

# Thickness Dependence of Critical Current Density in Thick MgB<sub>2</sub> Films

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**Abstract** - A study was performed to examine the  $J_c$  behavior as a function of thickness in MgB<sub>2</sub> films fabricated by *ex situ* annealing at 840°C of boron films, grown by chemical vapor deposition, in Mg vapor. The film thicknesses range between 300 nm and 10 μm. The values of  $J_c$  range from  $1.2 \times 10^7$  A/cm<sup>2</sup> for 300 nm to  $1.9 \times 10^5$  A/cm<sup>2</sup> for 10 μm film thickness at 20 K and self-field. The study shows that critical current density ( $J_c$ ) in MgB<sub>2</sub> films decrease with increasing film thickness, similar to that observed in YBCO coated conductors. The results were interpreted.

## I. INTRODUCTION

For coated-conductor applications, it is necessary to deposit thick films in order to maximize the critical current in the wire or tape and enhance the engineering critical current density. However, the main obstacle to higher current in thicker films is that the critical current density ( $J_c$ ) in the superconductor drops dramatically as the coating thickness is increased [1-7]. This limitation was first observed over 18 years ago in YBCO coated-conductors, yet only recently it became one of the most important remaining challenges in the coated conductors field as they are approaching the commercialization stage [5]. As MgB<sub>2</sub> coated conductors started hitting the road since the discovery of superconductivity in MgB<sub>2</sub> seven years ago [8], a study on  $J_c$  thickness dependence in MgB<sub>2</sub> films is deemed to be necessary.

## II. EXPERIMENTS

The films used in the study were fabricated by *ex situ* annealing of boron films in Mg vapor. Boron films were deposited on (0001) 6H-SiC substrates by CVD using a precursor gas of 5% B<sub>2</sub>H<sub>6</sub> in H<sub>2</sub>. The B film was then wrapped with a Nb foil and sealed in a low-carbon steel tube with high purity Mg pellets wrapped in a separate Nb foil and the tube was sealed in Ar atmosphere and sintered. The normal sintering procedure included a fast heating to a constant temperature of 840°C for all films in 30 min, followed by holding at that temperature for 2 to 6 h depending on the boron film thickness, and then quench cooling to room temperature in 10 min. The films experience a volume expansion of about 200% when changing from B to MgB<sub>2</sub>. For example, a ~2.5 μm thick B film results into a ~5 μm thick MgB<sub>2</sub> film after the reaction. Critical current densities were obtained using

magnetization measurements and determined from hysteresis loops using the Bean critical state model.

## III. RESULTS

Table 1 shows a list of films used in this study and their corresponding thicknesses as well as critical current density and critical current values at self field at temperatures of 5 K and 20 K. These values are higher than values previously reported in the literature for MgB<sub>2</sub> films fabricated by the *ex situ* annealing method [9-19]. The high  $J_c$  values reported here show the efficiency of the *ex situ* annealing method in the fabrication of MgB<sub>2</sub> films; and achieving them at such a low annealing temperature of 840°C is promising for low temperature industrial fabrication of resonance frequency (RF) MgB<sub>2</sub> cavities.

TABLE I  
MgB<sub>2</sub> FILMS STUDIED AND THEIR CORRESPONDING CRITICAL CURRENT DENSITY AND CRITICAL CURRENT VALUES

MgB <sub>2</sub> Film Thickness (μm)	$J_c$ (A/cm <sup>2</sup> ) @ 0T,5K	$J_c$ (A/cm <sup>2</sup> ) @ 0T,20K	$I_c$ (A/cm-w) @ 0T,5K	$I_c$ (A/cm-w) @ 0T,20K
0.3	$1.97 \times 10^7$	$1.2 \times 10^7$	591	360
0.5	$1.41 \times 10^7$	$7.42 \times 10^6$	705	371
1	$1.36 \times 10^7$	$7.28 \times 10^6$	1360	728
2	$2.1 \times 10^6$	$1.14 \times 10^6$	420	228
3	$1.79 \times 10^6$	$9.94 \times 10^5$	537	298
5	$1.03 \times 10^6$	$5.41 \times 10^5$	515	270
10	$3 \times 10^5$	$1.9 \times 10^5$	387	190

Figure 1 shows the MgB<sub>2</sub> films  $J_c$  thickness dependence curve at self field. It shows a clear decrease of  $J_c$  with increasing film thickness. This behavior of  $J_c$  decreasing with increasing MgB<sub>2</sub> film thickness is similar to the  $J_c$  thickness dependence behavior in YBCO coated conductors [1-7].

In an effort to understand the dependence of  $J_c$  on film thickness in MgB<sub>2</sub>, we used a model developed by Foltyn et al. which they used to understand the same behavior in YBCO. In their model, the value of  $j_c$  in YBCO is a function of the slice's distance from the substrate ( $z$ ), and  $j_c(z)$  is related to the average  $J_c$  for a film of thickness  $t$  by

$$J_c(t) = (1/t) \int_0^t j_c(z) dz.$$

The model suggests that there are two main values for  $J_c$ , the highest one near the film-substrate interface ( $j_{ci}$ ), which also corresponds to the one in very thin films, and the lowest one near the bulk value ( $j_{cb}$ ), which also corresponds to the one in thick films. Thus the model assumes that  $J_c$  decreases linearly from  $j_{ci}$  to  $j_{cb}$  over a range  $z_r$ , and then remains constant throughout the rest of the film, leaving  $z_r$  as the only adjustable parameter. By using that model, both calculated and measured  $J_c$  values were in excellent agreement for experimental results, where the best fit was obtained for a  $z_r$  of 0.65  $\mu\text{m}$  in YBCO. Applying the same model to  $\text{MgB}_2$  films show the same agreement. Figure 2 shows the experimental  $\text{MgB}_2$  confirmation of the conceptual incremental function  $j_c(z)$ . The continuous line in the figure is the incremental critical current density model used for the fit. The best fit is obtained when  $z_r$  is adjusted to 2  $\mu\text{m}$ . The distance from substrate ( $z$ ) in figure 2 corresponds directly to the  $\text{MgB}_2$  film thickness, where each thickness represents a new  $\text{MgB}_2$  film. The reasons for the  $J_c$  decrease with increasing film thickness can be the high density of flux pinning defects and dislocations near the film-substrate interface, the presence of vortices in the film, and/or the degradation of microstructure at higher thickness.

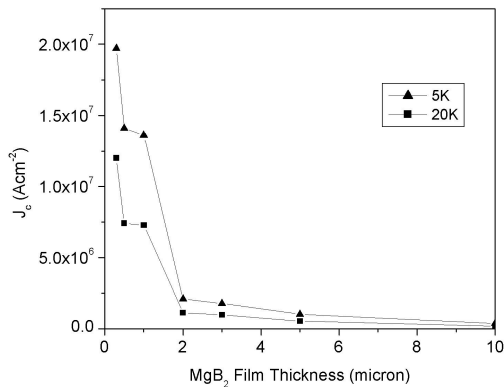


Fig. 1.  $\text{MgB}_2$  films  $J_c$  thickness dependence at self field.

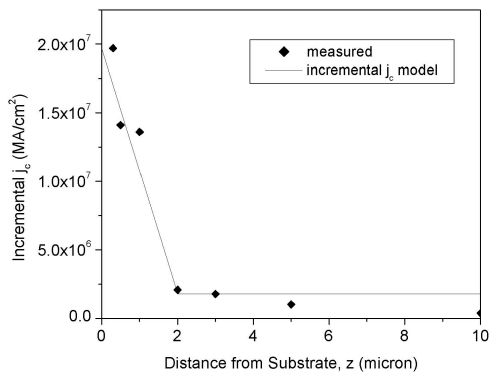


Fig. 2. Experimental confirmation of the conceptual incremental function  $j_c(z)$ . The best fit is obtained when  $z_r$  is adjusted to 2  $\mu\text{m}$   $\text{MgB}_2$  film thickness.

Figure 3 shows the  $I_c$  thickness dependence of  $\text{MgB}_2$  films at self field. The figure displays that, similar to YBCO coated conductors, critical current in  $\text{MgB}_2$  films drops beyond  $\sim 1 \mu\text{m}$  thickness, which is probably due to impurity diffusion during annealing and microstructural degradation for thicker films. This indicates that 1  $\mu\text{m}$   $\text{MgB}_2$  films can carry critical current that is higher than thicker  $\text{MgB}_2$  films, which is economical for industrial applications such as the fabrication of  $\text{MgB}_2$  superconducting cavities and coated-conductor wires and tapes.

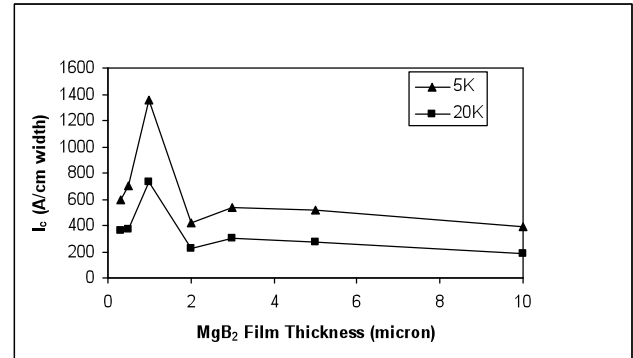


Fig. 3.  $\text{MgB}_2$  films  $I_c$  thickness dependence at self field.

#### ACKNOWLEDGMENT

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