Certificate

TÜVRheinland®

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

MDSAP
MEDICAL DEVICE SINGLE AUDIT PROGRAM

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.

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