Compositional and Microstructural Profiles across Nb₃Sn Filaments Produced by Different Fabrication Methods

P. J. Lee and D. C. Larbalestier

Abstract—The next generation of high field magnets for fusion and accelerator applications requires the development of Nb₃Sn strand with significantly higher critical current densities. Our recent work has shown that when the specific pinning force is normalized to the grain boundary density (the primary pinning site in this material), a strong increase in pinning force is observed for high Sn content strand, suggesting a strong compositional dependency for pinning in this material. In this study we use advanced FESEM techniques to quantify the compositional and microstructural variations to a sub-100 nm level. A stepped compositional and microstructural variation was observed across a bronze-processed Nb₃Sn filaments with sharp interfaces for both Cu(Sn)-Nb₃Sn and Nb₃Sn-Nb. Microstructural variation was observed on the scale of the original filaments in high Sn, MJR but not across the coalesced filament mass. A very high microstructural uniformity was observed in a PIT monofilament. Thus a major influence of Sn appears to be exerted through its influence on the composition gradient.

Index Terms—niobium-tin compounds, microstructure, critical current, composition gradient

I. INTRODUCTION

A significant increase in the critical current density of Nb₃Sn is required for both medium and high field VLHC options - are we already close enough to perfection that significant improvements can not be expected?

The high energy physics led development of Nb-Ti has resulted in a commodity, commercial product available in long lengths at low cost from a number of highly competitive companies. For Nb-Ti:

1. With the exception of a small region at the edge of the filaments the microstructure is homogeneous across each filament.
2. The micro-structure/chemistry is uniform from filament to filament.
3. The filaments are uniform in shape and size and are

Fig. 1. FESEM fractographs revealing a) Transverse cross-section, b) longitudinal cross-section and c) filament-surface microstructure in a bronze-process Vacuumschmelze ITER strand.
uncoupled.

4. The role of chemistry, heat treatment, and processing on the $J_c$ is well understood and well controlled.

The contrast between Nb$_3$Sn and Nb-Ti is quite striking:

1. The microstructure is inhomogeneous across each and every filament (e.g. [1]).
2. The microstructure/chemistry generally varies with filament position (e.g. [1]-[4]).
3. The filaments are generally irregular in shape and size and become more coupled as superconductor volume in the cross-section increases (e.g. [5]).
4. The role of chemistry, heat treatment and processing on $J_c$, $H_{c2}$, or $I_{c1}$ or $T_c$ is not well understood and is thus largely uncontrolled.

The task of understanding the influences of processing variables is complicated by the complexity of the Nb-Sn system and the inhomogeneity of the strands. In this paper we examine aspects of the microstructural and chemical inhomogeneity for bronze and internal tin MJR conductors.

II. EXPERIMENTAL PROCEDURE

A. Microstructure

The inhomogeneity of Nb$_3$Sn composites provides a considerable challenge to microstructural characterization because of variations within filaments and from filament to filament. The location within the strand cross-section and within the filament must be known and numerous analyses must be performed in order to assess this variation. Traditional analysis by Transmission Electron Microscopy, TEM is then exceptionally laborious and difficult because of the difficulty in preparing a full-filament, electron-transparent cross-section, while retaining location information of that filament. Recent advances in the resolution of Field Emission Scanning Electron Microscopes, however, have made it possible to reliably quantify the microstructure of A15 phases by fractography [1]. Fig. 1 shows FESEM fractographs of a bronze process ITER-VAC strand after a standard ITER heat treatment. The contrasts are quite striking between the transverse and longitudinal cross-sections and the grain size at the filament surface. In order to show the variation in microstructure with position we use a process of position normalization. Each analyzed microstructural feature is located with respect to its distance from the filament centroid and the closest reaction interface.

B. Composition

We have recently extended the position normalization technique to Backscattered Electron Imaging, BEI, in the FESEM, in order to observe chemical changes at a resolution below 100nm [6].

The potential for applying backscattered electron spectroscopy, BES, to reveal chemical composition changes with sub-micron spatial resolution has been explored by a number of authors [7]-[9]. Practical resolution appears to be only limited by the signal to noise ratio [10]. By breaking a conventionally obtained BEI image, into individual pixels and normalizing the positions of the pixels to the interfaces under study, we can combine signals from the entire image. For Nb$_3$Sn filaments with < 200 nm grain diameters, this approach also averages over multiple grain orientations, thus eliminating crystallographic back scatter yield variations. By Monte Carlo simulation [11] of the electron trajectories, the mean maximum depth of the backscattered electrons in Nb$_3$Sn is 40-50 nm for a 5 kV accelerating voltage. This rises

![Fig. 2 Simulated backscatter traverse across depleted 2 atomic % Sn bronze (left) to stoichiometric A15 (right) boundary using 50,000 electrons per point. 100,000 electron simulated levels for bulk Cu, Nb, 25 atomic % Sn Nb$_3$Sn and 20 atomic % Sn Nb$_3$Sn are also shown.

Fig. 2 Compositional variation across bronze process filament in a fully heat treated Vacuumschmelze ITER strand.

![Fig. 3 Compositional variation across bronze process filament in a fully heat treated Vacuumschmelze ITER strand.](https://example.com/fig3.png)
with accelerating voltage to over 180 nm at 12 kV. With the interfacial roughness indicated in Fig. 1c, it is important to keep the accelerating voltage as low as possible to obtain resolution on the sub-100 nm scale. A backscatter traverse across a depleted bronze to stoichiometric Nb$_3$Sn interface is shown in Fig. 2.

III. RESULTS

A. Bronze Process ITER strand.

Fig. 3 shows the 5 kV electron backscatter variation across a bronze process filament. The gradient suggests a decline in Sn composition across the radius of the filament. The interface is as sharp as that simulated (Fig. 2) for a planar depleted bronze to 25 atomic % Sn A15 interface. The backscatter intensity across the A15 phase is shown in detail in Fig. 4. A flat intensity is observed over the first ~300-400 nm of the filament radius, followed by a continuous decline until approximately 900 nm from the bronze-filament interface.

In Fig. 5 the change in grain morphology (aspect ratio and grain boundary density) across the filament radius is shown. The aspect ratio is low for the first ~300 nm, after which there is a steady increase. The grain boundary density is highest near the filament surface, after which there is a plateau extending to ~300 nm. After 300 nm there is a changeover which initially sees a spike in grain boundary density followed by a continuous decline.

The inverse grain size, 1/d, is often used as an indicator of $F_{\text{pmax}}$, in A15 superconductors, and allows comparison of this data with earlier work. In this work we calculate the diameter, d*, from the transverse cross-sectional area of the grain assuming it to be circular. The grain morphology has a relatively small impact on grain boundary density compared with d* and consequently the grain boundary density and 1/d* trends are almost identical even for filaments such as this with highly aspected grains. In Fig. 6, we contrast the distribution in 1/d for the outer, high-Sn 100 nm of the radius with that of the grains beyond 500 nm from the bronze, where BSE analysis indicates the Sn level is declining. As with the other measured microstructural features in Nb$_3$Sn (and Nb-Ti) the size distributions are best described as log-normal. The log-normal generated mean 1/d* for the outer 100 nm is 11.1 $\mu$m$^{-1}$, compared with a value of 6.7 $\mu$m$^{-1}$ for the lower

Fig. 7 FESEM fractograph detail from within the A15 layer in a high $J_c$, high Sn MJR strand. The original filaments are surrounded by a layer of large A15 grains. The highlighted filament is analyzed individually.
Sn interior (>500 nm from the bronze).

B. High Sn MJR internal Sn process strands.

Analysis of high-Sn, high-\(J_c\), internal-Sn, MJR composites is complicated by the growing together of the filaments into a single filament mass. In Fig. 7 an original filament within the coalesced A15 layer of a high-Sn MJR composite is outlined. The microstructural variation across this layer (Fig. 8b) is strong compared with that averaged over the coalesced A15 layer (Fig. 8a).

A position normalized backscatter profile across the outer 4 \(\mu \text{m}\) of the reacted A15 layer is shown in Fig. 9. Although there is variability at the individual filament scale, the overall trace is flat.

C. High Sn source PIT strands.

ECN-type PIT filaments characteristically have a small interior layer of large grains surrounded by the main layer of fine grains. In Fig. 10 we show the grain size variation across the fine-grained A15 region in a PIT monofilament manufactured by Supercon. There is only a shallow gradient in grain size over the entire 20 \(\mu \text{m}\) layer.

IV. SUMMARY

We now have the tools to provide a quantitative description of complex Nb\(_3\)Sn composites. These initial studies reveal important insights into the microstructural control of these strands. In a bronze-process strand we observe the onset of columnar growth simultaneously with a downturn in %Sn. In the high-Sn strand we observe that the principal variations are across the original filaments and not across the coalesced filament pack. In the PIT strand, with a very high Sn content powder core, a uniform, and fine grain size can be achieved over a very long distance.

ACKNOWLEDGMENT

Internal Sn strand was provided by Oremet Wah Chang and by Seung Hong of OF-ST. Reacted monofilament PIT Nb\(_3\)Sn was supplied by Terrence Wong of Supercon. Processing and metallography at the UW was performed by William Starch, Danica Christensen, and Walid Gabr-Rayan.

REFERENCES


