

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 oudited the quality system in recordance with MOD and found that the quality system meets the requirements of MOD.

Manufacture and Facility

Device(s)

Type(s)

Device(s)in

Relevant report(s)

Audit report

Medicinos linija, UAB

Karauciaus St.29 LT-78374

Siauliai, Lithuania

Dental materials

See Annex 1

Class Ia

44064/2023/C

M020/10-5



Marek Rudak

Valid until August 19, 2029

Issued on August 20, 2024

Manufacturer can affix the CE mark with number Notified Body only in case cavices are in comply with all relevant and effective Directives of European Parliament end of the Concil Sarvelliance audits according to Annex I will be held to validity of the Certificate.