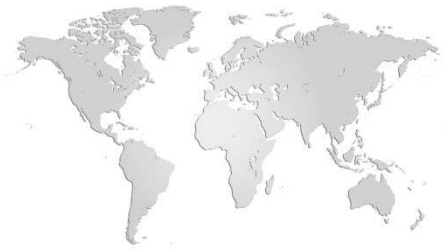
EU Certificate

for the assessment of the technical documentation



according to Medical Device Regulation (EU) 2017/745 Annex IX Chapter II

As a Notified Body of the European Union, DEKRA Certification GmbH certifies, that the manufacturer

PAJUNK GmbH Medizintechnologie

Single Registration Number (SRN): DE-MF-000005510
Karl-Hall-Straße 1, 78187 Geisingen, Germany

that the technical documentation of the product(s) described in the annex complies with the provisions of the Medical Device Directive (EU) 2017/745. The certificate is based on the results of the assessment of the technical documentation according to the Medical Devices Regulation (EU) 2017/745 Annex IX Chapter II, which are recorded in the report referred to in the annex.

Product: SPROTTE® STANDARD (LUER/ NRFit®) Anaesthesia: Cannula/ Needle

EU Certificate no.: 51268-61-I1

Certificate valid from: 2022-07-08
Certificate valid to: 2026-09-16

Previous certificate no. 51268-61-I0, valid from 2021-09-17 to 2022-07-07
Change to previous certificate: subsequent addition of single registration number (SRN)

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
BS-MDR-092

Natascha Jezyschek
DEKRA Certification GmbH, Stuttgart, 2022-07-08
Notified Body ID number: 0124