





Production Quality Assurance Certificate

This is to certify that: Kentec Electronics Limited

Unit 25-26

Fawkes Avenue Questor

Dartford DA1 1JQ United Kingdom

Holds Certificate Number: BSI/MED/PC/755657

In respect of:

MED/3.51

Fixed fire detection and fire alarm systems components for control stations, service spaces, accommodation spaces, cabin balconies, machinery spaces and unattended machinery spaces

(a) Control and Indication equipment

(b) Power Supply Equipment

On the basis that BSI carried out the relevant conformity to type based on quality assurance of the production process for the equipment identified above which was found to be in compliance with the Marine Equipment Directive (MED) 2014/90/EU and Regulation (EU) 2021/1158, subject to any conditions in the schedule attached hereto. The attached schedule of approval forms part of this certificate.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 2797):

Drs. Dave Hagenaars, Managing Director

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PRODUCTS RVA C 646



...making excellence a habit.™

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To check its validity telephone +31 20 3460780. An electronic certificate can be authenticated online.

Production Quality Assurance Certificate

No. BSI/MED/PC/755657

Schedule of Approval

Product Description:

Product	Model	Certificate	Issue	Notified
Designation		No.	Date	Body
MED/3.51 (a) & (b)	Syncro ASM M8000 Series - Analogue Addressable Fire Alarm Control Panel with General Emergency Alarm. Models - MA80161M2, MA80162M2, MH80161M2, MH80162M2	BSI/MED/3.51/755656	29/11/2021	2797

Certificate Amendment Record and BSI internal Review relating to this Certificate

Issue date	Comments	BSI Review No.
November 2021	First issue.	2797:11:3503509

Conditions of Certification

- i) This certificate remains valid unless cancelled or revoked, provided the conditions listed below are complied with and the equipment remains satisfactory in service
- ii) This certificate loses its validity if the manufacturer makes any changes or modifications to the approved quality system, which have not been notified to, and agreed with the notified body named on this certificate and/or after lapse of time, withdrawal or revocation of the Type Examination (Module B) Certificate.
- iii) The equipment detailed above is to be manufactured in accordance with Conformity to Type Based on Quality Assurance of the Production Process (Module D) Certificate of the Marine Equipment Directive.
- iv) If the specified standards are amended during the validity of this certificate, the product type are to be reapproved prior to it being supplied to vessels to which the amended standards apply.
- v) Production tests are to be conducted in accordance with the applicable requirements of the UK Marine Regulation and be recorded by the manufacturer in accordance with the approved Conformity to Type Based on Quality Assurance of the Production Process (Module D) Certificate of the Marine Equipment Directive.

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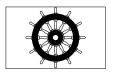
No. BSI/MED/PC/755657

Schedule of Approval

Conditions of Certification continued

- vi) This certificate authorises the manufacturer or his authorised representative established within the Community in conjunction with the EC Type Examination (Module B) Certificate of the equipment listed in the scope to affix the "Mark of Conformity".
- vii) Each equipment is to have the "Mark of Conformity" affixed and be issued with a "Declaration of Conformity".

Example for the Application of the "Mark of Conformity":



2797/YYYY

"Wheelmark" Format yyyy Last four digits of year mark affixed. 2797 Notified Body number undertaking surveillance module.

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