

High critical-current density and scaling of phase-slip processes in YBaCuO nanowires

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Abstract.

YBaCuO nanowires were reproducibly fabricated down to widths of 50 nm. A Au/Ti cap layer on YBCO yielded high electrical performance up to temperatures above 80 K in single nanowires. Critical current density of tens of MA/cm² at T = 4.2 K and of 10 MA/cm² at 77 K were achieved that survive in high magnetic fields. Phase-slip processes were tuned by choosing the size of the nanochannels and the intensity of the applied external magnetic field. Data indicate that YBCO nanowires are rather attractive system for the fabrication of efficient sensors, supporting the notion of futuristic THz devices.

1. Introduction

Superconducting nanostructures operating at the liquid nitrogen temperature are very attractive for several applications in superconducting electronics. In particular high critical-temperature superconductor (HTS) nanowires not only have the advantage of high temperature operation, but in principle can be functionally scaled to smaller sizes [1, 2] thanks to their extremely short coherence lengths ξ_0 . Other intrinsic properties of these materials can play a relevant role too. For instance, their characteristic fast relaxation times [3] offer higher counting rates in photo-detection experiments when compared to traditional superconductors [4].

In this context reliable top-down fabrication techniques allowing a more direct integration of the HTS nanostructures into hybrid systems are of much interest. Here we analyze the electrical properties of a number of nanofabricated wires down to 50 nm in width. Our measurements show very good performance for single YBCO nanowires both in terms of critical temperature (T_C) and critical-current density (J_C) with excellent potential for sensor applications. Critical-current densities up to 70 MA/cm² at $T = 4.2$ K and 10 MA/cm² at 77 K were achieved on YBCO capped by a thin protective layer of Au/Ti. These are among the best results reported for single nanowires [5, 6, 7, 8] (see below). Contrary to other works [2], where even smaller sizes were achieved for *arrays* of nanowires, in this case we isolated single nanowires, thus responding to a much wider range of requirements for circuit design.

2. Experimental details

We have employed YBCO thin films produced on various substrates through different deposition techniques (sputtering and reactive thermal co-evaporation [9]). For the aims of this work of demonstrating a reliable top-down technology to obtain single YBCO nanowires down to a width of 50 nm and the utility of using a Au cap layer, the properties of the thin films to privilege are rms roughness as low as possible, and absence of macroscopic impurities in the film. The initial values of T_C and J_C of unpatterned films are relevant but not crucial. In order to perform systematic experiments and reliably compare nanostructures realized on different samples, we have employed thin films produced on large areas by co-evaporation technique by Theva [11]. From the same wafer, it is possible to cut 20 samples of size of 5 mm x 5 mm with the same properties. In this work we focus on c-axis YBCO thin films (50 nm thick), grown on a CeO_2 seed layer (40 nm thick), and protected by a 20 nm thick Au layer [11]. The substrate is a large wafer of Yttria-stabilized zirconia (YSZ). The values of the original critical temperature and of the critical current density are high ($T_C \simeq 86.5$ K and $J_C = 2.2$ MA/cm² at $T = 77$ K) but not optimized to the highest possible values. The key-steps of the nanotechnology approach are based on the use of a 30 nm thick Ti mask patterned through e-beam lithography (ZEISS MERLIN), and on a low energy milling procedure keeping the sample at a temperature of about -150°C. Most of the procedure

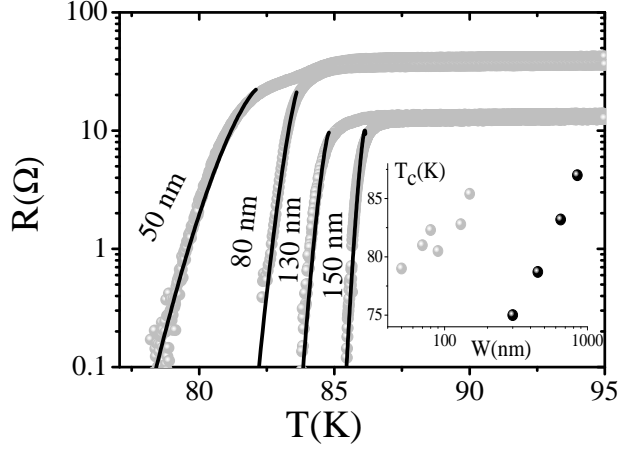


Figure 1. Resistive transitions of YBCO/Au/Ti nanowires of different widths indicated in the figure. The dashed lines are the fits achieved using the LAMH theory (see text). Inset: Dependence of the critical temperatures on the nanowire width. The grey dots refer to the T_c 's of YBCO/Au/Ti samples while the black points to bare YBCO nanowires.

is described in detail in [12]. All nanobridges described in this work are $1 \mu\text{m}$ long. A comparative study of bare and capped YBCO nanowires on samples produced in the same deposition run and only differing by the last fabrication step (bare nanowires undergo an additional ion milling step in order to remove the Au layer) sheds light on the role of Au layer protecting the nanowires. The Au/Ti layer makes easier the integration of the YBCO component into hybrid structures because of its better compatibility with other materials as opposed to HTS. If only genuine superconducting properties are required, the YBCO capped nanowires can perfectly replace YBCO nanowires for most functions. Nano-fabrication techniques are not restricted, for the resolutions specified above, to any specific choice of the substrates usually employed for HTS thin films, or thin film deposition conditions, as long as surface roughness is below some thresholds, for instance lower than 5nm on a total thickness of 50 nm.

Measurements were performed in liquid helium using a four probe configuration. The set up employed is equipped with two stages of cold filters (copper powder and RC Pi filters with a cut-off frequency of the order of a few GHz and 160 MHz respectively).

3. Results and Discussions

The critical temperature of various representative devices is reported as a function of wire width in the inset of Fig. 1. T_c scales linearly for both types of bridges. For the capped YBCO, T_c decreases from about 86 K for unpatterned films to about 85 (78) K for nanowires of width $w=140$ (50) nm respectively. For the bare YBCO samples, T_c of $1\mu\text{m}$ wide bridges is basically the same of the unpatterned films and linearly decreases down to 75 K for nanowires of width $w=300$ nm. Therefore the 'critical' region is for

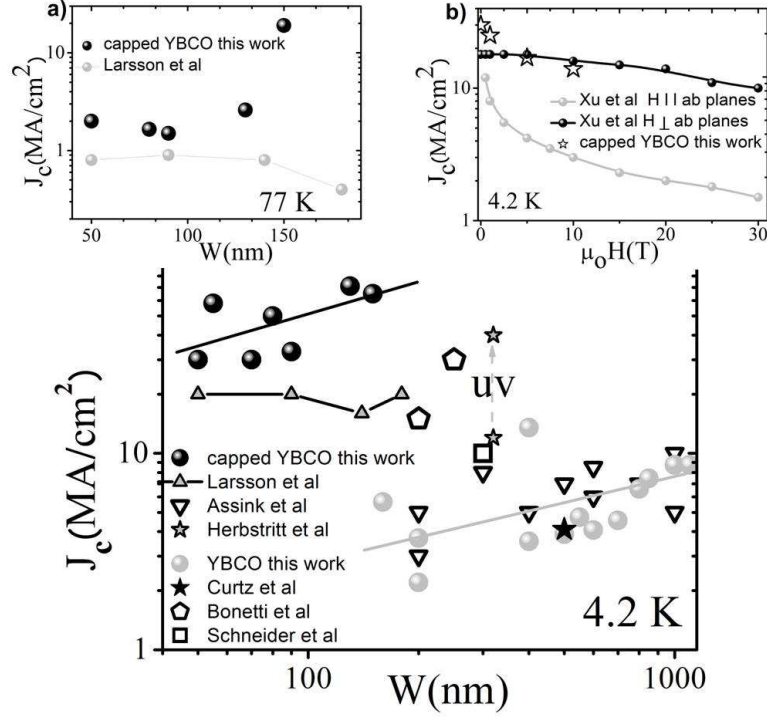


Figure 2. Values of critical current density J_C measured at 4.2K versus channel width (W). Data from this work (grey circles for YBCO nanowires, black circles for YBCO/Au/Ti nanowires) are compared with results available in literature [5, 6, 8, 7]. Uv stands for the increase of J_C obtained by ozone treatment [7]. Inset a): J_C vs W of the YBCO/Au/Ti nanowires measured at 77K are compared with [6]. Inset b): J_C of the 50 nm wide nanochannel (measured at $T = 4.2$ K) at different values of the magnetic field H are compared with typical data from literature [17].

widths below about $1 \mu\text{m}$. Superconductivity is quite robust also for widths less than 200 nm, but the properties of devices are less reproducible.

Not surprisingly, the bare YBCO thin films are more sensitive to ion milling and processing details. Figure 1 shows resistance vs temperature curves for the capped YBCO nanowires. Both bare and capped YBCO data can be well fitted in terms of the thermally-activated phase-slip model, i.e. the Langer- Ambegaokar- McCumber - Halperin (LAMH) model [13] commonly used to describe the broadening of $R(T)$ in superconducting nanostructures [14]. The total resistance of the wire is commonly expressed as the parallel [14] between the resistance activated by phase-slip processes R_{LAMH}

$$R_{LAMH} = R_Q \frac{\hbar \Omega}{k_B T} \exp \left(-\frac{\Delta F}{k_B T} \right) \quad (1)$$

and the normal resistance of quasiparticles $R_N(T) = R_N \exp(-\Delta F(T)/k_B T)$ [15]: $R(T)^{-1} = R_{LAMH}(T)^{-1} + R_N^{-1}(T)$. Here $\Omega = (L/\xi) \sqrt{(\Delta F/k_B T)} (1/\tau_{GL})$ is the phase slipping rate, $\tau_{GL} = (\pi \hbar/k_B (T_C - T))$, $R_Q = h/4e^2$ is the quantum resistance constant,

L is the sample length, k_B is the Boltzmann constant, and ΔF is the free energy where the superconducting phase "falls off". ΔF at $T=0$ K can be expressed as $8\sqrt{2}/(48\pi^2)A(\phi_0^2/(\mu_0\lambda^2\xi_0))$ (where A and λ are the cross section and the London penetration depth respectively) in analogy with other experiments on HTS nanowires [2]. The extracted values of the fitting parameters are $\xi_0 \sim 2nm$ and of $\lambda \sim 300nm$ which, are quite consistent with typical values of the cuprates [10]. ΔF is for all curves about 10^5 K.

Measured critical-current density values are reported as a function of the width (w) for $T = 4.2$ K in Fig. 2 and compared with results available in literature [5, 6, 7, 8]. J_C in capped YBCO nanowires is higher than 30 MA/cm^2 (for $w = 50 \text{ nm}$) and reaches a maximum of 70 MA/cm^2 for $W = 140 \text{ nm}$. These are among the highest values reported [5, 6, 8, 7] and only a few times lower than the theoretical depairing limit (300 MA/cm^2) [16]. J_C values at $T = 77$ K are reported in Fig. 2 inset a) for YBCO/Au/Ti nanowires and range from 2 MA/cm^2 for $w = 50 \text{ nm}$ to 20 MA/cm^2 for $W = 140 \text{ nm}$. The better performances offered by capped nanowires when compared with those of bare YBCO nanowires of this and previous works [6], reveal that the protecting Au layer is key for enhancing the quality of HTS nanowires. We believe that there are margins to further improve J_C performances especially at 77 K by employing films with higher values of the initial T_C .

Nanowires keep good transport properties also when an external magnetic field is applied. For instance, in the extreme limit of 50 nm , a magnetic field of 10 T reduces T_C by less than 10% , while J_C at 4.2 K is only halved. Superconductivity in 50 nm wide strips survives up to $H = 1 \text{ T}$ at 77 K . Figure 2 inset b) allows a comparison of the dependence of J_C on magnetic field at $T = 4.2 \text{ K}$ [17] in the present samples with other results and confirms the robustness against an externally applied magnetic field. The use of HTS for power applications is a field of research of great relevance and a 'smart' control of pinning centers is the key to increase J_C . In macroscopic samples this role is usually played by the introduction of nanodefects [18, 19], while in nanowires by the detailed structure of the edge barrier [20].

Figure 3a shows the experimental current-voltage (I-V) characteristics measured at $T = 4.2 \text{ K}$ for YBCO/Au/Ti nanowires 130 , 80 and 50 nm wide. The distinctive feature is the presence of steps accompanied by a very large hysteresis that exceeds 30% of the total current. Similar features were reported by several author in experiments involving nanostructures [14] and HTS nanowires [2, 22, 23]. These observations were interpreted in terms of phenomena driving the superconducting channel to the normal state, i.e. the development of phase-slip centers (PSC) or normal hot spots (HP) [25]. Because of their low carrier density and thus low superfluid rigidity, HTS systems are inherently prone to phase fluctuations and therefore more susceptible to PSC in reduced dimensions [21, 14]. I-V curves reported in Fig. 3 show that the amplitude and the position of the steps depends on the size of the nanochannels and on the magnetic field. By increasing the width of the channel, the switch occurs at higher voltage. This is consistent with the idea that in wider samples the transition to the fast vortex motion that leads to the step

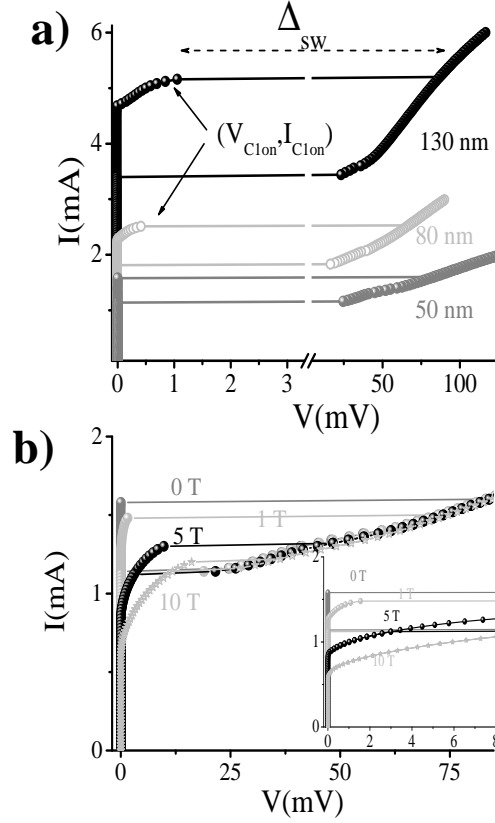


Figure 3. a) I-V curves at $T = 4.2$ K for YBCO/Au/Ti nanowires of widths 130 nm, 80 nm and 50 nm, respectively. b) For the 50 nm wide bridge, I-V curves are reported for different values of the magnetic field ($H=0, 1$ T, 5 T and 10 T).

structure occurs at a larger current [28]. As shown in Fig. 3b, an increase in H moves V_{1on} to higher voltages to a maximum of about 15 mV for $H = 10$ T. The voltage jump Δ_{SW} is also reduced from $\Delta_{SW} = 80$ mV at $H = 0$ T to about 10 mV at $H = 10$ T.

From the value of the critical voltage $V_{1on} = v_C H L$ [26, 28] it is possible to estimate the vortex critical velocity $v_C = 10^3$ m/s (at $H = 1$ T), and hence the inelastic electron-electron scattering time $\tau_S \approx 0.6$ ns on the basis of the expression $\tau_S = D[14\zeta(3)]^{0.5} (1 - T/T_C)^{0.5} / \pi v_C^2$ [27], where D is the quasiparticle diffusion constant and $\zeta(3) \sim 1.2$. This value is in agreement with other experiments [27] and estimates [28].

The scaling of the I-V curves and of the steps with the width of the nano-channels and the influence of H on the I-Vs reflects the presence of phase slip lines (PSL) [29, 28]. PSL are a 2-dimensional analogue of PSC and reveal some kind of phase transition in the vortex lattice at the instability point with regions of fast and slow vortex motion [29, 28]. When the width of the bridges is reduced to $w = 50$ nm, the step position ($V_{1on} = 0$) and the shape of the I-V indicate a transition from PSL to the more classical 1-dim PSC (at $H = 0$ T). In the narrowest nanowire, not more than one line of vortices at low fields can be hosted. When the magnetic field is increased, the size of inter-

vortex distance a ($a = \sqrt{\Phi_0/H}$) is decreased (for instance $a = 20$ nm at $H = 5$ T), and the entrance of additional rows of vortices is favored, with a consequent transition to PSL. This is documented by the steps at finite voltage in the presence of magnetic field ($V_{1on} > 0$). Vortex motion is characterized by oscillatory behavior of flux lines which cross the sample resulting in an emission of electromagnetic radiation $\nu = v_c/a$ that is on the order of 0.2×10^{11} Hz at $H=1$ T.

Studies on the dynamics of kinematic vortex-antivortex lines confirm that their individual velocity can be manipulated by applying magnetic field and current [30], supporting the notion of futuristic THz devices.

4. Conclusions

In conclusion single YBCO nanowires were fabricated through a Au/Ti cap layer to reach very high values of the critical current density also at 77 K that is very robust against an externally applied magnetic field H . The detailed structures of the I-V curves of these single nanowires can be tuned by choosing the size of the bridge and by setting the magnetic field intensity. Threshold mechanisms associated to phase slips processes and visible as steps in I-Vs pave the way to HTS nanowires as sensors for photodetection experiments.

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