



META BIOMED CO., LTD.

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EC Declaration of Conformity

**according to the Medical Devices Directive 93/42/EEC
as amended by Directive 2007/47/EC**

For the following medical device :

Product Name : Calcium Hydroxide Temporary Filling Material

Brand Name: Metapaste (detailed model names attached)

Classification : Class IIa by Rule 7 of Annex IX, Council Directive 93/42/EEC as amended by Directive 2007/47/EC

Manufacturer's Name : META BIOMED CO., LTD.

Manufacturer's Address : 270, Osongsaengmyeong1-ro, Osong-eup, Heungdeok-gu,
Cheongju-si, Chungcheongbuk-do, Korea

is herewith the conformity with the requirements set out in the Council Directive 93/42/EEC of 14 June 1993 as amended by Directive 2007/47/EC of the European Parliament and of the Council of 5 September 2007 concerning Medical Devices declared, For the evaluation of the conformity with this Directive, the following standards have been applied:

MEDDEV 2.12-1 Rev.8, MEDDEV 2.7.1 Rev.4, MEDDEV 2.12.2 Rev.2, Directive 93/42/EEC amended by Directive 2007/47/EC, FDA 21 CFR 820, EN ISO 6876:2012, EN ISO 10993-1:2009, EN ISO 10993-3:2014, EN ISO 10993-5:2009, EN ISO 10993-6:2009, EN ISO 10993-10:2023, EN ISO 10993-11:2018, EN 1041:2008, EN ISO 7405:2018, EN ISO 15223-1:2016, EN ISO 13485:2016, EN ISO 14971:2019, ISO/TR 24971:2020, EN 1641:2009, EN ISO 14644-1:2015, EN ISO 14644-2:2015, EN ISO 14644-3:2019, EN ISO 14644-4:2001, EN ISO 14644-5:2004, KS I ISO 14644-6:2017, EN ISO 14644-7:2004, EN ISO 14644-8:2022, EN ISO 14644-9:2022, EN ISO 14644-10:2022, ISO 14644-12:2018, EN ISO 14644-13:2017, EN ISO 14644-14:2016, EN ISO 14644-15:2017, EN ISO 14644-16:2019, EN ISO 14644-17:2021, EN ISO 14698-1:2003, EN ISO 14698-2:2003, BS EN 17141:2020, KS A 3011:2018, USP <1116>, ASTM F2847-17, ASTM G131-96(2016)e1, EN ISO 11737-1:2018/A1:2021, EN ISO/TS 17665-2:2009, Korean Pharmacopeia 12th, ISTA 1A 2022, ASTM F1980-21, EN 62366-1:2015, IEC TR 62366-2:2016

is subject to the procedure set out in Annex II (excluding sect-4) of Directive 93/42/EEC under the supervision of Notified Body Number 1639, SGS Belgium NV, Noorderlaan 87, BE-2030 Antwerpen, Belgium.

GMDN Code : 60275(Endodontic temporary cement)

EC Certificate Number : KR19/81826259

Responsible for making this declaration is the :

■ Manufacturer: META BIOMED CO., LTD.

European Representative is:

■ Meta Biomed Europe GmbH, Wiesenstr. 35, 45473 Mülheim an der Ruhr, Germany

■ Tel:+49-208-30991910, Fax: +49-208-30991999

We, the manufacturer's declaration, declare and ensure with sole responsibility, that the Medical Device meet the provisions of Council Directive 93/42/EEC (Medical Device Directive) which apply to them.

Person responsible for making this declaration:

Name, Surname : Suk Song Oh

Position/Title : PRESIDENT

Place: Cheongju-si **Date:** November 08, 2023

Suk Song Oh
(Company stamp and legal signature)



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(Attachment: Model name list)

No	Brand Name	Model Name
1	Metapaste	Metapaste