



Declaration of Conformity

as per Annex IV of the Directive 98/79/EC of the European Parliament and the Council of 27 October 1998 via BSI as the Notified Body (registration no. BSI-NL 2797)

Document No.: **DOC-2022-14**

Manufacturer: **Roche Molecular Systems, Inc.
1080 US Highway 202 South
Branchburg, NJ 08876
USA**

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

Name, Address and Identification number of the Notified Body: **BSI Group The Netherlands B.V.
Notified Body Number: 2797
Say Building, John M. Keynesplein 9, 1066 EP
Amsterdam, Netherlands**

Roche Molecular Systems, Inc. declares that the *in vitro* diagnostic medical device:

Product Name: **cobas**[®] HBV/HCV/HIV-1 Control Kit
*Positive control kit for use on the **cobas**[®] 5800/6800/8800 Systems*

P/N: 09040773190

Complies with the requirements of the Directive 98/79/EC of the European Parliament and of the Council of 27 October 1998 on *in vitro* diagnostic medical devices.

This declaration is supported by the following certificates:

EC Certificate – Full Quality Assurance: CE 707974, first issued 2019-03-26, valid until 2026-05-26
EC Design-Examination Certificate: CE 709228, first issued 2019-03-26, valid until 2025-05-26

The Manufacturer agrees to develop, implement, and maintain a documented post-production monitoring process, including the notification of reportable events, under the European Medical Device Vigilance System Guidelines. As a legal manufacturer, Roche Molecular Systems, Inc., is solely responsible for the product. This declaration is issued under the sole responsibility of Roche Molecular Systems, Inc.



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Place: Tucson, AZ

Date: 19-May-2022

Jeff Boone

Jeff Boone
Vice President, Quality Management

Place: Pleasanton, CA

Date: 17-May-2022

Rita Hoady

Rita Hoady
Network Lead Molecular Lab
Director, Global Regulatory Affairs