ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

MDSAP
MEDICAL DEVICE SINGLE AUDIT PROGRAM

TÜV SÜD America

CERTIFICATE

No. QS6 033098 0047 Rev. 02

Certificate Holder: **Kettenbach GmbH & Co. KG**
Im Heerfeld 7
35713 Eschenburg
GERMANY

Certification Mark:

Scope of Certificate: **Design and Development, Production and Distribution of Dental Products (Denture Soft Liner, Dental Impression Materials, Bite Registration Materials, Dental Restoration Materials for Crowns and Bridges, Dual-Curing Build Up Materials); Distribution of Dental Products (Mixing Units for Dental Products (Non-Active Medical Devices), Adhesives for Impression Materials, Impression Trays)**

Standard(s): **ISO 13485:2016**

Regulatory Authority(ies): **Australia TGA, Brazil ANVISA, Health Canada, USA FDA, MHLW / PMDA. See attached for listing of specific regulatory requirements.**

The Certification Body of TÜV SÜD America Inc. certifies that the quality management system of the manufacturer listed above has been audited against the stated criteria and found to conform to those criteria for the scope of certification listed. Validity of this certificate can be obtained by visiting the website www.tuvsud.com/ps-cert
TÜV SÜD America Inc. is an MDSAP Recognized Auditing Organization.

REPs Facility ID: **F001258**

Effective Date: **2022-02-23**

Expiry Date: **2025-01-28**

Date of Issue: 2022-02-25

(Michael Ogunleye)
Manager, US Certification Body,
Medical and Health Services

TÜV®

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

MDSAP
MEDICAL DEVICE SINGLE AUDIT PROGRAM

TÜV SÜD America

C E R T I F I C A T E

No. QS6 033098 0047 Rev. 02

Regulatory Requirements: **Audit/Certification Criteria**

Australia
Therapeutic Goods (Medical Devices) Regulations 2002
- Schedule 3, Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure

Brazil
- RDC ANVISA n. 16/2013
- RDC ANVISA n. 23/2012
- RDC ANVISA n. 67/2009

Canada
- Medical Device Regulations – Part 1- SOR 98/282

Japan
- MHLW Ministerial Ordinance 169, Article 4 to Article 68
- PMD Act

United States
- 21 CFR Part 803
- 21 CFR Part 806
- 21 CFR Part 807 – Subparts A to D
- 21 CFR Part 820

Facility(ies): Kettenbach GmbH & Co. KG
Im Heerfeld 7, 35713 Eschenburg, GERMANY

Facility Scopes: Design and Development, Production and Distribution of Dental Products (Denture Soft Liner, Dental Impression Materials, Bite Registration Materials, Dental Restoration Materials for Crowns and Bridges, Dual-Curing Build Up Materials); Distribution of Dental Products (Mixing Units for Dental Products (Non-Active Medical Devices), Adhesives for Impression Materials, Impression Trays)
REPs Facility ID: F001258

Date of Issue: 2022-02-25

(Michael Ogunleye)
Manager, US Certification Body,
Medical and Health Services

TÜV®

TÜV SÜD America, Inc. • 401 Edgewater Place Suite #500 • Wakefield • MA 01880 • USA • www.tuvsud.com