

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 Solifix Crepp LF white
see page 2
76300161A9101ZP

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: 4m stretched

A91010040.040	Wero Swiss Solifix Crepp LF 4cm x 4m white
A91010060.040	Wero Swiss Solifix Crepp LF 6cm x 4m white
A91010080.040	Wero Swiss Solifix Crepp LF 8cm x 4m white
A91010100.040	Wero Swiss Solifix Crepp LF 10cm x 4m white
A91010120.040	Wero Swiss Solifix Crepp LF 12cm x 4m white

Length: 20m stretched

A91010040.200	Wero Swiss Solifix Crepp LF 4cm x 20m white
A91010060.200	Wero Swiss Solifix Crepp LF 6cm x 20m white
A91010080.200	Wero Swiss Solifix Crepp LF 8cm x 20m white
A91010100.200	Wero Swiss Solifix Crepp LF 10cm x 20m white
A91010120.200	Wero Swiss Solifix Crepp LF 12cm x 20m white