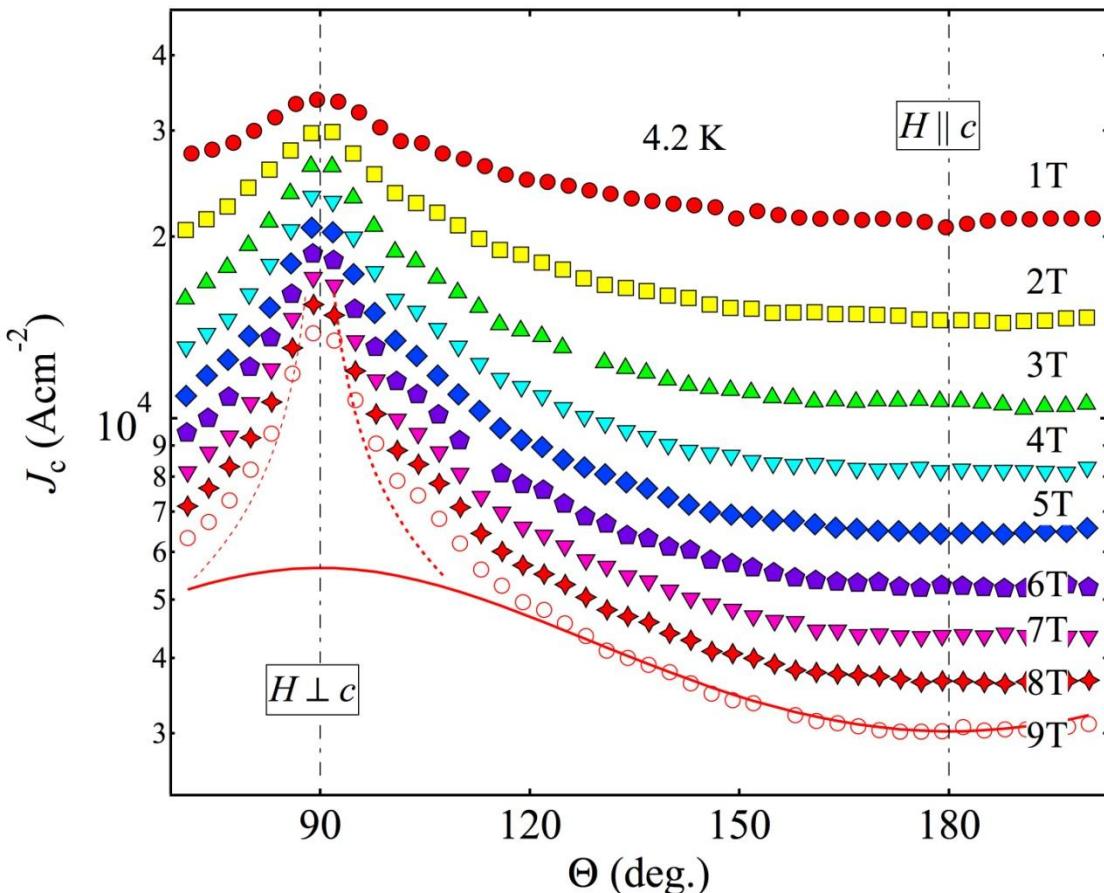
Ba-122: Resistive Measurements



- $R(T)$ static fields ≤ 9 T (PPMS)
- $R(B)$ pulsed fields ≤ 45 T
- T_c , H_{c2} defined at 90% $R(25\text{K})$
- Ba122 shows anisotropic H_{c2}
 $H_{c2}^{\parallel c} < H_{c2}^{\perp c}$

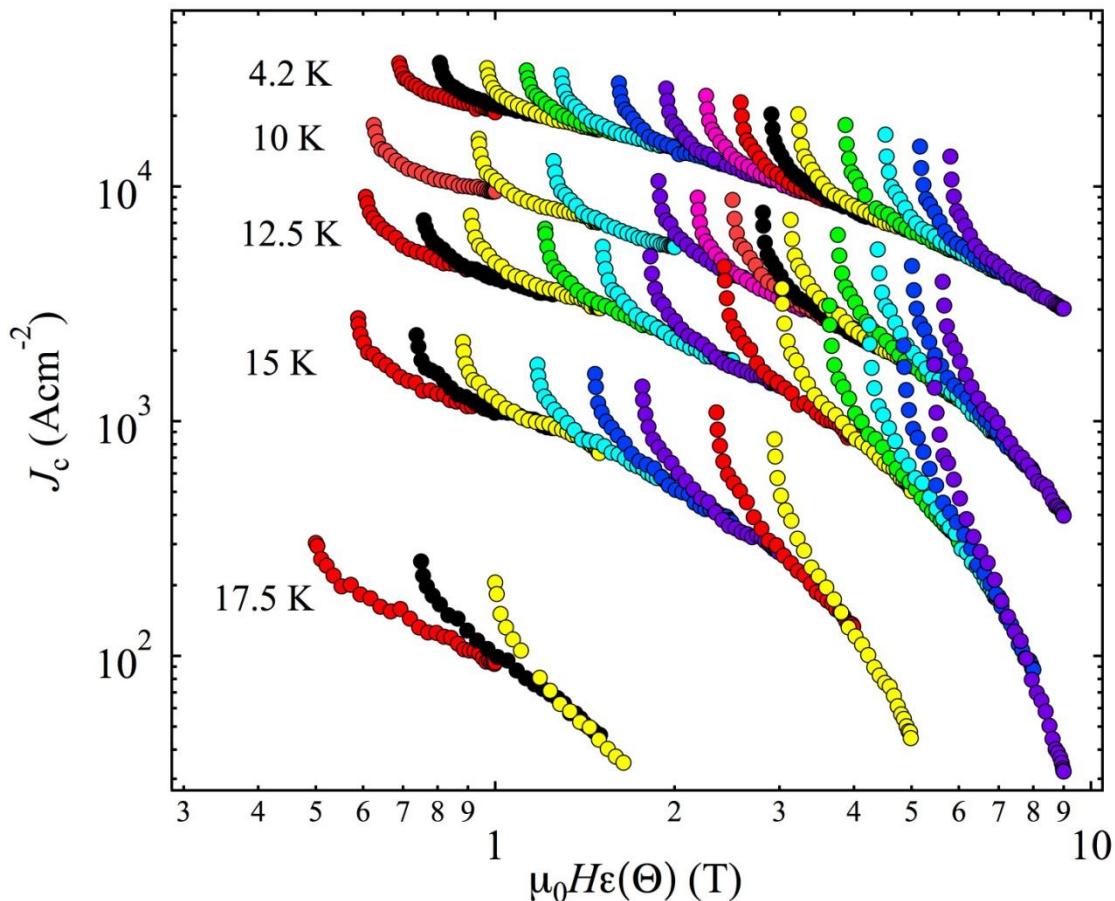


J_c anisotropy



- J_c from $J(E)$ curves,
 $E_c = 1$ μ V/cm,
 $w = 1$ mm, $l = 5$ mm
- Overall low absolute J_c values (clean structure without strong pinning centres)
- Maximum at $B \parallel ab$: due to
 1. electronic anisotropy
 2. pinning at occasional planar defects/surfaces
- No second peak for $B \parallel c$ (absence of correlated defects, cf. TEM)

J_c scaling



If:

- anisotropic Ginzburg-Landau theory holds
- only random and isotropic pinning centers
- angle θ well defined:
 - Large magnetic fields ($\kappa \gg B_{c2}/B$)
 - c-axis textured growth

Then:

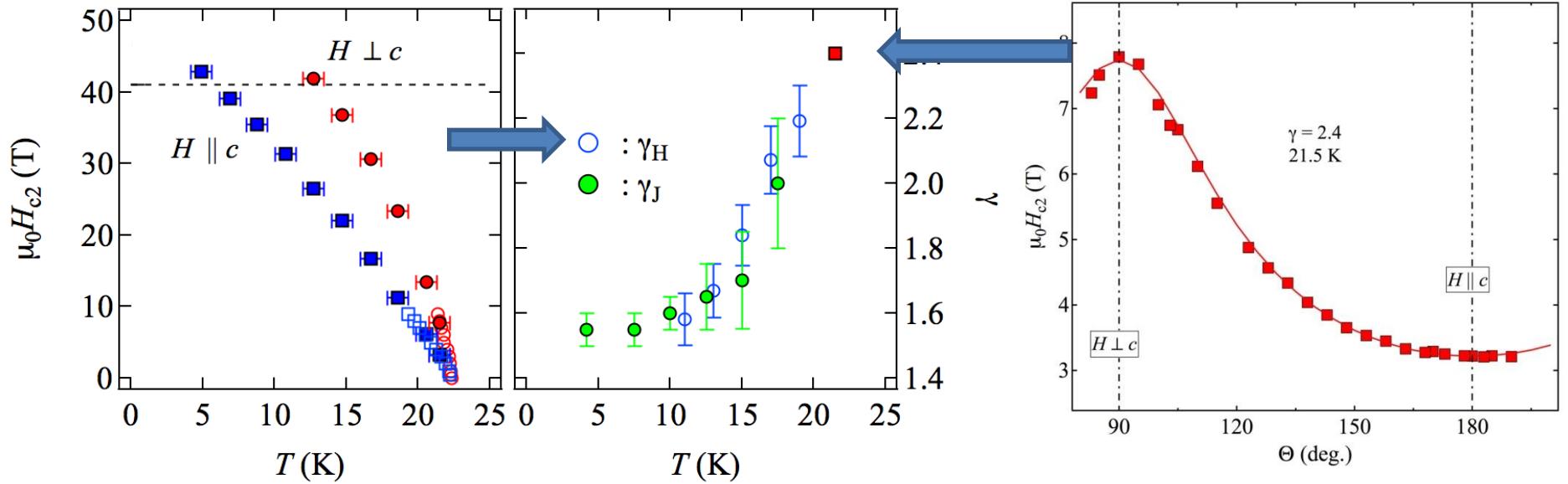
$J_c(\theta)$ can be scaled with:

$$J_c(H, \Theta) = J_c(\tilde{H}),$$

$$\text{where } \tilde{H} = \varepsilon(\Theta)H$$

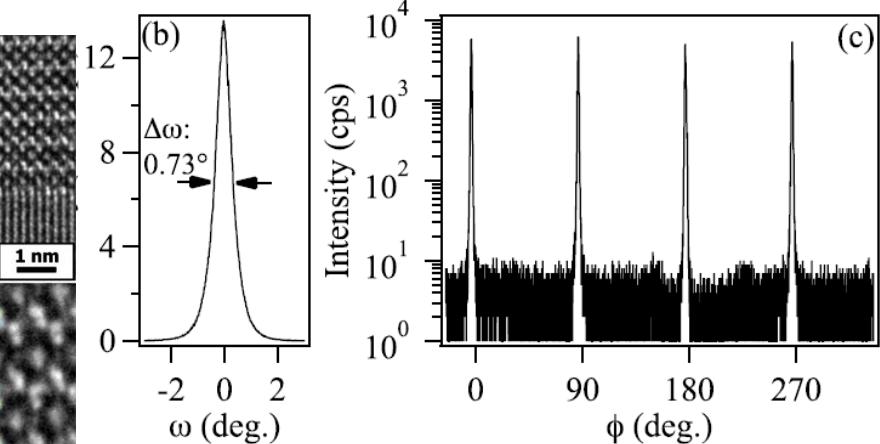
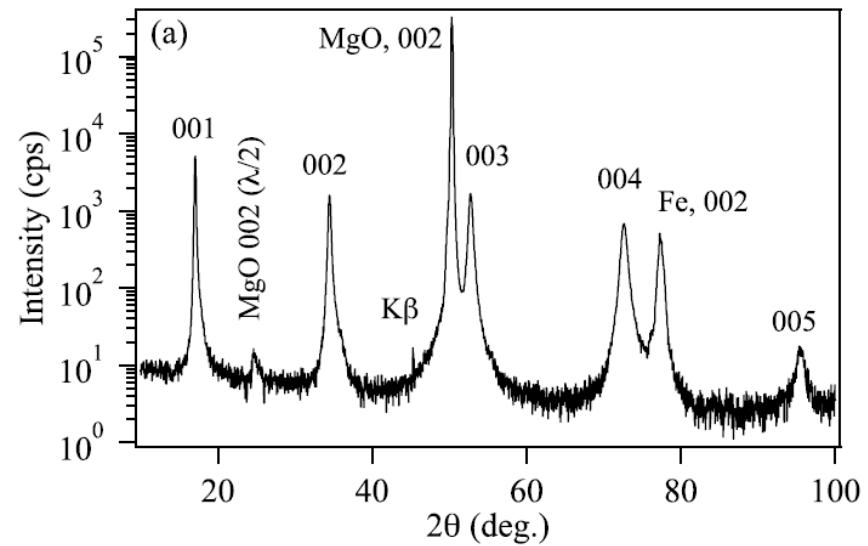
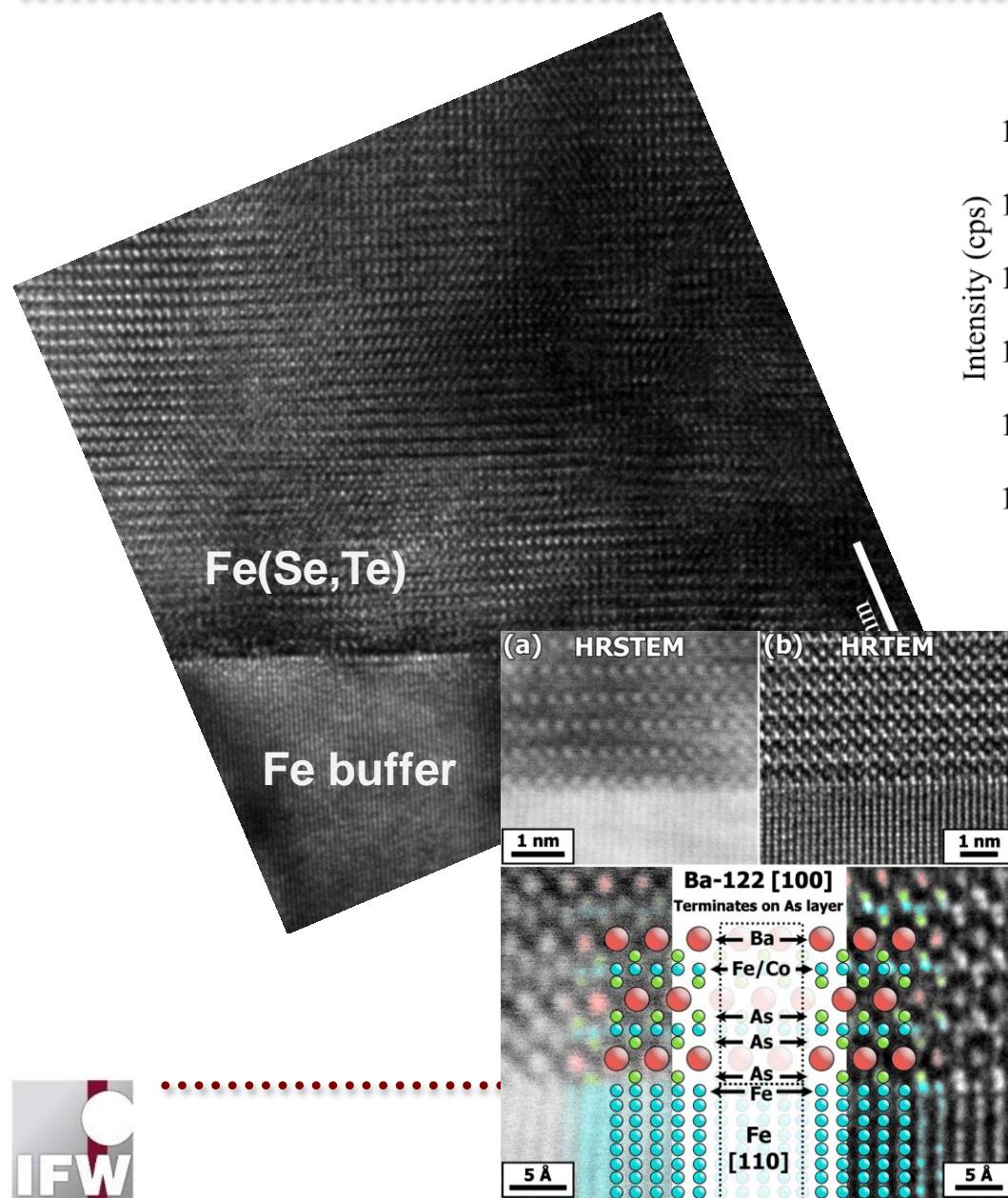
$$\varepsilon(\Theta) = \sqrt{\cos(\Theta)^2 + \gamma^{-2} \sin(\Theta)^2}$$

Electronic Anisotropy



- Low T : J_c scaling, medium T : $H_{c2}^{\perp c}/H_{c2}^{\parallel c}$, high T : full GL fitting of $H_{c2}(\theta)$
- H_{c2} anisotropy temperature dependent
→ evidence for multiband superconductivity (in 3D limit)
- J_c scaling probes anisotropy of upper critical field

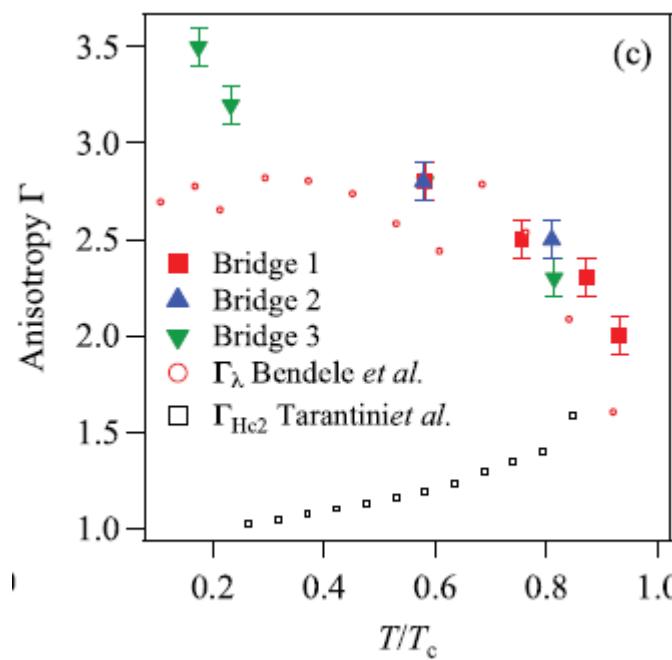
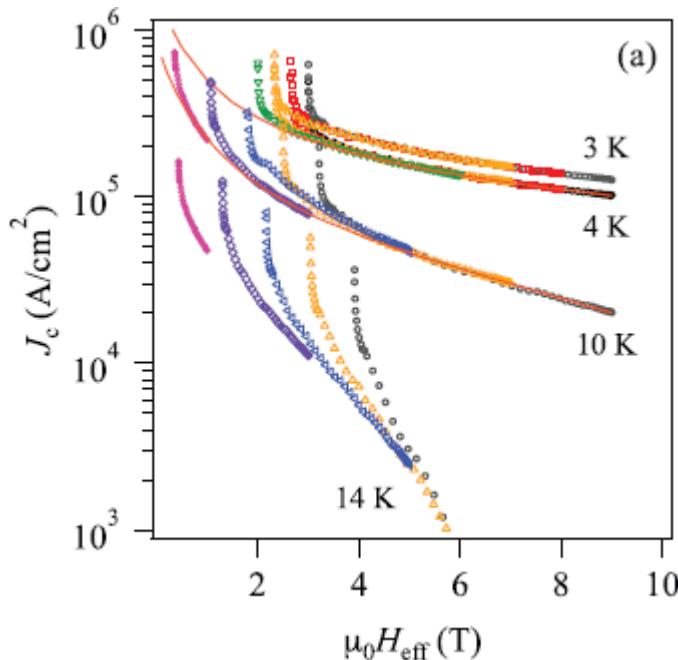
Fe(Se,Te): XRD, TEM



K. Iida et al., PRB 87, 104510 (2013)

T. Thersleff et al., APL 97, 022506 (2010)

Fe(Se,Te): J_c and N Scaling



$$H_{\text{eff}} = H\varepsilon(\Theta)$$

$$\epsilon(\Theta) = \sqrt{\cos^2(\Theta) + \Gamma^{-2} \sin^2(\Theta)}$$

Scaling parameter Γ follows Γ_λ
(in contrast to Ba-122, RE-1111)

Scaling with ξ or λ ?

In general, H_{irr} is extrinsic. But in absence of strong, extended defects, H_{irr} is governed by properties of the flux line lattice (intrinsic) rather than the defect structure (extrinsic).

Melting line:

3D:

$$B_m^J(T) = \frac{\pi^2 c_L^4}{G i} H_{c2}(0) \frac{T_c^2}{T^2} \left(1 - \frac{T^2}{T_c^2}\right)^2$$

Ba-122



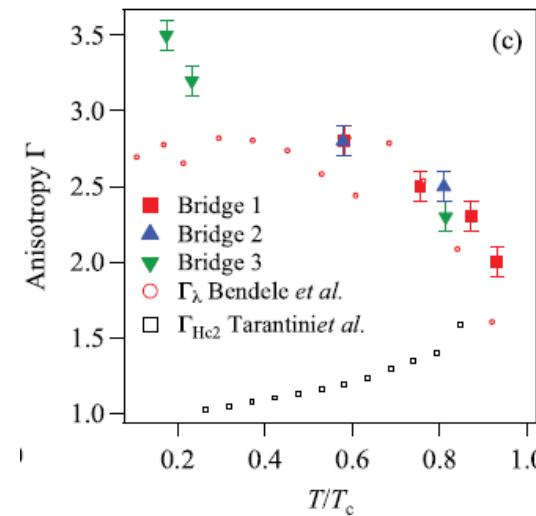
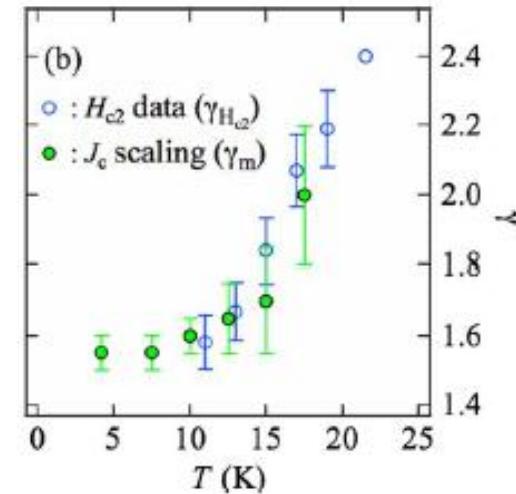
2D:

$$B_m^{\text{em}}(T) \approx \frac{c_L^2}{2\beta} B_\lambda \frac{\epsilon_0 d}{T}$$

Fe(Se,Te)

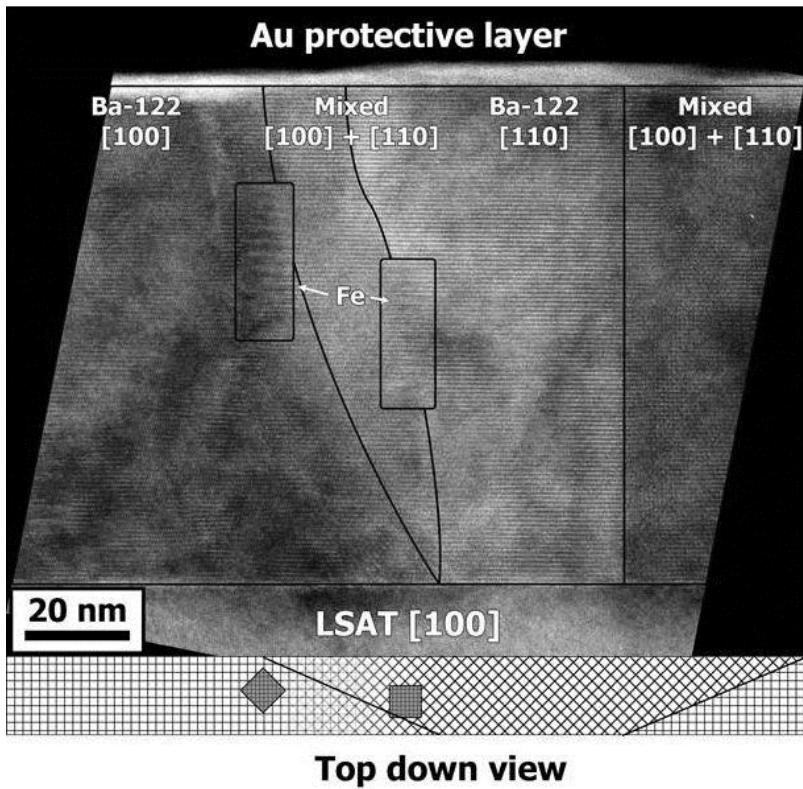
$$\bar{B}_\lambda = \Phi_0 / \lambda^2$$

(same holds for glass-liquid transition)

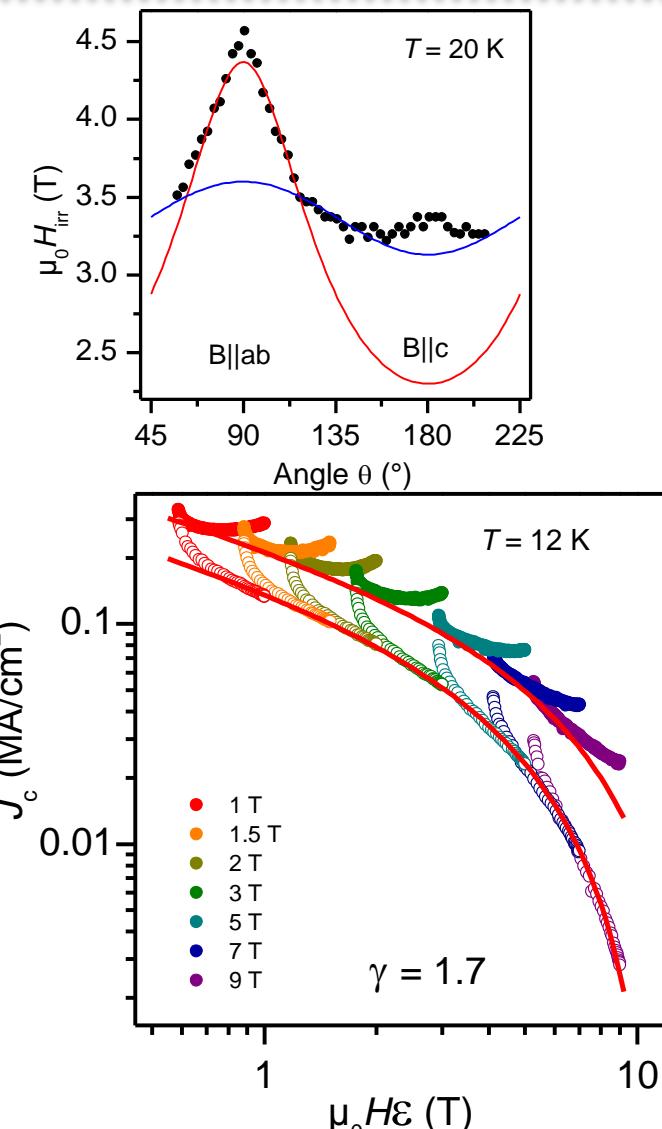


Effect of multiband + anisotropy!

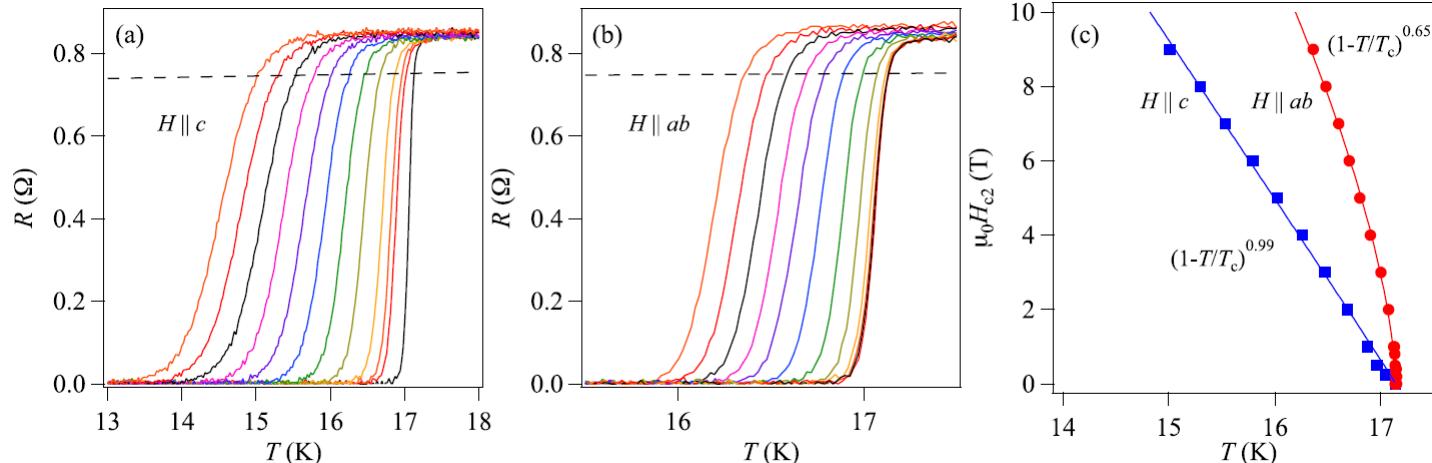
Scaling with extended defects



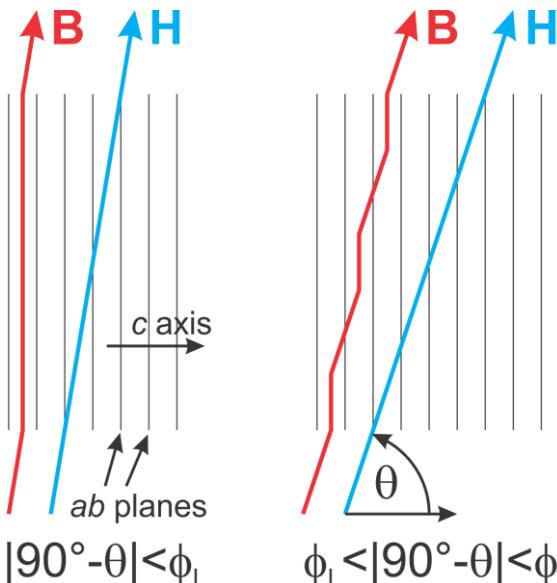
- Ba-122 with 45° c-axis tilt GBs + Fe particles → enhanced c-axis pinning
- Scaling with H_{c2} anisotropy only possible in small angular region



Fe(Se,Te): Upper critical field



Steep slope of $H_{c2} \parallel ab$ and negative curvature \rightarrow 2D behaviour



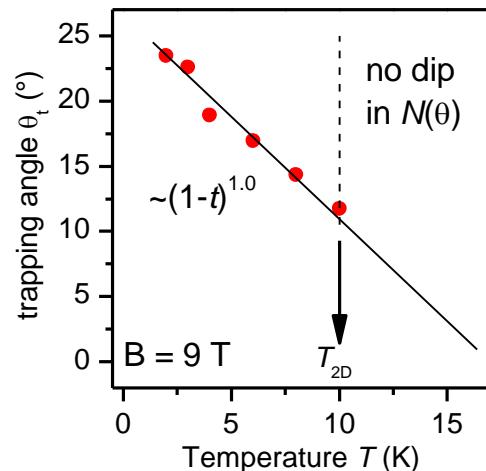
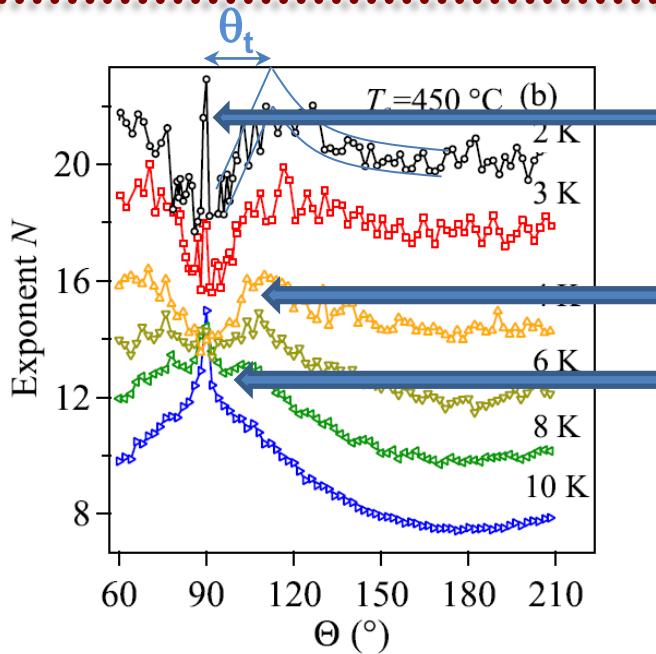
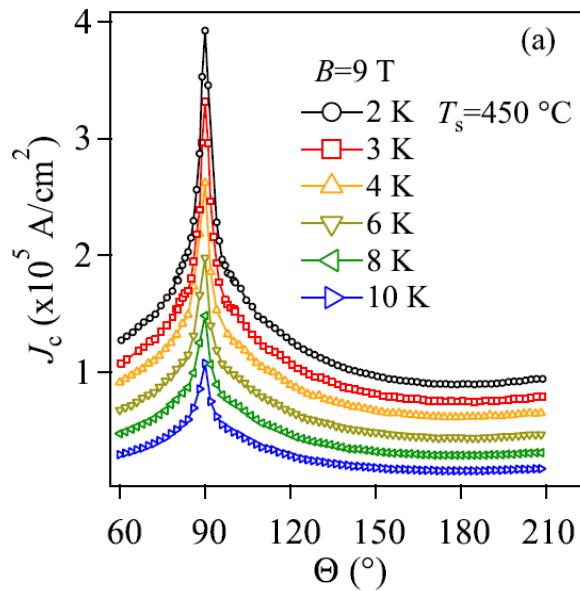
\rightarrow trapping and lock-in of flux lines

$$\varphi_t = \left[\frac{2\epsilon_{TP}}{\epsilon_l} \right]^{1/2}$$

$$\varphi_L = \frac{4\pi\epsilon_l}{\Phi_0 H} \varphi_t$$

- At angles smaller than the trapping angle ϕ_t vortices form staircases.
- At angles smaller than the lock-in angle ϕ_L vortices lock into the ab -planes.

Temperature Dependence

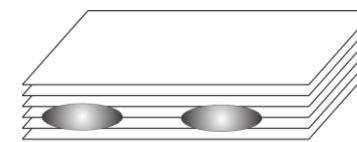


Transition between anisotropic 3D and quasi-2D:

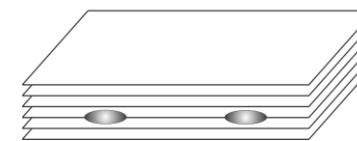
$$T_{\text{cr}} = (1 - \tau_{\text{cr}}) T_c < T_c$$

$$\tau_{\text{cr}} = 2\xi_c^2(0)/d^2$$

→ $\xi_c = 0.2\text{ nm}$



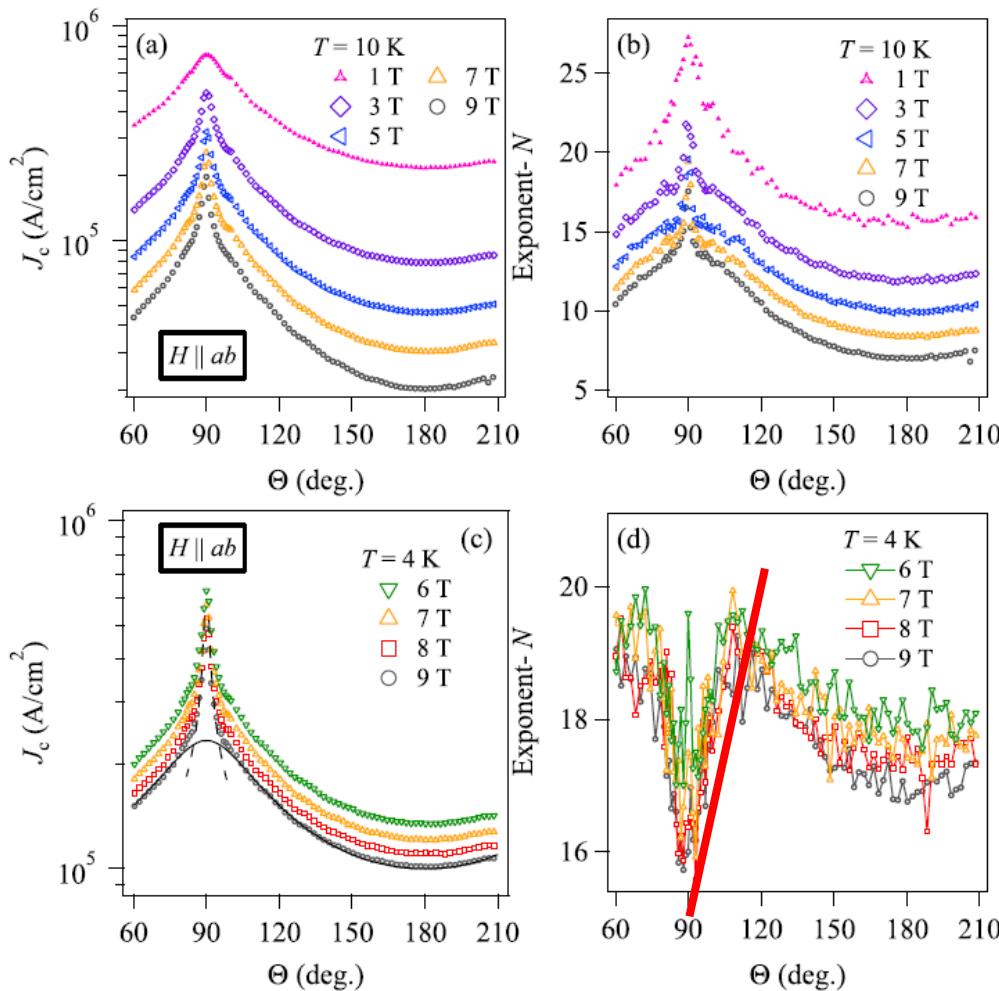
$$T > T^*$$



$$T < T^*$$

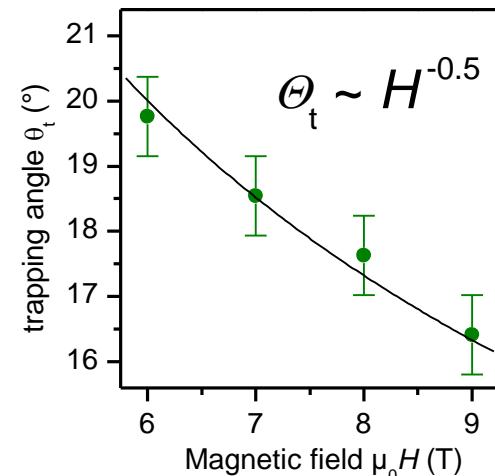
$$2\xi_c < d$$

Field Dependence

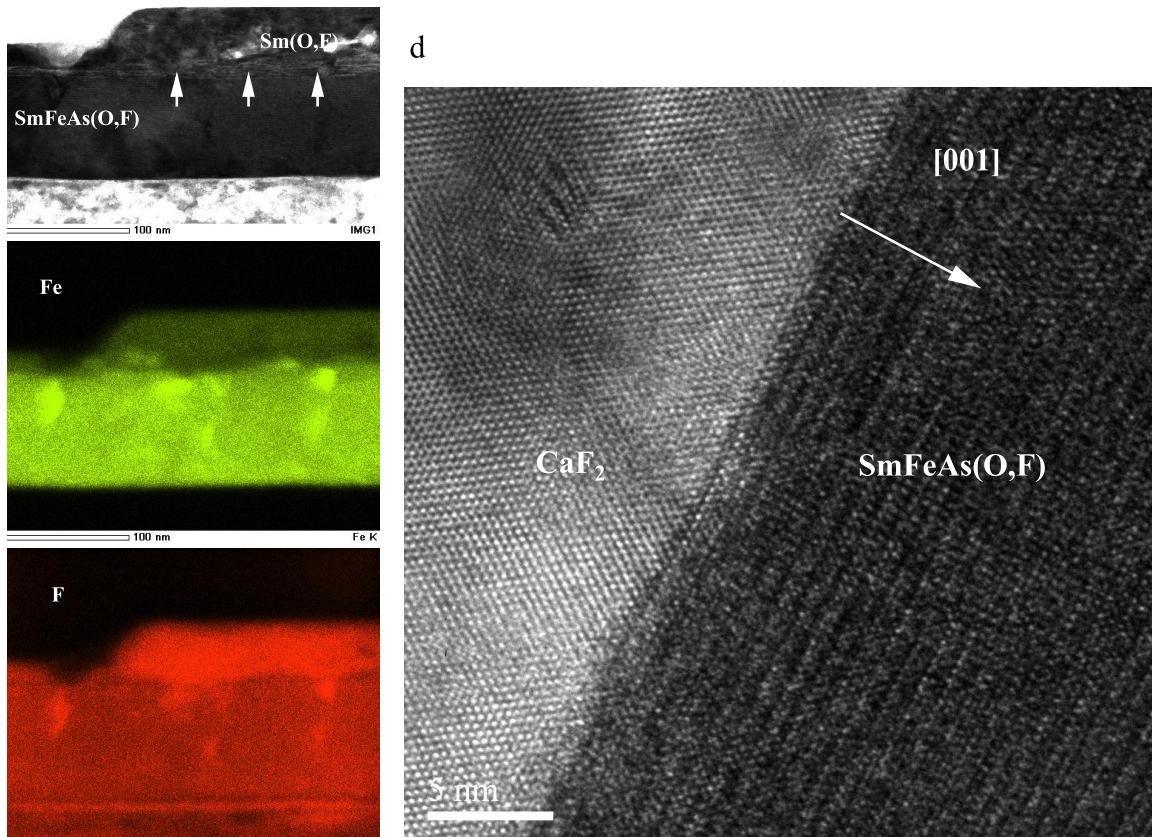


dN/dθ in trapping region:

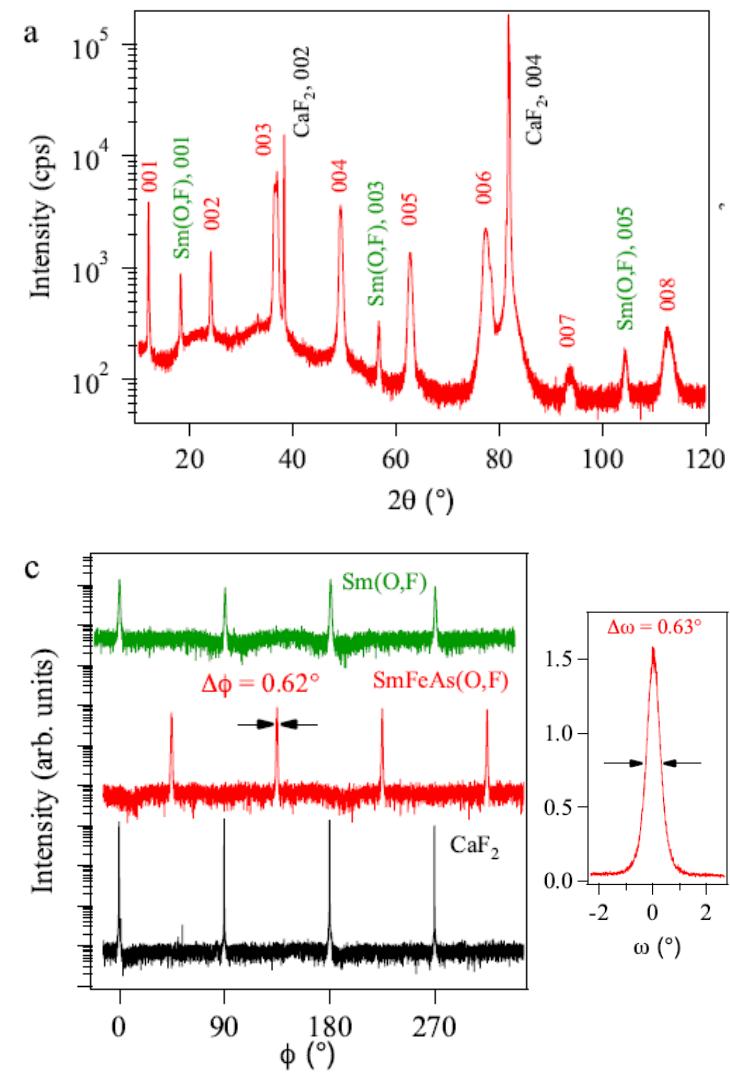
- + independent of B and T ,
- + depends on double-kink density per line



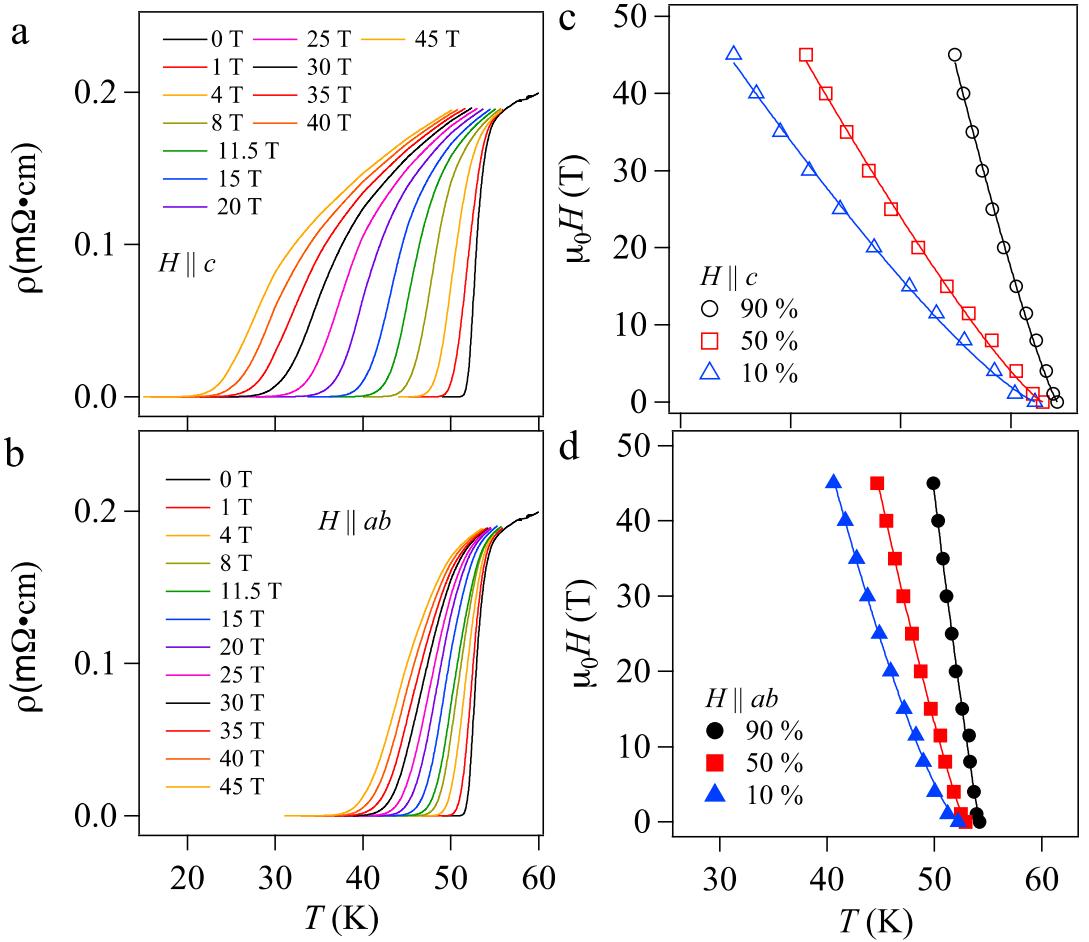
MBE Sm-1111: Microstructure



- Sharp biaxial texture
- Clean interface
- Occasional Fe-F particles



Sm-1111: resistive transition and H_{c2}

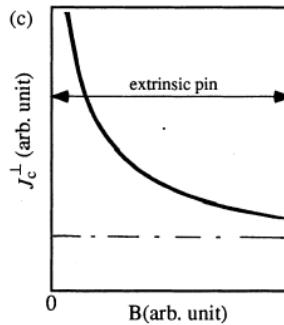
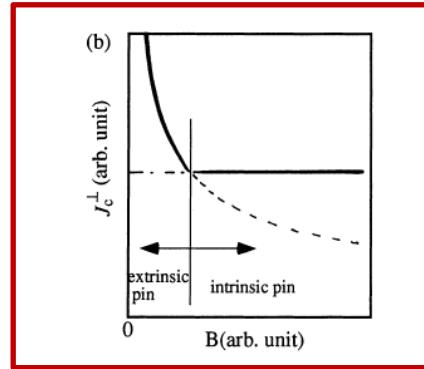
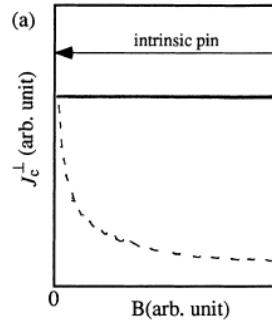
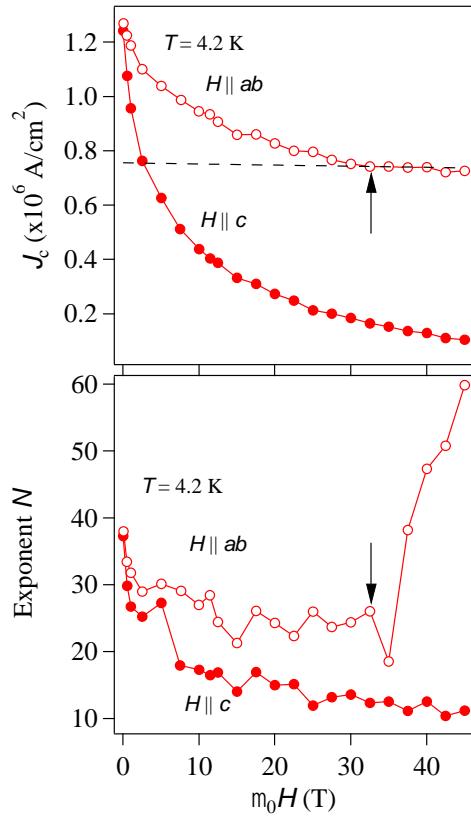


$$\left| \frac{d\mu_0 H_{c2}^{\parallel ab}}{dT} \right|_{T_c} = 11.4 \text{ T/K}$$

$$\xrightarrow{\quad} \xi_c = 0.28 \text{ nm} \\ < 0.858 \text{ nm} = d$$

**Extremely 2D
superconductor**

Sm-1111: $J_c(B)$



Above 35 T:

- Constant $J_c(B \parallel ab)$
- Sharply Increasing $N(B)$

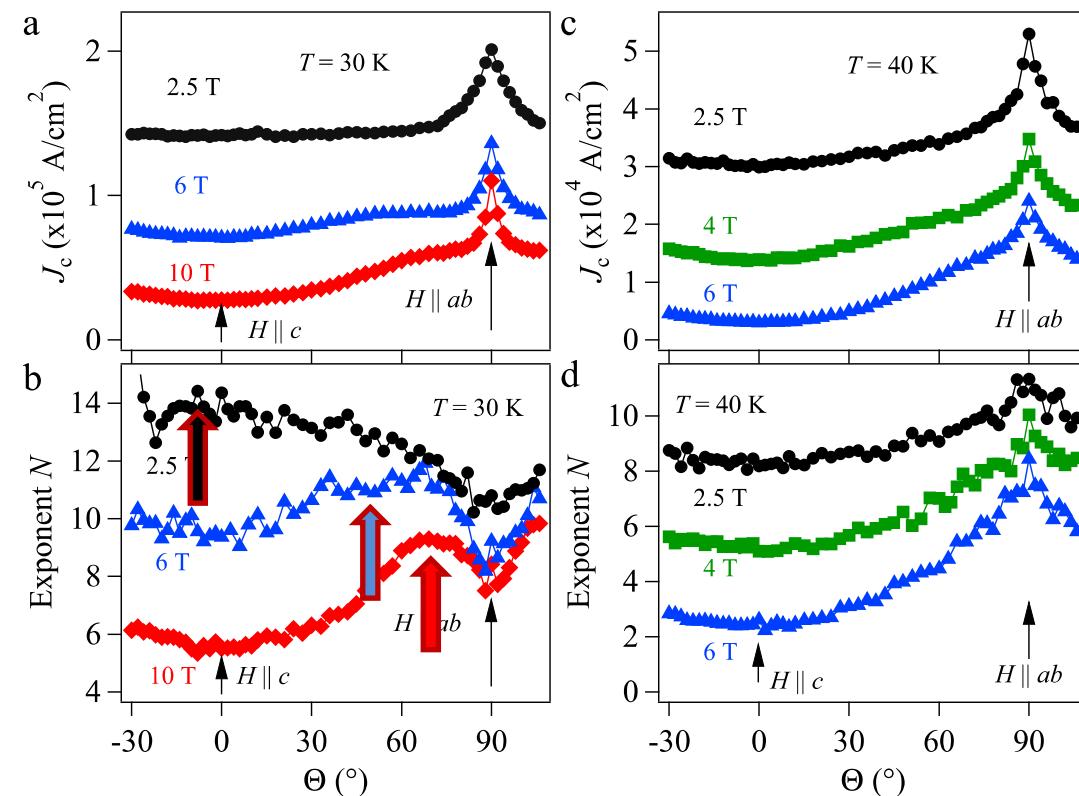
→ Intrinsic Pinning dominant

Below 35 T:

- J_c and N scale (as for $B \parallel c$)

→ Extrinsic Pinning dominant

Sm-1111: $J_c(\theta)$



$$T_{\text{cr}} = (1 - \tau_{\text{cr}}) T_c < T_c$$

$$\tau_{\text{cr}} = 2\xi_c^2(0)/d^2$$

Above 40 K:

- No dip in N observed

→ Extrinsic Pinning dominant

Below 30 K:

- Typical dip in N near ab

→ Intrinsic Pinning dominant

→ Large Lock-in angles
(due to large anisotropy)

→ 2D-3D transition at $T_{\text{cr}} \sim 35\text{ K}$
leads to

$$\xi_c \sim 0.35\text{ nm}$$

Conclusion

- Active and growing community for Fe-based sc thin films
- Basic investigations and evaluation of possible applications
- T_c strongly strain dependent
- J_c in clean films can be scaled with anisotropy of ξ (3D) or λ (2D)
- Intrinsic pinning clearly identified in Fe(Se,Te) and Sm-1111

