Certificate

The SQS herewith attests that the organisation named below has a management system that meets the requirements of the normative basis mentioned.

ruetschi

Ruetschi Technology AG
Fabrikstrasse 35
3286 Muntelier
Switzerland

Further sites according to appendix

Scope

Design, development, manufacturing and sale of medical devices, plastic components and precision parts
Assembling and packaging of non-sterile and sterile medical devices under clean room conditions

Normative base

EN ISO 13485:2016 Medical devices – Quality Management System

Reg. no. H60601

Validity 01.12.2021 – 30.11.2024
Issue 01.12.2021

A. Grisard, President SQS

F. Müller, CEO SQS

Swiss Association for Quality and Management Systems (SQS)
Bernstrasse 103, 3052 Zollikofen, Switzerland

sqs.ch

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Appendix

Appendix of main certificate Reg. no. H60601

Ruetschi Technology AG
Fabrikstrasse 35
3286 Muntelier
Switzerland

Central Function	*Scope*	*Norm / Revision*	*Reg. no.*	*Validity*
Ruetschi Technology AG Fabrikstrasse 35 3286 Muntelier Switzerland	Design, development, manufacturing and sale of medical devices, plastic components and precision parts Assembling and packaging of non-sterile and sterile medical devices under clean room conditions	EN ISO 13485:2016	H60601	01.12.2021 30.11.2024

Locations	*Scope*	*Norm / Revision*	*Reg. no.*	*Validity*
Ruetschi Technology AG Länggasse 13 3280 Murten Switzerland	Design, development, manufacturing and sale of medical devices, plastic components and precision parts Assembling and packaging of non-sterile and sterile medical devices under clean room conditions	EN ISO 13485:2016	H60601	01.12.2021 30.11.2024
Ruetschi Technology SA Rue des Prés-du-Lac 63 1400 Yverdon-les-Bains Switzerland	Design, development, manufacturing and sale of medical devices, plastic components and precision parts Development and manufacturing of injection molds Assembling and packaging of non-sterile and sterile medical devices under clean room conditions	EN ISO 13485:2016	H60601	01.12.2021 30.11.2024

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