IVF HARTMANN AG
Victor von Bruns-Strasse 28
8212 Neuhausen am Rheinfall
Switzerland

Telefon: +41 52 674 31 11
Fax: +41 52 672 74 41
info@ivf.hartmann.info
ivf.hartmann.info

HARTMANN
Helps. Cares. Protects.

EU Declaration of Conformity

Neuhausen, 2025-12-08

Product Group Number	3901 Verbandpäckchen 22 (IVP) (scope see table 1)
Manufacturer	IVF HARTMANN AG Victor-von-Bruns-Strasse 28 8212 Neuhausen, Switzerland
Authorized Representative	PAUL HARTMANN AG Paul-Hartmann-Straße 12 89520 Heidenheim, Germany

We herewith declare under our sole responsibility that the product listed above, first placed on the market by IVF HARTMANN AG, satisfy the applicable provisions of the following listed legislation. The conformity assessment procedures have been performed and the Technical Documentation is kept available.

Applied legislative act

Medical Device Regulation (EU) 2017/745

High Level Intended Purpose	Non-active, non-implantable devices for wound and skin care	
Classification	Risk Class	Rule
	Class Is	Rule 4, indent 1
Conformity Assessment Procedure	Article 52 (7) and Annex XI part A with respect to sterility	
Notified Body	TÜV SÜD Product Service GmbH, Ridlerstr. 65, 80339 München, Germany, ID-Nr. 0123	
EU Certificates	EU Quality Assurance Certificate (MDR) No. G26 047326 0014	
Single Registration Number	Manufacturer: CH-MF-000015962	
Swiss Single Registration Number	Manufacturer: CHRN-MF-20000305	
Basic UDI-DI	76116003901N3	
	Code	Term
EMDN	M030401	Elastic Compression Bandages
GMDN	47011	First aid absorbent pad/bandage

IVF HARTMANN AG

i.V.

Robert Hämmerl
Regulatory Affairs Manager

i.V.

Dr. Heinrich Giegrich
Head of RA & Quality

Valid until: 2030-12-07

Table 1: Scope

REF	Description
988671	IVP 22 15 cm