

Certificate

The Certification Body of
TÜV Rheinland Product Safety GmbH

hereby certifies that the organization

DRG Instruments GmbH
Frauenbergstr. 18
35039 Marburg
Deutschland

has established and applies a quality management system for medical devices
for the following scope:

**Design and Development, Manufacture and Distribution
of in vitro Diagnostics by Enzymeimmunoassays (ELISAS)
and Radioimmunoassays (RIAS) Technology**

Proof has been furnished that the requirements specified in

EN ISO 13485:2003 + AC:2007

are fulfilled. The quality management system is subject to yearly surveillance.

Certificate Registration No.: SX 60024027 0001

An audit was performed. Report No.: 21138422 002

This Certificate is valid until: 04.02.2014




Akkreditiert durch
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln
und Medizinprodukten
ZLG-ZQ-995.00.01-46

Cologne, 05.02.2009



Certification Body


Dr. H. Lüdemann

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