

EU-DECLARATION OF CONFORMITY

Manufacturer	Wernli AG Eggasse 4 4852 Rothrist Switzerland
Single Registration Number CH:	CHRN-MF-200000003
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