

EU DECLARATION OF CONFORMITY



Name of product: MATRIX MTA+

Manufacturer:

Przedsiębiorstwo Produkcyjno-Handlowe CERKAMED Wojciech Pawłowski
37-450 Stalowa Wola, ul. Kwiatkowskiego 1, POLAND

SRN (Single Registration Number):

PL-MF-000003211

Purpose and range of use:

MTA + dosing block is designated for forming precise doses of MTA+ product for the tooth cavity or root canal.

Medical device of class I, according to the rule 1 of Annex VIII MDR (EU) 2017/745.
Evaluation of conformity was conducted following the procedure relating to Annex II and III Regulation (EU) 2017/745.

BASIC UDI-DI:

590755302MATRIXJC

Harmonised standards:

EN ISO 14971:2012

Medical devices — Application of risk management to medical devices

EN 1041:2008+A1:2013

Information provided by the manufacturer of medical devices.

EN ISO 15223-1:2016

Medical devices — Symbols to be used with medical device labels, labelling and information to be supplied-Part 1: General requirements

Reference documents:

- Regulation (EU) 2017/745 of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC
- The Act of May 20, 2010 about medical devices with a changes

We declare with full responsibility that the manufactured product, which this statement refers to, complies with the reference documents mentioned above.

This declaration of conformity is issued under the sole responsibility of manufacturer.

Honorata Sołowiej,
Person responsible for regulatory compliance
On behalf of Wojciech Pawłowski,
Stalowa Wola

PRZEDSIĘBIORSTWO PRODUKCYJNO-HANDLOWE

CERKAMED

WOJCIECH PAWŁOWSKI

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signature, company stamp, date

28.12.2021

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H. Sołowiej