

EvpU



EC Certificate

Full Quality Assurance System

Directive 93/42/EES on Medical devices, as amended by 2007/4/EC of the European Parliament and of the Council (MOD) Annex II excluding (4) (Modele H) Slovak government No. 587/2008 Collection of Laws as amended

No. 44064/102/1/2023/CE

EVPU as Notified Body No.2293, has audited the quality system in accordance with MOD and found that the quality system meets the requirements of MOD.

Manufacture and Facility

Medicinos linija, UAB

Karaciaus St.29 LT-78374

Siauliai, Lithuania

Device(s)

Dental materials

Type(s)

See Annex 1

Device(s) in

Class Ia

Relevant report(s)

44064/2023/C

Audit report

M020/10-5



Marek Rudak

Issued on August 20, 2024

Valid until August 19, 2029

Manufacturer can affix the CE mark with number Notified Body only in case devices are in compliance with all relevant and effective Directives of European Parliament and of the Council. Surveillance audits according to Annex I will be held to validity of the Certificate.