Schedule of Accreditation

issued by

United Kingdom Accreditation Service

2 Pine Trees, Chertsey Lane, Staines-upon-Thames, TW18 3HR, UK



SUMMARY OF ACCREDITED SCOPE

Accredited to provide certification of the following Management Systems Standards and related Sector Schemes as detailed in this schedule:

• Medical Devices - Quality Management Systems (MD-QMS) to ISO 13485: 2016



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DEKRA Certification UK Ltd

Issue No: 005 Issue date: 19 December 2024

KEY LOCATION ADDRESS	SMD-QM
DEKRA Certification UK Ltd	
Stokenchurch House	\checkmark
Oxford Road	

Stokenchurch

This Certification Body has demonstrated to UKAS that it has the systems and processes in place to provide the capability to manage and issue accredited management systems certification only in the country in which it is established for the standards and scopes detailed on this schedule, unless specifically detailed in the individual Management System scope table (denoted by *).



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DEKRA Certification UK Ltd

Issue No: 005 Issue date: 19 December 2024

MEDICAL DEVICES QUALITY MANAGEMENT SYSTEMS In accordance with ISO/IEC 17021-1: 2015 ISO 13485: 2016 Certification **IAF Mandatory Scopes of Accreditation Technical Area** Scope Reference 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 Medical Devices (Non-General active medical devices Implantable) Devices for imaging Monitoring devices Devices for radiation therapy and thermo therapy Active (non-implantable) medical devices other than specified above



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MEDICAL DEVICES QUALITY MANAGEMENT SYSTEMS In accordance with ISO/IEC 17021-1: 2015		
ISO 13485: 2016 Certification		
1.4 In Vitro Diagnostic Medical Devices (IVD)	Reagents and reagent products, calibrators and control materials	
	for:	
	Clinical Chemistry	
	Immunochemistry (immunology)	
	Haematology /Haemostasis /Immunohematology	
	Microbiology	
	Infection Immunology	
	Histology/cytology	
	Genetic Testing	
	In vitro diagnostic instruments and software	
	IVD medical devices other than specified above	

1.5 Sterilisation Method for Medical Devices	Ethylene oxide gas sterilisation (EOG)
	Moist heat
	Aseptic processing
	Radiation sterilisation (e.g. gamma, x-ray, electron beam)
	Low temperature steam and formaldehyde sterilisation
	Thermic sterilisation with dry heat



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MEDICAL DEVICES QUALITY MANAGEMENT SYSTEMS In accordarue with ISO/IEC 17021-1: 2015 ISO 13485: 2016 Certification Sterilisation with hydrogen peroxide Sterilisation with hydrogen peroxide Sterilisation method other than specified above 1.6 Devices Incorporating/Utilising Specific Substances/Technologies Medical devices incorporating medicinal substances Medical devices utilising tissues of animal origin Medical devices incorporating derivatives of human blood Medical devices utilising micromechanics Medical devices utilising micromechanics Medical devices utilising biological active coatings and/or materials being wholly or mainly absorbed Medical devices utilising biological active coatings and/or materials being wholly or mainly absorbed END