

## EC DECLARATION OF CONFORMITY

<b>Manufacturer:</b>	Becton, Dickinson and Company Belliver Industrial Estate, Belliver Way, Roborough, Plymouth, PL6 7BP, United Kingdom																																												
<b>Manufacturing Site(s):</b>	Becton, Dickinson and Company Belliver Industrial Estate, Belliver Way, Roborough, Plymouth, PL6 7BP, United Kingdom																																												
<b>Products:</b>	<table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="text-align: left;">Catalogue number</th> <th style="text-align: left;">Device name</th> </tr> </thead> <tbody> <tr><td>362080</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>362081</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>362082</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>362091</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>362092</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>364660</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>365904</td><td>BD Vacutainer® SERUM/CAT Plus Blood Collection Tubes</td></tr> <tr><td>367837</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>367895</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>367896</td><td>BD Vacutainer® CAT (Clot Activator Tube) Plus Blood Collection Tubes</td></tr> <tr><td>368271</td><td>BD Vacutainer® SERUM/CAT Plus Blood Collection Tubes</td></tr> <tr><td>368492</td><td>BD Vacutainer® CAT (Clot Activator Tube) Plus Blood Collection Tubes</td></tr> <tr><td>368493</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>368813</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>368814</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>368815</td><td>BD Vacutainer® CAT (Clot Activator Tube) Plus Blood Collection Tubes</td></tr> <tr><td>368817</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>368863</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>368975</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> <tr><td>369032</td><td>BD Vacutainer® CAT (Clot Activator Tube) Plus Blood Collection Tubes</td></tr> <tr><td>369625</td><td>BD Vacutainer® CAT Plus Blood Collection Tubes</td></tr> </tbody> </table>	Catalogue number	Device name	362080	BD Vacutainer® CAT Plus Blood Collection Tubes	362081	BD Vacutainer® CAT Plus Blood Collection Tubes	362082	BD Vacutainer® CAT Plus Blood Collection Tubes	362091	BD Vacutainer® CAT Plus Blood Collection Tubes	362092	BD Vacutainer® CAT Plus Blood Collection Tubes	364660	BD Vacutainer® CAT Plus Blood Collection Tubes	365904	BD Vacutainer® SERUM/CAT Plus Blood Collection Tubes	367837	BD Vacutainer® CAT Plus Blood Collection Tubes	367895	BD Vacutainer® CAT Plus Blood Collection Tubes	367896	BD Vacutainer® CAT (Clot Activator Tube) Plus Blood Collection Tubes	368271	BD Vacutainer® SERUM/CAT Plus Blood Collection Tubes	368492	BD Vacutainer® CAT (Clot Activator Tube) Plus Blood Collection Tubes	368493	BD Vacutainer® CAT Plus Blood Collection Tubes	368813	BD Vacutainer® CAT Plus Blood Collection Tubes	368814	BD Vacutainer® CAT Plus Blood Collection Tubes	368815	BD Vacutainer® CAT (Clot Activator Tube) Plus Blood Collection Tubes	368817	BD Vacutainer® CAT Plus Blood Collection Tubes	368863	BD Vacutainer® CAT Plus Blood Collection Tubes	368975	BD Vacutainer® CAT Plus Blood Collection Tubes	369032	BD Vacutainer® CAT (Clot Activator Tube) Plus Blood Collection Tubes	369625	BD Vacutainer® CAT Plus Blood Collection Tubes
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<b>IVDD Classification:</b>	Non Annex II <i>In Vitro</i> Diagnostic Medical Device																																												
<b>IVDD Conformity Assessment Route:</b>	Annex III (excluding Annex III.6)																																												
<b>GMDN:</b>	42386																																												

We herewith declare that the above mentioned products meet the provisions of the Council Directive 98/79/EC for *in vitro* diagnostic medical devices. This declaration is based on conformity to Annex III (excluding Annex III.6). All supporting documentation is retained under the premises of the manufacturer.

<b>List of Harmonized Standards:</b>
<b>EN ISO 13485:2012</b> Medical devices — Quality management systems — Requirements for regulatory purposes <b>EN ISO 14971:2012</b> Medical Devices – Application of risk management to medical devices <b>EN 556-</b>

**1:2001** Sterilisation of medical devices – Requirements for medical devices to be designated “STERILE” – Part 1: Requirements for terminally sterilized medical devices **EN ISO 11137-1:2015** Sterilization of health care products - Radiation - Part 1: Requirements for development, validation and routine control of a sterilization process for medical devices **EN ISO 11137-2:2015** Sterilization of health care products — Radiation — Part 2: Establishing the sterilization dose. **EN ISO 11737-2:2009** Sterilization of medical devices - Microbiological methods Part 2: Tests of sterility performed in the definition, validation and maintenance of a sterilization process **EN 14820:2004** Single-use containers for human venous blood specimen collection **EN 62366:2008** Medical devices - Application of usability engineering to medical devices **EN ISO 18113-1: 2011** In vitro diagnostic medical devices – Information supplied by the manufacturer (Labelling). Part 1: Terms, definitions and general requirements (ISO 18113-1:2009) **EN ISO 18113-2: 2011** In vitro diagnostic medical devices – Information supplied by the manufacturer (Labelling). Part 2: In vitro diagnostic reagents for professional use (ISO 18113-2:2009) **EN ISO 15223-1:2016** Medical Devices – Symbols to be used with medical device labels, labelling and information to be supplied – Part 1: General Requirements

**List of Non-Harmonised Standards:**

**ISO 14001:2015** Environmental management systems - Requirements with guidance for use **EN ISO 11137-3:2017** Sterilisation of health care product – Radiation – part 3: guidance on dosimetric aspects of development, validation and routine control **EN ISO 11737-1:2018** Sterilization of medical devices - Microbiological methods - Part 1: Determination of a population of microorganisms on products **ISO 6710:1995** Single-Use Containers for Venous Blood Specimen Collection **EN ISO 14698-1:2003** Cleanrooms and associated controlled environments -- Biocontamination control — Part 1: General principles and methods **EN ISO 14698-2:2003** Cleanrooms and associated controlled environments -- Biocontamination control — Part 2: Evaluation and interpretation of biocontamination data **EN ISO 14644-1:2015** Cleanrooms and associated controlled environments - Part 1: Classification of air cleanliness **EN ISO 14644-2:2015** Cleanrooms and associated controlled environments -- Part 2: Monitoring to provide evidence of cleanroom performance related to air cleanliness by particle concentration **ISO 2859-1:1999** Sampling procedures for inspection by attributes - Part 1: Sampling schemes indexed by acceptance quality limit (AQL) for lot-by-lot inspection **ASTM D5276:1998 (R 2009)** Standard Test Method for Drop Test of Loaded Containers by Free Fall **ASTM D999: 2008 (R2015)** Standard Test Methods for Vibration Testing of Shipping Containers **ASTM D4169: 2014** Standard Practice for Performance Testing of Shipping Containers and Systems **ASTM D4728: 2006 (R2012)** Standard Test Method for Random Vibration Testing of Shipping Containers **ASTM D-775: 1980 (R 1986)** Standard Test Method for Drop Test for Loaded Boxes

SIGNED FOR AND ON BEHALF OF: Becton, Dickinson and Company

PLACE, DATE OF ISSUE: Plymouth, 18<sup>th</sup> September 2018

Signature: 

Brad Spring

Vice President, Regulatory Affairs

BD Life Sciences

Document Number: VR4310005



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### **VERSION HISTORY**

Current Version Prepared By: Joseph Statham

<b>REV.</b>	<b>Version Description</b>
A	Transferred from QDMS to ECC – Version number remained
B	Transfer into new IVD Declaration of Conformity Template (MED-RA-001D)
C	Update EN ISO 11737-1:2006 to EN ISO 11737-1:2018 as per CAPA 325553.