

# Superconductivity —

## Part 3: Critical current measurement — DC critical current of Ag- and/or Ag alloy-sheathed Bi-2212 and Bi-2223 oxide superconductors

The European Standard EN 61788-3:2006 has the status of a British Standard

ICS 17.220; 29.050

## National foreword

This British Standard was published by BSI. It is the UK implementation of EN 61788-3:2006. It is identical with IEC 61788-3:2006. It supersedes BS EN 61788-3:2001 which is withdrawn.

The UK participation in its preparation was entrusted to Technical Committee L/-/90, Super Conductivity.

A list of organizations represented on L/-/90 can be obtained on request to its secretary.

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This British Standard was published under the authority of the Standards Policy and Strategy Committee on 29 September 2006

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ISBN 0 580 49146 3

### Amendments issued since publication

Amd. No.	Date	Comments

English version

**Superconductivity**  
**Part 3: Critical current measurement -**  
**DC critical current of Ag- and/or Ag alloy-sheathed**  
**Bi-2212 and Bi-2223 oxide superconductors**  
(IEC 61788-3:2006)

Supraconductivité  
Partie 3: Mesure du courant critique -  
Courant critique continu des oxydes  
supraconducteurs Bi-2212 et Bi-2223  
avec gaine Ag et/ou en alliage d'Ag  
(CEI 61788-3:2006)

Supraleitfähigkeit  
Teil 3: Messen des kritischen Stromes -  
Kritischer Strom (Gleichstrom) von  
Ag- und/oder Ag-Legierung ummantelten  
oxidischen Bi-2212 und  
Bi-2223-Supraleitern  
(IEC 61788-3:2006)

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**CENELEC**

European Committee for Electrotechnical Standardization  
Comité Européen de Normalisation Electrotechnique  
Europäisches Komitee für Elektrotechnische Normung

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## Foreword

The text of document 90/184/FDIS, future edition 2 of IEC 61788-3, prepared by IEC TC 90, Superconductivity, was submitted to the IEC-CENELEC parallel vote and was approved by CENELEC as EN 61788-3 on 2006-06-01.

This European Standard supersedes EN 61788-3:2001.

Modifications made to EN 61788-3:2001 mostly involve wording and essentially include no technical changes.

Examples of technical changes introduced include the voltage lead diameter being smaller than 0,21 mm and the mode of expression for magnetic field accuracy being  $\pm 1\%$  and  $\pm 0,02\text{ T}$  instead of  $1\%$ . The expression for magnetic field precision has been changed in the same way.

The following dates were fixed:

- latest date by which the EN has to be implemented at national level by publication of an identical national standard or by endorsement (dop) 2007-03-01
- latest date by which the national standards conflicting with the EN have to be withdrawn (dow) 2009-06-01

Annex ZA has been added by CENELEC.

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## Endorsement notice

The text of the International Standard IEC 61788-3:2006 was approved by CENELEC as a European Standard without any modification.

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## INTRODUCTION

In 1986 J.G. Bednorz and K.A. Mueller discovered that some Perovskite type Cu-containing oxides show superconductivity at temperatures far above those which metallic superconductors have shown. Since then, extensive R & D work on high-temperature oxide superconductors has been and is being made worldwide, and its application to high-field magnet machines, low-loss power transmission, electronics and many other technologies is in progress [1].<sup>1)</sup>

Fabrication technology is essential to the application of high-temperature oxide superconductors. Among high-temperature oxide superconductors developed so far, BiSrCaCu oxide (Bi-2212 and Bi-2223) superconductors have been the most successful at being fabricated into wires and tapes of practical length and superconducting properties. These conductors can be wound into a magnet to generate a magnetic field of several tesla [2]. It has also been shown that Bi-2212 and Bi-2223 conductors can substantially raise the limit of magnetic field generation by a superconducting magnet [3].

In summer 1993, VAMAS-TWA16 started working on the test methods of critical currents in Bi-oxide superconductors. In September 1997, the TWA16 worked out a guideline (VAMAS guideline) on the critical current measurement method for Ag-sheathed Bi-2212 and Bi-2223 oxide superconductors. This pre-standardization work of VAMAS was taken as the base for the IEC standard, described in the present document, on the dc critical current test method of Ag-sheathed Bi-2212 and Bi-2223 oxide superconductors.

The test method covered in this International Standard is intended to give an appropriate and agreeable technical base to those engineers working in the field of superconductivity technology.

The critical current of composite superconductors like Ag-sheathed Bi-oxide superconductors depends on many variables. These variables need to be considered in both the testing and the application of these materials. Test conditions such as magnetic field, temperature and relative orientation of the specimen and magnetic field are determined by the particular application. The test configuration may be determined by the particular conductor through certain tolerances. The specific critical current criterion may be determined by the particular application. It may be appropriate to measure a number of test specimens if there are irregularities in testing.

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<sup>1)</sup> The numbers in brackets refer to the bibliography.

## SUPERCONDUCTIVITY –

### Part 3: Critical current measurement – DC critical current of Ag- and/or Ag alloy-sheathed Bi-2212 and Bi-2223 oxide superconductors

#### 1 Scope

This part of IEC 61788 covers a test method for the determination of the dc critical current of short and straight Ag- and/or Ag alloy-sheathed Bi-2212 and Bi-2223 oxide superconductors that have a monolithic structure and a shape of round wire or flat or square tape containing mono- or multicores of oxides.

This method is intended for use with superconductors that have critical currents less than 500 A and  $n$ -values larger than 5. The test is carried out with and without an applying external magnetic field. For all tests in a magnetic field, the magnetic field is perpendicular to the length of the specimen. In the test of a tape specimen in a magnetic field, the magnetic field is parallel or perpendicular to the wider tape surface (or one surface if square). The test specimen is immersed either in a liquid helium bath or a liquid nitrogen bath during testing. Deviations from this test method that are allowed for routine tests and other specific restrictions are given in this standard.

#### 2 Normative reference

The following referenced document is indispensable for the application of this document. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

IEC 60050-815:2000, *International Electrotechnical Vocabulary (IEV) – Part 815: Superconductivity*

#### 3 Terms and definitions

For the purposes of this document, the terms and definitions given in IEC 60050-815, several of which have been repeated here for convenience, and the following apply.

##### 3.1 critical current

$I_c$   
maximum direct current that can be regarded as flowing without resistance

NOTE  $I_c$  is a function of magnetic field strength and temperature.

[IEV 815-03-01]

##### 3.2 critical current criterion

$I_c$  criterion  
criterion to determine the critical current,  $I_c$ , based on the electric field strength,  $E$  or the resistivity,  $\rho$

NOTE 1  $E = 10 \mu\text{V/m}$  or  $E = 100 \mu\text{V/m}$  is often used as the electric field strength criterion, and  $\rho = 10^{-13} \Omega \cdot \text{m}$  or  $\rho = 10^{-14} \Omega \cdot \text{m}$  is often used as the resistivity criterion.

NOTE 2 For short high temperature oxide superconductor specimens, less sensitive criteria than those shown in Note 1 are sometimes used.

[IEV 815-03-02, modified]

### 3.3

**n-value** (of a superconductor)

exponent obtained in a specific range of electric field strength or resistivity when the voltage/current  $U$ - $I$  curve is approximated by the equation  $U \propto I^n$

NOTE In the case for high temperature oxide superconductors, the equation  $U \propto I^n$  does not hold in a wide range of  $U$ .

[IEV 815-03-10, modified]

### 3.4

**quench**

uncontrollable and irreversible transition of a superconductor or a superconducting device from the superconducting state to the normal conducting state

NOTE A term usually applied to superconducting magnets.

[IEV 815-03-11]

### 3.5

**Lorentz force** (on fluxons)

force applied to fluxons by a current

NOTE 1 The force per unit volume is given by  $\mathbf{J} \times \mathbf{B}$ , where  $\mathbf{J}$  is a current density, and  $\mathbf{B}$  is a magnetic flux density.

NOTE 2 "Lorentz force" is defined in IEC 60050-111-20.

[IEV 815-03-16]

### 3.6

**current transfer** (of composite superconductor)

phenomenon that a dc current transfers spatially from filament to filament in a composite superconductor, resulting in a voltage generation along the conductor

NOTE In the  $I_c$  measurement, this phenomenon appears typically near the current contacts where the injected current flows along the conductor from periphery to inside until uniform distribution among filaments is accomplished.

### 3.7

**constant sweep rate method**

a  $U$ - $I$  data acquisition method where a current is swept at a constant rate from zero to a current above  $I_c$  while continuously or frequently and periodically acquiring  $U$ - $I$  data

### 3.8

**ramp-and-hold method**

a  $U$ - $I$  data acquisition method where a current is ramped to a number of appropriately distributed points along the  $U$ - $I$  curve and held constant at each one of these points while acquiring a number of voltages and current readings

### 3.9

**Bi-2212 and Bi-2223 oxide superconductors**

oxide superconductors with layered structure containing  $\text{CuO}_2$  sheets and chemical formulae,  $\text{Bi}_2\text{Sr}_2\text{CaCu}_2\text{O}_x$  ( $x \sim 8$ ) and  $(\text{Bi,Pb})_2\text{Sr}_2\text{Ca}_2\text{Cu}_3\text{O}_x$  ( $x \sim 10$ ), respectively



## 4 Principle

The critical current of a composite superconductor is determined from a voltage ( $U$ ) - current ( $I$ ) characteristic measured at a certain value of a static applied magnetic field strength (magnetic field) and at a specified temperature in a liquid cryogen bath at a constant pressure. To get a  $U$ - $I$  characteristic, a direct current is applied to the superconductor specimen and the voltage generated along a section of the specimen is measured. The current is increased from zero and the  $U$ - $I$  characteristic generated is recorded. The critical current is determined as the current at which a specific electric field strength criterion (electric field criterion) ( $E_c$ ) or resistivity criterion ( $\rho_c$ ) is reached. For either  $E_c$  or  $\rho_c$ , there is a corresponding voltage criterion ( $U_c$ ) for a specified voltage tap separation.

## 5 Requirements

The target precision of this method is a coefficient of variation (standard deviation divided by the average of the critical current determinations) that is less than 5 % for the measurement at 0 T and near 4,2 K or 77 K.

The use of a common current transfer correction is excluded from this test method. Furthermore, if a current transfer signature is pronounced in the measurement, then the measurement shall be considered invalid.

It is the responsibility of the user of this standard to consult and establish appropriate safety and health practices and to determine the applicability of regulatory limitations prior to use. Specific precautionary statements are given below.

Hazards exist in this type of measurement. Very large direct currents with very low voltages do not necessarily provide a direct personal hazard, but accidental shorting of the leads with another conductor, such as tools or transfer lines, can release significant amounts of energy and cause arcs or burns. It is imperative to isolate and protect current leads from shorting. Also the energy stored in the superconducting magnets commonly used for the background magnetic field can cause similar large current and/or voltage pulses or deposit large amounts of thermal energy in the cryogenic systems, causing rapid boil-off or even explosive conditions. Under rapid boil-off conditions, cryogens can create oxygen-deficient conditions in the immediate area and additional ventilation may be necessary. The use of cryogenic liquids is essential to cool the superconductors to allow the transition into the superconducting state. Direct contact of skin with cold liquid transfer lines, storage Dewars or apparatus components can cause immediate freezing, as can direct contact with a spilled cryogen. If improperly used, liquid helium storage Dewars can freeze air or water in pressure vent lines and cause the Dewar to over-pressurize and fail despite the common safety devices. It is imperative that safety precautions for handling cryogenic liquids be observed.

## 6 Apparatus

### 6.1 Measurement holder material

The measurement holder shall be made from an insulating material or from a conductive non-ferromagnetic material that is either covered or not covered with an insulating layer.

The critical current may inevitably depend on the measurement holder material due to the strain induced by the differential thermal contraction between the specimen and the measurement holder.

The total strain induced in the specimen at the measuring temperature shall be minimized to be within  $\pm 0,1$  %. If there is an excess strain due to the differential thermal contraction of the specimen and the holder, the critical current shall be noted to be determined under an excess strain state by identification of the holder material.

Suitable measurement holder materials are recommended in A.3.1. Any one of these may be used.

When a conductive material is used without an insulating layer, the leakage current through the holder shall be less than 1 % of the total current when the specimen current is at  $I_c$  (see 9.5).

## 6.2 Measurement holder construction

The holder shall have a flat surface on which a straight specimen can be placed.

The current contact shall be rigidly fastened to the measurement holder to avoid stress concentration in the region of transition between the holder and the current contact. It is important to have no difference in level between the mounting surfaces of the current contacts and the specimen holder.

## 6.3 Measurement set up

The apparatus to measure the  $U-I$  characteristic of the superconductor specimen consists of a specimen probe, a test cryostat, a magnet system and a  $U-I$  measurement system.

The specimen probe, which consists of a specimen, a measurement holder and a specimen support structure, is inserted in the test cryostat filled with liquid cryogen. In some cases the cryostat contains a superconducting solenoid magnet and its support structure to apply a magnetic field to the specimen. The  $U-I$  measurement system consists of a dc current source, a recorder and necessary preamplifiers, filters or voltmeters, or a combination thereof. A computer assisted data acquisition system is also allowed.

## 7 Specimen preparation

### 7.1 Reaction heat treatment

Reaction heat treatment shall be carried out according to the manufacturer's specification which includes reaction temperature, period and atmosphere, oxygen partial pressure, specimen warming and cooling rates, specimen protection method against mechanical strain, examination of deformation and surface condition of specimen and error limits which shall not be exceeded. Temperature variations within the furnace shall be controlled such as not to exceed those limits.

Reaction heat treatment can be skipped when it has already been carried out by the manufacturer.

### 7.2 Specimen mounting for measurement

After the reaction heat treatment, the ends of the specimen shall be trimmed to suit the measurement holder.

When using resistivity criteria for the critical current determination, the total cross-sectional area  $S$  of the specimen shall be determined to an accuracy of 5 %.

The specimen shall be mounted to the flat surface of the holder and both ends shall be soldered to the current contact blocks (see Clause A.5 for solder material).

For the test in magnetic fields, a low-temperature adhesive (such as epoxy) shall be used to bond the specimen to the measurement holder to reduce specimen motion against the Lorentz force.

For a tape specimen, the bond shall be strong enough to keep the specimen in place against the Lorentz force, in the case where the applied magnetic field is perpendicular to the specimen surface.

The length of a specimen to be measured shall be defined as follows:

$$L_1 = 2 \times L_2 + L + 2 \times L_3 \geq 5 \times W \quad (1)$$

$$L_2, L, L_3 \geq W \quad (2)$$

where

- $L$  is the distance between the voltage taps;
- $L_1$  is the length of a specimen to be measured;
- $L_2$  is the length of the soldered part of the current contact;
- $L_3$  is the shortest distance from a current contact to a voltage tap;
- $W$  is the width or diameter of a specimen to be measured.

For a specimen with a large current-carrying capacity, narrow tape, or round wire,  $L_2$  shall be larger.  $L$  shall be larger for a measurement that needs high sensitivity and  $L_3$  shall be larger when current transfer voltage cannot be neglected.

In the case of a specimen that has a stainless steel or other high-resistivity material backing or jacket,  $L_2$  shall be longer than  $3W$ .

In the case of the wire specimen the angle between the specimen axis and the magnetic field shall be  $(90 \pm 9)^\circ$ . This angle shall be determined with an accuracy of  $\pm 2^\circ$ .

In the case of tape specimens, there are two options in addition to the requirement that the angle between the longitudinal specimen axis and the magnetic field shall be  $(90 \pm 9)^\circ$ . In one option, the magnetic field shall be perpendicular to the specimen surface, the angle deviation being within  $\pm 7^\circ$ . In the second option, the magnetic field shall be parallel to the specimen surface, the angle deviation being within  $\pm 3^\circ$ .

The voltage taps shall be placed in the central part along both the specimen length and the specimen width.

All soldering shall be conducted as quickly as possible so as not to cause thermal damage to the specimen. Voltage leads with a diameter less than 0,21 mm shall be used and twisted together before soldering.

The distance between the voltage taps,  $L$ , shall be measured to an accuracy of 5 %.

## 8 Measurement procedure

The specimen shall be immersed in cryogen for the data acquisition phase. The specimen may be cooled slowly in cryogen vapour and then inserted into the cryogen bath, or inserted slowly into the cryogen bath, or, in the case of cooling to the 4,2 K range, first slowly immersed in liquid nitrogen and then liquid helium. The specimen shall be cooled from room temperature to liquid helium (or liquid nitrogen) temperature over a time period of at least 5 min.

When measuring at more than one temperature or magnetic field angle, between each measuring temperature and/or each magnetic-field angle, the specimen shall be cooled in zero field, from a temperature above the critical temperature down to the measuring temperature, and then the field angle with respect to the conductor cross-section shall be fixed while the field is still zero. This procedural step can only be omitted if one of the following two conditions is met:

only zero applied field measurements will be made with monotonically increasing temperatures or the specimen has a demonstrated magnetic hysteresis of less than 2 % at the magnetic fields where  $I_c$  is to be reported (see Annex B).

The temperature of the cryogen bath shall be measured during each determination of  $I_c$ .

Unless a quench protection circuit or resistive shunt is used to protect the specimen from damage, the specimen current shall be kept low enough so that the specimen does not enter the normal state.

When using the constant sweep rate method, the time for the ramp from zero current to  $I_c$  shall be more than 30 s.

When using the ramp-and-hold method, the current sweep rate between current set points shall be lower than the equivalent of ramping from zero current to  $I_c$  in 3 s. Data acquisition at each set point shall be started as soon as the flow/creep voltage generated by the current ramp can be disregarded. The current drift during each current set point shall be less than 1 % of  $I_c$ .

The relation between the magnetic field and the magnet current shall be measured beforehand. The magnet current shall be measured before each determination of  $I_c$ .

If the magnetic field is parallel to the surface of the measurement holder, the relative direction of the current to the applied magnetic field shall result in the Lorentz force which pushes the specimen against the surface of the measurement holder. In the case of the applied magnetic field perpendicular to the measurement holder surface, either direction of the current relative to the field is possible, with the condition that the specimen is rigidly mounted to the measurement holder with appropriate adhesive.

Record the  $U$ - $I$  characteristic with increasing current and at monotonically increasing magnetic fields (see Annex B).

The baseline voltage of the  $U$ - $I$  characteristic shall be taken as the recorded voltage at zero current for the ramp-and-hold method or the average voltage at approximately 0,1  $I_c$  for the constant sweep rate method.

## 9 Precision and accuracy of the test method

### 9.1 Critical current

The current source shall provide a dc current having a maximum periodic and random deviation of less than  $\pm 2$  % at  $I_c$ , within the bandwidth 10 Hz to 10 MHz.

A four-terminal standard resistor, with an accuracy of at least 0,5 %, shall be used to determine the specimen current.

A recorder and the necessary preamplifiers, filters or voltmeters, or a combination thereof, shall be used to record the  $U$ - $I$  characteristic. The record of the  $U$ - $I$  characteristic shall allow the determination of  $U_c$  to a precision of 10 % and the corresponding current to an accuracy of 1 % and with a precision of 1 %.

### 9.2 Temperature

A cryostat shall provide the necessary environment for measuring  $I_c$  and the specimen shall be measured while immersed in liquid helium or liquid nitrogen. The liquid temperature shall be reported to an accuracy of  $\pm 0,1$  K, measured by means of a pressure sensor or an appropriate temperature sensor.

The difference between the specimen temperature and the bath temperature shall be minimized.

To convert the pressure observed in the cryostat into a temperature value, the phase diagram of helium or nitrogen shall be used. The pressure measurement shall be accurate enough to obtain the required accuracy of the temperature measurement.

### 9.3 Magnetic field

A magnet system shall provide the magnetic field to an accuracy better than the larger of  $\pm 1\%$  and  $\pm 0,02$  T and a precision better than the larger of  $\pm 0,5\%$  and  $\pm 0,02$  T.

When testing without a magnet, the background magnetic field shall be measured to an accuracy of  $\pm 0,0002$  T and a precision of  $\pm 0,0001$  T.

The magnetic field shall have a uniformity better than the larger of  $0,5\%$  and  $0,02$  T over the length of the specimen between the voltage contacts.

The maximum periodic and random deviation of the magnetic field shall be less than the larger of  $\pm 1\%$  and  $\pm 0,02$  T.

For critical current measurements at zero or very low magnetic field, the residual magnetic field in a superconducting magnet shall be minimized.

### 9.4 Specimen and holder support structure

The support structure shall provide adequate support for the specimen and the orientation of the specimen with respect to the magnetic field. The specimen support is adequate if it allows additional determinations of critical current with a precision of  $2\%$ .

### 9.5 Specimen protection

If a resistive shunt or a quench protection circuit is used in parallel with the specimen, then the current through the shunt or the circuit shall be less than  $1\%$  of the total current at  $I_c$ .

## 10 Calculation of results

### 10.1 Critical current criteria

The critical current  $I_c$  shall be determined by using an electric field criterion  $E_c$  or a resistivity criterion  $\rho_c$  where the total cross-sectional area  $S$  of the composite superconductor is preferred for the estimation of the resistivity (see Figures 1 and 2).

In the case of the electric field criterion, two values of  $I_c$  shall be determined at criteria of  $100\ \mu\text{V/m}$  and  $500\ \mu\text{V/m}$ . In the other case, two values of  $I_c$  shall be determined at the resistivity criteria of  $2 \times 10^{-13}\ \Omega\text{m}$  and  $10^{-12}\ \Omega\text{m}$ .

When it is difficult to measure the  $I_c$  properly at a criterion of  $500\ \mu\text{V/m}$ , an  $E_c$  criterion of less than  $500\ \mu\text{V/m}$  shall be substituted. Otherwise, measurements using the resistivity criterion are recommended.

The  $I_c$  shall be determined as the current corresponding to the point on the  $U-I$  curve where the voltage is  $U_c$  measured relative to the baseline voltage (see Figures 1 and 2):

$$U_c = L E_c \quad (3)$$

where

$U_c$  is the voltage criterion in microvolts ( $\mu\text{V}$ );

$L$  is the voltage tap separation in meters (m);

$E_c$  is the electric field criterion in microvolts/meter ( $\mu\text{V}/\text{m}$ );

or, when using a resistivity criterion:

$$U_c = I_c \rho_c L/S \quad (4)$$

where  $U_c$ ,  $I_c$  and  $\rho_c$  are the corresponding voltage, current and resistivity to the intersecting point of a straight line with the  $U$ - $I$  curve as shown in Figure 1, and  $S$  is the total cross-sectional area in square meters.

A straight line shall be drawn from the baseline voltage to the average voltage near  $0,5 I_c$  (see Figures 1 and 2). A finite slope of this line may be due to current transfer and/or local sample damage. A valid determination of  $I_c$  requires that the slope of the line be less than  $0,3 U_c/I_c$ , where  $U_c$  and  $I_c$  are determined at a criterion of  $100 \mu\text{V}/\text{m}$  or  $2 \times 10^{-13} \Omega\text{m}$ .

## 10.2 $n$ -value (optional)

The  $n$ -value shall be calculated as the slope of the plot of  $\log U$  versus  $\log I$  in the region where the  $I_c$  is determined.

The range of the criteria used to determine  $n$  shall be reported.

## 11 Test report

### 11.1 Identification of test specimen

The test specimen shall be identified, if possible, by the following:

- a) name of the manufacturer of the specimen;
- b) classification and/or symbol;
- c) lot number;
- d) raw materials and their chemical composition;
- e) shape and area of the cross-section of the wire, number of cores, diameter of cores, volume fractions of cores to Ag and/or Ag alloy sheath, and other components in the wire;
- f) manufacturing process technique.

### 11.2 Report of $I_c$ values

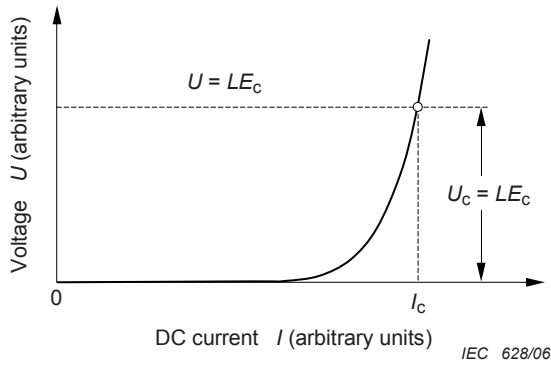
The  $I_c$  values, along with their corresponding criteria, and  $n$ -values (optional) shall be reported.

### 11.3 Report of test conditions

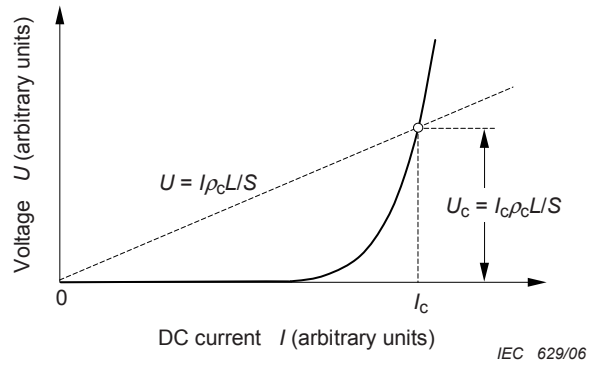
The following test conditions shall be reported:

- a) test magnetic field strength, and orientation, uniformity and accuracy of field;
- b) test temperature and accuracy of temperature;
- c) length between voltage taps and total specimen length;
- d) the shortest distance from a current contact to a voltage tap;
- e) soldered length of the current contacts;
- f) the specimen bonding method, including identification of the bonding material;

- g) Lorentz force direction;
- h) sample history with magnetic field sweep;
- i) sample history with temperature variation;
- j) sample history with current sweep and hold.



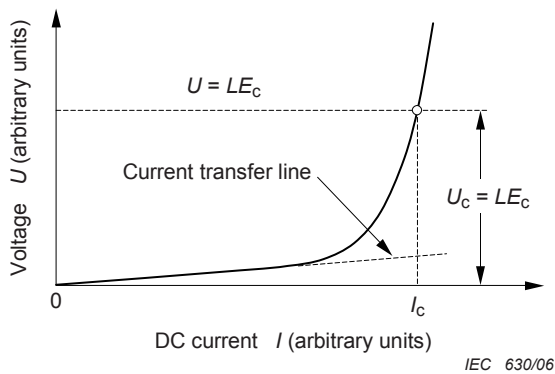
(a)



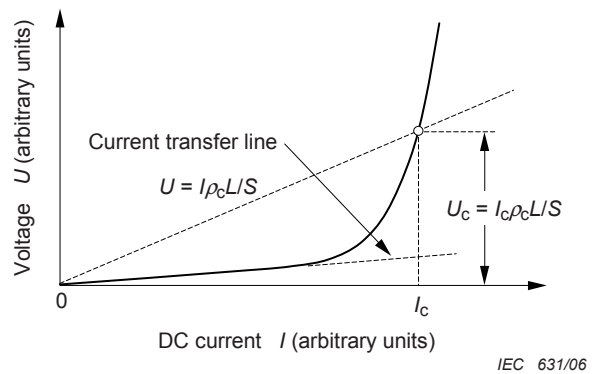
(b)

NOTE The application of the (a) electric field and (b) resistivity criteria to determine the critical current is shown above.

Figure 1 – Intrinsic  $U-I$  characteristic



(a)



(b)

NOTE The application of the (a) electric field and (b) resistivity criteria to determine the critical current on a  $U-I$  characteristic with a current transfer component exhibited as a linear region at low current is shown above.

Figure 2 –  $U-I$  characteristic with a current transfer component



## Annex A (informative)

### Additional information relating to Clauses 1 to 10

#### A.1 Scope

There are a large number of variables that have a significant effect on the measured value of critical current in Ag- and/or Ag alloy-sheathed Bi-2212 and Bi-2223 oxide superconductor wires and tapes. However, significant portions of the test method covered in this standard are common or similar to those for Nb<sub>3</sub>Sn composite superconductors (IEC 61788-2). Thus, only part of these variables will be addressed in this informative annex. For those variables that are not mentioned here, refer to IEC 61788-2.

Special features found for oxide superconductors may be classified into two groups. The first group is specific to oxide composite superconductors, including mechanical fragility, electro-magnetic weak links, cryogen gas bubble formation, aging degradation, magnetic flux flow and creep, large anisotropy, hysteresis in critical current with magnetic field sweep, etc. The second group is due to the short length of the specimen used in the standard. A critical current measurement on such a specimen may easily pick up different voltage signals due to thermal electromotive force, inductive voltage, thermal noise, current redistribution, specimen motion relative to the holder, etc. Current transfer voltages may be present due to the short distance from a current contact to a voltage tap. Short specimen length may reduce mechanical tolerance against the Lorentz force, for example, by promoting the formation of cryogen gas bubbles within the composite.

Superconductors with critical currents above 500 A could be measured with the present standard with an anticipated reduction both in accuracy and precision and a more significant self-field effect.

The reason for the restrictions in this test method is to obtain the necessary precision in the final definitive phase of long conductor qualification.

This standard assumes that measurements are made either in liquid helium or liquid nitrogen. However, it is generally accepted that, if these measurements can be made in liquid helium, then they can be made in other cryogens, such as liquid neon and liquid hydrogen, because the heat of vaporization of liquid helium is low compared to other cryogens. Thus, this standard should extend to measurements conducted in other cryogens.

Cryogens (including nitrogen, helium, neon, and hydrogen) are used at temperatures near boiling point for the normal atmospheric pressure of the test site. The use of cryogens at temperatures other than near boiling point, or measurements in a gas or a vacuum are not covered by the scope of this standard.

#### A.2 Requirements

The minimum total length of the tape specimen is five times the tape width ( $W$ ), which represents the sum of the following:

- the soldered length of current contacts ( $2 W$ );
- the shortest distance between current and voltage contacts ( $2 W$ );
- the minimum voltage tap separation ( $1 W$ ).



The target precision of the method described in this standard is defined by the results of an interlaboratory comparison. Results from the previous VAMAS interlaboratory comparisons (the regional and the general intercomparisons) were used in this test method to formulate the tolerances of the many variables that affect the precision of critical current measurements. The target precision, for an interlaboratory comparison, is a coefficient of variation (standard deviation divided by the average of critical current determination) that is less than 5 % for the measurement at 0 T and near 4,2 K or 77 K.

It is expected that the specimen mounting and the specimen cooling procedures in this test method may be one of the most significant contributors to the overall uncertainty of the critical current measurement.

In the case of routine tests where it is impractical to adhere to these specific restrictions, this standard can be used as a set of general guidelines with an anticipated reduction in precision.

The test method for determining the  $I_c$  values of Ag- and/or Ag alloy-sheathed Bi-oxide superconducting wires and tapes excluded from the present test method may be addressed in future documents.

In the case of measurements made in liquid hydrogen, additional safety precautions may be required since hydrogen leakage into air can result in accidental gas explosion.

### A.3 Apparatus

#### A.3.1 Measurement holder material

In this method, the specimen strain is controlled to a minimum (less than 0,1 %). A 0,1 % thermal contraction may result in no appreciable  $I_c$  deviation at 0 T and near 4,2 K or 77 K. One significant source of strain is the mismatch in thermal contraction rates between the measurement holder and the specimen when cooled to liquid helium or nitrogen temperature.

Based on the typical thermal contractions shown in table A.1, the following materials are suggested for the measurement holder material. For alternate holder materials, a carefully prepared qualification study should precede the routine tests.

Recommended holder material:

- fibreglass epoxy composite, with the specimen lying in the plane of the fabric wrap;
- ceramic dispersed epoxy;
- Ag/Ag alloy.

The leakage current through a conductive holder without an insulating layer can be estimated by making measurements under test conditions with and without a specimen on the holder. The measurement of the voltage drop from current contact to current contact without a specimen and under test conditions can be used to estimate the resistance of the leakage path including contact resistance. Then, measurement of the voltage drop from current contact to current contact with a specimen and under test conditions can be used to estimate leakage current.

#### A.3.2 Measurement holder construction

An example of a measurement holder is shown in Figure A.1. Typically, the current contacts are made from copper blocks, and the thickness of the contact should be determined so that there will be no difference in level between the holder surface and the contact surfaces; the contact blocks need to be rigidly affixed to the holder.

#### A.4 Specimen preparation

Extreme care must be taken in trimming and mounting the specimen after the reaction so as not to damage the specimen.

Specimen motion can damage the specimen and result in a premature quench (irreversible thermal runaway), voltage noise and ultimately a reduction in the repeatability of critical current.

It is recommended that a low-temperature adhesive, such as epoxy, be used to secure the specimen to the holder.

A rough and clean surface on the measurement holder and a clean surface on the specimen are needed for strong specimen bonding.

The low-temperature adhesive should withstand the heat from soldering.

The low-temperature adhesive should be a thin layer between the specimen and the holder surface.

After the low-temperature adhesive is applied, the specimen should be pressed to the holder with a pressure that does not damage the specimen.

The distance between the voltage taps is defined as the smallest distance between the soldered leads, irrespective of the sizes of the solder spots.

#### A.5 Measurement procedure

The specimen support structure is needed to hold the specimen in the centre of the background magnet in a liquid helium or nitrogen cryostat and to support current and voltage leads between room and measurement temperatures.

To reduce thermoelectric voltages on the specimen voltage leads, copper voltage leads are used which are continuous from the cryogen bath to room temperature where an isothermal environment for all room temperature joints or connections is provided. It should be noted that the joints or connections immersed in cryogen are isothermal.

As soldering material, Indium, Indium alloys or Bismuth alloys with a low melting temperature are preferable, although usual Pb-60 % Sn and Pb-70 % Sn are also applicable.

The specimen cooling rate may affect the measured critical current. The strength of the bond between the specimen and its holder changes during the cooling process when differential thermal contraction between the specimen and the holder is also occurring. This may result in disabling the low-temperature adhesive.

Oxygen impurities can cause boiling temperature variation of liquid nitrogen in the test cryostat if it is not tightly sealed. Liquid nitrogen stored in a non-pressurized Dewar for several weeks will condense enough oxygen impurities to shift the boiling point.

If the system noise is significant compared to the prescribed value of voltage, i.e.  $U_c$ , it is desirable to increase the time for the ramp from zero current to  $I_c$  to more than 150 s. In this case, care should be taken to increase the heat capacity and/or cooling surface of the current contacts enough to suppress the influence of heat generation due to the longer time required for the measurement. It should be noted that the ramp-and-hold method allows for averaging data which can be appropriately distributed along the  $U-I$  characteristic.

Ramping the specimen current can induce a positive or negative voltage on the voltage taps with time. This source of interfering voltage during the ramp can be identified by its proportional dependence on ramp rate. If this voltage is significant compared to  $U_c$ , then decrease the ramp rate, decrease the area of the loop formed by the voltage taps and the specimen between them, or else use the ramp-and-hold method.

Faster current ramp rates can be used for the ramp-and-hold method if the measurement system proves to yield consistent results with the specified ramp rate equivalent to ramping from zero to  $I_c$  in 3 s. It is possible to obtain consistent results with ramp rates as high as 500 A/s on a conductor with critical current from 10 A to 200 A.

Specimen motion can induce noise spikes. The specimen holder should be tightly mounted to the support structure, while the support structure should be held rigid against the magnetic force.

The baseline voltage may include thermoelectric, off-set, ground-loop and common-mode voltages. It is assumed that these voltages remain relatively constant for the time it takes to record each  $U-I$  characteristic. Small changes in thermoelectric and off-set voltages can be approximately removed by measuring the baseline voltage before and after the  $U-I$  curve measurement and assuming a linear change with time. If the change in the baseline voltage is significant compared to  $U_c$ , then corrective action to the experimental configuration should be taken.

For critical current measurements at zero or very low magnetic fields, the residual magnetic field in a superconducting magnet shall be completely minimized or the use of a superconducting magnet shall be avoided. This is especially important for the measurements near liquid nitrogen temperature.

## A.6 Precision and accuracy of the test method

An optional method for partially assessing the overall precision of a laboratory's critical current measurement system is to obtain and measure a superconductor simulator (Figure A.2) originally developed at the National Institute of Standards and Technology, Boulder, Colorado, USA [4].

## A.7 Calculation of results

### A.7.1 Critical current criteria

For some applications, the non-Ag (or non-Ag alloy) cross-sectional area is used in the resistivity criterion. It can be determined by using a graphical analysing method.

A larger separation between current and voltage connections may be necessary if a significant current transfer component exists relative to the criteria.

### A.7.2 $n$ -value

The superconductor's  $U-I$  characteristic curve near the  $I_c$  can usually be approximated by the empirical power-law equation:

$$U = U_0(I/I_0)^n \quad (\text{A.1})$$

where

$U$  is the specimen voltage in microvolts ( $\mu\text{V}$ );

$U_0$  is a reference voltage in microvolts ( $\mu\text{V}$ );

$I$  is the specimen current in amperes (A);

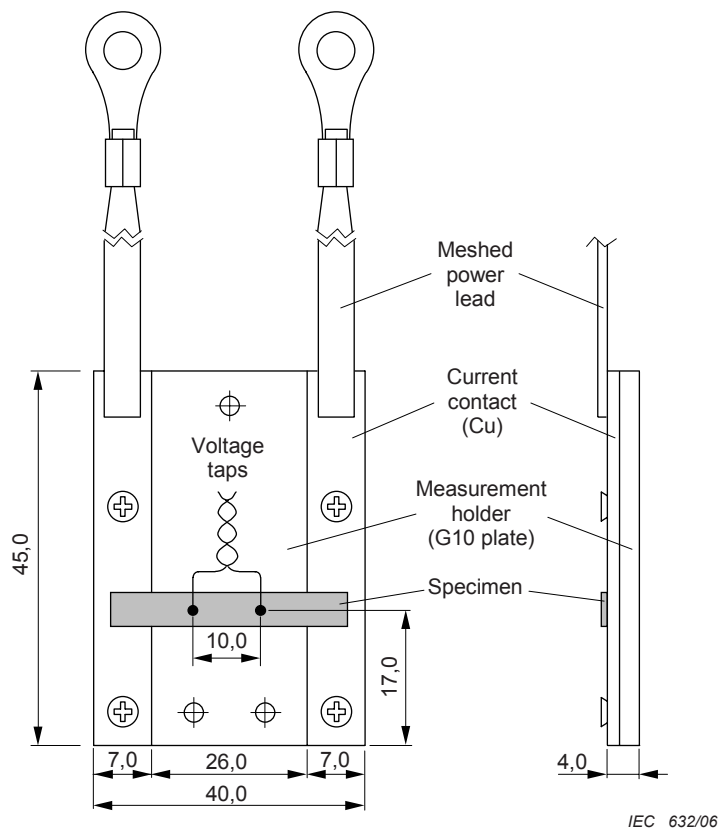
$I_0$  is a reference current in amperes (A).

The  $n$ -value (no units) reflects the general shape of the curve near the  $I_c$ .

**Table A.1 – Thermal expansion data of Bi-oxide superconductor and selected materials**

Material	Thermal expansion %							
	Temperature K							
	273	200	150	100	50	20	10	4
Bi-2223 Composite tape [5]	0	-0,09	-0,14	-0,17	-0,20	-0,20	-0,20	
1st cool-down								
2nd cool-down	0	-0,11	-0,18	-0,23	-0,27	-0,27	-0,27	
Silver [6]	0	-0,135	-0,221	-0,301	-0,360	-0,374	-0,375	-0,375
OFHC copper annealed [7]	0	-0,118	-0,18	-0,252	-0,288	-0,295	-0,295	
G10, warp [8]	0	-0,09	-0,13	-0,175	-0,205	-0,215	-0,220	-0,225
G10, normal [8]	0	-0,28	-0,428	-0,54	-0,62	-0,64	-0,65	-0,655
Stainless steel AISI SUS316 [7]	0	-0,111	-0,173	-0,23	-0,262	-0,265	-0,265	-0,265
Stainless steel AISI SUS304 [7]	0	-0,11	-0,172	-0,23	-0,261	-0,264	-0,264	-0,264

NOTE Thermal expansions are referred to zero at 273 K.



Dimensions in millimetres

Figure A.1 – Illustration of a measurement configuration for a short specimen of a few hundred A class conductors

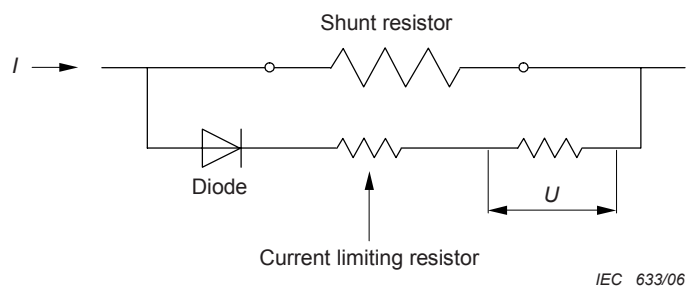


Figure A.2 – Illustration of superconductor simulator circuit

## Annex B (informative)

### Magnetic hysteresis of the critical current of high-temperature oxide superconductors

#### B.1 Magnetic hysteresis

The measured critical current ( $I_c$ ) of a high-temperature oxide superconductor can have a significant dependence on the history of the temperature, current, and applied magnetic-field strength and angle. As for magnetic-field strength,  $I_c$  at a given magnetic field of some multifilamentary Bi-2223 conductors can be as much as 100 % higher when they are measured with monotonically decreasing fields (from a higher field) compared to monotonically increasing fields. The  $I_c$  measured at zero magnetic field after a magnetic field sweep, can be lower than the initial  $I_c$  measured at zero field by as much as 40 %. The  $I_c$  hysteresis of some multifilamentary Bi-2212 conductors can be much less. It can be as low as 5 % to 10 % at low magnetic fields and as low as 2 % at high magnetic fields. This magnetic hysteresis of the  $I_c$  is due to weak links in a high-temperature oxide superconductor and can be removed by heating the superconductor above its critical temperature ( $T_c$ ) [9].

#### B.2 Sequence of measurement conditions

The sequence of conditions in a given application determines the appropriate sequence during the measurements. For the example of a tape conductor near the mid-plane of a simple solenoid magnet, the sequence of conditions would be: the conductor is cooled to the operating temperature and then the current and the magnetic field (parallel to the face of the tape) are ramped up together along the load-line of the magnet until the operating current or  $I_c$  is reached.

In a typical  $I_c$  measurement where the hysteresis with sample current can be regarded as small, this is simplified and the practice may be: the magnetic field and the sample current are ramped independently and the  $I_c$  is measured at a number of fixed and monotonically increasing magnetic fields.

The possible improvement in the  $I_c$  by first exposing the sample to higher magnetic fields is not a relevant sequence unless this is somehow to be accomplished in the application. Thus, in general, the relevant sequence of parameters is:

- a) the specimen is cooled in zero field, from a temperature above  $T_c$  down to the measuring temperature;
- b) the desired field angle with respect to the conductor cross-section is fixed;
- c) the  $I_c$  can be measured at various, monotonically increasing magnetic fields.

If  $I_c$  data at other field angles is needed, the specimen must be heated to a temperature above  $T_c$  and the above sequence repeated.

#### B.3 Reduction of magnetic hysteresis

The effects of magnetic hysteresis can be reduced by ramping the magnet to a lower or zero field and then back up to the test field. Using this technique, acceptably low hysteresis can be demonstrated on Bi-2223 conductors for the higher magnetic fields (for example 4 T at 4,2 K). If this can be demonstrated on a given conductor, then the step of warming and cooling in zero field can be omitted.

The effect of hysteresis on  $I_c$  may depend on many factors such as: material processing, magnetic field strength, field angle, temperature, criteria, and extent of field sweep. As these materials evolve, the magnitude of the hysteresis effects may be reduced.

#### **B.4 Magnetic field angle dependence**

Similar magnetic hysteresis of the  $I_c$  has been observed in magnetic-field angle dependence measurements on tape specimens. The  $I_c$  at a given magnetic-field angle of some multi-filamentary Bi-2223 conductors can be as much as 50 % higher when they are measured with an angle sweep direction that results in a decreasing normal magnetic-field component (field perpendicular to the face of the tape) compared to an angle sweep direction that results in an increasing normal magnetic-field component. As with field sweeps, angle sweeps result in fairly reproducible hysteresis loops, except the first point (the point that was arrived at with the relevant sequence of conditions) is not repeated. Comparisons of angle dependence measurements at fixed angle to angle sweeps show these data are related; however, if significant hysteresis exists, the data obtained from angle sweeps will be significantly different for some angles. These differences can be as large as 20 %.

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**Annex ZA**  
(normative)

**Normative references to international publications  
with their corresponding European publications**

The following referenced documents are indispensable for the application of this document. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

NOTE When an international publication has been modified by common modifications, indicated by (mod), the relevant EN/HD applies.

<u>Publication</u>	<u>Year</u>	<u>Title</u>	<u>EN/HD</u>	<u>Year</u>
IEC 60050-815	2000	International Electrotechnical Vocabulary (IEV) Part 815: Superconductivity	-	-

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