

Annex to declaration of accreditation (scope of accreditation)
Normative document: EN ISO/IEC 17021-1:2015
Registration number: **C 589**

of **DEKRA Certification B.V.**
Product Testing & Certification

This annex is valid from: **25-05-2023** to **01-03-2027** Replaces annex dated: **18-04-2023**
Voluntary withdrawal for the crossed out activities as of 18-04-2023

Location(s) where activities are performed under accreditation

Head Office

Meander 1051
6802 ED
Arnhem
The Netherlands

Location	Certification Scheme
Meander 1051 6802 ED Arnhem The Netherlands	ISO 9001 ISO 13485 Directive 2014/34/EU
1850 Gateway Blvd – Suite 925 Concord, CA, 94520 USA (West)	ISO 9001 ISO 13485 EN ISO 13485
Tzur 8. 4486200 Tzur Yigal Israel	ISO 9001 ISO 13485 EN ISO 13485
K.K., West Wing 7E, 1-28-10 Akebono-Cho, Tachikawa-shi, Tokyo, Japan	ISO 9001 ISO 13485 EN ISO 13485
5F, No. 250, Jiangchangsan Road, Shanghai, 200436, P. R. China	ISO 9001 ISO 13485 EN ISO 13485

This annex has been approved by the Board of the
Dutch Accreditation Council, on its behalf,

J.A.W.M. de Haas

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Standard / Normative document	Certification scheme ¹
ISO 9001	Quality management system for the scopes: (reference to IAF-codes and NACE Rev. 2 where relevant) 12 chemicals, chemical products and fibres 14 rubber and plastic products 17 basic metals and fabricated metal products 18 machinery and equipment 19 electrical and optical equipment 29 wholesale and retail trade, repair of motor vehicles, motorcycles and personal and household goods 34 engineering services 35 other services
ISO 13485	Medical devices — Quality management systems — Requirements for regulatory purposes for the scopes: 1.1 - Non-active medical devices - General non-active, non-implantable medical devices - Non-active implants - Devices for wound care - Non-active dental devices and accessories - Non-active medical devices other than specified above 1.2 - Active (non-implantable) medical devices - General active medical devices - Devices for imaging - Monitoring devices - Devices for radiation therapy and thermotherapy - Active (non-implantable) medical devices other than specified above 1.3 - Active implantable medical devices - General active implantable medical devices - Implantable medical devices other than specified above 1.4 - In Vitro Diagnostic Medical Devices - Reagents and reagent products, calibrators and control materials for: Clinical Chemistry Immunochemistry (Immunology) Haematology/Haemostasis/Immunohematology Microbiology Infectious Immunology Histology/Cytology Genetic Testing - In Vitro Diagnostic Instruments and software - IVD Medical Devices other than specified above 1.5 - Sterilization Methods for Medical Devices - Ethylene oxide gas sterilization (EOG)- including applicable

¹ If no date or version number is mentioned for a normative document, the accreditation concerns the most current version of the document or scheme.

¹ If there is a reference to a code starting with NAW, NAP, EA or IAF, this concerns a scheme mentioned on the [RvA-BR010-list](#).

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Standard / Normative document	Certification scheme ¹
	<ul style="list-style-type: none"> quality management system requirements within ISO 11135 - Moist heat - including applicable quality management system requirements within ISO 17665-1 - Aseptic processing - Radiation sterilization (e.g. gamma, x-ray, electron beam) - including applicable quality management system requirements within ISO 11137-1 -Low temperature steam and formaldehyde sterilization -Thermic sterilization with dry heat -Sterilization with hydrogen peroxide -Sterilization method other than specified above 1.6 - Devices incorporating/utilizing specific substances/technologies <ul style="list-style-type: none"> - Medical devices incorporating medicinal substances - Medical devices utilizing tissues of animal origin - Medical devices incorporating derivatives of human blood - Medical devices utilizing micromechanics - Medical devices utilizing nanomaterials - Medical devices utilizing biological active coatings and/or materials or being wholly or mainly absorbed 1.7 - Parts and Services <ul style="list-style-type: none"> - Raw materials - Components - Subassemblies - Calibrations services* - Distribution services - Maintenance services - Transportation services - Other services <p><small>*organizations providing calibration services should be accredited to ISO/IEC 17025</small></p>

Product / Product Group	Module / article	Conformity assessment procedure
Directive 2014/34/EU Equipment and protective systems intend for use in potentially explosive atmospheres <i>The accreditation for the specified activities is suitable for notification</i>		
Equipment category M-1 of equipment-group I- [electrical equipment] [and] [non-electrical equipment]	Conformity to type based on quality assurance of the production process (module D)	Annex IV

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Product / Product Group	Module / article	Conformity assessment procedure
Equipment category 1 of equipment-group II, explosive atmosphere due to gases, vapours or mists — electrical equipment] [and] [non-electrical equipment]	Conformity to type based on quality assurance of the production process (module D)	Annex IV
Equipment category 1 of equipment-group II, explosive atmosphere due to air/dust mixtures — electrical equipment] [and] [non-electrical equipment]	Conformity to type based on quality assurance of the production process (module D)	Annex IV
Equipment category M2 of equipment-group I — [electrical equipment] [and] [non-electrical equipment]	Conformity to type based on product quality assurance (module E)	Annex VII
Equipment category 2 of equipment-group II, explosive atmosphere due to air/dust mixture — [electrical equipment]]	Conformity to type based on product quality assurance (module E)	Annex VII
Equipment category 2 of equipment-group II, explosive atmosphere due to gases, vapours or mists — electrical equipment]	Conformity to type based on product quality assurance (module E)	Annex VII
Protective systems	Conformity to type based on quality assurance of the production process (module D)	Annex IV
Components	Conformity to type based on quality assurance of the production process (module D)	Annex IV
	Conformity to type based on product quality assurance (module E)	Annex VII