



CERTIFICATE



This is to certify that the company

Swiss Bionic Solutions Schweiz GmbH

Schulhausstrasse 17
8834 Schindellegi
Switzerland

with the organizational units/sites as listed in the annex

has implemented and maintains a **Quality Management System**.

Scope of certificate and applicable country-specific requirements:

Design and development, manufacturing, distribution and service of pulsed electromagnetic field therapy devices for relief of muscular aches and pains and to increase blood circulation.

-CND, USA (a,b,c,d)

Through an audit, documented in a report, performed by DQS Medizinprodukte GmbH, it was verified that the management system fulfills the requirements of the following standard:

ISO 13485 : 2016 (MDSAP Audit Model Edition 2)

including country-specific requirements as shown in the scope
(full references are listed in the annex)

Certificate registration no.	516451 MDSAP16
Certificate unique ID	170719492
Effective date	2018-12-07
Expiry date	2021-12-06
Frankfurt am Main	2018-12-07



DQS Medizinprodukte GmbH

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DQS Medizinprodukte GmbH is authorised under the Medical Devices Single Audit Program.

Visit <https://www.mydqs.com/en/customers/customer-database.html> to validate this certificate.



Annex to certificate

Certificate registration No.: 516451 MDSAP16

Certificate unique ID: 170719492

Effective date: 2018-12-07

Swiss Bionic Solutions Schweiz GmbH

Schulhausstrasse 17

8834 Schindellegi

Switzerland

Audited site

DUNS No., site scope and country-specific requirements

Swiss Bionic Solutions Schweiz GmbH

Schulhausstrasse 17

8834 Schindellegi

Switzerland

Design and development, manufacturing and distribution of pulsed electromagnetic field therapy devices for relief of muscular aches and pains and to increase blood circulation.

-CND, USA (a,b,c,d)

DUNS No.: 485825694

Swiss Bionic Solutions Schweiz GmbH

Gewerbecenter Gläntern 1,3,5

8864 Reichenburg

Switzerland

Manufacturing, distribution and service of pulsed electromagnetic field therapy devices for relief of muscular aches and pains and to increase blood circulation.

-CND, USA (a,b,c,d)

DUNS No.:480241315



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Swiss Bionic Solutions Schweiz GmbH

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Full references of country-specific requirements of MDSAP participating Regulatory Authorities

Abbreviation	Jurisdiction	Reference
AUS	Australia	(a) Therapeutic Goods (Medical Devices) Regulations 2002, Schedule 3, Part 1 – Full Quality Assurance Procedure (b) Therapeutic Goods (Medical Devices) Regulations 2002, Schedule 3, Part 4 – Production Quality Assurance Procedure
BRA	Brazil	RDC ANVISA n. 16/2013 RDC ANVISA n. 23/2012 RDC ANVISA n. 67/2009
CND	Canada	Medical Device Regulations SOR/98-282, Part 1
JPN	Japan	MHLW Ministerial Ordinance No. 169 (2004) as amended by MHLW Ordinance No. 128 (2014), Articles 4 to 68 Japan PMD Act (as applicable)
USA	United States	(a) 21 CFR Part 803 (b) 21 CFR Part 806 (c) 21 CFR Part 807 (d) 21 CFR Part 820 (e) 21 CFR Part 821