EG-Konformitätserklärung/*EC Declaration of Conformity*

gemäß Anhang IV der Richtlinie 98/79/EG des Europäischen Parlaments und des Rates vom 27. Oktober 1998 mit TÜV SÜD Product Service GmbH (Ridlerstraße 65, 80339 München, Germany) als Notified Body (Nr. 0123)

as per Annex IV of Directive 98/79/EC of the European Parliaments and Council of 27 October 1998 via TÜV SÜD Product Service GmbH (Ridlerstrasse 65, 80339 Munich, Germany) as the Notified Body (No. 0123)

Hersteller/*Manufacturer*:	Roche Diagnostics GmbH
Adresse/*Address:*	Roche Professional Diagnostics Sandhofer Straße 116 D-68305 Mannheim

Die Roche Diagnostics GmbH erklärt, dass das Produkt/die Produktfamilie (bei rezepturgleichen Produkten)
Roche Diagnostics GmbH declares that the product/the product line (in case of products manufactured by identical recipes)

Produktname/*Product name*:	**PreciControl Anti-HCV**
Art.-Nr./*Id. No.*:	03290379
Beschreibung/*Description*:	PreciControl Anti-HCV dient zur Qualitätskontrolle der Anti-HCV und Anti-HCV II Immunoassays an Elecsys und cobas e Immunoassay-Systemen. *PreciControl Anti-HCV is used for quality control of the Anti-HCV and Anti-HCV II immunoassays on the Elecsys and cobas e immunoassay analyzers.*

auf das/die sich diese Erklärung bezieht, den Forderungen der EG-Richtlinie 98/79/EG des Rates vom 27. Oktober 1998 (bzw. seine Umsetzung in nationales Recht der Mitgliedsstaaten in welchen das Produkt vermarktet werden soll) über In-vitro-Diagnostica entspricht.
to which this declaration relates fulfils the requirements of EC Directive 98/79/EC of the Council of 27 October 1998 (and its relevant transposition into the national laws of the Member States in which the device is intended to be placed on the market) concerning in-vitro diagnostic devices.

Mannheim, 23.Dezember 2011

Roche Diagnostics GmbH
ppa./*on behalf of the company*

Dr. M. Thein
Head of Quality
Roche Professional Diagnostics

i. V./*on behalf of the company*

Dr. B. Rauschel
Head of Quality Control Penzberg
Roche Diagnostics Global Operations

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ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Product Service

EC Certificate

EC Design-Examination Certificate

Directive 98/79/EC on In Vitro Diagnostic Medical Devices (IVDD), Annex IV (4) (List A)

No. V7 11 02 10283 243

Manufacturer: **Roche Diagnostics GmbH**
Sandhofer Str. 116
68305 Mannheim
GERMANY

Product: **Screening test for Hepatitis C marker**

Model(s): **Anti-HCV**
PreciControl Anti-HCV
Elecsys and cobas e analyzers

Parameters:

Model Name:	Model No.:
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Anti-HCV	03290352
PreciContol Anti-HCV	03290379
Elecsys and cobas e analyzers	

The Certification Body of TÜV SÜD Product Service GmbH declares that a design examination has been carried out on the respective devices in accordance with IVDD Annex IV (4). The design of the devices conforms to the requirements of this Directive. See also notes overleaf.

Report No.: 71380757-02

Valid from: 2011-02-22
Valid until: 2012-08-04

Date, 2011-02-23

Hans-Heiner Junker

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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