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Pinning force scaling of electrospun Bi-2212 nanowire networks

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Abstract

Flux pinning forces were determined on different network samples of superconducting Bi₂Sr₂CaCu₂O₈ (Bi-2212) nanowires prepared by the electrospinning technique. We employed magnetization data determined by SQUID magnetometry in a wide temperature range 10 K < T < 35 K, where a strong superconducting signal prevails. The scaling analysis of the pinning forces was applied to interpret the data obtained. Both pure and Li-doped Bi2212 nanowire networks exhibit a peak position of $h_0 \sim 0.11$, which is smaller than the expected value of $h_0 = 0.2$ indicating flux pinning at grain boundaries or extended defects. For the flowing currents through such a network, the crystallographic anisotropy and the percolation play an important role, resulting in reduced peak positions as compared to bulk samples.

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1. Introduction

Recently, superconducting nanowires of Bi₂Sr₂CaCu₂O₈ (Bi-2212) could successfully be prepared by employing the electrospinning technique [1, 2, 3]. The nanowires themselves were found to be polycrystalline, consisting of nanometer-sized grains of random orientation and exhibit diameters of about 250 nm, which is comparable to the London penetration depth, λ_0 . The networks of these nanowires formed by electrospinning and the subsequent heat treatment have an extremely low density and show numerous interconnects between the nanowires which enable a current flow through the entire network. The transport and magnetization measurements [2, 4] revealed that the currents can flow through the entire sample perimeter even at applied magnetic fields of 10 T. These properties of the nanowire network samples are very interesting from the underlying physics and for possible applications. Therefore, in this letter, we analyze the flux pinning forces obtained from M(H)measurements using the model of Dew-Hughes (DH) [5, 6] to gain a deeper understanding of the magnetic behavior of the nanowire networks, which may be interesting for certain applications whereever the weight of the samples counts, and we compare the resulting pinning force scaling diagrams to such of other types of Bi-2212 and Bi-2223 materials published in the

literature.

2. Experimental procedure

The Bi-2212 nanowire networks were fabricated using the electrospinning technique from an aqueous solution of polyvinylpyrrolidone (PVP, M_W 1,300,000), bismuth(III) acetate, strontium acetate hydrate, calcium acetate hydrate and copper(II) acetate monohydrate and followed by calcination treatment of the electrospun polymer/inorganic composite fibers including one oxygenation step in pure O₂. The chemicals were commercially available (Alfa Aesar GmbH & Co KG, purity above 99.99 %) and used as received without further purification. X-ray analysis confirmed that the samples are pure Bi-2212 phase with some residing carbon. For the sample with Li doping, 30 x.-% Li acetate was added to the precursor solution. Further details of the nanowire fabrication procedure are given elsewhere[3]. The density of our nanowire networks is found to be equal to 0.0459 g/cm³, which is considerably lower than the theoretical density of bulk Bi-2212 (6.4 g/cm^3).

The magnetization of the nanowire networks was measured using a SQUID magnetometer (Quantum Design MPMS3) with ± 7 T magnetic field applied perpendicular to the sample surface. The onset of superconductivity takes place at 76 K, which

is somewhat lower than the bulk T_c of the Bi-2212 phase (85 K) [2, 4].

3. Results and discussion



Figure 1: (a) Optical image of heat-treated nanowire network samples. (b) SEM image (magnification $3000 \times$) of the nanowire network after the final heat treatment.

The microstructure of the Bi-2212 nanowire networks are presented in Figs. 1 (a/b) after the final heat treatment. The nanowire networks (a) look like a thin fabric and are extremely brittle. Due to the numerous interconnects between the individual nanowires as shown in (b), transport currents can flow through the entire nanowire network sample.

Figure 2 shows magnetization hysteresis loops of the Bi2212 and Bi2212+Li samples measured at T = 15 K. The loops exhibit the typical asymmetric shape of polycrystalline materials [7]. Both loops show an additional magnetic signal overlaid on the superconducting loop. For pure Bi-2212, there is a diamagnetic signal, and for the Li-doped sample, the signal is paramagnetic.

The dependence of the pinning force density on the applied magnetic field was determined by $F_p(H) = \mu_0 H \times J_c(H)$. The $J_c(H)$ datapoints are obtained from the width of the experimental loops using the Bean formula. In Figs. 3, 4, the normalized



Figure 2: Magnetization hysteresis loops measured at T = 15 K.

pinning force $f_p = F_p/F_{p0}$ is plotted as a function of the reduced field $h = H/H_{irr}$ for different *T*, here F_{p0} is the maximal pinning force density at given *T*. For high- T_c superconductors, the values of H_{irr} are usually determined from a logarithmic plot. Here we see that noise appears in the $J_c(H)$ datapoints right at $H > H_{irr}$. The noise level was subtracted before the $f_p(h)$ dependencies were plotted. The resulting $f_p(h)$ dependencies at all temperatures are almost coinciding, thus the f_p scaling is well developed in both types of samples studied here.



Figure 3: Scaling of the pinning forces of the Bi2212 sample. The solid line represents the ECSM fitting curve with p = 0.37 and q = 3.

The pinning force scaling reveals a peak position, h_0 , of ~0.11. This value is smaller than $h_0 = 0.2$, which is expected for the flux pinning at 2D objects like grain boundaries [5] and was found for various Bi-based superconducting materials in the literature [8, 9, 10, 11, 12, 13, 14]. This small value of h_0 reflects the influence of anisotropy and percolation, which was described already in Refs. [9, 15]. The supercurrents



Figure 4: Scaling of the pinning forces of the Bi2212+Li sample. The solid line represents the ECSM fitting curve with p = 0.37 and q = 3.

flowing within an individual nanowire have to pass grains with many different orientations, so the anisotropy factor (i.e., the anisotropy of the upper critical fields) is important to describe the resulting current. According to Ref. [15] regarding the pinning force analysis in bulk MgB₂ samples, an anisotropy factor larger than 4, which is certainly fulfilled for Bi-2212, yields a position of $h_0 = 0.1$ for the grain boundary pinning which otherwise would give $h_0 = 0.2$. The influence of percolation effects on the peak position was found to be much smaller as described in Ref. [15]. However, the percolation effects were also found to reduce the peak positions as compared to the DH model. Therefore, one can safely conclude that the peak position $h_0 \sim 0.1$ describes grain boundary pinning in the case of an anisotropic material and percolating currents.

For a bulk type-II superconductor with sizes much higher than the London penetration depth λ_0 , the dependence of the critical current density J_c on the magnetic field H can be expressed by:

$$J_c(H) = J_{c0} \frac{1 - |H/H_{c2}|^{\alpha}}{1 + |H/(h_1 H_{c2})|^{\alpha}} \quad , \tag{1}$$

where $\alpha \leq 1$. This expression is a modification of the Kim-Anderson and Urban [16] dependencies. The extended critical state model [17, 18, 19] supports a faster decreasing of the $J_c(H)$ dependence due to an extension of the equilibrium surface. A type-II superconductor with size being comparable to λ_0 has the following $J_c(H)$ dependence:

$$J_c(H) = J_{c0} \frac{1 - |H/H_{c2}|^{\alpha}}{1 + |H/(h_1 H_{c2})|^{\alpha}} \left(1 - |H/H_{irr}|\right)^n \quad , \tag{2}$$

where *n* is the index defined by the geometry of the grain, n = 3 for a cylindrical sample and n = 2 for a thin plate[19].

We are interested in the behavior of the pinning force in the field range $H_p < H \ll H_{c2}$. Given $h_1 \ll 1$ and $H \ll H_{c2}$, one obtains the pinning force density from eq. (2):

$$F_p(H) \approx |H|^{1-\alpha} \left(1 - \frac{|H|}{H_{\rm irr}}\right)^n J_{\rm c0}(h_1 H_{c2})^{\alpha}$$
 (3)

After substitution for $\alpha = 1 - p$ and n = q, the well-known scaling law of Dew-Hughes [5] is resulted:

$$f_p(h) = \left(\frac{h}{h_0}\right)^p \left(\frac{1-h}{1-h_0}\right)^q \quad , \tag{4}$$

where h_0 is the position of the maximum, $h_0 = p/(p + q)$. The position of the pinning force maximum is easily determined from graphs such that one can find the value of α from $\alpha = 1 - n/(1/h_0 - 1)$. Given $h_0 = 0.11$ and n = 3 we obtain $\alpha = 0.63$. This corresponds to p = 0.37 and q = 3. These values differ from each other and are typical for anisotropic and polycrystalline superconductors [9, 15]. The solid line in Figs. 3, 4 represents the $f_p(h)$ dependence computed using eq. (3) and gives a good fit to the data in both cases.

Both types of nanowire networks (undoped and doped with Li) were found to show a good scaling of the flux pinning force data, and the peak, h_0 is located at ~0.11, which is clearly smaller than the expected value of 0.2 as found for Bi-based superconductors [8, 9, 10, 11, 12, 13, 14].

Here it should be noted that the high-*h* region of the F_p -scaling plots is influenced by several factors, (i) the magnetometer shows noise being close to its resolution, and (ii) the currents at high fields are very small and highly percolative. As a consequence, the irreversibility field is hard to determine from the plots. Nevertheless, the obtained scalung of the flux pinning forces turned out to be quite good in the temperature range investigated here, and the peak position of the f_p -plot is a good indicator of the active pinning mechanism, as flux creep effects do not play a role here.

It is interesting to note that the influence of anisotropy and percolation were only recently introduced to the pinning force analysis [15], when regarding the behavior of bulk, polycrystalline MgB₂ samples. In superconducting samples with a forced texture (bulk samples or tapes), the anisotropy does only play a minor role, and only recently, bulk, polycrystalline FeSe samples [20, 21] were found to behave similar to polycrystalline MgB₂ due to a well-developed coupling between the grains. According to Ref. [15], the changes in F_p due to changes in the percolation threshold are comparatively small since p_c is expected to vary only between 0.2 and 0.3 in three-dimensional systems [22]. The percolation threshold mainly depends on the number of connections between the grains; thus, clean dense samples have a comparatively small p_c and porous or dirty samples with a large amount of secondary phases have a larger p_c . For the present nanowire networks, the character of an individual nanowire is more 2D, as only a small number of grains form the entire nanowire. This number can be as small as 3-5 grains. Furthermore, the residing carbon in the nanowire samples may be located at the grain boundaries, and this effect is even enhanced in the case of the Li-doped samples, where we assume that Li is also found in the grain boundaries [23, 24]. Therefore, the nanowire network samples with their high porosity and the dirty grain boundaries should exhibit a stronger influence of p_c as compared to MgB₂ bulks analyzed in Ref. [15], which causes a move of the peak position in the $f_p(h)$ -diagram to lower values as observed here.

4. Conclusions

We have performed a pinning force scaling on the data obtained from electrospun Bi-2212 nanowire networks based on magnetization data. The analysis was performed using the Bean model and the ECSM model. The obtained scaling was found to quite good in the investigated temperature range 10 K < T < 35 K, and the peak position of the f_p -plots were found at ~0.11. Due to anisotropy and percolation, this value is reduced as compared to the value obtained for the grain boundary pinning in bulk materials.

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