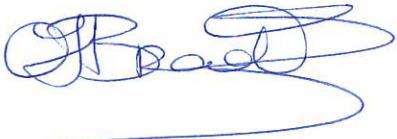




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|--|---|
| Manufacturer: | BD Kiestra B.V. Marconilaan 6 9207 JC Drachten The Netherlands |
| Conformity Assessment Procedure: | Annex III of the IVD Directive 98/79/EC |
| Products: | <p>BD Kiestra™ WCA</p> <p>Containing any of the following:</p> <p>447202 BD Kiestra™ Inoqula+™ (covered by its' own Declaration of Conformity (DOC) and not on the WCA DOC)</p> <p>447206 BD Kiestra™ ReadA™ Compact (covered by its' own DOC and not on the WCA DOC)</p> <p>447214 BD Kiestra™ Inoqula+™ Slide Preparation Module (covered by its' own DOC and not on the WCA DOC)</p> <p>447272 BD Kiestra™ Inoqula™ Magnetic Beads (covered by its' own DOC and not on the WCA DOC)</p> <p>447293 ProceedA WCA</p> <p>447322 Inoqula WCA upgrade kit</p> <p>447323 Inoqula WCA upgrade kit pre 1.2</p> |
| <p>We hereby declare that the above mentioned products comply with the European In Vitro Diagnostic Directive 98/79/EC and its relevant transposition into national laws of the member states into which we place the devices.</p> | |
| Signed in Drachten: | 14 November 2019 |
| Name and Authority: | Karin Brands Manager Regulatory Affairs |
| Signature: |  |