

Certificate

Certificate No.: MD 1676601 84938748-30

Manufacturer: **ChM Sp. z o.o.**
Lewickie 3b
16-061 Juchnowiec Kościelny
Poland

D-U-N-S No.: 367520157

Certification criteria ISO 13485:2016
Brazil RDC ANVISA n. 16/2013, RDC ANVISA n. 23/2012, RDC ANVISA n. 67/2009
Canada Medical Devices Regulations – Part 1 – SOR 98/282
United States 21 CFR 820, 21 CFR 821, 21 CFR 803, 21 CFR 806, 21 CFR 807 – Subparts A to D

Scope: Design and development, manufacture and distribution of non-sterile orthopaedic implants, non-sterile medical drivers, non-sterile attachments and non-active non-sterile surgical instruments in the area of orthopaedics.

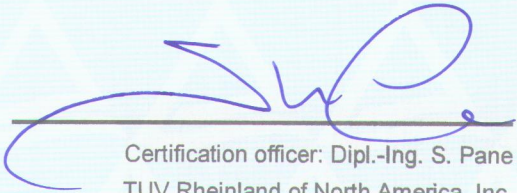
TUV Rheinland of North America, Inc., an MDSAP recognized Auditing Organization, certifies that the quality management system of the Manufacturer has been audited against and found to conform the Certification criteria for the Scope contained in this certificate. The quality management system is subject to annual surveillance audit(s).

Project No.: 26300273 012

Issue Date: 2019-10-10

Effective Date: 2019-10-10

Expiry Date: 2022-04-17



Certification officer: Dipl.-Ing. S. Pane
TUV Rheinland of North America, Inc.

The validity of the certificate can be verified on www.certipedia.com, via the QR code or calling 1-888-743-4652.