#### PAPER

## Contact resistance between two REBCO tapes: the effects of cyclic loading and surface coating

To cite this article: Jun Lu et al 2018 Supercond. Sci. Technol. 31 085006

View the article online for updates and enhancements.

### **Related content**

- <u>Contact resistance between two REBCO</u> <u>tapes under load and load cycles</u> Jun Lu, Robert Goddard, Ke Han et al.
- Metal-as-insulation variant of no-insulation HTS winding technique: pancake tests under high background magnetic field and high current at 4.2 K T Lécrevisse, A Badel, T Benkel et al.
- <u>Mechanism of notable difference in the</u> <u>field delay times of no-insulation laver-</u> <u>wound and pancake-wound REBCO coils</u> Y Suetomi, K Yanagisawa, H Nakagome et al.

## IOP ebooks<sup>™</sup>

Bringing you innovative digital publishing with leading voices to create your essential collection of books in STEM research.

Start exploring the collection - download the first chapter of every title for free.

Supercond. Sci. Technol. 31 (2018) 085006 (8pp)

# Contact resistance between two REBCO tapes: the effects of cyclic loading and surface coating

#### Jun Lu<sup>®</sup>, Jeremy Levitan, Dustin McRae and Robert Walsh

National High Magnetic Field Laboratory, Tallahassee, FL 32310, United States of America

E-mail: junlu@magnet.fsu.edu

Received 23 April 2018 Accepted for publication 18 June 2018 Published 10 July 2018



#### Abstract

No-insulation (NI) superconducting REBCO magnets have advantages of self-quenchprotection, a very high engineering current density and high mechanical strength, and the potential to reach very high magnetic fields. However, NI REBCO magnets have drawbacks of a long magnet charging time and high field ramp losses. These can be mitigated by controlling the turn-to-turn contact resistivity ( $R_c$ ). In an effort to control  $R_c$ , we consider two approaches. One is coating a REBCO conductor with various resistive thin films, and the other is to use a stainless steel (SS) tape as an interlayer which is also coated with different metallic films. We present experimental results of  $R_c$  of an as-received sample under cyclic contact pressure of 2.5–25 MPa up to 30 000 cycles. After an initial increase in  $R_c$  for the first 10–20 cycles,  $R_c$  decreases to about one tenth of its initial value after a few hundred cycles. A warm-up and cool-down thermal cycle does not significantly change the low  $R_c$  resulting from a previously high number of load cycles. We also studied  $R_c$  of REBCO tapes that are coated with different resistive layers and interlayers. In order to increase  $R_c$ , we experimented with electro- and electroless plating of Ni, Cr, and Ni–P. We also measured  $R_c$  with a thin metallic interlayer as a coil co-winding material which included Cu, SS, and SS plated with Ni and Cu. A SS interlayer increases  $R_c$  by about three orders of magnitude; while the Cu plated SS interlayer only increases  $R_c$  by one order of magnitude. Finally, we treated the as-received REBCO surface by oxidation using an Ebonol® C solution. This controlled oxidation allowed the  $R_c$  to be controlled over a wide range.

Keywords: coating, contact resistance, no-insulation, REBCO, cyclic loading

(Some figures may appear in colour only in the online journal)

#### 1. Introduction

No-insulation (NI) REBCO pancake magnet coils have several advantages over conventional insulated coils [1–3]. Due to low turn-to-turn electrical resistance, when a magnet quench happens in a NI REBCO coil, the quench current automatically bypasses the normal zone, and a hot spot is avoided. This self-quench-protection ability eliminates the need for a quench detection and protection system that can be very challenging and costly in a large high field REBCO magnet [4]. Consequently, a NI REBCO coil is very stable which allows for a thinner copper stabilizer to be used on the REBCO conductor. The thinner stabilizer and the elimination of insulation make the engineering critical current density and the effective mechanical strength of the coil very high. These suggest that this technology enables very compact magnets to reach a very high magnetic field.

There are, however, drawbacks with the current state of the NI REBCO coil technology. NI coils typically have long charging delay times [5–7] as well as high field ramp losses [8] which is a concern especially for cryo-cooled magnets where the cooling power is limited. These issues are directly related to low turn-to-turn contact resistivity ( $R_c$ ) which is defined as contact resistance multiplied by contact area, and can be mitigated by somehow increasing  $R_c$ . However, very high  $R_c$  compromises the turn-to-turn current transfer

#### Supercond. Sci. Technol. 31 (2018) 085006

Table 1. REBCO coating methods.

Coating	Electrolyte	<i>T</i> (°C)	Current density (A $dm^{-2}$ )	Growth rate $(\mu m \min^{-1})$
Ni	Caswell Ni	40	1	0.12
Ni–P	Caswell electroless Ni-P	90		0.5
Cr	$Cr_2O_3$ : $H_2SO_4$ : $H_2O = 45$ : 200: 0.45	22	2.5	0.03
Cu on SS	$CuSO_4 \cdot 5H_2O : H_2O = 7 : 250 \text{ (add } H_2SO_4 \text{ until } pH = 1)$	22	0.3	0.05

capability therefore jeopardizes the coil's self-protection ability. In principle,  $R_c$  can be optimized to achieve a relatively short charging delay time and low ramp losses without jeopardizing the coil's self-protection ability. Many theoretical analyses have been done to predict the quench behaviors of NI magnet coils and the effect of  $R_c$  [9–12]. These studies suggest that  $R_c$  values ranging from 1000 to  $100\,000\,\mu\Omega$  cm<sup>2</sup> might be suitable, which is significantly higher than about  $70 \,\mu\Omega$  cm<sup>2</sup> in the current NI coils [6]. Therefore it is critical to develop a technology to control  $R_c$  to this desirable level. To this end, the effect of 'insulation' by a metallic co-winding interlayer is also explored [13–15]. In addition, it is proposed that the conductor with graded  $R_c$  is used to reduce the ramp loss [8] as well as to improve stability and recovery speed after a quench [16] in large NI REBCO magnets.

Evidently  $R_c$  is a very critical parameter in the development of NI coil technology. In our previous paper on this subject, we reviewed REBCO  $R_c$  studies in the literature and reported our measurements and analysis of  $R_c$  between two SuperPower REBCO samples as a function of temperature and contact pressure for several load cycles [17]. In this paper, considering the fact that a practical superconducting magnet typically experiences a few thousand charge-discharge cycles in its lifetime, we study the effect of a large number of pressure cycles on  $R_c$ . Moreover, we performed a large number of experiments in order to develop a technology to control  $R_c$ . Here we present the effects on  $R_c$  of a thin layer of metal deposited on REBCO, various metallic interlayers, and oxidation of REBCO tape. Based our experimental results, we propose a practical method to control  $R_c$ . The implications of our results on NI REBCO coil applications will be discussed.

#### 2. Experimental

#### 2.1. Samples

Samples used in this experiment are REBCO conductors made by SuperPower (SCS4050AP) and SuNAM, both are surrounded by electroplated copper stabilizers. The Super-Power conductor is 4 mm wide with overall thickness of 95  $\mu$ m which includes a 20  $\mu$ m copper stabilizer on each side. The nominal critical current in the self-field at 77 K is 80 A. The resistivity ratio between room temperature and 4.2 K for the copper stabilizer layer is measured on similar SuperPower conductors to be about 50. Since SuperPower conductors were used for the majority of our experiments, the data in this paper are all from SuperPower SCS4050AP samples unless stated otherwise. The SuNAM conductor is 4 mm wide with an overall thickness of 135  $\mu$ m which includes a 30  $\mu$ m laminated copper stabilizer. The nominal critical current in the self-field at 77 K is 105 A. One SuNAM sample has 1  $\mu$ m stainless steel (SS) cladding.

#### 2.2. Thin film coating and surface oxidation

In order to control  $R_c$ , various kinds of surface coatings were made. The surface coating methods chosen in this experiment are relatively low cost, and easy to scale up to high volume production. Due to these reasons, a more sophisticated process such as physical vapor deposition (PVD) was not considered. It should be noted, however, a REBCO conductor manufacturer is typically equipped with a PVD system with reel-to-reel operation capabilities. Therefore, for a REBCO manufacturer to coat a thin film using PVD on a REBCO tape may not be very expensive.

Cr, Ni and Ni-P films were deposited on REBCO conductors directly. Cr film was electroplated on REBCO samples using an electrolyte made of 200 ml deionized water, 45 g Cr<sub>2</sub>O<sub>3</sub>, and 0.45 g sulfuric acid. Ni film was electroplated using a commercial nickel plating solution (Caswell Inc., USA). Ni-P with 5%-7% phosphorus is a high resistivity material with a high hardness and good wear resistance, and all these properties potentially lead to high contact resistance. Therefore several samples were plated with Ni-P on Ni plated REBCO using a commercial electroless Ni-P plating solution (Caswell Inc., USA). All these plating processes resulted in dense, uniform and adherent metallic coatings. In addition, Ni and Cu were plated on 316 SS tape, which was then used as an interlayer between two REBCO tapes. For Cu electroplating, a CuSO<sub>4</sub> solution was used. The detailed electrolytes and other conditions used for these plating processes are summarized in table 1.

In addition, we experimented with oxidation of the REBCO surface. In the oxidation experiments, we used Ebonol® C, a mixture of sodium hydroxide and sodium chromate [18] and dissolved it in deionized water at a weight ratio of Ebonol® C: water = 18: 80. The REBCO surface was oxidized in Ebonol® C solution at 98 °C for various durations. The thicknesses of the oxide layer on the copper strips were measured by weight loss after removing the oxide layer with a solution of HCl:  $H_2O = 1$ : 10, and assuming the density of the oxide is 6.31 g cm<sup>-3</sup> for CuO. As shown in figure 1, the oxide thickness increases almost linearly with the logarithm of time. Depending on the thickness of the oxides, the color of the oxidized REBCO surface changed from dark brown after 10 s to black after 30 s.



**Figure 1.** CuO layer thickness versus Ebonol® C oxidation time. Ebonol® C concentration is about 18 wt%, temperature was 98 °C.

## 2.3. Transverse load cycling and R<sub>c</sub> measurement at cryogenic temperatures

For  $R_c$  measurement, a probe used in our previous experiment [17] was modified to fit to a Material Test System tensile machine which enables fast load cycling at a rate up to 10 Hz at 77 K or 4.2 K, load cycles of 2.5–25 MPa being applied. More details of the experimental setup are described in our previous paper [17].  $R_c$  was measured during the load cycling by applying  $\pm$  1.00 A current from a Keithley 2400 bipolar DC current source, and measuring voltage with a Keithley 2010 digital multimeter. The instrument error for resistance measurements was estimated to be less than 0.1  $\mu\Omega$  cm<sup>2</sup>, obtained from a measurement of a REBCO solder joint sample using otherwise the same experimental conditions.

#### 3. Results and discussions

#### 3.1. R<sub>c</sub> variation due to surface conditions

It is known that surface conditions such as slight oxidation or other contamination, as well as roughness, have a significant effect on contact resistance. Since the surface condition of a commercial REBCO conductor is usually not specified, it is expected that  $R_c$  from different manufacturers and different production batches varies widely. We investigated this variation by measuring  $R_c$  of REBCO with different surface conditions at 25 MPa contact pressure at 77 K. As shown in table 2,  $R_c$  varied drastically from sample to sample. We observed a systematic difference in  $R_c$  between SuperPower and SuNAM conductors, which might be contributed to their respective surface flatness and roughness. Because while a SuperPower conductor has a rough finish with increased thickness at the center and the edges [17], a SuNAM conductor has a smooth and shiny finish with a relatively uniform thickness. After slightly polishing the surface of the Super-Power conductors with an abrasive (Scotch-Brite) or chemical etching it with a solution of HCl:  $H_2O = 1$ : 1,  $R_c$  was reduced by a factor of about ten. When the conductor surface was handled by ungloved hands,  $R_c$  increased by a factor of two to six. When a consistent surface cleaning method was applied, e.g. by wiping with an ethanol soaked tissue, the measured  $R_c$  values became much more consistent. Variation along the conductor length of the same piece was also observed.

#### 3.2. Effect of contact load cycling

We reported in our previous paper [17] that  $R_c$  increases with contact load cycling up to 14 cycles of 140 MPa at 77 K. In order to simulate a user magnet with frequent field ramps, in this paper we measured  $R_c$  of the SuperPower conductor with a larger number of load cycles between 2.5 and 25 MPa. This contact pressure range was chosen because it seems to be a reasonable range for radial stresses in a coil. Admittedly, the radial stress in a coil which depends on winding tension, thermal stress, and electromagnetic load during the coil's operation, is not uniform and very difficult to calculate precisely.

Figure 2(a) shows a series of curves of  $R_c$  versus contact pressure P measured after different numbers of load cycles at 77 K. Each  $R_c$  (P) curve shows the expected behavior where  $R_c$  decreases with transverse load. It is also evident that  $R_c$ increases within the first 10-20 cycles then decreases quickly. After a few hundred cycles,  $R_c$  levels off to a value only about one tenth of its initial value. This trend is presented more clearly in figure 2(b) where  $R_c$  at 25 MPa pressure is plotted against load cycles in a semi-logarithm scale. After only about 200 cycles,  $R_c$  decreases to about  $4 \mu \Omega$  cm<sup>2</sup> from the initial value of 40  $\mu\Omega$  cm<sup>2</sup> then levels off. This experiment was repeated a few times, and similar results were obtained. Since we are interested in a high field REBCO magnet operating at 4.2 K, the load cycling effect was studied at 4.2 K as well. The results are presented in figure 3 and compared with the 77 K data. Despite the difference between the 77 K and 4.2 K data which can be partly attributed to the variation from sample to sample, a similar trend was observed. The initial higher  $R_c$  at 4.2 K is likely due to random variations in surface conditions. The significant decrease in  $R_c$  seems to occur after a few hundred cycles, somewhat later than that at 77 K. At the end of 20 000 load cycles at 4.2 K,  $R_c$  is only 0.85  $\mu\Omega$  cm<sup>2</sup> which is significantly lower than 4  $\mu\Omega$  cm<sup>2</sup> at 77 K. The lower asymptotic  $R_c$  value at 4.2 K is likely due to the lower resistivity of copper, consistent with [17]. After these load cycling experiments, the sample surfaces were examined with a light microscope up to  $1000 \times$  magnification. No discernable surface morphology changes were observed.

This significant decrease in  $R_c$  upon a large number of contact pressure cycles was not discovered in our previous experiment [17] because the number of cycles was limited to 14. In order to understand this load cycling effect, we review the Holm theory of contact resistance where  $R_c$  is related to mechanical and electrical properties of the contacting surface by [17]

$$R_C = 0.886\rho_{\sqrt{\frac{HV}{PN}}} + \frac{\rho_f dHV}{P}$$
(1)

Sample ID	Surface	Number of samples	$R_c$ at 25 MPa ( $\mu\Omega$ cm <sup>2</sup> )
SuNAM	Ethanol wiped	2	5.9–7.5
SuperPower-1	Ethanol wiped, beginning of the spool	13	16–58
SuperPower-2	Ethanol wiped, end of the spool	7	47-180
SuperPower-3	Ethanol wiped then finger handled	1	100
SuperPower-4	Scotch-Brite wiped	5	0.8–7.3
SuperPower-5	HCl etched	6	3.4–22



**Figure 2.**  $R_c$  of as-received SuperPower SCS4050 at 77 K. (a)  $R_c$  versus contact pressure curves for different numbers of load cycles. (b)  $R_c$  measured at 25 MPa as a function of number of load cycles.

where  $\rho$  and  $\rho_f$  are resistivity of the contact material (Cu in our case) and the surface contamination film (probably Cu oxides) respectively, *HV* the Vickers hardness of the contact material, *N* the number density (in m<sup>-2</sup>) of contact spots, and *d* the thickness of the oxide film. According to equation (1), under a given contact pressure, the harder the contacting material, the smaller contact spot radius, therefore higher  $R_c$ .



**Figure 3.**  $R_c$  measured at 25 MPa as a function of number of load cycles at 77 K and 4.2 K.

So we speculate that initially the surface cold-work hardened the material and increased its resistivity, and both led to higher  $R_c$  as explained in our previous paper [17]. However, the cold-working effect reached its saturation at a few tens of cycles. The quick decrease in  $R_c$  after that might be explained by gradual wear-out of the oxide layer on the copper surface. The fact that there is a ten-fold difference in  $R_c$  between asreceived and freshly HCl etched samples seems to be consistent with the existence of a thin resistive oxide layer.

This appreciable load cycling effect has very important implications for NI magnet coil technology. Because a user magnet usually undergoes at least a few thousand charging/ discharging cycles in its lifetime.  $R_c$  of a magnet coil measured without significant load cycling might considerably underestimate the charging delay time, which, in some cases, could be ten times longer than its initial value after only a few hundred charging cycles. It should also be noted that during operation the contact pressure in a NI coil is not uniform and depends on the specific magnet design. Therefore the result from these experiments may not fully represent the situation in a real magnet coil.

#### 3.3. Effect of thermal cycling

In its lifetime, a superconducting magnet is usually warmed up to room temperature many times. Therefore it is very



**Figure 4.** Effect of thermal cycling.  $R_c$  versus number of load cycles at 4.2 K measured after first and second cool-down. During warming up to 273 K, the contact pressure was kept at 2.5 MPa.

relevant to investigate the effect of thermal cycling on  $R_c$ . In the following experiment,  $R_c$  of a sample was tested for up to 20 000 load cycles at 4.2 K. Then the sample was allowed to naturally warm up inside the cryostat overnight under a 2 MPa constant pressure to 273 K. The sample was cooled down to 4.2 K again and  $R_c$  was measured again. The measured  $R_c$  versus number of cycles for both the first and the second cool-downs are shown in a log–log plot in figure 4. As shown in the figure, thermal cycling does not reset  $R_c$  to its initial value before load cycling. After thermal cycling,  $R_c$ only increases moderately from 0.8  $\mu\Omega$  cm<sup>2</sup> to about 3.1  $\mu\Omega$ cm<sup>2</sup>, still much less than the initial value of nearly 100  $\mu\Omega$ cm<sup>2</sup>. This suggests that in a magnet coil the drastic decrease in  $R_c$  due to load cycling cannot be reset by warming the coil up to room temperature.

#### 3.4. Control of R<sub>c</sub>

For NI magnet applications, it is highly desirable to control  $R_c$  so the coil charging delay time is relatively short without compromising the coil's self-protection ability. Our study on the load cycling effect suggests that it is also important for  $R_c$  to be insensitive to load cycling. To this end, two approaches are considered. The first one is coating the conductor surface with a thin resistive film so  $R_c$  is increased and may be controlled by either the chemical composition or the thickness of the film. The second one is winding the coil with a resistive co-winding tape, and coating the co-winding tape with various resistive films to control  $R_c$ . The results of these efforts are presented in the following subsections.

3.4.1. Metallic cladding. One of the earliest efforts to control  $R_c$  was made by the REBCO manufacturer SuNAM [19]. REBCO conductors were coated with up to 1  $\mu$ m SS cladding (MC-REBCO). The charging delay of the test NI coils using MC-REBCO conductors reduced considerably. We measured  $R_c$  of a SuNAM MC-REBCO as a function of load cycles at



**Figure 5.**  $R_c$  versus load cycles of the SuNAM metallic cladded sample as compared with the SuperPower as-received sample.

both 77 K and 4.2 K. The results are plotted in figure 5 together with data from as-received SuperPower samples. Evidently SuNAM's MC-REBCO sample has higher  $R_c$  than the as-received SuperPower sample. Furthermore,  $R_c$  does not decrease drastically with load cycles especially at 77 K. It might be possible to further control  $R_c$  by changing the thickness of the SS cladding layer.

Encouraged by this result, we experimented with depositing metallic layers on REBCO by electroplating, which is more economical and easier to scale up compared with the magnetron sputtering method used by SuNAM [10]. In these experiments, short samples of SuperPower conductor were plated with Ni or Cr films of different thicknesses. However, the plated Ni thin films did not seem to increase  $R_c$ significantly as shown in table 3, where  $R_c$  under first load at 25 MPa for different surface coatings are listed. Cr plating and the native oxide layer on the Cr surface seem to have the potential to significantly increase  $R_c$ , but our results on three Cr plated samples vary widely, therefore are not presented in table 3. Further investigation is needed to understand the formation of the native Cr oxides and their effect on  $R_c$ . On the other hand, the environmental concerns of the hexavalent Cr plating makes it a less attractive option.

Ni5-7wt%P is a highly electrical resistive, wear and corrosion resistant material that can be plated to metal surfaces by an electroless process [20]. Therefore in order to increase the  $R_c$ , Ni–P of various thicknesses was plated on SuperPower samples which were already plated with a 1  $\mu$ m pure Ni layer. The 77 K  $R_c$  measurements on Ni–P plated samples show a near linear increase in  $R_c$  with Ni–P thickness, as shown in figure 6(a). This demonstrates a certain level of  $R_c$  control by varying the Ni–P thickness. However, the Ni–P sample seems to be sensitive to load cycling. As shown in figure 6(b), after only a few hundred load cycles,  $R_c$  of a 1  $\mu$ m Ni–P plated sample drops down almost two orders of magnitude and becomes lower than that of an as-received uncoated sample before load cycling.

Sample	Surface	Number of samples	$R_c$ at 25 MPa ( $\mu\Omega$ cm <sup>2</sup> )
SuNam	$1 \mu m$ SS cladding	2	180, 224
SuperPower	$1 \mu m$ Ni plating	3	19–64
SuperPower	Graphite sprayed	1	180



**Figure 6.** Effect of Ni–P plating. Ni–P is plated on REBCO with a 1  $\mu$ m pure Ni buffer layer (a)  $R_c$  versus Ni–P thickness. (b)  $R_c$  versus load cycles for 1  $\mu$ m Ni–P plated sample with an as-received sample for comparison.

3.4.2. Co-wind tape and its coating. Co-winding a REBCO pancake coil with 316 SS tape is another attractive option for a NI coil. Here the co-winding 316 SS tape is used as mechanical reinforcement as well as a metallic interlayer to increase  $R_c$ . The properties of 316 SS co-wound REBCO coils have been analyzed [21] and experimental results reported in [14, 22, 23]. In order to verify the  $R_c$  values reported in these works, we measured  $R_c$  of two SuperPower REBCO conductors which have a 50  $\mu$ m thick 316 SS tape placed in between them. As shown in figure 7,  $R_c$  is in the order of 10 000–100 000  $\mu\Omega$  cm<sup>2</sup> which is much higher than



**Figure 7.** Effect of co-wind tape,  $R_c$  versus load cycles for REBCO inserted with SS tape, Cu tape, Ni plated SS tape and Cu plated SS tape.

 $\sim 1000 \ \mu\Omega \ \mathrm{cm}^2$  reported in [14] and about three orders of magnitude higher than measured without the 316 SS interlayer. Since the resistance of  $50 \,\mu\text{m}$  SS tape in its thickness direction is only in the order of  $1 \ \mu\Omega \ cm^2$  at 77 K, the high  $R_c$  is likely due to the native oxide layer on the SS surface as well as its appreciable hardness compared with copper. This is consistent with equation (1) where high native oxide resistivity  $\rho_{\rm f}$  and high hardness HV make  $R_c$  higher. With such high  $R_c$ , the self-protection ability of such an NI coil becomes questionable, especially for the coils with large stored energy. So in an effort to reduce and control such high  $R_c$ , about 1  $\mu$ m of pure nickel or copper was plated on the SS tape. As also shown in figure 7, these thin films of pure metals on SS tape reduced  $R_c$  dramatically. The remarkable effect of these thin films on  $R_c$  demonstrates the importance of the native oxide resistivity and contact material hardness. This moderation of  $R_c$  by a thin film on SS co-wind tape seems to be a viable option for controlling  $R_c$  in a NI magnet coil. It is also very interesting to see that with a 125  $\mu$ m thick Cu tape between REBCO tapes, as plotted in figure 7,  $R_c$  is similar to that with the Cu plated SS. This further demonstrates the dominant role surface materials play in controlling  $R_c$ .

3.4.3. Surface treatment by Ebonol® C oxidation. Since electroplating pure metals on REBCO seems to have limited potential in controlling  $R_c$ , an alternative is to deposit a layer of highly resistive material on REBCO, so  $R_c$  may be controlled reliably by changing the thickness of the deposited layer. Copper oxide (CuO), being a semiconductor, is an obvious option. We used a commercial copper blackening agent Ebonol® C to oxidize the Cu surface of REBCO tapes



**Figure 8.**  $R_c$  versus load cycles for REBCO oxidized by Ebonol® C for 10, 30, and 60 s.



**Figure 9.** A summary of all the  $R_c$  versus load cycle curves measured at 77 K from samples of different surface and interlayer conditions.

and calibrated the oxide thickness as a function of oxidation time (figure 1). Figure 8 shows the  $R_c$  as function of load cycle for samples oxidized for 10, 30 and 60 s. The  $R_c$  of these samples ranges for two orders of magnitude and are significantly higher than for the as-received REBCO. Furthermore, they do not seem to decrease too much with load cycling. Therefore by controlling the oxidation time,  $R_c$ can be easily controlled and be tailored to different NI coils for different applications. This Ebonol® C oxidation process is also economical and easy to be scaled up to have a reel-toreel surface treatment process which allows industry to manufacture REBCO tapes with customized  $R_c$  at a low cost. Figure 9 summarizes all 77 K  $R_c$  versus load cycle curves presented previously. This covers large variety of  $R_c$ from different surface conditions. Among them, the most controllable cases are the Ebonol® C treated surfaces.

Finally, it should be noted that our  $R_c$  measurements are on small stamp samples. Although our technique has the advantages of being versatile and able to perform fast measurements under a large number of load cycles, as well as the ability to reveal the underlying physics of the contact resistance, the results of these experiments need to be confirmed with a coil test such as those performed in [1]. This is because, for the applications in magnet coils, the contact pressure varies from turn to turn, so the effect of the load cycling cannot be predicted precisely. An  $R_c$  test by a small coil would bring us one step closer to the real situation in a magnet coil. For these reasons, we are developing a reel-to-reel Ebonol® oxidation system, so we can oxidize the surface of long length of REBCO tapes in order to make small coils for  $R_c$  tests.

#### 4. Conclusions

The contact resistance  $R_c$  between two REBCO tapes was measured at 77 K and 4.2 K under mechanical load between 2.5 and 25 MPa up to 30 000 cycles. We found that  $R_c$  reaches a maximum at 10-20 cycles and then decreases. After about 1000 cycles,  $R_c$  decreased to about one tenth of its initial value. After 20 000 load cycles, thermal cycling was performed by warming the sample from 4.2 K to 273 K under 2.5 MPa pressure and cooling down to 4.2 K again. This increased  $R_c$  slightly which was still much lower than its value before load cycling. Both electroplating of Ni, Cr, and Ni-P directly on REBCO, and using a Cu or Ni plated SS cowind tape were able to control  $R_c$  to a certain degree. A SS interlayer increases  $R_c$  by about three orders of magnitude; while the Cu plated SS interlayer only increases  $R_c$  by one order of magnitude. Controlled REBCO surface oxidation by an Ebonol<sup>®</sup> C solution results in an easily controlled  $R_c$  that covers a wide range by varying oxidation time. This oxidation technique can be easily scaled up to meet the needs of large NI magnet coils.

#### Acknowledgments

We thank Mr Kevin Jiang for his contribution to Ni and Ni–P plating, and Dr Chris Ray for providing Ebonol® C. We thank Dr Rongmei Niu for her assistance in experiments. Mr Kevin Jiang is supported through the NHMFL summer intern program. The NHMFL is supported by NSF through NSF-DMR-1157490 and the State of Florida.

#### **ORCID** iDs

Jun Lu https://orcid.org/0000-0001-8521-489X

#### References

- [1] Hahn S et al 2011 IEEE Trans. Appl. Supercond. 21 1592-5
- [2] Yoon S et al 2016 Supercond. Sci. Technol. 29 04LT04
- [3] Kim K et al 2017 Supercond. Sci. Technol. 30 065008

- [4] Weijers H et al 2016 IEEE Trans. Appl. Supercond. 26 4300807
- [5] Hahn S et al 2014 IEEE Trans. Appl. Supercond. 24 4602705
- [6] Wang X et al IEEE Trans. Appl. Supercond. 25 4601805
- [7] Yanagisawa K et al IEEE Trans. Appl. Supercond. 26 4602304
- [8] Wang Y et al 2017 J. Appl. Phys. 122 053902
- [9] Markiewicz W D et al 2016 Supercond. Sci. Technol. 29
- 025001 [10] Katsumata K *et al* 2016 *IEEE Trans. Appl. Supercond.* **27** 4602005
- [11] Oki T et al 2016 IEEE Trans. Appl. Supercond. 26 4702905
- [12] Wang T et al 2015 IEEE Trans. Appl. Supercond. 25 4603409
- [13] Kwanglok K et al 2017 Supercond. Sci. Technol. 30 075001

- [14] Wang Y et al 2017 IEEE Trans. Appl. Supercond. 27 3700105
- [15] Sohn M-H et al 2016 IEEE Trans. Appl. Supercond. 26 4701304
- [16] Chan W K et al 2017 Supercond. Sci. Technol. 30 074007
- [17] Lu J et al 2017 Supercond. Sci. Technol. 30 045005
- [18] Mayer W R 1944 US Patent 2460896
- [19] Kim J et al 2016 IEEE Trans. Appl. Supercond. 26 4601906
- [20] Sudagar J et al 2013 J. Alloys Compounds 571 183
- [21] Noguchi S et al 2017 IEEE Trans. Appl. Supercond. 27 7700505
- [22] Gupta R et al 2016 IEEE Trans. Appl. Supercond. 26 4100705
- [23] Lécrevisse T and Iwasa Y 2016 IEEE Trans. Appl. Supercond. 26 4700405