



BSI Standards Publication

## Medical electrical equipment

Part 4-3: Guidance and interpretation —  
Considerations of unaddressed safety aspects  
in the third edition of IEC 60601-1 and  
proposals for new requirements

**National foreword**

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The UK participation in its preparation was entrusted by Technical Committee CH/62, Electrical Equipment in Medical Practice, to Subcommittee CH/62/1, Common aspects of Electrical Equipment used in Medical Practice.

A list of organizations represented on this committee can be obtained on request to its secretary.

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# TECHNICAL REPORT

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**Medical electrical equipment –  
Part 4-3: Guidance and interpretation – Considerations of unaddressed safety  
aspects in the third edition of IEC 60601-1 and proposals for new requirements**

INTERNATIONAL  
ELECTROTECHNICAL  
COMMISSION

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## CONTENTS

FOREWORD.....	5
INTRODUCTION.....	7
1 Scope and object.....	8
1.1 Scope .....	8
1.2 Object.....	8
2 Normative references .....	8
3 Recommendations .....	9
3.1 Template used for recommendations prepared by SC 62A/WG 14 .....	9
3.2 Recommendation sheets.....	10
3.2.101 Total PATIENT LEAKAGE CURRENT of a ME SYSTEM.....	10
3.2.102 Pollution degree for MOPP .....	10
3.2.103 Transients on d.c. mains.....	11
3.2.104 Altitude factor for DEFIBRILLATION-PROOF APPLIED PARTS.....	12
3.2.105 Defibrillation energy protection for MOOP / MOPP.....	13
3.2.106 Overvoltage categories III and IV.....	13
3.2.107 Pollution degree related to different micro/macro environments .....	13
3.2.108 Warnings versus ALARM SIGNALS .....	14
3.2.109 Single Y1 capacitor for MOPP .....	14
3.2.110 WORKING VOLTAGE > 14 140 V peak .....	15
3.2.111 CREEPAGE DISTANCE and AIR CLEARANCE for dental equipment.....	15
3.2.112 Short circuiting of one constituent part of DOUBLE INSULATION .....	16
3.2.113 Instability in transport position .....	16
3.2.114 When to conduct leakage current tests after humidity preconditioning treatment.....	17
3.2.115 DEFIBRILLATION-PROOF TYPE B APPLIED PARTS .....	17
3.2.116 Instability excluding transport position .....	18
3.2.117 DIELECTRIC STRENGTH of two serial MOPP barrier parts.....	18
3.2.118 Overheating transformer .....	19
3.2.119 Test equipment for recurrent tests according to IEC 62353 testing used within IEC 60601-1 type approval testing.....	20
3.2.120 Tolerances of apparatus .....	22
3.2.121 FUNCTIONAL EARTH CONDUCTOR and ESSENTIAL PERFORMANCE .....	23
3.2.122 A.c. motors .....	24
3.2.123 Operational insulation.....	25
3.2.124 WORKING VOLTAGE measurement .....	25
3.2.125 Defibrillation test.....	26
3.2.126 Oil containers for moving parts .....	27
3.2.127 PERMANENTLY INSTALLED ME EQUIPMENT in the HOME HEALTHCARE ENVIRONMENT .....	27
3.2.128 Polystyrene plate for LEAKAGE CURRENT tests .....	30
3.2.129 Push buttons .....	31
3.2.130 Temperature limit at the ENCLOSURE in SINGLE FAULT CONDITION.....	31
3.2.131 Optic coupler requirements.....	33
3.2.132 Eye-verification of tester before legibility test.....	35
3.2.133 End stops to prevent overtravel .....	36
3.2.134 MOPP barrier with low WORKING VOLTAGE r.m.s. and high WORKING VOLTAGE peak.....	37

3.2.135	Labeling: spare parts vs. detachable parts vs. ACCESSORIES .....	38
3.2.136	Protective earth impedance of ME SYSTEM >200 mΩ .....	41
3.2.137	Ball pressure test.....	42
3.2.138	Magnesium alloy ENCLOSURE.....	43
3.2.139	Instability with initial movement .....	44
3.2.140	Ball pressure test.....	45
3.2.141	DIELECTRIC STRENGTH test values .....	47
3.2.142	SECONDARY CIRCUITS .....	48
3.2.143	LEAKAGE CURRENTS in SINGLE FAULT CONDITION and during component faults .....	48
3.2.144	Impedance of a PROTECTIVE EARTH CONDUCTOR within a DETACHABLE POWER SUPPLY CORD.....	49
3.2.145	Time delay of the 100 VA limit .....	50
3.2.146	Test voltage multiplied by factor 1,6 .....	51
3.2.147	Overflow, spillage, .....	51
3.2.148	DIELECTRIC STRENGTH test of transformers without accessible frame .....	52
3.2.149	Expected voltage on SIP/SOPs .....	52
3.2.150	Flammability rating for transformer bobbin .....	53
3.2.151	COMPONENT WITH HIGH-INTEGRITY CHARACTERISTICS.....	54
3.2.152	Peak and r.m.s. WORKING VOLTAGES .....	55
3.2.153	Critical components .....	56
3.2.154	LEAKAGE CURRENT test for ME EQUIPMENT with multiple APPLIED PARTS.....	56
3.2.155	DIELECTRIC STRENGTH test value for extruded and spirally wrapped multi-layer wires .....	57
3.2.156	DIELECTRIC STRENGTH test after thermal cycling test.....	57
3.2.157	Required MOOP values higher than MOPP values .....	58
3.2.158	Optocouplers .....	58
3.2.159	Impact test.....	59
3.2.160	Spillage test in NORMAL CONDITION and in SINGLE FAULT CONDITION .....	60
3.2.161	TYPE B APPLIED PART connected to ACCESSIBLE PARTS .....	61
3.2.162	Current/power labeling.....	62
3.2.163	Separate power supply part of ME EQUIPMENT or ME SYSTEM .....	62
3.2.164	Specification of the allowed power supply.....	63
3.2.165	Mains transients for opposite polarity on the secondary side or battery pole to pole barrier .....	64
3.2.166	Keep dry and umbrella symbol.....	65
3.2.167	MOBILE and STATIONARY ME EQUIPMENT with wheels .....	66
3.2.168	Varistors installed in the MAINS PART .....	67
3.2.169	Using Y2 capacitors for MOPP .....	67
3.2.170	Overtravel end stops – specification of the speed .....	68
3.2.171	CREEPAGE DISTANCE and AIR CLEARANCE between input and output of fuse contacts .....	69
3.2.172	Examples of SINGLE FAULT CONDITION .....	69
3.2.173	Examples of ME SYSTEMS .....	70
3.2.174	Cross sectional area of POWER SUPPLY CORD for rated input current > 63 A.....	70
3.2.175	Biocompatibility for quasi APPLIED PARTS .....	71
3.2.176	Floating reference earth.....	71
3.2.177	SINGLE FAULT CONDITION in OXYGEN RICH ENVIRONMENT.....	72
3.2.178	Laser requirements .....	74

3.2.179	Flammability rating of insulated wires .....	74
3.2.180	Infrared lamps .....	75
3.2.181	Identification of internal fuses .....	76
3.2.182	Chargers for ME EQUIPMENT used at home .....	77
3.2.183	CLASS II ME EQUIPMENT with FUNCTIONAL EARTH CONDUCTOR .....	78
3.2.184	Symbol D2-2 on MSO .....	78
3.2.185	PATIENT leads connectors .....	79
3.2.186	Rationale for IP2X .....	80
3.2.187	Battery – limited power .....	80
3.2.188	TYPE B APPLIED PART separated from ACCESSIBLE PARTS.....	81
3.2.189	Protective earth test >25A .....	81
3.2.190	Reference to IEC 62304:2006.....	82
3.2.191	The SIP/SOP pin to earth TOUCH CURRENT.....	82
3.2.192	Overbalancing .....	84
3.2.193	MAINS VOLTAGE on APPLIED PART.....	85
Annex A (informative) Overview of the recommendations developed by IEC/SC 62A/WG 14 .....		86
Bibliography.....		91

Table A.1 – Cross-reference of recommendations by subclause of IEC 60601-1:2005 and IEC 60601-1:2005/AMD1:2012 (1 of 5).....	86
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## INTERNATIONAL ELECTROTECHNICAL COMMISSION

**MEDICAL ELECTRICAL EQUIPMENT –****Part 4-3: Guidance and interpretation – Considerations  
of unaddressed safety aspects in the third edition of IEC 60601-1  
and proposals for new requirements**

## FOREWORD

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IEC TR 60601-4-3, which is a technical report, has been prepared by subcommittee 62A: Common aspects of electrical equipment used in medical practice, of IEC technical committee 62: Electrical equipment in medical practice.

The text of this technical report is based on the following documents:

Enquiry draft	Report on voting
62A/951/DTR	62A/973A/RVC

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

This publication has been drafted in accordance with the ISO/IEC Directives, Part 2.

Terms used throughout this technical report that have been defined in Clause 3 of IEC 60601-1:2005 and IEC 60601-1:2005/AMD 1:2012 are printed in SMALL CAPITALS.

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

- reconfirmed,
- withdrawn,
- replaced by a revised edition, or
- amended.

A bilingual version of this technical report may be issued at a later date.

## INTRODUCTION

At the Sydney meeting in August 1994, IEC subcommittee (SC) 62A established a procedure under which working group (WG) 14 would develop recommendations regarding problems of interpretation or application of IEC 60601-1. WG 14 is made up of experts with particular expertise in testing according to the requirements of IEC 60601-1. Many of the experts on WG 14 are employed by test laboratories with a long history of applying IEC 60601-1 to MEDICAL ELECTRICAL EQUIPMENT. While the National Committee members of SC 62A nominate these experts, their recommendations were not to be formally adopted through any official voting procedure. To reinforce this process, the Subcommittee specifically directed that the following note appear on every page of the resulting informational circular:

**IMPORTANT NOTE:** Per the 62A decision at Sydney (see RM3755/SC62A, August 1994), the 62A Secretary is circulating this recommendation, prepared by 62A/WG 14, regarding problems of interpretation or application of IEC 60601-1 to all P-Member NCs.

This recommendation/interpretation is the result of considerations by a group of nominated experts and has not been formally adopted through any National Committee voting procedure. Distribution is only for information.

At the November 2000 meeting of SC 62A in Tokyo, the subcommittee discussed ways and means for achieving a wider distribution of the WG 14 recommendations. At the conclusion of this discussion, the subcommittee instructed the Secretariat to develop a technical report (TR) based on the published recommendations of WG 14. This technical report is intended to convey the results of WG 14's work to interested parties such as MANUFACTURERS and test laboratories while retaining the informative nature of the material.

This first edition of IEC TR 60601-4-3 contains 93 recommendations, numbered 101 to 193. All these recommendations are based upon IEC 60601-1:2005 and IEC 60601-1:2005/AMD1:2012.

The numbering starts with 101 instead of just 1 to ensure that these WG 14 recommendations (101 to 193) will not accidentally be confused with previous issued WG 14 recommendations 1 to 63, which are based on the second edition of IEC 60601-1 and published in IEC TR 62296.

This technical report may be amended from time to time as WG 14 prepares additional recommendations.

## MEDICAL ELECTRICAL EQUIPMENT

### Part 4-3: Guidance and interpretation – Considerations of unaddressed safety aspects in the third edition of IEC 60601-1 and proposals for new requirements

#### 1 Scope and object

##### 1.1 Scope

This technical report contains a series of recommendations developed by an expert working group of IEC subcommittee 62A in response to questions of interpretation of the third edition of IEC 60601-1.

This technical report is primarily intended to be used by:

- MANUFACTURERS of MEDICAL ELECTRICAL EQUIPMENT;
- test laboratories and others responsible for assessment of compliance with IEC 60601-1:2005 and IEC 60601-1:2005/AMD1:2012, and
- those developing subsequent editions of IEC 60601-1.

The recommendations in the first edition of IEC/TR 62296 were considered in preparing the third edition of IEC 60601-1. Similarly it is expected that these recommendations within IEC 60601-4-3 will be considered when preparing a future revision of IEC 60601-1.

##### 1.2 Object

The object of this technical report is to make the recommendations/interpretations developed by the experts in IEC/SC 62A/WG 14 available to those interested in the application of the third edition of IEC 60601-1.

The reader is reminded that, although a majority of the National Committee members of IEC/SC 62A have approved publication of this technical report, the contents remain the opinion of the expert members of WG 14. These recommendations/interpretations are the result of considerations by this group of nominated experts and have not been formally adopted through any National Committee voting procedure. Distribution is only for information.

#### 2 Normative references

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

IEC 60601-1:2005, *Medical electrical equipment – Part 1: General requirements for basic safety and essential performance*  
IEC 60601-1:2005/AMD1:2012

IEC 60601-1-8:2006, *Medical electrical equipment – Part 1-8: General requirements for basic safety and essential performance – Collateral standard: General requirements, tests and guidance for alarm systems in medical electrical equipment and medical electrical systems*  
IEC 60601-1-8:2006/AMD1:2012

IEC 60601-1-11:2010, *Medical electrical equipment – Part 1-11: General requirements for basic safety and essential performance – Collateral standard: Requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment*