EU Declaration of Conformity according to the Medical Devices Directive 93/42/EEC

The manufacturer:
Ampri Handelsgesellschaft mbH
Benzstr. 16
21423 Winsen (Luhe)
Germany
declares under its own responsibility that

art. no.
0988101, 0988102, 0988103, 0988104 SoreProtect Sensplast
elastic non-woven fixation

Complies with the requirements of Annex VII of Directive 93/42/EEC.

This product is a Class 1 medical device according to the classification in Annex IX.

Technical documentation is available to prove this is accordance with the requirements.

Winsen, 26.05.2020

i.V.

ppa. Stephan Welzin
Head of Quality Management & Operational Purchasing

This Declaration of Conformity is valid until 25.05.2021

revision 03