



EC- Declaration of Conformity

According to Annex III of the Directive 98/79/EC of the European Parliament and of the Council of 27. October 1998

Manufacturer: Hitachi High-Technologies Corporation
1-24-14 Nishi-Shinbashi
Minato-ku, Tokyo 105-8717, Japan

Authorised Representative: Roche Diagnostics GmbH
Sandhofer Straße 116
D-68305 Mannheim, Germany

Roche Diagnostics GmbH declares that the in vitro diagnostic medical instrument

Product name: cobas® 6000 system

Configuration information: Composed of 1 to 3 modules of cobas c 501 and/or cobas e 601 in any combination with the exception of 3 x cobas c 501 or 3 x cobas e 601

Description: Integrated system for clinical and immunological tests, intended for in-vitro determination of a wide range of analytes.

relating to this declaration, complies with the requirements of EC Directive 98/79/EC of the Council of October 27, 1998 concerning in-vitro diagnostic medical devices.

Mannheim, June 8, 2006

Roche Diagnostics GmbH

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