



BSI Standards Publication

Medical electrical system — Input interface for haemodialysis equipment for use of external alarming device

National foreword

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INTERNATIONAL ELECTROTECHNICAL COMMISSION

**MEDICAL ELECTRICAL SYSTEM – INPUT INTERFACE FOR
HAEMODIALYSIS EQUIPMENT FOR USE OF EXTERNAL
ALARMING DEVICE**

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IEC PAS 63023 has been processed by subcommittee 62D: Electromedical equipment, of IEC technical committee 62: Electrical equipment in medical practice

The text of this PAS is based on the following document:

This PAS was approved for publication by the P-members of the committee concerned as indicated in the following document

Draft PAS	Report on voting
62D/1273/PAS	62D/1281/RVD

Following publication of this PAS, which is a pre-standard publication, the technical committee or subcommittee concerned may transform it into an International Standard.

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INTRODUCTION

The effort to standardize an INPUT INTERFACE is based on the request to connect an EXTERNAL ALARMING DEVICE to the HAEMODIALYSIS EQUIPMENT in order to stop the extracorporeal circuit.

This PAS establishes a unique interface for connection of an EXTERNAL ALARMING DEVICE.

In extracorporeal treatment different hazardous situations may occur. One major concern is blood loss to the environment. In 12.4.4.104.1 of IEC 60601-2-16:2012, different technical solutions are given.

The relevant excerpt concerning this issue from 201.12.4.4.104.1 of IEC 60601-2-16:2012 reads as follows:

“a) The HAEMODIALYSIS EQUIPMENT shall include a PROTECTIVE SYSTEM to protect the PATIENT from extracorporeal blood loss to the environment that may cause a HAZARD.

NOTE 1 Today no system has been developed that can totally be relied upon to detect blood loss to the environment. The following recommendation is the best known system to detect blood loss to the environment.

If a PROTECTIVE SYSTEM is utilizing measurement of the VENOUS PRESSURE, the OPERATOR should have at least the possibility to adjust the lower ALARM LIMIT manually as closely as possible to the current measurement value. The single needle treatment mode needs additional measures.

b) The HAEMODIALYSIS EQUIPMENT shall include a PROTECTIVE SYSTEM to protect the PATIENT from extracorporeal blood loss to the environment caused by a rupture or separation in the EXTRACORPOREAL CIRCUIT due to excessive pressure, unless this is prevented by inherent safe design.

NOTE 2 This is not related to separation of the PATIENT CONNECTION or access needle but related to the potential pressure that can be generated by the pump which could cause tubing rupture or joint separation in the EXTRACORPOREAL CIRCUIT.

c) Operation of the PROTECTIVE SYSTEM shall achieve the following safe condition:

- activation of an audible and visual alarm signal (see 208.6.3.1, 208.6.3.3.2, 208.6.3.3.101);
- stoppage of the blood flow to the environment caused by the HAEMODIALYSIS EQUIPMENT, even under SINGLE FAULT CONDITION;
- in the case of HAEMOFILTRATION or HAEMODIAFILTRATION, stoppage of the SUBSTITUTION FLUID flow.”

Concerning Subclause 201.12.4.4.104.1 a) of IEC 60601-2-16:2012

Monitoring of the VENOUS PRESSURE is not always suitable for detecting a blood loss in time, in case the venous puncture cannula slips out. The VENOUS PRESSURE is determined mainly by the hydraulic resistance of the venous puncture cannula, particularly with today's usual high blood flow rates of up to 500 ml/min. Hence a VENOUS PRESSURE ALARM SYSTEM is not always able to detect whether or not the puncture cannula slips out.

If dialysis is performed in the single-needle mode with only one blood pump ("single-needle single pump", "SN click-clack"), the VENOUS PRESSURE measurement is an integral part of the control system. An error in this control system (e.g. pressure sensor stuck to low value) might lead to the upper changeover point of the VENOUS PRESSURE never being reached. As a result, the pressure becomes too high, the tubing system may burst, and the PATIENT may lose a great amount of blood. This may require a PROTECTIVE SYSTEM that is independent of the control system, e.g. monitoring of the phase duration by an independent microprocessor.

Inherent safe design is e.g. a pump rotor that is spring-mounted so smoothly that bursting of the tubing is not possible. However, in this case the HAZARD of haemolysis may exist.

Other measures for prevention of overpressure are holders for the EXTRACORPOREAL CIRCUIT lines and the DIALYSER which make kinking sufficiently unlikely.

Blood loss to the environment caused by disconnections or faults in the EXTRACORPOREAL CIRCUIT cannot be prevented by any PROTECTIVE SYSTEM. The PROTECTIVE SYSTEM should be designed so that blood loss is detected and major blood loss is prevented. Most reported cases of fatal blood loss are caused by blood access cannulas slipping from the fistula or graft. This cannot be prevented by the HAEMODIALYSIS EQUIPMENT. Traditionally, VENOUS PRESSURE monitors have been used for protection of blood loss to the environment. These sensors detect a drop of the pressure in the return bloodline. In case of a bloodline rupture or disconnection of the bloodline from the blood access device (cannula or central venous catheter) the pressure will drop considerably because of the high flow resistance in the blood access device. When the venous cannula slips from a fistula the pressure change is usually too low to be detected by the VENOUS PRESSURE monitor. The pressure drops only by the amount of the fistula pressure, which is typically 5 mmHg – 20 mmHg. To avoid frequent nuisance alarms caused by PATIENT movement the difference between the actual VENOUS PRESSURE and the lower pressure ALARM LIMIT is usually adjusted to 10 mmHg – 20 mmHg.

Monitors employing pressure pulses or other parameters may offer greater sensitivity but may also require up to a minute to detect the fault condition and switch off the blood pump. With high blood flow this may cause blood losses of 500 ml, which are usually not fatal for adults.

The effects of haemorrhage are described by [1]¹.

Concerning Subclause 201.12.4.4.104.1 c) of IEC 60601-2-16:2012

As alarm reaction, the stopping of an occluding blood pump is considered as sufficient. The additional closing of the safety clamp adds only little value, because a rupture will occur most likely at the point of highest pressure, which normally is between the blood pump and the DIALYSER. In this case “retrograde” blood loss via the venous bloodline is negligible compared to the direct blood loss through the arterial bloodline.

If staff is not present (e.g. home PATIENT) or delayed for a long period, in the case of venous puncture cannula slippage, the blood loss from the venous access (backwards) may become hazardous to the PATIENT.

An EXTERNAL ALARMING DEVICE could detect blood loss to the environment from venous needle disconnection. The alarm reaction of the HAEMODIALYSIS EQUIPMENT via the input interface avoids further blood loss.

The solution in this PAS is given by hardware and software with regard to the behaviour of the HAEMODIALYSIS EQUIPMENT in case of activation by an EXTERNAL ALARMING DEVICE.

It is the understanding of the standard committee that initially as little as possible should be standardized. It is hoped that a quasi-standard evolves, which can then be standardized.

The design of the socket and plug is under the responsibility of the manufacturers.

¹ Numbers in square brackets refer to the Bibliography.

MEDICAL ELECTRICAL SYSTEM – INPUT INTERFACE FOR HAEMODIALYSIS EQUIPMENT FOR USE OF EXTERNAL ALARMING DEVICE

1 Scope

This PAS establishes a unique INPUT INTERFACE for connection of an EXTERNAL ALARMING DEVICE to HAEMODIALYSIS EQUIPMENT.

The INPUT INTERFACE of the HAEMODIALYSIS EQUIPMENT is designed as a simple solution, which takes a SINGLE FAULT CONDITION of the INPUT INTERFACE into account, to stop the extracorporeal blood flow in case of needles slipping out from the fistula or graft detected by the EXTERNAL ALARMING DEVICE.

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 60601-2-16:2012, *Medical electrical equipment – Part 2-16: Particular requirements for the basic safety and essential performance of haemodialysis, haemodiafiltration and haemofiltration equipment.*

3 Terms and definitions

For the purposes of this document, the terms and definitions given in IEC 60601-2-16:2012 as well as the following terms and definitions apply:

Additions:

3.1

EXTERNAL ALARMING DEVICE

ACCESSORY that detects ALARM CONDITIONS

3.2

INPUT INTERFACE

part of HAEMODIALYSIS EQUIPMENT providing the possibility of access to an EXTERNAL ALARMING DEVICE.

3.3

INTERNAL SIGNAL PROCESSING

part of HAEMODIALYSIS EQUIPMENT intended to process signals

3.4

SIGNAL PLUG

TERMINAL DEVICE of the EXTERNAL ALARMING DEVICE for the connection to the HAEMODIALYSIS EQUIPMENT SIGNAL SOCKET

3.5

SIGNAL SOCKET

TERMINAL DEVICE of the INPUT INTERFACE

3.6**HAEMODIALYSIS EQUIPMENT GROUND**

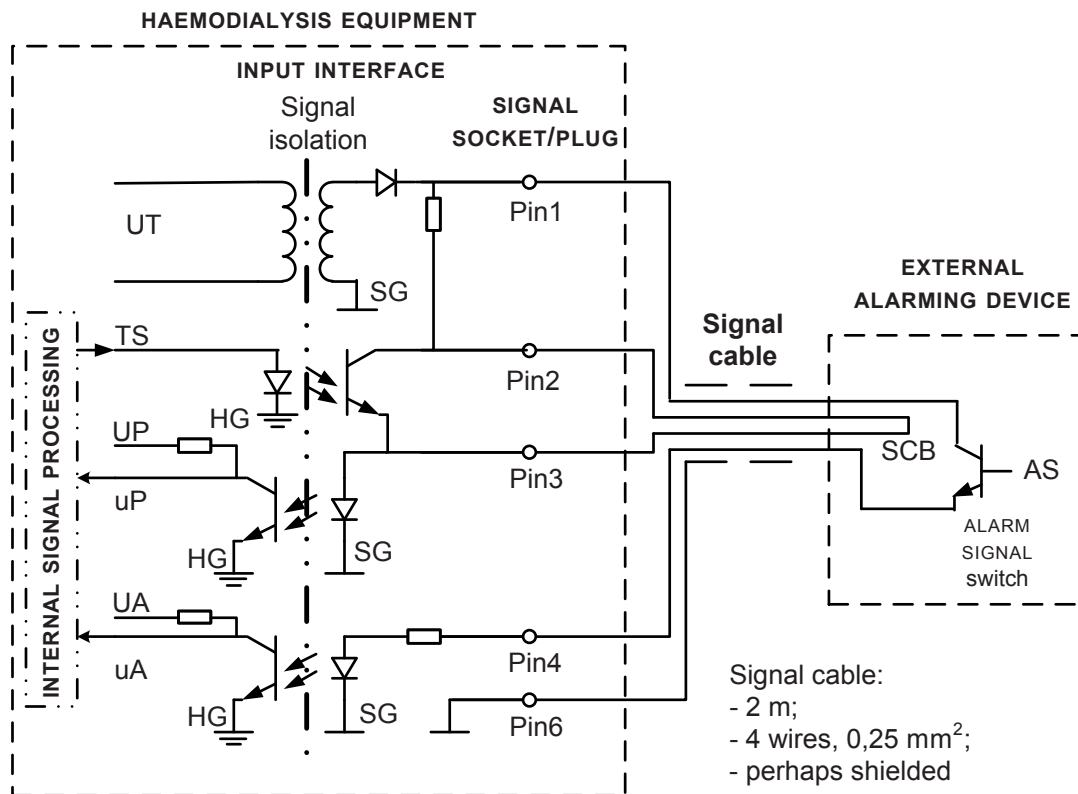
grounding terminal connected to conductive parts for INTERNAL SIGNAL PROCESSING

3.7**SIGNAL GROUND**

grounding terminal connected to conductive parts for external signal processing

4 Application**4.1 Simplified circuit diagram**

The simplified circuit diagram in Figure 1 provides a hardware solution for an INPUT INTERFACE.



UT: Transformer input voltage
 TS: Test signal
 UP: Plug voltage
 uP: SIGNAL PLUG connected
 UA: ALARM SIGNAL voltage
 uA: ALARM SIGNAL
 HG: HAEMODIALYSIS EQUIPMENT GROUND

Pin1: ALARM SIGNAL switch voltage
 Pin2: SIGNAL PLUG connected output
 Pin3: SIGNAL PLUG connected input
 Pin4: EXTERNAL ALARMING DEVICE signal
 Pin6: Signal cable shield
 SG: Isolated SIGNAL GROUND
 SCB: Signal cable bridge

- TS Imp1 Pulse train test signal: No definition
- AS Imp2 Pulse train ALARM SIGNAL: duty cycle (pulse duration/period duration) = 0,33
- AS Imp3 Pulse train No ALARM SIGNAL: Inverted pulse train ALARM SIGNAL
- Pulse time duration ALARM SIGNAL: 0,2 s ± 10 %; duration of one period 0,6 s ± 10 %
 - Switching voltage ALARM SIGNAL switch: minimal 6 V; maximal 30 V
 - Switching power ALARM SIGNAL switch: 200 mW

Figure 1 – Simplified circuit diagram

4.2 Periodic functional check

A failure of the INPUT INTERFACE of the HAEMODIALYSIS EQUIPMENT should become obvious to the OPERATOR at least once per day when in operational use.

The check of the INPUT INTERFACE is done with the test signal (TS). For this purpose, the test signal is set to operation voltage (OV) and the test results are monitored by the INTERNAL SIGNAL PROCESSING unit.

The expected test condition signals and test results are listed in Table 1.

Table 1 – Periodic functional check of the INPUT INTERFACE

Test condition	uP	uA	Test result
Test signal (TS) = operation voltage (OV)	HG	UA	No error
	HG	HG	Error
	HG	Imp2	No error
	HG	Imp3	No error
	Imp1	UA or HG or Imp2 or Imp3	Error
	UP	UA or HG or Imp2 or Imp3	Error

The test results from Table 1 lead to the expected reactions of the HAEMODIALYSIS EQUIPMENT, which are listed in Table 2.

Table 2 – Reaction of HAEMODIALYSIS EQUIPMENT

Test result	Reaction HAEMODIALYSIS EQUIPMENT
No error	Display status: test INPUT INTERFACE passed
Error	Display status: test INPUT INTERFACE failed

4.3 Condition of INPUT INTERFACE

The signals of the connected SIGNAL PLUG (uP) and ALARM SIGNAL (uA) lead to the results listed in Table 3.

Table 3 – Signal result of signal input to INTERNAL SIGNAL PROCESSING unit

uP	uA	Signal result
Imp1	UA	No EXTERNAL ALARMING DEVICE connected
Imp1	HG	INPUT INTERFACE defective
Imp1	Imp2	EXTERNAL ALARMING DEVICE or INPUT INTERFACE defective
Imp1	Imp3	EXTERNAL ALARMING DEVICE or INPUT INTERFACE defective
UP	UA	INPUT INTERFACE defective
UP	HG	INPUT INTERFACE defective
UP	Imp2	INPUT INTERFACE defective
UP	Imp3	INPUT INTERFACE defective
HG	UA	EXTERNAL ALARMING DEVICE or INPUT INTERFACE defective
HG	HG	EXTERNAL ALARMING DEVICE or INPUT INTERFACE defective
HG	Imp2	No ALARM SIGNAL from EXTERNAL ALARMING DEVICE
HG	Imp3	ALARM SIGNAL from EXTERNAL ALARMING DEVICE

4.4 Reaction of HAEMODIALYSIS EQUIPMENT

During the treatment, the signal result from Table 3 leads to the expected reactions of the HAEMODIALYSIS EQUIPMENT, which are listed in Table 4.

Table 4 – Reaction of HAEMODIALYSIS EQUIPMENT during the treatment

Signal result	Reaction HAEMODIALYSIS EQUIPMENT during the treatment
No EXTERNAL ALARMING DEVICE connected	Display status: no EXTERNAL ALARMING DEVICE connected
INPUT INTERFACE defective	<ul style="list-style-type: none"> – Visual ALARM SIGNAL – Audible ALARM SIGNAL – Display status: INPUT INTERFACE defective
EXTERNAL ALARMING DEVICE or INPUT INTERFACE defective	<ul style="list-style-type: none"> – Visual ALARM SIGNAL – Audible ALARM SIGNAL – Display status: EXTERNAL ALARMING DEVICE or INPUT INTERFACE defective
No ALARM SIGNAL from EXTERNAL ALARMING DEVICE	Display status: EXTERNAL ALARMING DEVICE is connected
ALARM SIGNAL from EXTERNAL ALARMING DEVICE	<ul style="list-style-type: none"> – Stoppage of blood flow – Visual ALARM SIGNAL – Audible ALARM SIGNAL – Display status: EXTERNAL ALARMING DEVICE in ALARM CONDITION

Bibliography

Guyton, AC. Circulatory Shock and Physiology of Its Treatment. In Guyton, AC, editor, *Textbook of Medical Physiology*, Eight Edition. W.B. Saunders Company, Philadelphia, 1991: pp 263-71

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