## **EC CERTIFICATE**

## **Full Quality Assurance System**

Directive 93/42/EEC on Medical Devices, Annex II excluding (4)

No. 5-869-200-1907

The NEOEMKI National Medical Device Conformity Assessment and Certification LLC. certifies that the manufacturer:

Genuine Technologies, razvoj in proizvodnja, d.o.o.

Slovenska cesta 54 1000 Liubliana Slovenia

Manufacturing site: Jamova 39 1000 Ljubljana Slovenia

for the product:

## RS+ dental root filler

applies a quality system which meets the requirements of Directive 93/42/EEC concerning medical devices, Annex II.

Registry number of the related audit report: NE/1025/2020

This certificate is valid until 2024-05-26 supposed that the results of the regular yearly surveillance audits are satisfactory.

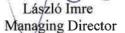
Issued by NEOEMKI LLC as a Notified Body with identification number 1011.

Issue: 3

First issued by the Directorate of Device Testing and Clinical Engineering (EMKI) on 29 July 2019.

Budapest, 2020-07-09







**EMKI 2396** 

The authenticity and validity of the certificate are verifiable at NEOEMKI LLC.

neoEMKI Nemzeti Orvostechnikai Eszköz Megfelelőségértékelő és Tanúsító Kft. neoEMKI National Medical Device Conformity Assessment and Certification LLC.

