



# CERTIFICATE



This is to certify that the company

## METOXIT AG

Emdwiesenstrasse 6  
8240 Thayngen  
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, development and production of ceramic blanks and corresponding dyeing liquids for the production of dental-prosthetic restorations.

**-CND, JPN, 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.	066501 MDSAP16
Certificate unique ID	170706341
Effective date	2018-10-16
Expiry date	2021-10-15
Frankfurt am Main	2018-10-16



### DQS Medizinprodukte GmbH

Sigrid Uhlemann  
Managing Director

Szymon Kurdyn  
Product Manager

August-Schanz-Straße 21, 60433 Frankfurt am Main,  
Tel. +49 (0) 69 95427-300, [medical.devices@dqs-med.de](mailto:medical.devices@dqs-med.de)

**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.: 066501 MDSAP16**  
**Certificate unique ID: 170706341**  
**Effective date: 2018-10-16**



## **METOXIT AG**

Emdwiesenstrasse 6  
8240 Thayngen  
Switzerland

### **Audited site**

**METOXIT AG**  
Emdwiesenstrasse 6  
8240 Thayngen  
Switzerland

### **DUNS No., site scope and country-specific requirements**

Design, development and production of ceramic  
blanks and corresponding dyeing liquids for the  
production of dental-prosthetic restorations.  
**-CND, JPN, USA (a,b,c,d)**  
**DUNS No.: 481080828**



**Annex to certificate**  
**Certificate registration No.: 066501 MDSAP16**  
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### **Full references of country-specific requirements of MDSAP participating Regulatory Authorities**

<b>Abbreviation</b>	<b>Jurisdiction</b>	<b>Reference</b>
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