

SLOVENSKI STANDARD oSIST prEN IEC 61557-16:2022

01-julij-2022

Električna varnost v nizkonapetostnih razdelilnih sistemih za izmenične napetosti do 1 000 kV in enosmerne napetosti do 1 500 kV - Oprema za preskušanje, merjenje ali nadzorovanje zaščitnih ukrepov - 16. del: Oprema za preskušanje učinkovitosti zaščitnih ukrepov električne opreme oziroma medicinske električne opreme

Electrical safety in low voltage distribution systems up to 1 000 V a.c. and 1 500 V d.c -Equipment for testing, measuring or monitoring of protective measures - Part 16: Equipment for testing the effectiveness of the protective measures of electrical equipment and/or medical electrical equipment

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Sécurité électrique dans les réseaux de distribution basse tension de 1 000 V c.a. et 1 500 V c.c. - Dispositifs de contrôle, de mesure ou de surveillance de mesures de protection - Partie 16: Équipement pour les essais de bon fonctionnement des mesures de protection de l'équipement électrique et/ou de l'équipement médical électrique

Ta slovenski standard je istoveten z: prEN IEC 61557-16:2022

<u>ICS:</u>

17.220.20	Merjenje električnih in magnetnih veličin	Measurement of electrical and magnetic quantities
29.080.01	Električna izolacija na splošno	Electrical insulation in general
29.240.01	Omrežja za prenos in distribucijo električne energije na splošno	Power transmission and distribution networks in general

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85/831/CDV

COMMITTEE DRAFT FOR VOTE (CDV)

PROJECT NUMBER:	
IEC 61557-16 ED2	
DATE OF CIRCULATION:	CLOSING DATE FOR VOTING:
2022-06-03	2022-08-26
SUPERSEDES DOCUMENTS:	
85/789/CD, 85/804A/CC	

IEC TC 85 : MEASURING EQUIPMENT FOR ELECTRICAL AND ELECTROMAGNETIC QUANTITIES		
SECRETARIAT:	SECRETARY:	
China	Ms Guiju HAN	
OF INTEREST TO THE FOLLOWING COMMITTEES:	PROPOSED HORIZONTAL STANDARD:	
SC 62A,TC 66		
	Other TC/SCs are requested to indicate their interest, if any, in this CDV to the secretary.	
FUNCTIONS CONCERNED:		
EMC Environment	QUALITY ASSURANCE SAFETY	
SUBMITTED FOR CENELEC PARALLEL VOTING	NOT SUBMITTED FOR CENELEC PARALLEL VOTING	
Attention IEC-CENELEC parallel voting		
The attention of IEC National Committees, members of CENELEC, is drawn to the fact that this Committee Draft for Vote (CDV) is submitted for parallel voting. The CENELEC members are invited to vote through the CENELEC online voting system.	<u>61557-16:2022</u> ards/sist/6fb48900-d286-4db2-8541- en-iec-61557-16-2022	

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TITLE:

Electrical safety in low voltage distribution systems up to 1 000 V a.c. and 1 500 V d.c -Equipment for testing, measuring or monitoring of protective measures - Part 16: Equipment for testing the effectiveness of the protective measures of electrical equipment and/or medical electrical equipment

PROPOSED STABILITY DATE: 2028

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54	IE	C CDV 61557-16 © IEC 2022 4 INTERNATIONAL ELECTROTECHNICAL COMMISSION	85/831/CDV
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56 57 58 59		ELECTRICAL SAFETY IN LOW VOLTAGE DISTRIBUTION SYST TO 1 000 V AC AND 1 500 V DC – EQUIPMENT FOR TESTIN MEASURING OR MONITORING OF PROTECTIVE MEASURE	EMS UP NG, S –
60 61 62 63		Part 16: Equipment for testing the effectiveness of the prote measures of electrical equipment and/or medical electrical equ	ective uipment
64		FOREWORD	
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97 98	Int Me	ternational Standard IEC 61557-16 has been prepared by IEC technical co easuring equipment for electrical and electromagnetic quantities.	mmittee 85:
99 100	Th co	nis second edition cancels and replaces the first edition published in 2014. Institutes a technical revision.	This edition
101 102	Th ed	his edition includes the following significant technical changes with respect to lition:	the previous
103 104	a)	splitting of uncertainty requirements for medical and non-medical electrical e	equipment in
105	b)	addition of a definition of ranges with defined uncertainty in 4.1.2 up to 4.1.7	
106	c)	addition of an optional measuring device (MD) for non-medical devices in 4.2.	Ι.
107 108	d)	addition of a limitation of the maximum intrinsic uncertainty for medical ap leakage current in 4.2.1	plications at

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- e) change of 4.2.3 from test sockets to sockets for service purposes; 109
- f) addition of a warning in the operating instructions; 110
- g) integration of former 6.3 into 6.2; 111
- h) update of Table 1; 112
- i) alignment of the structure with that of the whole IEC 61557 series. 113
- 114
- The text of this standard is based on the following documents: 115

FDIS	Report on voting

116

Full information on the voting for the approval of this standard can be found in the report on 117 voting indicated in the above table. 118

- This publication has been drafted in accordance with the ISO/IEC Directives, Part 2. 119
- This part of IEC 61557 shall be used in conjunction with IEC 61557-1. 120

A list of all parts in the IEC 61557 series, published under the general title Electrical safety in 121 low voltage distribution systems up to 1 000 V AC and 1 500 V DC - Equipment for testing, 122 measuring or monitoring of protective measures, can be found on the IEC website. 123

The committee has decided that the contents of this publication will remain unchanged until the 124 stability date indicated on the IEC web site under "http://webstore.iec.ch" in the data related to 125 the specific publication. At this date, the publication will be 126

- 127 •
- reconfirmed, https://standards.iteh.ai/catalog/standards/sist/6fb48900-d286-4db2-8541-
- withdrawn. 128 •
- replaced by a revised edition, or 129 •
- amended. 130 •
- 131
- 132

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INTRODUCTION

This part of IEC 61557 defines performance requirements of measuring equipment intended for testing the effectiveness of the protective measures of electrical equipment and/or medical electrical equipment (in accordance with IEC 62353). It is the intention of this standard to achieve comparable measuring results, additional safety for the testing person, and nondamaging electrical stress for the unit under test.

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ELECTRICAL SAFETY IN LOW VOLTAGE DISTRIBUTION SYSTEMS UP TO 1 000 V AC AND 1 500 V DC- EQUIPMENT FOR TESTING, MEASURING OR MONITORING OF PROTECTIVE MEASURES -

Part 16: Equipment for testing the effectiveness of the protective measures of electrical equipment and/or medical electrical equipment

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149 **1 Scope**

This part 16 of the IEC 61557 series specifies the requirements applicable to the performance for test and measurement equipment in order to determine the effectiveness of the protective measures for electrical equipment and/or medical electrical equipment described in IEC 62353.

153 **2** Normative references

The following documents are referred to in the text in such a way that some or all of their content
 constitutes requirements 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.

- 158 IEC 60529, Degrees of protection provided by enclosures (IP Code)
- 159 IEC 60601-1:2005, Medical electrical equipment Part 1: General requirements for basic
- 160 safety and essential performance
- 161 IEC 60601-1:2005/AMD1:2012
- 162 IEC 60601-1:2005/AMD2:2020 ai/catalog/standards/sist/6fb48900-d286-4db2-8541-
- 163 IEC 61000-4-8, *Electromagnetic compatibility (EMC) Part 4-8: Testing and measurement* 164 *techniques – Power frequency magnetic field immunity test*
- 165 IEC 61010-1:2010, Safety requirements for electrical equipment for measurement, control, and 166 laboratory use – Part 1: General requirements

167 IEC 61010-031, Safety requirements for electrical equipment for measurement, control and
 168 laboratory use – Part 031: Safety requirements for hand-held probe assemblies for electrical
 169 measurement and test

170 IEC 61010-2-030, Safety requirements for electrical equipment for measurement, control, and 171 laboratory use – Part 2-030: Particular requirements for testing and measuring circuits

172 IEC 61010-2-032, Safety requirements for electrical equipment for measurement, control and 173 laboratory use – Part 2-032: Particular requirements for hand-held and hand-manipulated 174 current sensors for electrical test and measurement

175 IEC 61326-1, Electrical equipment for measurement, control and laboratory use – EMC 176 requirements – Part 1: General requirements

177 IEC 61326-2-2, Electrical equipment for measurement, control and laboratory use – EMC 178 requirements – Part 2-2: Particular requirements – Test configurations, operational conditions 179 and performance criteria for portable test, measuring and monitoring equipment used in low-180 voltage distribution systems

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181 IEC 61557-1:2019, Electrical safety in low voltage distribution systems up to 1 000 V AC. and
 182 1 500 V DC – Equipment for testing, measuring or monitoring of protective measures – Part 1:
 183 General requirements

184 IEC 61557-2:2019, Electrical safety in low voltage distribution systems up to 1 000 V AC and
 1 500 V DC – Equipment for testing, measuring or monitoring of protective measures – Part 2:
 186 Insulation resistance

187 IEC 61557-4:2019, Electrical safety in low voltage distribution systems up to 1 000 V AC and
 188 1 500 V DC – Equipment for testing, measuring or monitoring of protective measures – Part 4:
 189 Resistance of earth connection and equipotential bonding

IEC 61557-10, Electrical safety in low voltage distribution systems up to 1 000 V AC and
 1 500 V DC – Equipment for testing, measuring or monitoring of protective measures – Part 10:
 Combined measuring equipment for testing, measuring or monitoring of protective measures

IEC 61557-13:20xx, Electrical safety in low voltage distribution systems up to 1 000 V AC and
 1 500 V DC – Equipment for testing, measuring or monitoring of protective measures – Part 13:
 Hand-held and hand-manipulated current clamps and sensors for measurement of leakage
 currents in electrical distribution systems

197 IEC 62353:2014, Medical electrical equipment – Recurrent test and test after repair of medical
 198 electrical equipment

3 Terms and definitions

For the purposes of this document, the terms and definitions given in IEC 61557-1:2019, IEC 61557-2:2019, IEC 61557-4:2019, IEC 61557-10:2013 and IEC 61557-13:2011, if applicable, and the following apply.

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- ISO and IEC maintain terminological databases for use in standardization at the following
 addresses:
- 205 IEC Electropedia: available at http://www.electropedia.org/
- 206 ISO Online browsing platform: available at http://www.iso.org/obp
- 207 **3.1**

208 test socket outlet

socket outlet on the test equipment for the unit under test, separated from the active parts ofthe mains circuit by double insulation

- 211 **3.2**
- 212 mains socket outlet
- socket outlet on the test equipment used to supply mains to the equipment under test
- 214 **3.3**

215 combined test-mains socket outlet

- socket outlet on the test equipment that can be switched to either test-condition and/or supply condition
- 218 **3.4**

219 service socket

socket outlet on the test equipment to supply mains to further test equipment or additional

221 equipment

- 222 **3.5**
- 223 test terminal
- terminal used independently, in parallel or in combination, with the test socket
- 225 **3.6**

226 measuring circuit MD

- 227 electric circuit with defined components and defined frequency characteristic
- 228 **3.7**
- 229 peak factor
- ratio of the maximum absolute value of an alternating quantity to its RMS value
- 231 [SOURCE: IEC 60050-103:2009, 103-06-15]
- 232 **3.8**

233 medical electrical equipment

234 **ME equipment**

electrical equipment having an applied part or transferring energy to or from the patient or detecting such energy transfer to or from the patient and which is:

- a) provided with not more than one connection to a particular supply means, and
- b) intended by its manufacturer to be used:
- 239 in the diagnosis, treatment, or monitoring of a patient, or
- 240 for compensation or alleviation of disease, injury or disability
- 241 Note 1 to entry: ME equipment includes those accessories as defined by the manufacturer that are necessary to 242 enable the normal use of the ME equipment.
- 243Note 2 to entry: Not all electrical equipment used in medical practice falls within this definition (e.g. some *in vitro*
diagnostic equipment).244OSIST prEN IEC 61557-16:2022
- Note 3 to entry: The implantable parts of active implantable medical devices can fall within this definition, but they are excluded from the scope of IEC 60601-1.
- 247 [SOURCE: IEC 60601-1:2005, 3.63]

248 **3.9**

249 protective earth resistance

- resistance between any accessible part which has to be connected for safety purposes to the protective earth terminal and the
- 252 protective conductor of the mains plug, or
- 253 protective conductor of the appliance inlet, or
- 254 protective conductor permanently connected to the supply mains;
- resistance between protective connectors at each end of a detachable power supply cord
- 256 (SOURCE: IEC 62353: 2014, 3.34)

257 **4 Requirements**

258 4.1 General requirements

In addition to the requirements of IEC 61557-1:2019, Clause 4, the following requirements shallapply.

- The requirements of IEC 62353:2014, Annex C shall apply for measuring equipment intended
- for medical electrical equipment testing.