Roche

Diagnostics

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: Roche Instrument Center AG
Forrenstrasse
CH-6343 Rotkreuz, Switzerland

Roche Instrument Center AG declares that the in-vitro diagnostic medical instrument

Product name: **cobas c 111 with ISE**

Description: In-vitro diagnostic analyzer performing clinical chemistry, specific protein and electrolyte tests. Analytes are measured photometrically or turbidimetrically or with ion sensitive electrodes

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.

Rotkreuz, October 18, 2006

Roche Instrument Center AG

J. Brand
Head of Instrumentation

Dr. A. Bayer
Head of Quality Management

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