



## Declaration of Conformity

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

<b>MANUFACTURER</b>		
Name of Company	Address	SRN
Bio/Data Corporation	155 Gibraltar Road, Horsham, PA 19044 U.S.A.	US-MF-000026991
<b>UK RESPONSIBLE PERSON</b>		
Name of Company	Address	Telephone/email
Alpha Laboratories Ltd	40 Parham Drive, Eastleigh, Hampshire SO50 4NU UNITED KINGDOM	+44 2380 483000 – phone <a href="mailto:quality@alphalabs.co.uk">quality@alphalabs.co.uk</a>
<b>PRODUCT IDENTIFICATION</b>		
Product / Trade Name	Product Code / Catalog Number	Basic UDI-DI
vW Abnormal Control Plasma 3 x 0.5mL	101270	++G0561012703E
Intended Purpose	Photo	
See Instructions for Use	See website <a href="http://www.biodatacorp.com">www.biodatacorp.com</a>	
Abnormal Control Plasma is prepared from a pool of normal infection negative human plasma which has been partially depleted of vW Factor and then lyophilized. It is used to verify the performance and sensitivity of the Ristocetin CoFactor Activity test.		
<b>IVDR RISK CLASS / COMMON SPECIFICATIONS</b>		
Device Classification		Common Specifications
Class	General IVD (Self Certified)	Sections 1-5 (as modified by Part III of Schedule 2A to the UK MDR 2002)
Rule	Part IV of the UK MDR 2002, Annex III	

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An ISO 13485 Registered Company

**Bio/Data Corporation, the Manufacturer, declares that the above-mentioned product meets the provision of the following UK legislation:**

Medical Devices Regulations 2002 (SI 2002 No 618, as amended) (UK MDR 2002)

**Conformity Statement:**

Bio/Data Corporation declares that the above-mentioned products meet the provision of the Medical Devices Regulations 2002 (SI 2002 No 618, as amended) (UK MDR 2002) for In Vitro Diagnostic Medical Devices.

**Notified Body (NB) Identification, conformity assessment procedure performed and identification if the certificate or certificates issued:**

Not applicable. The conformity assessment procedure for General IVD (non-Annex II) Devices should be carried out, as a general rule, under the sole responsibility of manufacturers, since such devices pose a low risk to patients.

(Self-Declaration)

**COMPANY REPRESENTATIVE:** William M. Trolio

**SIGNATURE:** \_\_\_\_\_



**TITLE/FUNCTION:** Director of Quality Assurance & Regulatory Affairs for Bio/Data Corporation

**PLACE:** Bio/Data Corporation, Horsham, PA. 19044 U.S.A.

**DATE:** 1 April 2023