Schweizerische Eidgenossenschaft
Confédération suisse
Confederazione Svizzera
Confederaziun svizra

Swiss Confederation

Federal Department of Economic Affairs,
Education and Research EAER

State Secretariat for Economic Affairs SECO
Swiss Accreditation Service SAS

SCESm Directory

Accreditation number: SCESm 0047

International standard:	ISO/IEC 17021-1:2015
Swiss standard:	SN EN ISO/IEC 17021-1:2015

QS Zürich AG
Postfach 6335
8050 Zürich

QS Zürich AG
Branch Basel
Erlenstrasse 31
4106 Therwil

Head:	Lukas Beljean
Responsible for MS:	Lukas Beljean
Telephone:	+41 44 35 04 665
E-Mail:	qs-zuerich@quality-service.ch
Internet:	www.quality-service.ch
Initial accreditation:	24.03.1998
Current accreditation:	24.03.2018 to 23.03.2020
Scope of accreditation see:	www.sas.admin.ch (Accredited bodies)

Scope of accreditation as of 17.06.2019

Certification body for management systems in the domain of quality, environment and security

Standards	Approved technical scopes	Remarks
ISO 9001:2015 ISO 14001:2015		
		IAF Code
	Mining and quarrying	2
	Food products, beverages and tobacco	3
	Textiles and textile products	4 (only for ISO 9001:2015)
	Publishing companies	8
	Printing companies	9
	Chemicals, chemical products and fibres	12
	Pharmaceuticals	13
	Rubber and plastic products	14
	Concrete, cement, lime, plaster etc.	16

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Standards	Approved technical scopes	Remarks
	Basic metals and fabricated metal products	17
	Machinery and equipment	18
	Electrical and optical equipment	19
	Other transport equipment	22
	Manufacturing not elsewhere classified	23
	Recycling	24
	Water supply	27
	Construction	28
	Wholesale and retail trade; repair of motor vehicles, motorcycles and personal and household goods	29
	Hotels and restaurants	30
	Transport, storage and communication	31
	Financial intrmediation, real estate, renting	32
	Information technology	33
	Engineering services	34
	Other services	35
	Public administratioin	36
	Education	37
	Health and social work	38
	Other social services	39
ISO 45001:2018 BS OHSAS 18001:2007	Occupational health and safety management systems	Already granted certificates according to the standard BS OHSAS 18001:2007 keep their validity until 11.03.2021, at the latest.
		IAF Code
	Food products, beverages and tobacco	3
	Publishing companies	8
	Printing companies	9
	Chemicals, chemical products and fibres	12
	Pharmaceuticals	13

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Standards	Approved technical scopes	Remarks
	Rubber and plastic products	14
	Concrete, cement, lime, plaster etc.	16
	Basic metals and fabricated metal products	17
	Machinery and equipment	18
	Electrical and optical equipment	19
	Other transport equipment	22
	Manufacturing not elsewhere classified	23
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	Other services	35
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	Other social services	39
Management system certification (all international standards of the series ISO 13485)		As a general rule, the following applies to certification in the field of medical devices: • Only certification according to the norms ISO 13485 without requirements to MDO, 93/42/ EEC and (EU) 2017/745. • The norm SN EN ISO 13485: 2012 is valid until at least 27.02.2019.

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Standards	Approved technical scopes	Remarks
SN EN ISO 13485:2012 SN EN ISO 13485:2016	**MD 0000 Non Active Medical Devices**	
SN EN ISO 13485:2012 SN EN ISO 13485:2016	**MD 0100 General non-active, non-implantable medical devices** - MD 0101 Non-active devices for anaesthesia emergency and intensive care - MD 0102 Non-active devices for injection, infusion, transfusion and dialysis - MD 0103 Non-active orthopaedic and rehabilitation devices - MD 0104 Non-active medical devices with measuring function, reusable instruments - MD 0105 Non-active ophthalmologic devices - MD 0106 Non-active instruments - MD 0107 Contraceptive medical devices, single-use medical devices - MD 0108 Non-active medical devices for disinfecting, cleaning, rinsing - MD 0110 Non-active medical devices for ingestion	
SN EN ISO 13485:2012 SN EN ISO 13485:2016	**MD 0200 Non-active implants** - MD 0202 Non-active orthopaedic implants - MD 0203 Non-active functional implants	
SN EN ISO 13485:2012 SN EN ISO 13485:2016	**MD 0300 Devices for wound care** - MD 0301 Bandages and wound dressings - MD 0302 Suture material and clamps - MD 0303 Other medical devices for wound care	

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Standards	Approved technical scopes	Remarks
SN EN ISO 13485:2012 SN EN ISO 13485:2016	**MD 0400 Non-active dental devices and accessoires** - MD 0401 Non-active dental equipment and instruments - MD 0402 Dental materials - MD 0403 Dental implants	
SN EN ISO 13485:2012 SN EN ISO 13485:2016	**MD 1000 Active Medical Devices**	
SN EN ISO 13485:2012 SN EN ISO 13485:2016	**MD 1100 General active medical devices** - MD 1106 Active dental devices - MD 1109 Active devices for patient positioning and transport	
SN EN ISO 13485:2012 SN EN ISO 13485:2016	**MD 1200 Devices for imaging** - MD 1202 Imaging devices utilising non-ionizing radiation	
SN EN ISO 13485:2012 SN EN ISO 13485:2016	**MD 1300 Monitoring devices** - MD 1301 Monitoring devices of non-vital physiological parameters	
SN EN ISO 13485:2012 SN EN ISO 13485:2016	**MDS 7000 Specifics of Medical Devices** - MDS 7001 Medical devices incorporating medicinal substances, according to Directive 2001/83/EC - MDS 7004 Medical devices referencing the Directive 2006/42/EC on machinery - -MDS 7006 Medical devices in sterile condition - -MDS 7009 Medical devices utilizing biological active coatings and/or materials or being wholly or mainly absorbed	
SN EN ISO 3834-2:2006 (ISO 3834-2:2005)	Quality requirements for welding, Comprehensive quality requirements	Combined with a certification based on ISO 9001 (Replaces EN 729-2)

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Standards	Approved technical scopes	Remarks
ISO 22000:2005	Food safety management systems Cluster / categories 2 - 6	The certification body fulfils the requirements of ISO/TS 22003:2013 for the sectors
	2. Food and Feed Processing (C + D)	C Food Manufacturing D Animal Feed Production
	3. Catering (E)	E Catering
	4. Retail, transport and storage (F + G)	F Distribution G Provision of Transport and Storage Services
	5. Auxiliary Services (H + I + J)	H Services I Production of Food Packaging and Packaging Materials J Equipment Manufacturing
ISO 50001:2011	Energy Management Systems - Industry – light to medium - Industry – heavy - Mining - Energy supply - Buildings - Building complexes - Transport - Agriculture	The certification body fulfils the requirements of the standard ISO 50003:2014

Abbreviation	Signification
IAF Code	IAF ID1:2014
(EU) 2017/745	Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC; 5 May 2017
EnMS	Energy Management System
MDO	Swiss Medical Device Ordinance, SR 812.213
93/42/EEC	Council directive 93/42/EEC of 14 June 1993 concerning medical devices

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