

EC Declaration of Conformity

I, the undersigned, hereby declare that the products specified below conform to Annex III of Directive 98/79/EC of the European Parliament and Council of 27 October 1998, on in-vitro diagnostic medical devices (and its relevant transposition into the national laws of the Member States in which the device is intended to be placed on the market); and

to Annex VI of Directive 2011/65/EU of the European Parliament and Council of 8 June 2011 on the restriction of the use of certain hazardous substances according Annex II (lead, mercury, hexavalent chromium, cadmium, polybrominated biphenyls and polybrominated diphenyl ethers) in electrical and electronic equipment (and its relevant transposition into the national laws of the Member States in which the device is intended to be placed on the market).

<u>Manufacturer:</u>	Ventana Medical Systems, Inc.
<u>Address:</u>	1910 E. Innovation Park Drive Tucson, AZ USA 85755
<u>European Authorized Representative:</u>	ROCHE DIAGNOSTICS GmbH Sandhofer Strasse 116 D-68305 Mannheim Germany
<u>Manufacturing Site:</u>	Tucson, AZ USA
<u>Product name:</u>	BENCHMARK XT
<u>Ventana P/N:</u>	750-700
<u>Roche P/N:</u>	05265231001
<u>Classification:</u>	General IVD
<u>Technical Data File:</u>	TDF 0040
<u>Date of First CE Marking:</u>	04-Dec-2003

Name of Authorized Signatory: Deepshikha Bhandari
Vice President, Regulatory Affairs

Signature: _____



Date: _____

22-JULY-2016