

EU RO Mutual Recognition Technical Requirements

SWITCHES	Version	0.4
	Adoption Date	1 April 2016
	Application Date	1 October 2016
	Tier	1
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1. PRODUCT DESCRIPTION

1.a General description of the product

Description of product according to IEC IEV 441-14-10:

Switch (mechanical): is a mechanical switching device capable of making, carrying and breaking currents under normal circuit conditions which may include specified operating overload conditions and also carrying for a specified time currents under specified abnormal circuit conditions such as those of short circuit.

Note: a switch may be capable of making but not breaking short circuit currents.

1.b Application limitations

- a) Restricted to LV mechanical switches according IEC 60947-3 where the rated voltage does not exceed 1000 V a.c. or 1500 V d.c. and without electronic devices;
- b) Onboard ships with exceptions as per SOLAS Ch. I, Reg. 3;
- c) Installation on board ships within locations with climatic, biological, chemically active, mechanically active and mechanical environmental conditions not exceeding those for which performance has been proved according to IEC 60721-3-6 (1993) + A2 (1997).

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1.c Intended use

- a) Power supply system characteristics as per IEC 60092-101; 2.8.;
- b) Environment category 6K4, 6B2, 6C2, 6S2 and 6M3 according to IEC 60721-3-6 (1993) + A2 (1997).

1.d System context

See 1.c.

2. DESIGN EVALUATION

2.a Engineering evaluation requirements

2.a i. Technical Requirements

- a) Type, ratings and characteristics of switches for intended applications shall be evaluated;
- b) In general, IEC 60947 series shall be observed.

2.a.ii. Technical documents to be submitted

IMPORTANT: The English Language shall be used for all submitted documents.

Prior to tests:

- a) Proposed test program and test schedule;
- b) Description of the test specimens and explanation of the selected test sample(s) providing evidence that the selected sample meets the most rigorous and demanding requirements;
- c) Product descriptions, manuals, data sheets, assembly drawings, dimension drawings, etc. clearly identifying the product.;
- d) Complete accreditation certificate of the Test Laboratory;
- e) Details of production sites;
- f) Product specification;
- g) Application, working area;
- h) Instructions on fitting, assembly and operation;
- i) QM-certificate according to ISO 9001.

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After completion of tests:

- j) The test report with an identification number shall contain all relevant data and test results including place and date of the tests, the names of the responsible personnel carrying out the test;
- k) Type references and serial numbers of the products tested;
- l) Test reports and test records must be signed by the personnel members in charge of the test and are to be confirmed by the EU RO by signing and marking the test report;
- m) Certificate of conformity to IEC Standards issued by one of the Certification Body under the ILAC or LOVAG or IECEE mutual recognition agreement.

2.b Type testing requirements

- a) All type tests according to IEC 60947-3 sub-clause 8.2 and 8.3 to be performed in accredited test laboratories recognized either by ILAC or LOVAG or IECEE. In cases where the tests are conducted at Nationally Accredited Laboratories, the presence of the EU RO surveyor may be omitted†.
- b) Test laboratory shall be authorised to conduct type tests according to the relevant IEC standards;
- c) Test specimens shall be taken from the production line or from stocks†.
- d) Special test according to IEC 60947-3 clause 8 and IEC 60947-1 Annex Q Special tests –Damp heat, salt mist, vibration and shock; (Category D).All tests to be performed on the agreed test samples;

† For further clarification of witnessing of tests and sampling the test specimen(s), refer to paragraphs 6, 7 and 8 of the EU RO "Design Evaluation Scheme" procedure (Appendix V of EU RO Framework Document for the Mutual Recognition of Type Approval found on

<http://www.euromr.org/Guidance%20for%20Mutual%20Recognition>).

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3. PRODUCTION REQUIREMENTS

3.a. General Requirements

- a) Refer to EU RO "Design Evaluation Scheme" procedure (Appendix V of EU RO Framework Document for the Mutual Recognition of Type Approval);
- b) Refer to EU RO "Product Quality Assurance (PQA)" procedure (Appendix VI of EU RO Framework Document for the Mutual Recognition of Type Approval).

3.b. Special Requirements

- a) Routine test according to IEC 60947-3 sub-clause 8.1.3;
- b) Production certification according to ISO 9001 by accredited certification bodies;
- c) QM/QS audit (annual) to be submitted to EU RO for review;
- d) Production of the equipment is limited to those facility listed on EU RO certificate;
- e) Changes to the product will void the EU RO certification;
- f) EU RO shall be granted access to all manufacturing and testing facilities, and to be provided with all the information necessary to performance its duties;
- g) General terms and conditions of the EU RO shall be observed.

4. MARKING REQUIREMENTS

Marking as required by IEC 60947-3 (sub-clause 5.2).

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5. TYPE APPROVAL CERTIFICATE CONTENT

The EU RO MR Type Approval Certificate shall contain the minimum information as defined in the EU RO Framework Document for the Mutual Recognition of Type Approval - see Appendix I EU RO MR Type Approval Certificate Information.

The following information is specifically applicable to products relevant to this technical requirement and shall be included on the relevant EU RO MR Type Approval Certificate:

- a) Technical data according to IEC marking;
- b) Validity according to "EU RO Product Quality Assurance, validity";
- c) RO certificate is void if IEC standards are up-dated or withdrawn;
- d) Reference to approved technical documents;
- e) Application and limitations.

6. APPROVAL DATE AND REVISION NUMBER

Date	Revision	Comment
8 July 2012	0.0	Accepted by Advisory Board
19 September 2012	0.1	Errors corrected
31 January 2014	0.2	Added reference to the EU RO Framework Document for the Mutual Recognition of Type Approval.
31 January 2015	0.3	CRF018 – Revision to par. 2.a.ii - Technical documents to be submitted in English; CRF020 – Revision to par. 5 - 'Type Approval Certificate Content'.
1 April 2016	0.4	CRF025 – Updated to new MR TR document format incl. par. 8; CRF026/026a – Witness testing & control of test specimen; CRF028 – addition of 6 month application clause.

7. BACKGROUND INFORMATION / REFERENCES

- a) IEC 60092-101; 2.8;
- b) IEC 60721-3-6 (1993) + A2 (1997);
- c) IEC 60947;
- d) IEC IEV 441-14-10;
- e) ISO 9001;
- f) SOLAS Ch. 1, Reg. 3;
- g) EU RO Framework Document for the Mutual Recognition of Type Approval.

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8. MAINTENANCE & CLARIFICATION OF TECHNICAL REQUIREMENTS

Anyone wishing to propose changes to this document or request clarification of technical issues should contact the EU RO MR Group Secretariat in the first instance: Secretariat@euomr.org.

Review and approval of change requests shall follow the EU RO MR Maintenance Process detailed in the EU RO Framework Document for the Mutual Recognition of Type Approval: <http://www.euomr.org/Guidance%20for%20Mutual%20Recognition>

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