

INTERNATIONAL STANDARD

NORME INTERNATIONALE



Medical electrical equipment –

**Part 2-31: Particular requirements for the basic safety and essential performance
of external cardiac pacemakers with internal power source**

Appareils électromédicaux –

**Partie 2-31: Exigences particulières pour la sécurité de base et les performances
essentielles des stimulateurs cardiaques externes à source d'énergie interne**



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performances essentielles des stimulateurs cardiaques externes à source
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CONTENTS

FOREWORD	4
INTRODUCTION	7
201.1 Scope, object and related standards	8
201.2 Normative references	10
201.3 * Terms and definitions	10
201.4 General requirements	12
201.5 General requirements for testing ME EQUIPMENT	13
201.6 Classification of ME EQUIPMENT and ME SYSTEMS	13
201.7 ME EQUIPMENT identification, marking and documents	13
201.8 Protection against electrical HAZARDS from ME EQUIPMENT	17
201.9 Protection against MECHANICAL HAZARDS of ME EQUIPMENT and ME SYSTEMS	23
201.10 Protection against unwanted and excessive radiation HAZARDS	23
201.11 Protection against excessive temperatures and other HAZARDS	23
201.12 Accuracy of controls and instruments and protection against hazardous outputs	24
201.13 HAZARDOUS SITUATIONS and fault conditions for ME EQUIPMENT	29
201.14 PROGRAMMABLE ELECTRICAL MEDICAL SYSTEMS (PEMS)	29
201.15 Construction of ME EQUIPMENT	29
201.16 ME SYSTEMS	29
201.17 Electromagnetic compatibility of ME EQUIPMENT and ME SYSTEMS	29
202 * ELECTROMAGNETIC DISTURBANCES – Requirements and tests	29
Annexes	31
Annex I Identification of IMMUNITY pass/fail criteria	31
Annex AA (informative) Particular guidance and rationale	32
Bibliography	52
Index of defined terms used in this particular standard	53
Figure 201.101 – Test waveform V_{test} implemented by example RCL circuit using $C = 120 \mu\text{F}$, $L = 25 \mu\text{H}$, $RL + R = 1 \Omega$	18
Figure 201.102 – Example circuit of defibrillation test voltage generator for generating a decaying exponential waveform	19
Figure 201.103 – Test setup for a SINGLE CHAMBER external CARDIAC PACEMAKER	20
Figure 201.104 – Test setup for a DUAL CHAMBER external CARDIAC PACEMAKER	20
Figure 201.105 – Test setup for a triple chamber external CARDIAC PACEMAKER, e.g. bi-ventricular external CARDIAC PACEMAKER	21
Figure 201.106 – Timing sequence	21
Figure 201.107 – Measuring circuit for the PATIENT AUXILIARY CURRENT for ME EQUIPMENT with an INTERNAL ELECTRICAL POWER SOURCE	23
Figure 201.108 – Measuring circuit for the MAXIMUM TRACKING RATE	26
Figure 201.109 – Initial oscilloscope display when measuring MAXIMUM TRACKING RATE	27
Figure AA.1 – Simple model of a SINGLE CHAMBER EXTERNAL PACEMAKER during defibrillation	39

Figure AA.2 – First proposal for a defib-protection test of SINGLE CHAMBER EXTERNAL PACEMAKER.....	41
Figure AA.3 – Circuit for a defibrillation test generator for defibrillation test according to conditions during open heart surgery	42
Figure AA.4 – Defibrillation PULSE generated by the defibrillation test generator from Figure AA.3	43
Figure AA.5 – Rise times of a defibrillation PULSE according to the circuit proposed in Figure AA.3	47
Table 201.101 – Distributed ESSENTIAL PERFORMANCE requirements	12
Table 201.102 – DUAL CHAMBER connector terminal marking.....	14
Table 201.103 – ME EQUIPMENT parameters.....	25
Table 202.101 – Static discharge requirements.....	30
Table AA.1 – EXTERNAL PACEMAKER HAZARD inventory	33
Table AA.2 – PULSE energies calculated for $C = 120 \mu\text{F} \pm 5 \%$	44
Table AA.3 – PULSE energies calculated for $C = 122 \mu\text{F} \pm 5 \%$	45
Table AA.4 – PULSE energies calculated for $C = 126,32 \mu\text{F} \pm 5 \%$	46

INTERNATIONAL ELECTROTECHNICAL COMMISSION

MEDICAL ELECTRICAL EQUIPMENT –**Part 2-31: Particular requirements for the basic safety and essential performance of external cardiac pacemakers with internal power source**

FOREWORD

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International standard IEC 60601-2-31 has been prepared by a Joint Working Group of IEC subcommittee 62D: Electromedical equipment, of IEC technical committee 62: Electrical equipment in medical practice, and ISO subcommittee SC6: Active implants, of ISO technical committee 150: Implants for surgery.

This publication is published as a double logo standard.

This third edition cancels and replaces the second edition published in 2008 and Amendment 1:2011. This edition constitutes a technical revision.

This edition includes the following significant technical changes with respect to the previous edition:

- a) The requirement for testing for energy reduction has been removed;
- b) The test for exposure to external defibrillation has been completely revised;



- c) The exclusion for testing ESD immunity only with respect to air discharges has been removed;
- d) Alignment with the latest edition of ISO 14708-2 for pacemakers, as well as the associated EMC standard ISO 14117;
- e) Additional rationale for all changes.

The text of this International Standard is based on the following documents of IEC:

FDIS	Report on voting
62D/1719/FDIS	62D/1732A/RVD

Full information on the voting for the approval of this International Standard can be found in the report on voting indicated in the above table. In ISO, the standard has been approved by 10 P members out of 10 having cast a vote.

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

In this document, the following print types are used:

- requirements and definitions: roman type;
- *test specifications: italic type;*
- informative material appearing outside of tables, such as notes, examples and references: in smaller type. Normative text of tables is also in a smaller type;
- TERMS DEFINED IN CLAUSE 3 OF THE GENERAL STANDARD, IN THIS PARTICULAR STANDARD OR AS NOTED: SMALL CAPITALS.

In referring to the structure of this document, the term

- "clause" means one of the seventeen numbered divisions within the table of contents, inclusive of all subdivisions (e.g. Clause 7 includes subclauses 7.1, 7.2, etc.);
- "subclause" means a numbered subdivision of a clause (e.g. 7.1, 7.2 and 7.2.1 are all subclauses of Clause 7).

References to clauses within this document are preceded by the term "Clause" followed by the clause number. References to subclauses within this particular standard are by number only.

In this document, the conjunctive "or" is used as an "inclusive or" so a statement is true if any combination of the conditions is true.

The verbal forms used in this document conform to usage described in Clause 7 of the ISO/IEC Directives, Part 2. For the purposes of this document, the auxiliary verb:

- "shall" means that compliance with a requirement or a test is mandatory for compliance with this document;
- "should" means that compliance with a requirement or a test is recommended but is not mandatory for compliance with this document;
- "may" is used to describe a permissible way to achieve compliance with a requirement or test.

An asterisk (*) as the first character of a title or at the beginning of a paragraph or table title indicates that there is guidance or rationale related to that item in Annex AA.

A list of all parts of the IEC 60601 series, published under the general title *Medical electrical equipment*, can be found on the IEC website.

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

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

NOTE The attention of users of this document is drawn to the fact that equipment manufacturers and testing organizations may need a transitional period following publication of a new, amended or revised IEC publication in which to make products in accordance with the new requirements and to equip themselves for conducting new or revised tests. It is the recommendation of the committee that the content of this publication be adopted for implementation nationally not earlier than 3 years from the date of publication.

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INTRODUCTION

The minimum safety requirements specified in this particular standard are considered to provide for a practical degree of safety in the operation of EXTERNAL PACEMAKERS with an internal power source.

Basically, CARDIAC PACEMAKERS treat cardiac arrhythmias. Such arrhythmias reduce cardiac output and can lead to confusion, dizziness, loss of consciousness and death. The objective of pacing is to restore cardiac rhythm and output appropriate to the PATIENT's physiological needs.

There are two distinct families of CARDIAC PACEMAKERS, implantable PACEMAKERS and EXTERNAL PACEMAKERS. EXTERNAL PACEMAKERS are used to pace PATIENTS temporarily prior to implanting an implantable PACEMAKER as well as for temporary pacing related to other medical PROCEDURES, e.g. open heart surgery.

CARDIAC PACEMAKERS differ in the various ways in which they maintain and monitor cardiac activity in different circumstances. The simplest model stimulates the atrium or ventricle independently of the cardiac activity; others detect atrial or ventricular activity and stimulate the atrium or ventricle as and when this is necessary; others, more complex, detect the spontaneous heart activity and stimulate appropriately the atrium and/or the ventricle. Certain PACEMAKERS work on preset frequency values, amplitudes and impulse duration. Others can have several values for parameters.

Standards for EXTERNAL PACEMAKERS require attention to information which will aid in developing and applying these devices. It is through these aspects of standardization that the central role of clinical experience should be, or has been, acknowledged. The ability to predict how a PACEMAKER will perform in a specific PATIENT based on testing of a device to a set of technical criteria is limited.

This particular standard does not take into consideration the specific safety aspects of EXTERNAL PACEMAKERS that are connected to a SUPPLY MAINS while simultaneously connected to the PATIENT.

This particular standard amends and supplements IEC 60601-1:2005 and IEC 60601-1:2005/AMD1:2012, *Medical electrical equipment – Part 1: General requirements for basic safety and essential performance*, hereinafter referred to as the general standard.

The requirements are followed by specifications for the relevant tests.

Following the decision taken by subcommittee 62D at the meeting in Washington in 1979, a "General guidance and rationale" section giving some explanatory notes, where appropriate, about the more important requirements is included in Annex AA.

Clauses or subclauses for which there are explanatory notes in Annex AA are marked with an asterisk (*).

An inventory of the PATIENT's safety posed by EXTERNAL PACEMAKERS and a rationale for the safety requirements contained in this particular standard are given in Annex AA. It is considered that knowledge of the reasons for these requirements will not only facilitate the proper application of this particular standard but will, in due course, expedite any revision necessitated by changes in clinical practice or as a result of developments in technology. However, Annex AA does not form part of the requirements of this document.

MEDICAL ELECTRICAL EQUIPMENT –

Part 2-31: Particular requirements for the basic safety and essential performance of external cardiac pacemakers with internal power source

201.1 Scope, object and related standards

Clause 1 of the general standard¹ applies, except as follows:

201.1.1 * Scope

Replacement:

This part of IEC 60601 applies to the BASIC SAFETY and ESSENTIAL PERFORMANCE of EXTERNAL PACEMAKERS powered by an INTERNAL ELECTRICAL POWER SOURCE, hereafter referred to as ME EQUIPMENT.

This document applies to PATIENT CABLES as defined in 201.3.209, but does not apply to LEADS as defined in 201.3.206.

HAZARDS inherent in the intended physiological function of ME EQUIPMENT within the scope of this document are not covered by specific requirements in this document except in 7.2.13 and 8.4.1 of the general standard.

NOTE See also 4.2 of the general standard.

This document does not apply to the implantable parts of ACTIVE IMPLANTABLE MEDICAL DEVICES covered by ISO 14708-1. This document does not apply to EXTERNAL PACEMAKERS which can be connected directly or indirectly to a SUPPLY MAINS.

This document does not apply to transthoracic and oesophageal pacing ME EQUIPMENT and antitachycardia ME EQUIPMENT.

201.1.2 Object

Replacement:

The object of this particular standard is to establish particular BASIC SAFETY and ESSENTIAL PERFORMANCE requirements for EXTERNAL PACEMAKERS as defined in 201.3.205.

201.1.3 Collateral standards

Addition:

This particular standard refers to those applicable collateral standards that are listed in Clause 2 of the general standard and Clause 201.2 of this particular standard.

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

IEC 60601-1-2:2014 applies as modified in Clause 202. IEC 60601-1-3 does not apply. All other published collateral standards in the IEC 60601-1 series apply as published.

201.1.4 Particular standards

Replacement:

In the IEC 60601 series, particular standards may modify, replace or delete requirements contained in the general standard and collateral standards as appropriate for the particular ME EQUIPMENT under consideration, and may add other BASIC SAFETY and ESSENTIAL PERFORMANCE requirements.

A requirement of a particular standard takes priority over the general standard.

For brevity, IEC 60601-1:2005 and IEC 60601-1:2005/AMD1:2012 are referred to in this particular standard as the general standard. Collateral standards are referred to by their document number.

The numbering of clauses and subclauses of this particular standard corresponds to that of the general standard with the prefix "201" (e.g. 201.1 in this document addresses the content of Clause 1 of the general standard) or applicable collateral standard with the prefix "20x" where x is the final digit(s) of the collateral standard document number (e.g. 202.4 in this particular standard addresses the content of Clause 4 of the IEC 60601-1-2 collateral standard, 203.4 in this particular standard addresses the content of Clause 4 of the IEC 60601-1-3 collateral standard, etc.). The changes to the text of the general standard and applicable collateral standards are specified by the use of the following words:

"Replacement" means that the clause or subclause of the general standard or applicable collateral standard is replaced completely by the text of this particular standard.

"Addition" means that the text of this particular standard is additional to the requirements of the general standard or applicable collateral standard.

"Amendment" means that the clause or subclause of the general standard or applicable collateral standard is amended as indicated by the text of this particular standard.

Subclauses, figures or tables which are additional to those of the general standard are numbered starting from 201.101. However, due to the fact that definitions in the general standard are numbered 3.1 through 3.147, additional definitions in this document are numbered beginning from 201.3.201. Additional annexes are lettered AA, BB, etc., and additional items aa), bb), etc.

Subclauses, figures or tables which are additional to those of a collateral standard are numbered starting from 20x, where "x" is the number of the collateral standard, e.g. 202 for IEC 60601-1-2, 203 for IEC 60601-1-3, etc.

The term "this document" is used to make reference to the general standard, any applicable collateral standards and this particular standard taken together.

Where there is no corresponding clause or subclause in this particular standard, the clause or subclause of the general standard or applicable collateral standard, although possibly not relevant, applies without modification; where it is intended that any part of the general standard or applicable collateral standard, although possibly relevant, is not to be applied, a statement to that effect is given in this particular standard.

201.2 Normative references

NOTE Informative references are listed in the Bibliography.

Clause 2 of the general standard applies, except as follows:

Replacement:

IEC 60601-1-2:2014, *Medical electrical equipment – Part 1-2: General requirements for basic safety and essential performance – Collateral Standard: Electromagnetic disturbances – Requirements and tests*

Addition:

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

ISO 14117:2019, *Active implantable medical devices – Electromagnetic compatibility – EMC test protocols for implantable cardiac pacemakers, implantable cardioverter defibrillators and cardiac resynchronization devices*

ISO 14708-2:2019, *Implants for surgery – Active implantable medical devices – Part 2: Cardiac pacemakers*

201.3 * Terms and definitions

For the purposes of this document, the terms and definitions specified in IEC 60601-1:2005 and IEC 60601-1:2005/AMD1:2012, ISO 14117:2019, and ISO 14708-2:2019 and the following apply.

ISO and IEC maintain terminological databases for use in standardization at the following addresses:

- IEC Electropedia: available at <http://www.electropedia.org/>
- ISO Online browsing platform: available at <http://www.iso.org/obp>

NOTE An index of defined terms is found beginning on page 53.

Addition:

201.3.201

ACTIVE IMPLANTABLE MEDICAL DEVICE

active medical device which is intended to be totally or partially introduced, surgically or medically, into the human body or by medical intervention into a natural orifice, and which is intended to remain in place after the procedure

[SOURCE: ISO 14708-1:2014, 3.2, modified – The words "in place" have been added to the definition, and the note to entry has been deleted.]

201.3.202

BATTERY DEPLETION INDICATOR

means of indicating when the battery should be replaced

201.3.203

CARDIAC PACEMAKER

ME EQUIPMENT intended to treat bradyarrhythmias

201.3.204**DUAL CHAMBER**

relating to both atrium and ventricle

201.3.205**EXTERNAL PACEMAKER**

CARDIAC PACEMAKER consisting of a NON-IMPLANTABLE PULSE GENERATOR and PATIENT CABLE(S) (if used)

201.3.206**LEAD**

flexible tube enclosing one or more insulated electrical conductors, intended to transfer electrical energy along its length between the EXTERNAL PACEMAKER and the PATIENT'S heart

[SOURCE: ISO 14708-1:2014, 3.13, modified – The words "between the EXTERNAL PACEMAKER and the PATIENT'S heart" have been added to the definition, and the note to entry has been deleted.]

201.3.207**MAXIMUM TRACKING RATE**

maximum PULSE RATE at which the multi-chamber NON-IMPLANTABLE PULSE GENERATOR will respond on a 1:1 basis to a sensed atrial signal

[SOURCE: ISO 14708-2:2019, 3.30, modified – The word "IMPLANTABLE" has been replaced by "multi-chamber NON-IMPLANTABLE" and the word "triggering" by "sensed atrial".]

201.3.208**NON-IMPLANTABLE PULSE GENERATOR**

ME EQUIPMENT with an INTERNAL ELECTRICAL POWER SOURCE which is intended for use outside the body and which produces a periodic electrical PULSE intended to stimulate the heart through a LEAD (or combination of a LEAD and PATIENT CABLE)

201.3.209**PATIENT CABLE**

cable used to extend the distance between the NON-IMPLANTABLE PULSE GENERATOR and the pacing LEAD

201.3.210**POST-VENTRICULAR ATRIAL REFRACTORY PERIOD****PVARP**

refractory period in atrial channel after paced or sensed event in ventricular channel, used in DUAL CHAMBER modes

Note 1 to entry: This note applies to the French language only.

201.3.211**PRIMARY BATTERY**

one or more cells, which are not designed to be electrically recharged, that are fitted with devices necessary for use, for example case, terminals, marking and protective devices

[SOURCE: IEC 60050-482:2004, 482-01-04, modified – The word "primary" has been added to the term, and the words "which are not designed to be electrically recharged" have been added to the definition.]

201.3.212**SINGLE CHAMBER**

relating to either atrium or ventricle

201.3.213

BASIC RATE

PULSE RATE of a NON-IMPLANTABLE PULSE GENERATOR, either atrial or ventricular, unmodified by sensed cardiac or other electrical influence

[SOURCE: ISO 14708-2:2019, 3.26, modified – The words "an implantable" have been replaced by "a NON-IMPLANTABLE".]

201.3.214

ESCAPE INTERVAL

time elapsing between the sensing of a spontaneous BEAT and the succeeding non-triggered PULSE of a NON-IMPLANTABLE PULSE GENERATOR

[SOURCE: ISO 14117:2019, 3.128, modified – The words "an implantable" have been replaced by "a NON-IMPLANTABLE".]

201.3.215

INTERFERENCE PULSE RATE

PULSE RATE with which the NON-IMPLANTABLE PULSE GENERATOR responds when it senses electrical activity that it recognizes as interference

[SOURCE: ISO 14117:2019, 3.129, modified – The words "an implantable" have been replaced by "NON-IMPLANTABLE".]

201.4 General requirements

Clause 4 of the general standard applies, except as follows:

201.4.3 ESSENTIAL PERFORMANCE

Additional subclause:

201.4.3.101 Additional ESSENTIAL PERFORMANCE requirements

Additional ESSENTIAL PERFORMANCE requirements are found in the subclauses listed in Table 201.101.

Table 201.101 – Distributed ESSENTIAL PERFORMANCE requirements

Requirement	Subclause
BATTERY DEPLETION INDICATOR	201.11.8
ME EQUIPMENT parameter stability	201.12.1.101
PULSE AMPLITUDE stability	201.12.1.102
Disabling runaway rate protection	201.12.4.1
Deliberate action required to change settings	201.12.4.101
Parameter stability at onset of the BATTERY DEPLETION INDICATOR	201.12.4.102
Runaway protection	201.12.4.103
Interference reversion in the presence of sensed ELECTROMAGNETIC DISTURBANCE or electrical energy sources	201.12.4.104
Limit at which the ventricle is paced in response to sensed atrial activity	201.12.4.105

201.4.10.1 Source of power for ME EQUIPMENT

Replacement:

ME EQUIPMENT shall be powered by a PRIMARY BATTERY.

Compliance is checked by inspection of the ACCOMPANYING DOCUMENTS.

201.4.10.2 SUPPLY MAINS for ME EQUIPMENT and ME SYSTEMS

This subclause of the general standard does not apply.

201.4.11 * Power input

This subclause of the general standard does not apply.

201.5 General requirements for testing ME EQUIPMENT

Clause 5 of the general standard applies.

201.6 Classification of ME EQUIPMENT and ME SYSTEMS

Clause 6 of the general standard applies, except as follows:

201.6.2 * Protection against electric shock

Replacement:

ME EQUIPMENT shall be classified as INTERNALLY POWERED ME EQUIPMENT.

ME EQUIPMENT shall be recognized as INTERNALLY POWERED only if no external connections to an electrical power source are provided.

APPLIED PARTS shall be classified as TYPE CF APPLIED PARTS. APPLIED PARTS shall be classified as DEFIBRILLATION-PROOF APPLIED PARTS.

201.7 ME EQUIPMENT identification, marking and documents

Clause 7 of the general standard applies, except as follows:

201.7.2 Marking on the outside of ME EQUIPMENT or ME EQUIPMENT parts

Additional subclauses:

201.7.2.101 ME EQUIPMENT intended for SINGLE CHAMBER application

If the ME EQUIPMENT is intended for SINGLE CHAMBER applications, the connector terminals (if used) shall be conspicuously marked positive (+) and negative (–).

201.7.2.102 * ME EQUIPMENT intended for DUAL CHAMBER application

If the ME EQUIPMENT is intended for DUAL CHAMBER application, the connector terminals (if used) shall be marked according to Table 201.102. If colour is used to differentiate between channels in a DUAL CHAMBER application, then the ventricular channel should be marked with the colour white and the atrial channel should be marked with a contrasting colour.

Table 201.102 – DUAL CHAMBER connector terminal marking

Channel	Symbol		Terminal label
	Positive terminal	Negative terminal	
Atrial channel	A+	A–	ATRIUM
Ventricular channel	V+	V–	VENTRICLE

201.7.2.103 Bipolar connectors

When bipolar connectors are used, they shall have keyways that prevent inadvertent polarity reversal.

201.7.2.104 * Battery compartment

The means of access to the battery compartment shall be easily identifiable. The battery compartment shall be clearly and permanently marked with the IEC battery nomenclature, the voltage and type. The battery compartment shall be clearly and permanently marked to show the correct orientation of the battery or batteries.

201.7.4 Marking of controls and instruments

Additional subclauses:

201.7.4.101 * Control or indicator for pacing output

If constant current output is used, the control for selecting pacing output or the relevant indicating means shall be marked in terms of current in milliamperes (mA) through a resistive load of $500\ \Omega \pm 1\%$. If a constant voltage output is used, the pacing output or the relevant indicating means shall be marked in terms of volts (V) across a resistive load of $500\ \Omega \pm 1\%$.

201.7.4.102 * Control or indicator for PULSE RATE

The control for selecting PULSE RATE or the relevant indicating means shall be marked in terms of reciprocal minutes.

201.7.4.103 * Control for selecting pacing mode

If a means of selecting the pacing mode is provided, the ME EQUIPMENT shall indicate, as well as the mode selected, the possible pacing modes using the codes described in Annex C of ISO 14708-2:2019.

201.7.9 ACCOMPANYING DOCUMENTS**201.7.9.2.2 * Warning and safety notices**

Replacement:

The instructions for use shall include all warning and safety notices.

General warnings and safety notices should be placed in a specifically identified section of the instructions for use. A warning or safety notice that applies only to a specific instruction or action should precede the instruction to which it applies.

The instructions for use shall provide the OPERATOR or RESPONSIBLE ORGANIZATION with warnings regarding any significant RISKS of reciprocal interference posed by the presence of the ME EQUIPMENT during specific investigations or treatments.

The instructions for use shall include the following.

- a) * Warnings regarding potential changes in the behaviour of the NON-IMPLANTABLE PULSE GENERATOR caused by ELECTROMAGNETIC DISTURBANCE sources (e.g. communication transmitters in hospitals, emergency transport vehicles, cellular telephones, etc.) and the effects of therapeutic and diagnostic energy sources (e.g. external cardioversion, diathermy, transcutaneous electrical nerve stimulators [TENS] devices, high-frequency surgical equipment, magnetic resonance imaging or similar sources) on the NON-IMPLANTABLE PULSE GENERATOR. This shall include advice on recognizing when the behaviour of the NON-IMPLANTABLE PULSE GENERATOR is being influenced by external ELECTROMAGNETIC DISTURBANCE or electrical energy sources and steps to be taken to avoid interference.
- b) * A warning about the danger of inadvertently introducing LEAKAGE CURRENT into the heart if SUPPLY MAINS-operated equipment is connected to the LEAD system.
- c) * A warning that the PATIENT CABLE shall be connected to the NON-IMPLANTABLE PULSE GENERATOR before the pacing LEADS are connected to the PATIENT CABLE.
- d) * A warning that when handling indwelling LEADS, the terminal pins or exposed metal are not to be touched nor be allowed to contact electrically conductive or wet surfaces.
- e) * A warning regarding the HAZARDS of using PRIMARY BATTERIES other than those recommended by the MANUFACTURER (for example, short BATTERY life after the indication of low BATTERY condition, degraded ME EQUIPMENT performance, overall reduced BATTERY life, and erratic or no pacing).
- f) * A warning that, before handling the NON-IMPLANTABLE PULSE GENERATOR, the PATIENT CABLE or indwelling LEADS, steps shall be taken to equalize the electrostatic potential between the user and the PATIENT, for example by touching the PATIENT at a site remote from the pacing LEADS.
- g) * A caution that, when clinically indicated, supplemental monitoring of the PATIENT should be considered.

201.7.9.2.4 * Electrical power source

Replacement:

The instructions for use shall contain advice on removal of the PRIMARY BATTERY if the ME EQUIPMENT is to be stored or when a long period of disuse is anticipated.

The instructions for use shall state the recommended PRIMARY BATTERY specification.

The instructions for use shall contain the estimated service time from a fully charged BATTERY at 20 °C ambient temperature when operating under specified conditions.

The instructions for use shall contain the estimated service time following activation of the BATTERY DEPLETION INDICATOR when operating under specified conditions.

The instructions for use shall contain the information (including a reference to the appropriate PRIMARY BATTERY specified in IEC 60086-2 [1] ² giving the identity of the PRIMARY BATTERIES to be used.

201.7.9.2.5 * ME EQUIPMENT description

Addition:

The instructions for use shall include the following.

² Numbers in square brackets refer to the Bibliography.

- a) * A general description, explanation of function available, and a description of each heart/PULSE GENERATOR interaction for each available pacing mode. See Clause C.3 of ISO 14708-2:2019 for a description of pacing modes.
- b) * The connector configuration, the geometry and/or dimensions of the receiving connectors and instructions for connecting the LEAD(S) or PATIENT CABLE(S) to the NON-IMPLANTABLE PULSE GENERATOR.
- c) * The electrical characteristics (including tolerances where applicable) with $500\ \Omega \pm 1\%$ load, unless otherwise stated, as follows:
 - ranges of BASIC RATE, ESCAPE INTERVAL, MAXIMUM TRACKING RATE and INTERFERENCE PULSE RATES (as applicable);
 - PULSE AMPLITUDE(S);
 - PULSE DURATION(S);
 - the SENSITIVITY range for both positive and negative polarities (if a sensing function is provided);
 - sensing amplifier blanking period(s) (if a sensing function is provided);
 - the REFRACTORY PERIOD(S) (pacing and sensing) and A-V INTERVAL(S) (as applicable);
 - mode of operation in the presence of sensed ELECTROMAGNETIC DISTURBANCE or electrical energy sources;
 - the rate limit (runaway protection), in reciprocal minutes.
- d) * The electrical characteristics listed below and as reported in 201.7.9.2.5 c) upon activation of the BATTERY DEPLETION INDICATOR, unless these are unchanged from the values previously reported:
 - BASIC RATE or equivalent PULSE INTERVAL;
 - PULSE AMPLITUDES(S);
 - PULSE DURATION(S);
 - SENSITIVITY (if a sensing function is provided);
 - mode change (if applicable).

201.7.9.2.8 * Start-up PROCEDURE

Addition:

The instructions for use shall contain any environmental limitations regarding storing the ME EQUIPMENT immediately prior to use.

201.7.9.2.13 * Maintenance

Addition:

The instructions for use shall contain details for replacing the PRIMARY BATTERY and the means of ascertaining when replacement is required.

The instructions for use shall contain information calling the RESPONSIBLE ORGANIZATION'S attention to the need for periodic maintenance, as well as to the need for maintenance after any malfunction or accident of the ME EQUIPMENT irrespective of usage, especially:

- cleaning and disinfection of reusable PATIENT CABLES;
- cleaning and disinfection of the NON-IMPLANTABLE PULSE GENERATOR;
- inspection of cables and connections for possible defects, for example, loosening of connections and other wear and tear from such causes as PATIENT movement;
- inspection of the NON-IMPLANTABLE PULSE GENERATOR and PATIENT CABLE for signs of physical damage or contamination, in particular damage or contamination that can have a detrimental effect on the electrical isolation properties of the ME EQUIPMENT;

- functional checks, calibration, activation of keys, switches, etc., especially if the ME EQUIPMENT has suffered severe shock, for example, by being dropped.

201.8 Protection against electrical HAZARDS from ME EQUIPMENT

Clause 8 of the general standard applies, except as follows:

201.8.5.5 DEFIBRILLATION-PROOF APPLIED PARTS

201.8.5.5.1 * Defibrillation protection

Replacement:

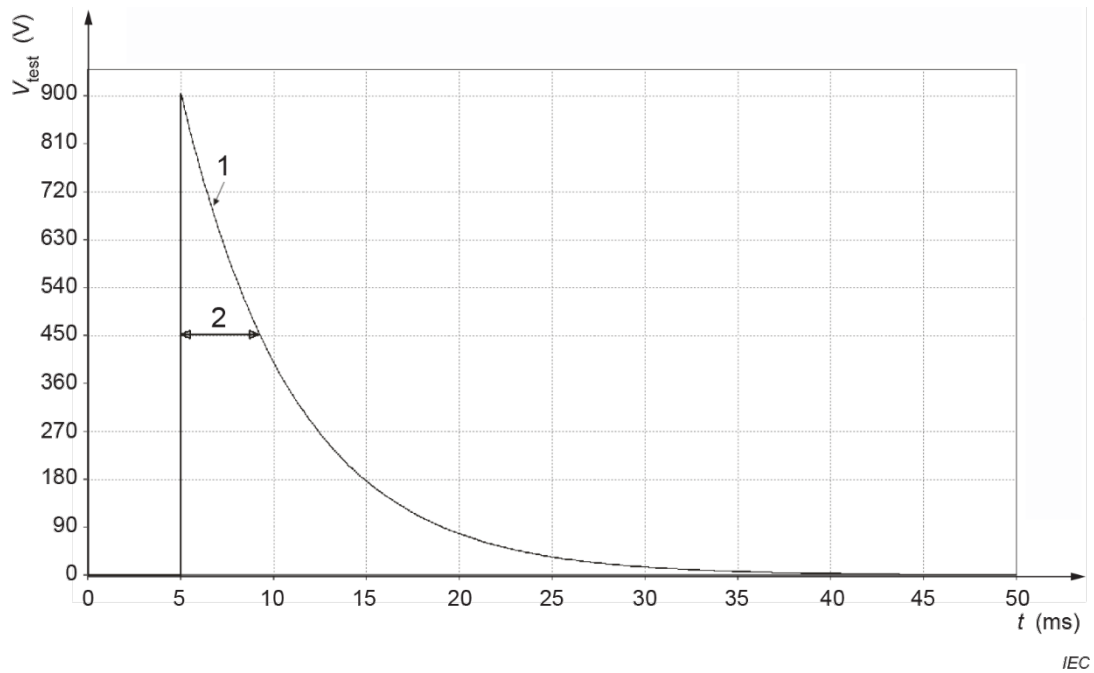
201.8.5.5.1.1 General requirements

The ME EQUIPMENT shall be designed so that defibrillation of the PATIENT using an internal defibrillator will not permanently affect the ME EQUIPMENT, provided that the internal defibrillator electrodes (e.g. paddles) are placed according to the ME EQUIPMENT MANUFACTURER's recommendations.

201.8.5.5.1.2 Test of defibrillation protection

Test equipment: Use a defibrillation test voltage generator providing a decaying exponential waveform presented in Figure 201.101 with the following characteristics: a maximum voltage V_{test} of $900 \text{ V}^{+2,5\%}_{-0}$ and an energy of 50 J to 55 J, having $T_{w50} = 4,05 \text{ ms}$ to $4,6 \text{ ms}$, where T_{w50} is the time interval during which the test voltage is above 50 % of the maximum value V_{test} , if discharged into a load resistance $R_{\text{heart}} = 50 \Omega \pm 1 \%$.

NOTE 1 The resistance $R_{\text{heart}} = 50 \Omega \pm 1 \%$ simulates the resistance of the heart seen by a defibrillator during open heart surgery.



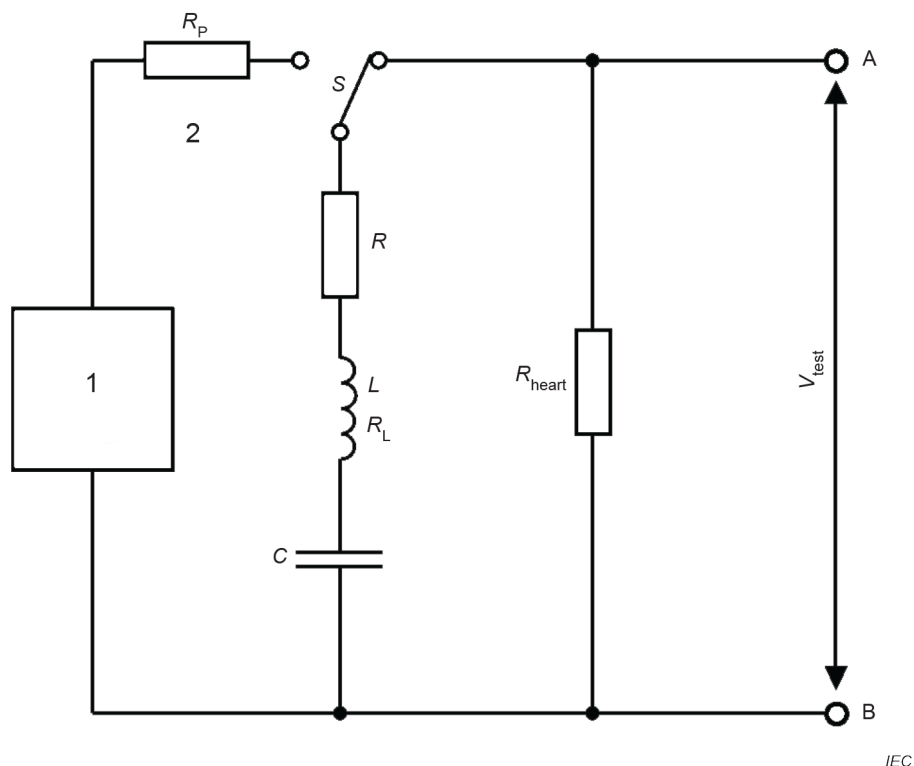
Key

- 1 decaying exponential test waveform
- 2 $T_{w50} = 4,05 \text{ ms to } 4,6 \text{ ms}$: time interval during which the test voltage is above 50 % of the maximum value V_{test}

Figure 201.101 – Test waveform V_{test} implemented by example RCL circuit using $C = 120 \mu\text{F}$, $L = 25 \mu\text{H}$, $RL + R = 1 \Omega$

Figure 201.102 illustrates an example schematic of such a defibrillation test generator with $C = 120 \mu\text{F} \pm 5\%$, $L = 25 \mu\text{H} \pm 5\%$, $R_L + R = 1 \Omega \pm 5\%$, where R_L is the resistance of the inductance L , and R is the output resistance of the defibrillation test voltage generator.

NOTE 2 The current limiting resistor R_p can be used to protect the voltage generator during capacitor charging.

**Key**

- 1 voltage generator
- 2 current limiting resistor
- S switch for applying the test voltage
- L $25 \mu\text{H} \pm 5\%$ inductance
- R_L resistance of inductor
- R resistor in series with L and C
- C $120 \mu\text{F} \pm 5\%$ capacitor
- R_{heart} $50 \Omega \pm 1\%$ load resistance, which simulates the resistance of the heart seen by a defibrillator during open heart surgery
- V_{test} test voltage
- A,B output terminals of the defibrillation test voltage generator

Figure 201.102 – Example circuit of defibrillation test voltage generator for generating a decaying exponential waveform

Test PROCEDURE: Connect the output V_{test} with its terminals A and B to the EXTERNAL PACEMAKER as described below.

The ME EQUIPMENT shall be categorized into one or more of two groups of EXTERNAL PACEMAKERS as appropriate and connected as indicated:

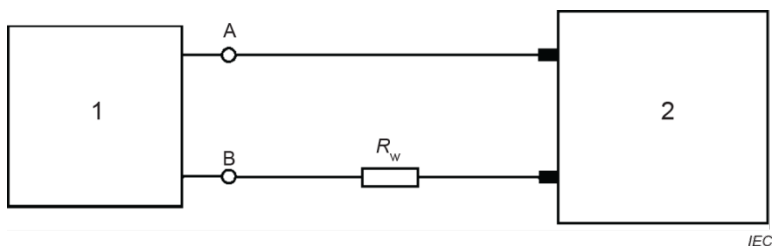
– SINGLE CHAMBER EXTERNAL PACEMAKER:

The test voltage V_{test} is applied between one EXTERNAL PACEMAKER PATIENT CONNECTION connected to the terminal A of the defibrillation test voltage generator and the second EXTERNAL PACEMAKER PATIENT CONNECTION connected via an 80Ω resistor to the terminal B of the defibrillation test voltage generator. The test setup is shown in Figure 201.103.

– Multi-chamber EXTERNAL PACEMAKER:

The test voltage V_{test} is applied to each EXTERNAL PACEMAKER PATIENT CONNECTION, being connected to the terminal A of the defibrillation test voltage generator, in turn with all the remaining EXTERNAL PACEMAKER PATIENT CONNECTIONS being connected together via $80\ \Omega$ resistors to the terminal B of the defibrillation test voltage generator. Figure 201.104 provides the test setup for a DUAL CHAMBER EXTERNAL PACEMAKER, whereas Figure 201.105 shows the test setup for a triple chamber EXTERNAL PACEMAKER (e.g. bi-ventricular).

NOTE 3 The $80\ \Omega$ resistors shown in the test setups below simulate the impedance of applied heart wires and the tissue between the internal defibrillator and the EXTERNAL PACEMAKER terminals.



Key

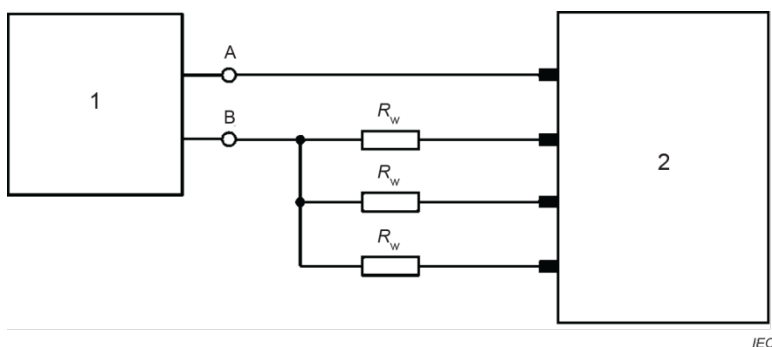
1 defibrillation test voltage generator

2 SINGLE CHAMBER EXTERNAL PACEMAKER

R_w $80\ \Omega \pm 1\%$, resistor simulating the impedance of heart wires and tissue between defibrillator and EXTERNAL PACEMAKER

A, B output terminals of the defibrillation test voltage generator

Figure 201.103 – Test setup for a SINGLE CHAMBER external CARDIAC PACEMAKER



Key

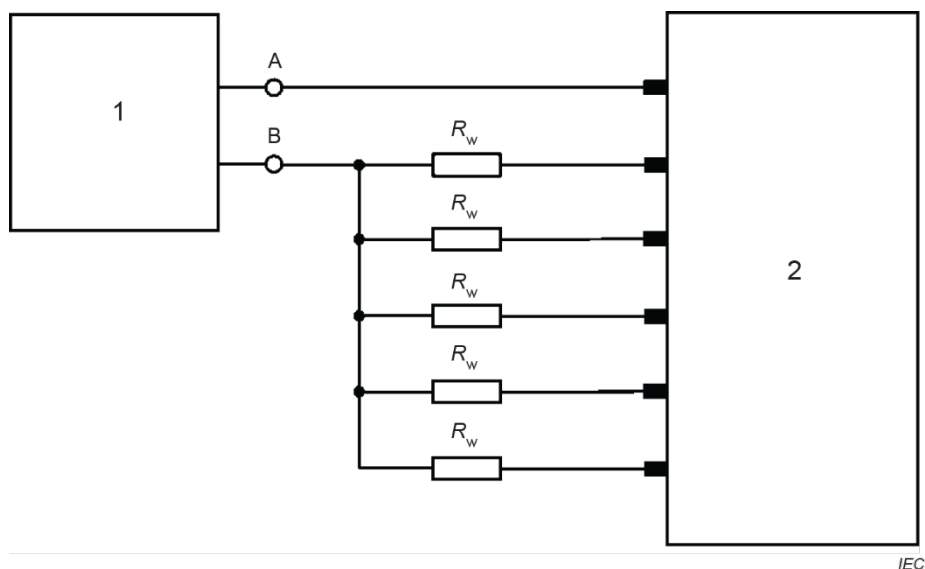
1 defibrillation test voltage generator

2 DUAL CHAMBER EXTERNAL PACEMAKER

R_w $80\ \Omega \pm 1\%$, resistor simulating the impedance of heart wires and tissue between defibrillator and EXTERNAL PACEMAKER

A, B output terminals of the defibrillation test voltage generator

Figure 201.104 – Test setup for a DUAL CHAMBER external CARDIAC PACEMAKER

**Key**

- 1 defibrillation test voltage generator
- 2 multi CHAMBER EXTERNAL PACEMAKER, e.g. bi-ventricular EXTERNAL PACEMAKER providing three pacing CHAMBERS
- R_w $80\ \Omega \pm 1\%$, resistor simulating the impedance of heart wires and tissue between defibrillator and EXTERNAL PACEMAKER
- A, B output terminals of the defibrillation test voltage generator

Figure 201.105 – Test setup for a triple chamber external CARDIAC PACEMAKER, e.g. bi-ventricular external CARDIAC PACEMAKER

Test by applying a sequence of three voltage PULSES of positive polarity at intervals of 20 s to 25 s. Then, after an interval of 60 s (minimum), repeat the test with PULSES of negative polarity (see Figure 201.106).

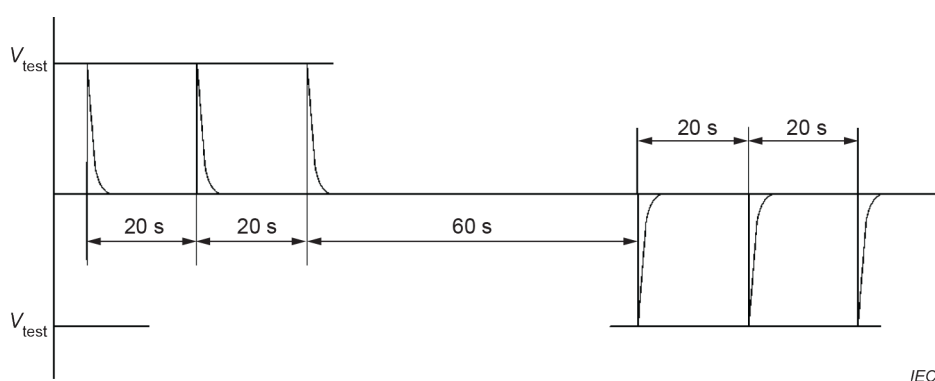


Figure 201.106 – Timing sequence

Compliance shall be confirmed if, after completing the test PROCEDURE, the ME EQUIPMENT continues to provide BASIC SAFETY and ESSENTIAL PERFORMANCE.

201.8.5.5.2 * Energy reduction test

This subclause of the general standard does not apply.

201.8.7.3 * Allowable values

Amendment:

In Table 3 of IEC 60601-1:2005, replace the values for PATIENT AUXILIARY CURRENT for TYPE CF APPLIED PARTS for DC by 1 μ A in NORMAL CONDITION (NC) and 5 μ A in SINGLE FAULT CONDITION (SFC).

201.8.7.4 Measurements

201.8.7.4.1 * General

Addition:

- aa) * The NON-IMPLANTABLE PULSE GENERATOR output should be disabled during LEAKAGE CURRENT testing if possible. If the output is to be active, its contribution should not be considered part of the LEAKAGE CURRENT.

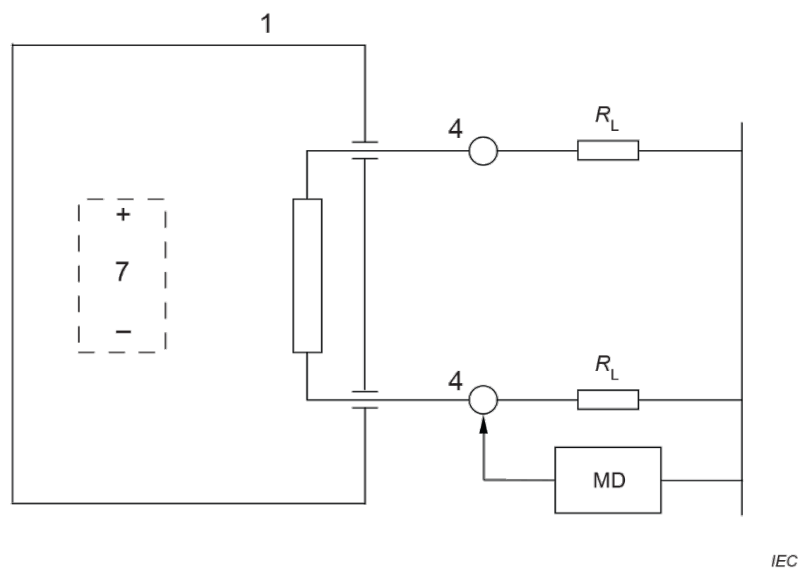
201.8.7.4.8 * Measurement of the PATIENT AUXILIARY CURRENT

Replacement:

For measurement of the PATIENT AUXILIARY CURRENT, the ME EQUIPMENT is connected as shown in Figure 201.107. Each PATIENT CONNECTION is connected to a common bus through a $500\ \Omega \pm 1\%$ load resistor (R_L). Using a measuring device (MD) consisting of a DC voltmeter, resolution better than 2 μ V, fed through a low pass filter with a time constant of at least 10 s, measure the average direct voltage across each load resistor. Steady state condition shall be reached before the measurement is made.

The NON-IMPLANTABLE PULSE GENERATOR shall be set to the nominal settings recommended by the MANUFACTURER (i.e., the factory recommended settings) but with the PULSE AMPLITUDE and PULSE DURATION programmed to the highest available settings.

The low pass filter can be implemented by a four-element RC filter with elements built from 1 M Ω resistors and 10 μ F metalized polypropylene capacitors. The input resistance of the DC voltmeter should be $\geq 400\ \text{M}\Omega$.

**Key**

1	ME EQUIPMENT ENCLOSURE
4	PATIENT CONNECTIONS
7	INTERNAL ELECTRICAL POWER SOURCE
R_L	load resistor
MD	measuring device (see 201.8.7.4.8)

Figure 201.107 – Measuring circuit for the PATIENT AUXILIARY CURRENT for ME EQUIPMENT with an INTERNAL ELECTRICAL POWER SOURCE

Additional subclause:

201.8.101 High-frequency surgical ME EQUIPMENT protection

The ME EQUIPMENT shall comply with 6.1.2 of ISO 14117:2019.

201.9 Protection against MECHANICAL HAZARDS of ME EQUIPMENT and ME SYSTEMS

Clause 9 of the general standard applies.

201.10 Protection against unwanted and excessive radiation HAZARDS

Clause 10 of the general standard applies.

201.11 Protection against excessive temperatures and other HAZARDS

Clause 11 of the general standard applies, except as follows:

201.11.6.5 * Ingress of water or particulate matter into ME EQUIPMENT and ME SYSTEMS

Replacement:

The ME Equipment shall be so constructed that the ingress of liquids (accidental wetting), shall not result in an unacceptable RISK.

Compliance is checked by the following test.

The ME EQUIPMENT is placed in the least favourable position of NORMAL USE with the PATIENT CABLE attached. The ME EQUIPMENT is subjected to a spill of 400 ml of 9 g/l saline solution from a height of 30 cm. The entire 400 ml is poured over the ME EQUIPMENT in less than 5 s. Following the spill, the ME EQUIPMENT is not to be resting in a depth of more than 5 mm of saline solution.

Immediately after 30 s of exposure, the ME EQUIPMENT is removed from the saline solution and visible moisture on the outside of the ENCLOSURE is removed.

The ME EQUIPMENT is to operate within specification during and after the spill.

After at least 24 h have passed, the ME EQUIPMENT is to operate within specification. The ME EQUIPMENT is then disassembled and inspected. Any evidence that liquid has entered the electronic compartment constitutes a failure.

201.11.8 * Interruption of the power supply / SUPPLY MAINS to ME EQUIPMENT

Replacement:

The ME EQUIPMENT shall be equipped with a BATTERY DEPLETION INDICATOR which clearly indicates when the power source is to be replaced.

Compliance is checked by inspection and by functional test.

201.12 Accuracy of controls and instruments and protection against hazardous outputs

Clause 12 of the general standard applies, except as follows:

201.12.1 Accuracy of controls and instruments

Replacement:

201.12.1.101 * ME EQUIPMENT PARAMETERS

The measured values of the ME EQUIPMENT parameters shown in Table 201.103 shall be within the MANUFACTURER'S published tolerance. The MANUFACTURER shall insure that measurement equipment accuracy is sufficient to support the stated tolerances for the parameters being measured within 201.12.1.101 and stated by the MANUFACTURER when measured at PULSE RATE settings of 60 and 120 PULSES per minute with a fully charged BATTERY. If 60 or 120 PULSES per minute are not within the range for PULSE RATE settings for the ME EQUIPMENT, then the test shall be conducted at the minimum or maximum allowable settings.

Compliance is checked by either the appropriate methods described below and in 6.1 of ISO 14708-2:2019, or by any other method provided it can demonstrate an accuracy which is sufficient to support the stated tolerances. In case of dispute, the test described below and in 6.1 of ISO 14708-2:2019 shall apply.

Table 201.103 – ME EQUIPMENT parameters

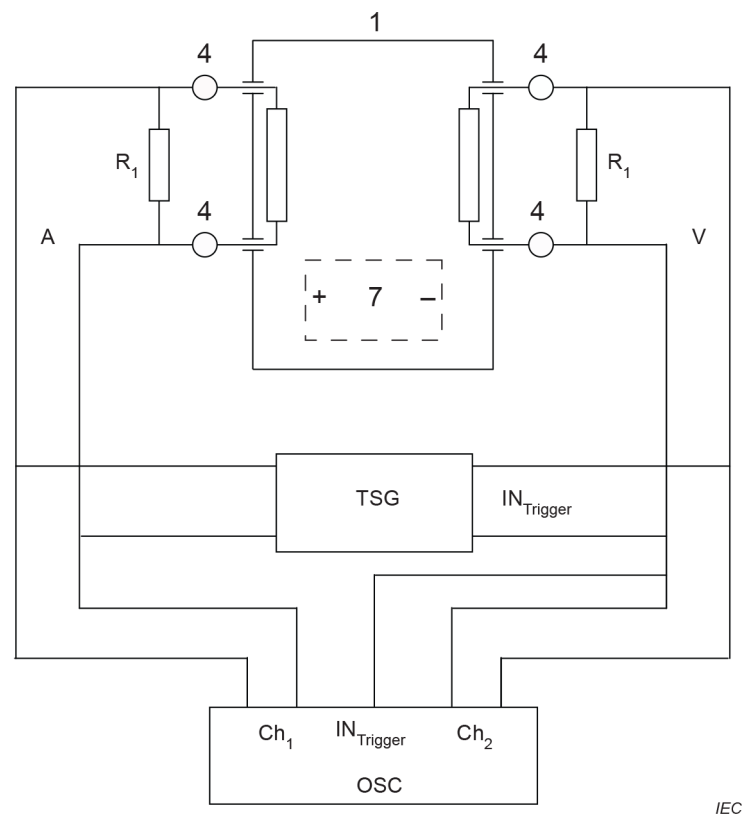
Measurement
PULSE AMPLITUDE
PULSE DURATION
PULSE RATE
SENSITIVITY (if applicable)
ESCAPE INTERVAL
REFRACTORY PERIOD(S) (if applicable)
A-V INTERVAL (if applicable)
MAXIMUM TRACKING RATE (if applicable)

Measurement of MAXIMUM TRACKING RATE is made using the following test.

With a fully charged battery and the NON-IMPLANTABLE PULSE GENERATOR in an A-V sequential mode with sensing and pacing in both chambers (DDD), the ME EQUIPMENT is connected according to Figure 201.108. The test apparatus is described in 6.1.3 of ISO 14708-2:2019. Adjust the signal generator until the amplitude of the test signal is approximately $2e_{pos}$ or $2e_{neg}$ as determined in 6.1.3 of ISO 14708-2:2019.

The delay from the triggering of the signal generator to the production of the test signal is designated D. Adjust the signal generator so that D is slightly greater than the POST-VENTRICULAR ATRIAL REFRACTORY PERIOD (PVARP). Slowly increase D until the ventricular pacing PULSE just begins to track the additional delay as observed on Channel 2 of the oscilloscope. Measure the interval between sequential pacing PULSES on Channel 2 in milliseconds. Designate that as interval T. Adjust the oscilloscope so that the display illustrated in Figure 201.109 is obtained.

Calculate the MAXIMUM TRACKING RATE [PULSES per minute] = $60\,000/T$ [ms].



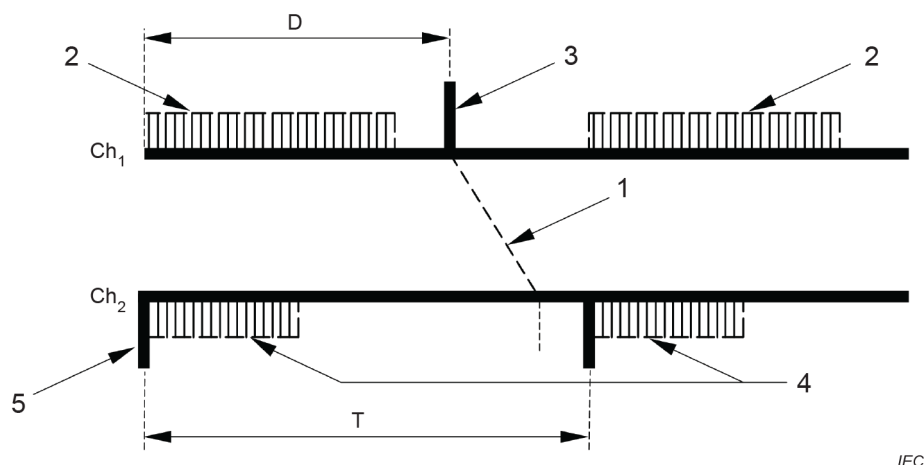
IEC

Key

R_1	$R_1 = 500 \, \Omega \pm 1 \, \%$
A	ATRIAL CHANNEL
V	VENTRICULAR CHANNEL
$IN_{Trigger}$	TRIGGER INPUT
CH_1	Channel 1
CH_2	Channel 2
TSG	TRIGGERABLE SIGNAL GENERATOR
OSC	OSCILLOSCOPE
①	ME EQUIPMENT ENCLOSURE
④	PATIENT CONNECTIONS
⑦	INTERNAL ELECTRICAL POWER SOURCE

See also Table 5 of IEC 60601-1:2005 and IEC 60601-1:2005/AMD1:2012.

Figure 201.108 – Measuring circuit for the MAXIMUM TRACKING RATE

**Key**

①

A-V Interval

②

POST-VENTRICULAR ATRIAL REFRACTORY PERIOD (PVARP)

③

test signal

④

ventricular refractory period

⑤

ventricular pacing PULSE

Ch1 Channel 1

Ch2 Channel 2

 D delay from the triggering of the signal generator to the production of the test signal ($D > \text{PVARP}^1$) T maximum tracking interval¹ Atrial refractory period minus the A-V delay. See [2].MAXIMUM TRACKING RATE [PULSES per minute] = $60\,000/T$ [ms].**Figure 201.109 – Initial oscilloscope display when measuring MAXIMUM TRACKING RATE****201.12.1.102 * PULSE AMPLITUDE**

The PULSE AMPLITUDE expressed either as voltage or current shall not vary from the indicated value by more than the percentage listed in the MANUFACTURER'S published specifications when the load is varied from $200\ \Omega$ to $1\,000\ \Omega$, at a pacing rate of 70 PULSES per minute with a fully charged battery.

Compliance is checked by using the basic test method described in 6.1.2 of ISO 14708-2:2019 with test loads of $200\ \Omega \pm 1\%$ and $1\,000\ \Omega \pm 1\%$ in order to determine how PULSE AMPLITUDE changes as a function of resistance.

201.12.4 Protection against hazardous output

201.12.4.1 * Intentional exceeding of safety limits

Replacement:

If the ME EQUIPMENT incorporates features which require PULSE RATES above the rate limit (see 201.12.4.103), the runaway rate protection may be disarmed when the feature is in use. The means for disarming the runaway rate protection shall require the OPERATOR to engage the activating mechanism continuously.

Compliance is checked by inspection and by a functional test.

Additional subclauses:

201.12.4.101 * Protection against accidental change of controls and tampering

Means shall be provided so that a deliberate action is required to change settings.

Compliance is checked by inspection.

201.12.4.102 * Protection against a low battery condition

Upon activation of the BATTERY DEPLETION INDICATOR, the measured values of the ME EQUIPMENT parameters listed in 201.7.9.2.5 d) shall be within the MANUFACTURER'S published tolerance when measured with the NON-IMPLANTABLE PULSE GENERATOR with $500\ \Omega \pm 1\%$ load.

Compliance is checked by either the appropriate methods described in 6.1 of ISO 14708-2:2019, or by any other method provided it can demonstrate an accuracy sufficient to support the MANUFACTURER'S published tolerances. In case of dispute, the test described in 6.1 of ISO 14708-2:2019 shall apply.

201.12.4.103 * Rate limit (runaway protection)

Means shall be provided to limit PULSE RATE, in the event of a SINGLE FAULT CONDITION, to the value specified by the MANUFACTURER.

Compliance is checked by inspection of the MANUFACTURER'S data.

201.12.4.104 * Interference reversion

In the presence of sensed electrical ELECTROMAGNETIC DISTURBANCE or electrical energy sources, the NON-IMPLANTABLE PULSE GENERATOR shall revert to a pacing mode and PULSE RATE specified by the MANUFACTURER.

Compliance is checked by inspecting the MANUFACTURER'S data.

201.12.4.105 * MAXIMUM TRACKING RATE

In DUAL CHAMBER modes incorporating atrial-synchronous ventricular pacing, a means shall be provided to set a limit at which the ventricle is paced in response to sensed atrial activity. The ME EQUIPMENT shall respond to sensed atrial activity above the MAXIMUM TRACKING RATE in a manner stated by the MANUFACTURER.

Compliance is checked by inspection and functional test.

201.13 HAZARDOUS SITUATIONS and fault conditions for ME EQUIPMENT

Clause 13 of the general standard applies.

201.14 PROGRAMMABLE ELECTRICAL MEDICAL SYSTEMS (PEMS)

Clause 14 of the general standard applies.

201.15 Construction of ME EQUIPMENT

Clause 15 of the general standard applies, except as follows:

Additional subclauses:

201.15.101 * Output indicator

The ME EQUIPMENT shall incorporate a means to indicate when the ME EQUIPMENT has emitted a pacing PULSE.

Compliance is checked by inspection and functional test.

201.15.102 * Input indicator

If a sensing function is provided, the ME EQUIPMENT shall incorporate a means of indicating that the ME EQUIPMENT has detected signals and has responded to them as if they are associated with the electrical activity of the heart, and that it is reacting to the signals as specified by the MANUFACTURER for the selected pacing mode and other operational characteristics.

Compliance is checked by inspection and functional test.

201.16 ME SYSTEMS

Clause 16 of the general standard does not apply.

201.17 Electromagnetic compatibility of ME EQUIPMENT and ME SYSTEMS

Clause 17 of the general standard applies.

202 * ELECTROMAGNETIC DISTURBANCES – Requirements and tests

IEC 60601-1-2:2014 applies except as follows:

202.8.9 * IMMUNITY TEST LEVELS

Replacement:

ME EQUIPMENT shall comply with the requirements of 8.9 of IEC 60601-1-2:2014 except as modified below for ELECTROSTATIC DISCHARGE.

The IMMUNITY TEST LEVELS specified in Table 202.101 shall be used. For this requirement, the following conditions associated with BASIC SAFETY and ESSENTIAL PERFORMANCE shall apply:

- no permanent degradation, or loss of function which is not recoverable shall be observed at any IMMUNITY TEST LEVEL, as a result of

- damage of ME EQUIPMENT (components), or
 - software corruption, or
 - loss of data;
- no inappropriate delivery of energy to the PATIENT shall occur at any IMMUNITY TEST LEVEL;
 - at IMMUNITY TEST LEVELS 1 and 2, the ME EQUIPMENT shall maintain ESSENTIAL PERFORMANCE within the specification limits;
 - at IMMUNITY TEST LEVELS 3 and 4, the temporary degradation or loss of function or performance which requires OPERATOR intervention is acceptable.

Table 202.101 – Static discharge requirements

IMMUNITY TEST LEVEL ^a	Test voltage (air discharge) kV	Test voltage (contact discharge) kV
1	2	2
2	4	4
3	8	6
4	15	8

^a The IMMUNITY test LEVELS for static discharge are defined in Table 1 of IEC 61000-4-2:2008 [3].

Compliance is checked by application of the tests in 8.9 of IEC 60601-1-2:2014, Tables 4, 7 and 8. Evaluate the response of the ME EQUIPMENT or ME SYSTEM during and after these tests in accordance with 8.9 of IEC 60601-1-2:2014 as modified above, considering each discharge individually.

Annexes

The annexes of the general standard apply except as follows:

Annex I

Identification of IMMUNITY pass/fail criteria

Annex I of the general standard does not apply.

Annex AA (informative)

Particular guidance and rationale

AA.1 General guidance

Annex AA explains the reason for the provisions of this particular standard in the IEC 60601 family, as useful background in reviewing, applying and revising the standard.

The rationale is directed towards those familiar with the subject of this particular standard but who have not participated in its development. Where the reason for a requirement is considered self-evident to such persons, reasons are not given. An understanding of the reasons for the main requirements is considered to be essential for proper application of this particular standard. Furthermore, as clinical practice and technology change, changes in this particular standard can be made with an understanding of previous concerns.

- **RISK ANALYSIS**

EXTERNAL PACEMAKERS are used to treat PATIENTS who have symptomatic or acute bradycardia as well as for temporary pacing related to other medical procedures. PATIENT safety is affected by the medical procedure involved, by the understanding of ME EQUIPMENT function by the clinician and by ME EQUIPMENT function. The requirements as specified in this particular standard are considered to provide for an acceptable RISK.

As a basis for establishing safety, an inventory of the RISKS to the PATIENT's safety posed by EXTERNAL PACEMAKERS was developed. The results of that analysis are summarized in Table AA.1. To facilitate the review of this document, a reference to the clause(s) in this particular standard where the action is described has been added to the table.

The tentative conclusion based on clinical experience is that failure to pace is the most probable occurrence of those HAZARDS listed.

Table AA.1 – EXTERNAL PACEMAKER HAZARD inventory

HAZARD	Cause	Action	Reference in this particular standard
FAILURE TO PACE	Low battery	Battery indicator	201.11.8
	Poor connection	Test of connection	201.7.9.2.13
	Threshold rise	Clinical technique	Not applicable
	Oversensing	Clinical technique	Not applicable
	Fault	Stability of parameters with battery and depletion	201.7.9.2.5 d) and 201.12.4.102
		Input indicator	201.15.102
		Defibrillator equipment protection	201.8.5.5.1
		High frequency surgical equipment protection	201.8.101
		Spillage protection	201.11.6.5
		Static electric discharges	202.8.9
		Maintenance	201.7.9.2.13
		Output indicator	201.15.101
	Maladjustment	Protective means	201.12.4.101
	Electrode dislodgement	Marking of controls	201.7.4
		Clinical technique	Not applicable
High rate	Fault	Rate limit (runaway protection)	201.12.4.103
	Maladjustment	Protective means	201.12.4.101
	Temporary high rate	Protective means	201.12.4.1
	Atrial tachyarrhythmia	MAXIMUM TRACKING RATE	201.12.4.105
Unwanted stimulation	Undersensing	Clinical technique	Not applicable
		Stability of parameters with battery depletion	201.7.9.2.5 d) and 201.12.4.102
		Input indicator	201.15.102
	Low battery	Battery indicator	201.11.8
	Maladjustment	Protective means	201.12.4.101
	Noise	Marking of controls	201.7.4
		Noise reversion	201.7.9.2.5 c) and 201.12.4.104
		Warnings	201.7.9.2.2 a)
	Poor connection (LEAD or battery)	Test of connection	201.7.9.2.13
	Fault	See failure to pace	
Micro/Macro shock	LEAKAGE CURRENT	LEAKAGE CURRENT limit	201.6.2 and Clause 201.8
	Injection current	Warning	201.7.9.2.2 a), 201.7.9.2.2 b), 201.7.9.2.2 c), 201.7.9.2.2 d), and 201.7.9.2.2 f)
Tissue/electrode damage	PATIENT AUXILIARY CURRENT	PATIENT AUXILIARY CURRENT limit	201.8.7.3

AA.2 Rationale for particular clauses and subclauses

The following are rationales for specific clauses and subclause in this particular standard, with clause and subclause numbers parallel to those in the body of the document. The numbering is, therefore, not consecutive.