

EU-DECLARATION OF CONFORMITY

Manufacturer: Wernli AG
Eggasse 4
4852 Rothrist
Switzerland

Single Registration Number CH: CHRN-MF-20000003

EC Representative: Wero Swiss Med Kft.
Ipartelep utca 6
4220 Hajdúböszörmény
Hungary

Single Registration Number HU: HU-AR-000049642

We declare under our sole responsibility that

the medical device
with reference no
with Basic-UDI-DI (product code)

Wero Swiss Forte
see page 2
76300161A17PB

Risk Class: **class 1, unsterile**
According to Annex VIII, Rule 1

meets all the provisions of the Regulation (EU) 2017 /745 which apply to it.

Applied harmonised standards, national standards or other normative documents: EN ISO 10993-1:2021 and applicable parts of the EN ISO 10993 Series
EN ISO 15223-1:2021
EN ISO 14971:2019 / A11:2021

Conformity assessment procedures: Regulation (EU) 2017 /745 according to MDR Article 10 and MDR Article 52 (7) Annex I and Annex II

Place, date: Rothrist, 09.03.2026

Name and function



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Felix Schoenle, CEO

Reference Numbers:

Length: 7m stretched

A1720060.070	Wero Swiss Forte 6cm x 7m brown
A1720080.070	Wero Swiss Forte 8cm x 7m brown
A1720100.070	Wero Swiss Forte 10cm x 7m brown
A1720120.070	Wero Swiss Forte 12cm x 7m brown
A1720150.070	Wero Swiss Forte 15cm x 7m brown