



EU Declaration of Conformity

2625 Patton Road
Roseville, MN 55113
651-294-2200
www.ablenetinc.com

This declaration of conformity is issued under the sole responsibility of the manufacturer. The device covered by the present declaration is in conformity with all regulations below, including compliance with related Essential Requirements.

Object of the declaration:

Product Name:	<i>Specs IC</i>
Product Model Designator:	Specs IC
Product Part Number(s):	10000114
Basic UDI-DI <required for MDR>	00850011150382
Control Indicator:	Date codes starting 2025W26
Global Medical Device Nomenclature Code (GMDN) and Description	
Product Options/Accessories:	N/A

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The object of the declaration described above is in conformity with the following regulations:

EU Regulation	Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices
Device Classification:	Class I based on Annex VIII and Rule 13
Conformity Assessment Path	<i>Not Applicable – Class I device</i>
Name/Address/ID of Notified Body:	<i>Not Applicable – Class I device</i>
Standards and Common Specifications	<p>The products listed above have been tested in a typical configuration as described in the Manufacturer's accompanying documentation and are compliant with the product standards listed below.</p> <p><i>BS EN ISO 14971:2012 – Medical Devices – Application of risk management to medical devices</i> <i>BS EN ISO 13485:2016 – Medical Devices – Quality management systems – Requirements for regulatory purposes</i> <i>Information Technology Equipment-Safety IEC 60950-1:2005+A1:2009</i></p>



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Additional information:

EU Authorized Representative:	<i>EUCEREP Roald Dahllaan 33 5629MC – Eindhoven The Netherlands</i>
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Signature (signed for and on behalf of AbleNet):	Date of Issue: <DD Month YYYY> 04 June 2025
Printed Name: Jesse Boyle	Place of Issue: AbleNet Inc – Roseville, MN
Title: Manager, Regulatory & Compliance	Document Number: DoC_10000114_SpecsIC_20250604