

Declaration of Conformity

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

MANUFACTURER			
Name of Company		Address	SRN
Bio/Data Corporation		155 Gibraltar Road,	US-MF-000026991
		Horsham, PA 19044 U.S.A.	
UK RESPONSIBLE PERSON			
Name of Company		Address	Telephone/email
Alpha Lab	ooratories Ltd	40 Parham Drive, Eastleigh,	+44 2380 483000 – phone
		Hampshire SO50 4NU UNITED KINGDOM	guality@alphalabs.co.uk
PRODUCT IDENTIFICATION			
Product / Trade Name		Product Code / Catalog Number	Basic UDI-DI
vW Select Ristocetin Cofactor Assay System		106730	++G0561067304W
Intended Purpose			Photo
See Instructions for Use			See website www.biodatacorp.com
The vW Select Ristocetin Cofactor Assay System is optimized to elicit the agglutination of lyophilized platelets in the presence of the patient's Platelet Poor Plasma and Ristocetin.			
		ION SPECIFICATIONS	
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		

155 Gibraltar Road, Horsham, PA 19044 U.S.A. Worldwide: (215) 441-4000 U.S.A.: (800) 257-3282 Fax Worldwide: (215) 443-8820 www.biodatacorp.com e-mail: customer.service@biodatacorp.com An ISO 13485 Registered Company



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

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