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Declaration of Conformity

This European Declaration of Conformity is issued under the sole responsibility of the manufacturer.

MANUFACTURER					
Name of Company				SRN	
Helmer Scientific DBA Helmer Inc.			Blvd Noblesville	US-MF-000003326	
AUTHORIZED REPRESENTATIVE					
Address			SRN	Pl	hone/email
-		The Hague	NL-AR-000	000116 +3	31.70.345.8570
The Netherlands				Eı	mergoEurope@ul.com
PRODUCT IDENTIFICATION					
Code / Catalog Number					
	DH2, DH4, DH8				
Intended Purpose				Basic UDI-DI	
Intended to by used by blood banks, hospitals and clinics to decrease the thaw time for fresh frozen plasma. It is NOT INTENDED to warm the plasma prior to infusion.				081639402TFR0036N	
RISK CLASS FOR DEVICES					
Device Classification Common Specifications / Standards					
1	EN61010-1 2010 Safety Requirements for Electrical Equipment for Measurement, Control, and Laboratory Use				
1	EN ISO 15223-2:2012 Medical devices — Symbols to be used with medical device labels				
	EN 62366:2012 Medical Devices – Application of Usability Engineering				
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	EN60601-1 Medical electrical equipment – Part 1-2: – Collateral Standard: Electromagnetic disturbances EN1041 Information to be supplied by the manufacturer with medical device				
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Helmer Scientific declares that the above-mentioned products meet the provision of the following EU legislation:

• Medical Devices Regulation (EU) 2017/745

• RoHS Recast Directive 2011/65/EU including the amendment to Annex II described in Commission Delegated Directive (EU) 2015/863.

COMPANY REPRESENTATIVE: Renee Schultz SIGNATURE:

TITLE: Director of Regulatory Affairs

PLACE: 14400 Bergen Blvd, Noblesville In USA **DATE:** 01 January 2022

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