



EU Declaration of Conformity

as per Annex IV of the Regulation EU 2017/746 on in-vitro diagnostic medical devices

Manufacturer: Roche Diagnostics GmbH
Address: Sandhofer Strasse 116
 68305 Mannheim
 Germany

Single Registration Number: DE-MF-000006260

Roche Diagnostics GmbH, under the sole responsibility, declares that the product/the product line

Product Name	Cat. No.	Basic UDI-DI
cobas h 232	04901126190	761333601340A7
cobas h 232	04901126041	761333602325AJ
cobas h 232	04901126026	761333602326AL
cobas h 232	04901126016	761333602327AN
cobas h 232	04901126005	761333602328AQ
cobas h 232	04901126045	761333602329AS
cobas h 232	04901126037	761333602330AB
cobas h 232	04901126057	761333602331AD
cobas h 232	04901126060	761333602332AF
cobas h 232	04901126134	761333602333AH
cobas h 232	04901126039	761333602334AK
cobas h 232	04901126051	761333602335AM
cobas h 232	04901126053	761333602336AP
cobas h 232	04901142190	761333601341A9
cobas h 232	04901142041	761333602337AR
cobas h 232	04901142026	761333602338AT
cobas h 232	04901142016	761333602339AV
cobas h 232	04901142005	761333602340AE
cobas h 232	04901142037	761333602341AG
cobas h 232	04901142057	761333602342AJ
cobas h 232	04901142013	761333602343AL
cobas h 232	04901142245	761333602344AN
cobas h 232	04901142060	761333602345AQ
cobas h 232	04901142134	761333602346AS
cobas h 232	04901142039	761333602347AU
cobas h 232	04901142051	761333602348AW
cobas h 232	04901142220	761333602349AY
cobas h 232	04901142053	761333602350AH

**Intended Use:**

The cobas h 232 is an in vitro diagnostic instrument intended for the quantitative evaluation of immunoassays using the Roche CARDIAC test strips in human whole blood. The cobas h 232 is intended for near-patient testing.

Not for self-testing.

Risk Class: ☒ A ☐ B ☐ C ☐ D

Conformity Route: ☒ Self-Declaration of Conformity (Class A)
☐ Technical Documentation Assessment Class B/C – Annex IX
☐ Technical Documentation Assessment Class B/C/D for Self-Testing – Annex IX
☐ Technical Documentation Assessment Class B/C/D for Near-Patient Testing – Annex IX
☐ Technical Documentation Assessment Class C/D for Companion Diagnostics – Annex IX

Certificates: ☐ EU QM Certificate No.:
☐ EU Technical Documentation Assessment Certificate No. (Near-Patient Testing, Self-Testing and Companion Diagnostics):

Other: ☐ Common Specifications:

Notified Body (NB) Name: N/A
 NB Address:

NB Ident. No.: N/A

to which this declaration relates fulfils the requirements of Regulation EU 2017/746 on in-vitro diagnostic medical devices

and

- fulfills the requirements of DIRECTIVE 2011/65/EU of the European Parliament and of the Council of 8 June 2011 (RoHS) on the restriction of the use of certain hazardous substances in electrical and electronic equipment.

and

- fulfills the requirements of Directive 2014/53/EU of the European Parliament and Council of 16 April 2014 (RED) relating to the making available on the market of radio equipment.



Mannheim, 3 July 2023

Roche Diagnostics GmbH

i.V./on behalf of the company

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ppa./on behalf of the company

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