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).

Manufacturer:	Ventana Medical Systems, Inc.
Address:	1910 E. Innovation Park Drive Tucson, AZ USA 85755
European Authorized Representative:	ROCHE DIAGNOSTICS GmbH Sandhofer Strasse 116 D-68305 Mannheim Germany
Manufacturing Site:	Tucson, AZ USA
Product name:	Reticulum II Staining Kit
Ventana P/N: *Roche P/N*:	860-024 05279399001
Classification: *Technical Data File:* *Date of First CE Marking:*	General IVD TDF-0056 16-May-2007

Name of Authorized Signatory: Peter Martin
Vice President, Regulatory Affairs

Signature: _____ Date: 22-Aug-2018

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