BD Declaration of Conformity

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:	BD Kiestra™ WCA Containing any of the following: 447202 BD Kiestra™ InoqulA+™ (covered by its' own Declaration of Conformity (DOC) and not on the WCA DOC) 447206 BD Kiestra™ ReadA™ Compact (covered by its' own DOC and not on the WCA DOC) 447214 BD Kiestra™ InoqulA+™ Slide Preparation Module (covered by its' own DOC and not on the WCA DOC) 447272 BD Kiestra™ InoqulA™ Magnetic Beads (covered by its' own DOC and not on the WCA DOC) 447293 ProceedA WCA 447322 InoqulA WCA upgrade kit 447323 InoqulA WCA upgrade kit pre 1.2

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.

Signed in Drachten:	14 November 2019
Name and Authority:	Karin Brands Manager Regulatory Affairs
Signature: