# *Declaration of Conformity*

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|---|---|
| **Manufacturer:** | Becton, Dickinson and Company<br>7 Loveton Circle<br>Sparks, Maryland 21152, USA<br>Tel: +1.410.316.4000<br>Fax: +1.410.316.4499 |
| **Authorized Representative:** | Benex Limited<br>Pottery Road, Dun Laoghaire<br>Co. Dublin, Ireland<br>Tel. : +353.1.202.5222<br>Fax : +353.1.202.5388 |
| **Conformity Assessment Procedure:** | IVD Directive 98/79/EC, of the European Parliament and of the Council, Annex III of Directive 98/79/EC |

**Product:**

| REF | Product Name |
|---|---|
| 256122 | BD Veritor™ System for Rapid Detection of SARS-CoV-2 & Flu A+B -30 pk |
| 256090 | SARS-CoV-2 & Flu A+B Control Swab Set – 30 pk |

**We hereby declare that the above mentioned products comply with the European In Vitro Diagnostic Medical Devices Directive and its relevant transposition into national laws of the member states into which we place the devices. This declaration is issued under the sole responsibility of Becton, Dickinson and Company.**

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| **Date:** | 13 MAY 2022 |
| **Name and Authority:** | Anne Zavertnik<br>Vice President Regulatory Affairs,<br>Integrated Diagnostic Solutions |
| **Signature:** |  |

Technical File Number: BDDSTF256122

| RECORD REVISION HISTORY TABLE | |
|---|---|
| **Revision** | **Description of Changes** |
| 01 | New Product |