

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 RESP | ONSIBLE PERSON | V | |
| Name of | Company | Address | Telephone/email |
| Alpha Lab | oratories 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 |
| MagneTube | | 107146 | *+G0561071461 |
| Intended Purpose | | | Photo |
| See Instructions for Use | | | See website www.biodatacorp.com |
| Disposable Micro Test Tubes 7.25 x 55 mm containing Micro Stir Bars | | | |
| are for use with the PAP-8E aggregometer or PAP-4 series | | | |
| aggregometer using micro volume adapters. | | | |
| IVDR RIS | K CLASS / COMN | NON 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 | | |



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