

## **Declaration of Conformity**

This European 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.                                   |                 |                                 |  |  |
| AUTHOR   | RIZED REPR            | ESEN'                          | TATIVE   |                 |                                 |  |  |
| Name of Company Add  |                       | Addı                           | ress SRN   |                 | Telephone/email                 |  |  |
| mdi Euro   | pa GmbH               | Lang                           | enhagener Str. 71  | DE-AR-000006218 | +49-511-3908 9531 – phone       |  |  |
|  |                       | D-30855 Langenhagen<br>GERMANY |  |                 | info@mdi-europa.com             |  |  |
|  |                       |                                |  |                 |                                 |  |  |
| PRODUCT IDENTIFICATION                                       |                       |                                |  |                 |                                 |  |  |
| Product / Trade Name   |                       | Product Code / Catalog Number  |  | Basic UDI-DI    |                                 |  |  |
| Arachidonic Acid   |                       | 101297                         |  | ++G05610129742  |                                 |  |  |
| Intended   | l Purpose             |                                |  |                 | Photo                           |  |  |
| See Instructions for Use                                     |                       |                                |  |                 | See website www.biodatacorp.com |  |  |
| Arachidonic Acid reagent is for routine use in demonstrating |                       |                                |  |                 |                                 |  |  |
| thrombo  | xane A2 a             | ctivati                        | on response in Pla   |                 |                                 |  |  |
| samples.   |                       |                                |  |                 |                                 |  |  |
| IVDR RISK CLASS / COMMON SPECIFICATIONS                      |                       |                                |  |                 |                                 |  |  |
| Device Classification  |                       |                                | Common Specifications                                      |                 |                                 |  |  |
| Class  | A non-ste             | erile                          | No relevant common specifications have been published yet. |                 |                                 |  |  |
| Rule   | 5a per An             | inex                           |  |                 | •                               |  |  |
|  | VIII of IV<br>2017/74 | DR                             |  |                 |                                 |  |  |



## www.biodatacorp.com customer.service@biodatacorp.com

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

In Vitro Diagnostic Medical Device Regulation (IVDR) (EU) 2017/746

## **Conformity Statement:**

Bio/Data Corporation confirms that the device covered by this declaration is in conformity with the (EU) IVDR 2017/746 Regulation and, if applicable, with any other relevant Union Legislation that provides for the issuing of an EU Declaration of Conformity.

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

Not applicable. The conformity assessment procedure for Class A 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: _        | C da ho              |
|---|---------------------|----------------------|
|   |                     |                      |
|   |                     | D: /D                |
| TITLE/FUNCTION: Director of Quality Assurance & Reg | ulatory Affairs for | Bio/Data Corporation |
|   |                     |                      |
| PLACE: Bio/Data Corporation, Horsham, PA. 19044 U   | S.A. DATE: 1        | L April 2023         |