



a member of the Roche Group

Ventana Medical Systems, Inc.  
1910 E. Innovation Park Drive  
Tucson, Arizona 85755  
Phone: (520) 887-2155  
Toll Free: (800)-227-2155  
[www.ventana.com](http://www.ventana.com)

## Declaration of Conformity to 98/79/EC

Manufacturer: Ventana Medical Systems, Inc.  
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

	Ventana	Roche
	[REF]	
Product name/	CONFIRM anti-MSH6 (44)	790-4455
Catalogue No.	Mouse Monoclonal Primary Antibody	05929911001
Technical Data File:	TDF-0412	
Classification:	General IVD	
Conformity Assessment:	98/79/EC Annex III	

Ventana Medical Systems, Inc. declares that the product(s) listed is/are in conformity with the essential requirements of Annex I of Council Directive 98/79/EC. All supporting documentation is retained under the premise of the Manufacturer.

Ventana Medical Systems, Inc. maintains a quality system based on EN ISO 13485:2012 / EN ISO 13485:2012 + AC:2012 as certified by TÜV Rheinland LGA Products GmbH (Notified Body No. 0197).

Place of Issue: Tucson, AZ USA 85755

Name of Authorized Signatory: Troy Quander  
Vice President, Regulatory Affairs

Signature:

Date: 2-Apr-15 - 2015

# EU Declaration of Conformity

In accordance with EU Regulation 2017/746 of the European Parliament and of the Council of 5 April 2017 on *in vitro* diagnostic medical devices.

Manufacturer: **Ventana Medical Systems Inc.  
1910 E Innovation Park Drive  
Tucson, AZ 85755, USA**

Single Registration Number (SRN) **US-MF-000016993**  
Manufacturer:

Authorized Representative: **Roche Diagnostics GmbH  
Sandhofer Strasse 116  
68305 Mannheim  
Germany**

Single Registration Number (SRN) **DE-AR-000006262**  
Authorized Representative:

This declaration is issued under the sole responsibility of Ventana Medical Systems Inc.

## Product Information

Part Number:	Product Name:	Basic UDI-DI:
06695248001 (alternative P/N: 805-4713)	CINtec® p16 Histology	761333601839BF
06695256001 (alternative P/N: 825-4713)		

**Intended Purpose:** CINtec p16 Histology is an immuno-histochemistry assay for the qualitative detection of the p16INK4a protein on formalin-fixed, paraffin-embedded tissue sections prepared from cervical biopsies. It is indicated to be used in conjunction with H&E stained slides prepared from the same cervical tissue specimen as an aid to increase diagnostic accuracy and inter-observer agreement in the diagnosis of high grade cervical intraepithelial neoplasia.

**Risk Class:** Class C

**Common Specifications:** Not applicable as no Common Specifications exist for the concerned device.

**Name, Address and  
Identification number of  
the Notified Body:** TÜV SÜD Product Service GmbH (No. 0123)  
Ridlerstraße 65  
80339 MÜNCHEN  
Germany

Conformity Assessment was established through the procedure described in Annex IX of EU Regulation 2017/746, including an assessment of the Technical Documentation as described in Annex IX, Chapter II. This declaration is supported by the following certificate(s):

EU Quality Management System Certificate: V12 096981 0003

Conformity of the product with EU Regulation 2017/746 and other applicable EU legislation has been established.

On behalf of Ventana Medical Systems Inc.

Place: Tucson, AZ 85755, USA

Date:

DocuSigned by:

  
Jeff Boone

D9C50B8025B5B4D8...

Site Head of Quality Function

Place: Tucson, AZ 85755, USA

Date:

DocuSigned by:

  
Benjamin Curson

0E08FEB02001421...

Site Head of Regulatory Affairs Function



# EU Declaration of Conformity

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68305 Mannheim  
Germany**

Single Registration Number (SRN) **DE-AR-000006262**  
Authorized Representative:

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## Product Information

Part Number:	Product Name:	Basic UDI-DI:
05267102001 (alternative P/N: 760-2542)	Anti-p53 (Bp53-11) Primary Antibody	761333601186AM

**Intended Purpose:** Anti-p53 (Bp53-11) Primary Antibody is intended for laboratory use in the qualitative immunohistochemical detection of p53 protein by light microscopy in sections of formalin-fixed, paraffin-embedded tissue on a BenchMark IHC/ISH instrument.

**Risk Class:** Class C

**Common Specifications:** Not applicable as no Common Specifications exist for the concerned device.

**Name, Address and Identification number of the Notified Body:** TÜV SÜD Product Service GmbH (No. 0123)  
Ridlerstraße 65  
80339 MÜNCHEN  
Germany

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EU Quality Management System Certificate: V12 096981 0003



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On behalf of Ventana Medical Systems Inc.

Place: Tucson, AZ 85755, USA

22-Jun-2022

Date:

*Jeff Boone*

**Jeff Boone**

Site Head of Quality Function

Place: Tucson, AZ 85755, USA

21-Jun-2022

Date:

*Ben Curson*

**Benjamin Curson**

Site Head of Regulatory Affairs Function



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In accordance with EU Regulation 2017/746 of the European Parliament and of the Council of 5 April 2017 on *in vitro* diagnostic medical devices.

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Manufacturer:

Authorized Representative: **Roche Diagnostics GmbH  
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68305 Mannheim  
Germany**

Single Registration Number (SRN) **DE-AR-000006262**  
Authorized Representative:

This declaration is issued under the sole responsibility of Ventana Medical Systems Inc.

## Product Information

Part Number:	Product Name:	Basic UDI-DI:
05867061001 (alternative P/N: 790-4509)	VENTANA anti-p63 (4A4) Mouse Monoclonal Primary Antibody	761333600840AR

**Intended Purpose:** VENTANA anti-p63 (4A4) Mouse Monoclonal Primary Antibody is intended for laboratory use in the qualitative immunohistochemical detection of the p63 protein by light microscopy in sections of formalin-fixed, paraffin-embedded tissue stained on a BenchMark IHC/ISH instrument.

**Risk Class:** Class C

**Common Specifications:** Not applicable as no Common Specifications exist for the concerned device.

**Name, Address and Identification number of the Notified Body:** TÜV SÜD Product Service GmbH (No. 0123)  
Ridlerstraße 65  
80339 MÜNCHEN  
Germany

Conformity Assessment was established through the procedure described in Annex IX of EU Regulation 2017/746, including an assessment of the Technical Documentation as described in Annex IX, Chapter II. This declaration is supported by the following certificate(s):

EU Quality Management System Certificate: V12 096981



Conformity of the product with EU Regulation 2017/746 and other applicable EU legislation has been established.

On behalf of Ventana Medical Systems Inc.

Place: Tucson, AZ 85755, USA

Date: 22-Feb-2022

DocuSigned by:

A handwritten signature in black ink, appearing to read "Jeff Boone".

DOC56B8025BB4D8...

**Jeff Boone**

Site Head of Quality Function

Place: Tucson, AZ 85755, USA

Date: 01-Mar-2022

A handwritten signature in black ink, appearing to read "Ben Curson".

**Benjamin Curson**

Site Head of Regulatory Affairs Function



# EU Declaration of Conformity

In accordance with EU Regulation 2017/746 of the European Parliament and of the Council of 5 April 2017 on *in vitro* diagnostic medical devices.

Manufacturer: **Ventana Medical Systems Inc.  
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Manufacturer:

Authorized Representative: **Roche Diagnostics GmbH  
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68305 Mannheim  
Germany**

Single Registration Number (SRN) **DE-AR-000006262**  
Authorized Representative:

This declaration is issued under the sole responsibility of Ventana Medical Systems Inc.

## Product Information

Part Number:	Product Name:	Basic UDI-DI:
05277990001 (alternative P/N: 790-2223) 50 tests	CONFIRM anti-Progesterone Receptor (PR) (1E2) Rabbit Monoclonal Primary Antibody	761333601236AB
05278392001 (alternative P/N: 790-4296) 250 tests		

**Intended Purpose:** CONFIRM anti-Progesterone Receptor (PR) (1E2) Rabbit Monoclonal (IgG) Primary Antibody is intended for laboratory use in the qualitative detection of progesterone receptor (PR) antigen in sections of formalin-fixed, paraffin-embedded tissue on a VENTANA automated slide stainer with VENTANA detection kits and ancillary reagents. CONFIRM anti-PR (1E2) antibody is directed against an epitope present on human progesterone receptor protein located in the nucleus of PR positive normal and neoplastic cells. CONFIRM anti-PR (1E2) antibody is indicated as an aid in the management, prognosis, and prediction of therapy outcome of breast carcinoma. This product should be interpreted by a qualified pathologist in conjunction with histological examination, relevant clinical information, and proper controls. Prescription use only. This antibody is intended for *in vitro* diagnostic (IVD) use.

**Risk Class:** Class C

**Common Specifications:** Not applicable as no Common Specifications exist for the concerned devices.



**Name, Address and  
Identification number of  
the Notified Body:**

TÜV SÜD Product Service GmbH (No. 0123)  
Ridlerstraße 65  
80339 MÜNCHEN  
Germany

Conformity Assessment was established through the procedure described in Annex IX of EU Regulation 2017/746, including an assessment of the Technical Documentation as described in Annex IX, Chapter II. This declaration is supported by the following certificate(s):

EU Quality Management System Certificate: V12 096981

Conformity of the product with EU Regulation 2017/746 and other applicable EU legislation has been established.

On behalf of Ventana Medical Systems Inc.

Place: Tucson, AZ 85755, USA

Date: 26-Jan-2022

DocuSigned by:

A handwritten signature of "Jeff Boone" is enclosed in a blue rectangular box.

Jeff Boone

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Site Head of Quality Function

Place: Tucson, AZ 85755, USA

Date: 31-Jan-2022

DocuSigned by:

A handwritten signature of "Ben Curson" is enclosed in a blue rectangular box.

Benjamin Curson

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Site Head of Regulatory Affairs Function

# EU Declaration of Conformity

In accordance with EU Regulation 2017/746 of the European Parliament and of the Council of 5 April 2017 on *in vitro* diagnostic medical devices.

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Manufacturer:

Authorized Representative: **Roche Diagnostics GmbH  
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68305 Mannheim  
Germany**

Single Registration Number (SRN) **DE-AR-000006262**  
Authorized Representative:

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## Product Information

Part Number:	Product Name:	Basic UDI-DI:
05266939001 (alternative P/N: 760-2506)	CONFIRM anti-Prostate Specific Antigen (PSA) Rabbit Polyclonal Primary Antibody	761333601234A7

**Intended Purpose:** CONFIRM anti-Prostate Specific Antigen (PSA) Rabbit Polyclonal Primary Antibody is intended for laboratory use in the qualitative immunohistochemical detection of PSA by light microscopy in sections of formalin-fixed, paraffin-embedde tissue stained on a BenchMark IHC/ISH instrument.

**Risk Class:** Class C

**Common Specifications:** Not applicable as no Common Specifications exist for the concerned device.

**Name, Address and Identification number of the Notified Body:** TÜV SÜD Product Service GmbH (No. 0123)  
Ridlerstraße 65  
80339 MÜNCHEN  
Germany

Conformity Assessment was established through the procedure described in Annex IX of EU Regulation 2017/746, including an assessment of the Technical Documentation as described in Annex IX, Chapter II. This declaration is supported by the following certificate(s):

EU Quality Management System Certificate: V12 096981 0003

Conformity of the product with EU Regulation 2017/746 and other applicable EU legislation has been established.

On behalf of Ventana Medical Systems Inc.

Place: Tucson, AZ 85755, USA

Date:

DocuSigned by:

Jeff Boone

09C56B8025B5B4D8...

**Jeff Boone**

Site Head of Quality Function

Place: Tucson, AZ 85755, USA

Date:

DocuSigned by:

Benjamin Curson

0E68FEB8A2C01421...

**Benjamin Curson**

Site Head of Regulatory Affairs Function



# EU Declaration of Conformity

In accordance with EU Regulation 2017/746 of the European Parliament and of the Council of 5 April 2017 on *in vitro* diagnostic medical devices.

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Single Registration Number (SRN) **US-MF-000016993**  
Manufacturer:

Authorized Representative: **Roche Diagnostics GmbH  
Sandhofer Strasse 116  
68305 Mannheim  
Germany**

Single Registration Number (SRN) **DE-AR-000006262**  
Authorized Representative:

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## Product Information

Part Number:	Product Name:	Basic UDI-DI:
05267072001 (alternative P/N: 760-2523)	CONFIRM anti-S100 (Polyclonal) Primary Antibody	761333601207A4

**Intended Purpose:** CONFIRM anti-S100 (Polyclonal) Primary Antibody is a rabbit polyclonal antibody intended for laboratory use in the qualitative immunohistochemical detection of S100 protein by light microscopy in sections of formalin-fixed, paraffin-embedded tissue on a BenchMark IHC/ISH instrument.

**Risk Class:** Class C

**Common Specifications:** Not applicable as no Common Specifications exist for the concerned device.

**Name, Address and Identification number of the Notified Body:** TÜV SÜD Product Service GmbH (No. 0123)  
Ridlerstraße 65  
80339 MÜNCHEN  
Germany

Conformity Assessment was established through the procedure described in Annex IX of EU Regulation 2017/746, including an assessment of the Technical Documentation as described in Annex IX, Chapter II. This declaration is supported by the following certificate(s):

EU Quality Management System Certificate: V12 096981 0003



Conformity of the product with EU Regulation 2017/746 and other applicable EU legislation has been established.

On behalf of Ventana Medical Systems Inc.

Place: Tucson, AZ 85755, USA

22-Jun-2022

Date:

*Jeff Boone*

**Jeff Boone**

Site Head of Quality Function

Place: Tucson, AZ 85755, USA

21-Jun-2022

Date:

*Ben Curson*

**Benjamin Curson**

Site Head of Regulatory Affairs Function



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Single Registration Number (SRN) **US-MF-000016993**  
Manufacturer:

Authorized Representative: **Roche Diagnostics GmbH  
Sandhofer Strasse 116  
68305 Mannheim  
Germany**

Single Registration Number (SRN) **DE-AR-000006262**  
Authorized Representative:

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## Product Information

Part Number:	Product Name:	Basic UDI-DI:
05266688001 (alternative P/N: 760-2018)	Protease 1	761333601904AZ

**Intended Purpose:** Protease 1 is an endopeptidase (alkaline protease) of the serine protease family and cleaves proteins in the tissue section, allowing primary antibodies to recognize and bind epitope(s). The reagent is intended for enzymatic digestion of sections of routine formalin-fixed, paraffin-embedded tissue on a Benchmark IHC/ISH instrument.  
This reagent is intended for *in vitro* diagnostic (IVD) use.

**Risk Class:** Class A

**Common Specifications:** Not applicable as no Common Specifications exist for the concerned device.

Conformity of the product with EU Regulation 2017/746 and other applicable EU legislation has been established.

On behalf of Ventana Medical Systems Inc.

Place: Tucson, AZ 85755, USA

Date: 29-Mar-2022

*Jeff Boone*

**Jeff Boone**

Site Head of Quality Function

Place: Tucson, AZ 85755, USA

Date: 28-Mar-2022

*Ben Curson*

**Benjamin Curson**

Site Head of Regulatory Affairs Function



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Manufacturer:

Authorized Representative: **Roche Diagnostics GmbH  
Sandhofer Strasse 116  
68305 Mannheim  
Germany**

Single Registration Number (SRN) **DE-AR-000006262**  
Authorized Representative:

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## Product Information

Part Number:	Product Name:	Basic UDI-DI:
05279771001 (alternative P/N: 950-102)	EZ Prep Concentrate (10X)	761333601901AT

**Intended Purpose:** Ventana Medical Systems' (Ventana) EZ Prep Concentrate (10X) solution (EZ Prep) is used for paraffin removal from tissue samples during immunohistochemistry and in situ hybridization reactions, and to dilute 2X SSC during stringency washes during in situ hybridization reactions carried out on Ventana automated slide stainers.  
This product is designed for use on BenchMark Series automated slide stainers.  
This product is intended for in vitro diagnostic (IVD) use.

**Risk Class:** Class A

**Common Specifications:** Not applicable as no Common Specifications exist for the concerned device.

Conformity of the product with EU Regulation 2017/746 and other applicable EU legislation has been established.

On behalf of Ventana Medical Systems Inc.

Place: Tucson, AZ 85755, USA

Date: 29-Mar-2022

*Jeff Boone*

**Jeff Boone**

Site Head of Quality Function

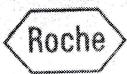
Place: Tucson, AZ 85755, USA

Date: 28-Mar-2022

*Ben Curson*

**Benjamin Curson**

Site Head of Regulatory Affairs Function



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**Tucson, AZ 85755, USA**

Single Registration Number (SRN) **US-MF-000016993**  
 Manufacturer:

Authorized Representative: **Roche Diagnostics GmbH**  
**Sandhofer Strasse 116**  
**68305 Mannheim**  
**Germany**

Single Registration Number (SRN) **DE-AR-000006262**  
 Authorized Representative:

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### Product Information

Part Number:	Product Name:	Basic UDI-DI:
05264839001 (alternative P/N: 650-010)	LCS (Predilute)	7613336012209U

**Intended Purpose:** LCS (Predilute) is a prediluted coverslip solution intended for laboratory use as a barrier between the aqueous reagents and the air. This barrier prevents evaporation, thereby providing a stable aqueous environment for the immunohistochemistry, immunocytochemistry, or *in situ* hybridization reactions on BenchMark GX and BenchMark XT instruments. This reagent is intended for *in vitro* diagnostic use.

**Risk Class:** Class A

**Common Specifications:** Not applicable as no Common Specifications exist for the concerned device.

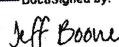
Conformity of the product with EU Regulation 2017/746 and other applicable EU legislation has been established.

On behalf of Ventana Medical Systems Inc.

Place: Tucson, AZ 85755, USA

Date: 22-Feb-2022

DocuSigned by:

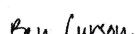
  
 Jeff Boone

Jeff Boone

Site Head of Quality Function

Place: Tucson, AZ 85755, USA

Date: 01-Mar-2022

  
 Ben Curson

Benjamin Curson

Site Head of Regulatory Affairs Function





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Authorized Representative: **Roche Diagnostics GmbH  
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68305 Mannheim  
Germany**

Single Registration Number (SRN) **DE-AR-000006262**  
Authorized Representative:

This declaration is issued under the sole responsibility of Ventana Medical Systems Inc.

## Product Information

Part Number:	Product Name:	Basic UDI-DI:
05446724001 (alternative P/N: 780-003)	ultraView Silver Wash II	761333601900AR

**Intended Purpose:** ultraView Silver Wash II solution is intended for laboratory use to provide an appropriate aqueous environment prior to silver detection chemistry steps on a BenchMark IHC/ISH instrument. This product is intended for *in vitro* diagnostic (IVD) use.

**Risk Class:** Class A

**Common Specifications:** Not applicable as no Common Specifications exist for the concerned device.

Conformity of the product with EU Regulation 2017/746 and other applicable EU legislation has been established.

On behalf of Ventana Medical Systems Inc.

Place: Tucson, AZ 85755, USA

Date: 22-Feb-2022

DocuSigned by:  
  
**Jeff Boone**

Site Head of Quality Function

Place: Tucson, AZ 85755, USA

Date: 01-Mar-2022

DocuSigned by:  
  
**Benjamin Curson**

Site Head of Regulatory Affairs Function



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www.ventana.com

## Declaration of Conformity to 98/79/EC

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Tucson, AZ USA 85755

European Authorized  
Representative: ROCHE DIAGNOSTICS GmbH  
Sandhofer Strasse 116  
D-68305 Mannheim  
Germany

Manufacturing Site: Tucson, AZ USA

Ventana	Roche
REF	

Product name/ Catalogue No. ultraView Universal Alkaline Phosphatase Red Detection Kit 760-501 05269814001

Technical Data File: D018734

Classification: General IVD

Conformity Assessment: 98/79/EC Annex III

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Place of Issue: Tucson, AZ USA 85755

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

Signature: R. Bonner

Date: 11 May 2016

*Roxane Bonner for  
Deepshikha Bhandari*



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Tucson, AZ USA 85755

European Authorized  
Representative: ROCHE DIAGNOSTICS GmbH  
Sandhofer Strasse 116  
D-68305 Mannheim  
Germany

Manufacturing Site: Tucson, AZ USA

	Ventana	Roche
Product name/	Hematoxylin	760-2021
Catalogue No.		05266726001
Technical Data File:	TDF-0043	
Classification:	General IVD	
Conformity Assessment:	98/79/EC Annex III	

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Place of Issue: Tucson, AZ USA 85755

Name of Authorized Signatory: Troy Quander  
Vice President, Regulatory Affairs

Signature:

Date: 5 - March - 2015



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European Authorized  
Representative: ROCHE DIAGNOSTICS GmbH  
Sandhofer Strasse 116  
D-68305 Mannheim  
Germany

Manufacturing Site: Tucson, AZ USA

	Ventana	Roche
	<b>REF</b>	
Product name/	Bluing Reagent	760-2037
Catalogue No.		05266769001
Technical Data File:	TDF 0046	
Classification:	General IVD	
Conformity Assessment:	98/79/EC Annex III	

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Ventana Medical Systems, Inc. maintains a quality system based on EN ISO 13485:2012 / EN ISO 13485:2012 + AC:2012 as certified by TÜV Rheinland LGA Products GmbH (Notified Body No. 0197).

Place of Issue: Tucson, AZ USA 85755

Name of Authorized Signatory: Troy Quander  
Vice President, Regulatory Affairs

Signature:

Date: 10- April 2015



# EU Declaration of Conformity

In accordance with EU Regulation 2017/746 of the European Parliament and of the Council of 5 April 2017 on *in vitro* diagnostic medical devices.

Manufacturer: **Ventana Medical Systems Inc.  
1910 E Innovation Park Drive  
Tucson, AZ 85755, USA**

Single Registration Number (SRN) **US-MF-000016993**  
Manufacturer:

Authorized Representative: **Roche Diagnostics GmbH  
Sandhofer Strasse 116  
68305 Mannheim  
Germany**

Single Registration Number (SRN) **DE-AR-000006262**  
Authorized Representative:

This declaration is issued under the sole responsibility of Ventana Medical Systems Inc.

## Product Information

Part Number:	Product Name:	Basic UDI-DI:
05277965001 (alternative P/N: 790-2208)	Hematoxylin II	761333601224A4

**Intended Purpose:** Hematoxylin II is a modified Mayer's hematoxylin intended for laboratory use in staining cellular nuclei on slides containing cells from frozen tissue, or formalin fixed, paraffin-embedded tissue on a BenchMark IHC/ISH instrument. This reagent is intended as a counterstain to immunohistochemistry, and in situ hybridization applications. This reagent is intended for *in vitro* diagnostic use.

**Risk Class:** Class A

**Common Specifications:** Not applicable as no Common Specifications exist for the concerned device.

Conformity of the product with EU Regulation 2017/746 and other applicable EU legislation has been established.

On behalf of Ventana Medical Systems Inc.

Place: Tucson, AZ 85755, USA

Date: 22-Feb-2022

DocuSigned by:

  
Jeff Boone

D9C56B8025BB4D8

**Jeff Boone**

Site Head of Quality Function

Place: Tucson, AZ 85755, USA

Date: 01-Mar-2022

Ben Curson

**Benjamin Curson**

Site Head of Regulatory Affairs Function

## Declaration of Conformity

as per Annex III of Directive 98/79/EC of the European Parliaments and Council of 27 October 1998

Manufacturer:

Roche Diagnostics GmbH  
Sandhofer Strasse 116  
68305 Mannheim  
Germany

Roche Diagnostics GmbH declares that the product/the product line

*Product name:* VENTANA PD-L1 (SP263) Assay

*Roche Id.-No.:* 07419821001

*Ventana Id.-No.:* 741-4905

to which this declaration relates fulfils the requirements 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).

Mannheim, 28 April 2022

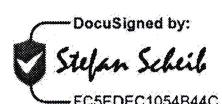
Roche Diagnostics GmbH

ppa./on behalf of the company

DocuSigned by:  
  
Ralf Zielenksi

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Ralf Zielenksi  
Head Q&R Compliance, PRRC RDG  
Centralised and Point of Care Solutions

ppa./on behalf of the company

DocuSigned by:  
  
Stefan Scheib  
FC5EDEC1054B44C...

Dr. Stefan Scheib  
Network Lead Core Lab, Global Regulatory Affairs  
Centralised and Point of Care Solutions

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