Electro-Dynamics of strongly disordered superconducting TiN films

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Why TiN for KIDs?

- High normal state resistivity (~ 100 $\mu\Omega$ cm)
 - Efficient far-IR absorption with 20-50 nm thick films
 - Reasonable area filling fraction
 - High kinetic inductance fraction
- T_c varies with stoichiometry (0 4.5 K)
- Reasonable quasiparticle lifetime
 - Maximum lifetime varies as $\sim T_c^2$
- Extremely high quality factors (> 10⁷)



• Improved sensitivity / figure of merit: $\mathcal{F} = \alpha_{\rm sc} \tau_{\rm max} Q_{i,{\rm max}} / N_0 V_{\rm sc}$

Leduc *et al*, APL **97**, 102509 (2010)



TiN

ŤUDelft



B. Sacépé *et.al.*, Nature Physics **7** 239 (2011)

 $k_{F}l$

¹² £

Electronic inhomogeneities

- Superconducting properties become inhomogeneous on a mesoscopic scale
 - Spatial fluctuation of spectral gap Δ
 - Formation of SC islands
 - Not directly linked to microscopic disorder (e.g. grain boundaries)





TUDelft

B. Sacépé et al, PRL 101, 157006 (2008)

How do we describe the electrodynamics of strongly disordered SC TiN films?

- To whish degree is TiN (NbTiN, NbN) a well-behaving textbook BCS superconductor like Al?
- To what extent is Mattis-Bardeen applicable for TiN?
- How can we justify the use of a large broadening parameter?
- What kind of problems can the high resistivity in the normal state of TiN films (that we whish to use) pose on the development of KIDs?







Jiansong Gao's formulas (JLTP 2008)

$$Q = \frac{2}{\alpha\beta} \frac{\sigma_2}{\sigma_1}$$

$$\frac{\delta Q}{Q} = -\left(\frac{\delta\sigma_1}{\sigma_1} - \frac{\delta\sigma_2}{\sigma_2}\right)$$

$$\delta\left(\frac{1}{Q}\right) = \frac{\alpha\beta}{2}\frac{\delta\sigma_1}{\sigma_2}$$

$$\frac{\delta f_0}{f_0} = \frac{\alpha\beta}{4} \frac{\delta\sigma_2}{\sigma_2}$$



Complex conductivity response: $\sigma_1 - i\sigma_2$ Mattis-Bardeen, Phys. Rev. **111** (1958)

$$\frac{\sigma_1(\omega)}{\sigma_N} = \frac{2}{\hbar\omega} \int_{\Delta}^{\infty} dE [f(E) - f(E + \hbar\omega)] \left(1 + \frac{\Delta^2}{E(E + \hbar\omega)}\right) N_S(E) N_S(E + \hbar\omega)$$

$$\frac{\sigma_2(\omega)}{\sigma_N} = \frac{1}{\hbar\omega} \int_{\max(\Delta - \hbar\omega, -\Delta)}^{\Delta} dE [1 - 2f(E + \hbar\omega)] \frac{1}{i} \left(1 + \frac{\Delta^2}{E(E + \hbar\omega)} \right) N_S(E) N_S(E + \hbar\omega)$$

$$N_{S}(E) = \frac{E}{\sqrt{E^{2} - \Delta^{2}}}$$
$$\Delta_{0}^{0} - \Delta \approx 2 \int_{\Delta}^{\infty} dE \frac{1}{\sqrt{E^{2} - \Delta^{2}}} f(E)$$

 $E \rightarrow E - i\Gamma$ or $\Delta \rightarrow \Delta - i\Gamma$

Broadening of the BCS density of states

Results

- TiN (Atomic Layer Deposition)
 - Low Disorder: $\rho = 191 \ \mu\Omega \cdot cm$, $T_c = 3.2 \ K$, $t = 55 \ nm$, $k_F / = 6.7$
 - High Disorder: $\rho = 309 \ \mu\Omega \cdot cm$, $T_c = 2.1 \ K$, $t = 11 \ nm$, $k_F / = 3.4$
- NbTiN (Sputtering)
 - Low Disorder: $\rho = 141 \ \mu\Omega \cdot cm$, $T_c = 13.6 \ K$, $t = 50 \ nm$
 - High Disorder: $\rho = 506 \ \mu\Omega \cdot cm$, $T_c = 11.8 \ K$, $t = 50 \ nm$



TiN ALD – series of: films with decreasing thickness















Explanations for Γ

- Lifetime broadening (but what determines the lifetime?)
- Electronic inhomogeneities
- Phase fluctuations (breakdown of long range superconducting coherence)





Conclusion

• Low disorder \rightarrow

MB theory + broadening of the BCS density of states

- For increasing disorder \rightarrow broadening parameter increases
- Broadening is directly related to the increase in disorder
- In agreement with numerical simulations of disordered superconductors

 - Ghosal et al, PRL 81, 3940 (1998)
 Bouadim et al, arXiv 1011.3275v1 (2010)



Outlook

 Microwave properties of a complete series of TiN film with increasing disorder

- ALD TIN
- Sputtered TiN (stoichiometry)
- NbTiN
- Lifetime measurements
- Complete characterization (DC and normal state properties of TiN)



Lifetime measurements TiN Low Disorder







TiN High Disorder two decay times! $r_1^* \exp[-t/T_1] + r_2^* \exp[-t/T_2]$

40

40



TUDelft

0

x 10⁻⁴

20

20

3

0

0

phase 1

0.0001

0.00001

Complete characterization

	Measured					Calculated				
Film ID	d	T _c	ρ	D	n @10 K	N(0)	ξ	<i>m</i> *	ł	k F •l
	[nm]	[K]	[µΩcm]	[cm²/s]	[cm⁻³]	[eV ⁻¹ cm ⁻³]	[nm]	[m _e]	[Å]	[-]
ALD1	55	3.2	191	0.94	2.9E+22	3.5E+22	6.0	2.8	7.0	6.7
ALD2	11	1.9	360	0.36	3.2E+22	4.8E+22	4.8	3.7	3.5	3.4
S2	250	4.7	100	1.60	4.4E+22	3.9E+22	6.4	2.7	10.2	11.1
S 3	320	4.7	365	0.44	2.5E+22	3.9E+22	3.4	3.2	4.1	3.7
S4	400	4.4	584	0.49	1.7E+22	2.2E+22	3.7	2.1	3.3	2.6



Questions



Homogeneously disordered versus granular disordered



FIG. 3: Resistance variations with temperature in Pb films upon increasing their thickness (from top to bottom) [12]. (a) Superconductor-insulator transition in finely dispersed quasihomogeneous films deposited on an SiO surface over an intermediate thin layer of amorphous Ge. In the superconducting region, the R(T) curves demonstrate a correlation between the normal resistance and the superconducting transition temperature. (b) Superconductor-insulator transition in granular films deposited directly onto the SiO surface. In such a method of deposition, the lead atoms coalesce into granules. The temperature of the superconducting transition in the film becomes constant at a film thickness exceeding the critical one.

Gantmakher *et al*, arXiv1004.3761v1 (2010)



Complex conductivity response: $\sigma_1 - i\sigma_2$ Mattis-Bardeen, Phys. Rev. **111** (1958)

$$\frac{\sigma_1(\omega)}{\sigma_N} = \frac{2}{\hbar\omega} \int_{\Delta}^{\infty} dE \frac{E^2 + \Delta^2 + \hbar\omega E}{\sqrt{E^2 - \Delta^2}\sqrt{(E + \hbar\omega)^2 - \Delta^2}} [f(E) - f(E + \hbar\omega)]$$

$$\frac{\sigma_2(\omega)}{\sigma_N} = \frac{1}{\hbar\omega} \int_{\max(\Delta - \hbar\omega, -\Delta)}^{\Delta} dE \frac{E^2 + \Delta^2 + \hbar\omega E}{\sqrt{\Delta^2 - E^2}\sqrt{(E + \hbar\omega)^2 - \Delta^2}} [1 - 2f(E + \hbar\omega)]$$

$$N_{S}(E) = \frac{E}{\sqrt{E^{2} - \Delta^{2}}}$$

$$E \rightarrow E - i\Gamma \quad \text{or} \quad \Delta \rightarrow \Delta - i\Gamma$$
Broadening of the
BCS density of states



TiN High Disorder (ALD250)











Atomic Layer Deposition



Why Atomic Layer Deposition?

- Very thin layers of TiN
- High resistance per square
- High uniformity
- Possibly superior properties compared to sputtered materials





Roughness and grain size



30

Relative roughness compared to resistivity







ALD021: bring it ON the carpet

-Fitting the amplitude instead of the phase will give a different decay time, roughly a factor of 2 smaller.

-The scatter in the lifetime deduced from the amplitude is very large, although the fits are reasonable.

-Resonator 4 gives a large scattering in the lifetime (phase and amplitude), although the fits are reasonable.

-We cannot really speak of an amplitude or phase response since the data is not properly normalized.

٥...

18_□

16

14

lifetime (µs)

6

4

2

0∟ 300

350

400

450

500

T (mK)

T







ALD250: additional plots

- I have only data of one resonator
- Further, I have only analyzed the 'phase' data and no proper normalization was possible









• How close to localization are our TiN films?

• Anderson localization at $k_F \ell \sim 1$



Blue Oxford Cryostat

- Base temperature 1.5 K
- Magnetic field up to ~13.5 T



Dynes et al., PRL 53 (1984)



FIG. 3. The density of states N(E) deconvoluted from the data of Fig. 2 (solid line). The dashed line is a BCS density of states broadened by the value of Γ shown in the figure for each case.

