

EC – DECLARATION OF CONFORMITY CE-IVD MARKING (In-Vitro-Diagnostic-Devices)

Hereby and with the CE-sign it is confirmed that following products, manufactured by AHN Biotechnologie GmbH, Uthleber Weg 14, D-99734 Nordhausen, Germany, meets the fundamental regulations of the directive of the council of the European community to the adjustment of legal rules of the member states regarding "In-vitro-Diagnostics" 98/79/EG, 27.10.1998:

Products:	AHN pipet4u Digital Micropipettes, Production Lot 2010
Serial Numbers:	800001 – 899999
Products:	AHN Racks for pipet4u Micropipettes, Production Lot 2010
Size:	Racks for 4 and 6 Micropipettes, Racks for 8-channel Pipette
Products:	AHN pipette tips and filter tips, Production Lot 2010
Volume ranges:	0.1-10 µl, 0.5-10/20 µl, 1-200 µl, 5-300 µl, 100-1000 µl, 1-5 ml, 1-5,5 ml, 1-10 ml
Products:	AHN PCR-tubes/strips and qPCR-Capillaries, Production Lot 2010
Volume ranges:	0.2 and 0.5 ml Volume
Products:	AHN Microtubes and Cryotubes, Production Lot 2010
Volume ranges:	0.5 ml, 1.5 mm, 1.8 mm and 2.0 ml Volume
Products:	AHN Spin Columns and Filterplates, Production Lot 2010
Volume ranges:	0.8 ml and 1 ml Volume
Products:	AHN Cryobox and Storage Box, Production Lot 2010
Size:	100-places
Products:	AHN Gelloader tips, Production Lot 2010
Volume ranges:	1-200 µl Volume
Products:	AHN Gelloader filter tips, Production Lot 2010
Volume ranges:	30 µl, 100 µl Volume
Products:	AHN Rackboxes, Production Lot 2010
Sizes:	60-places and 96-places
Products:	AHN ULTRAPrep Miniprep DNA-Purification Kits, Production Lot 2010
Sizes:	50 reactions and 250 reactions
Products:	AHN ULTRAPrep Miniprep RNA-Purification Kits, Production Lot 2010
Sizes:	50 reactions and 250 reactions

To evaluate the conformity of the products, the following standards were taken into consideration:

European Standard ISO 8655, part 1, part 2 and part 6, piston operated volumetric apparatus and pipette tips. General obligation to notify pursuant to §§ 25 and 30 (2) German Medical devices Act. European Standard 98/79/EU for In-Vitro-Diagnostics Medical Devices. The standards used for the measurements are traceable to the national standards of the Federal Republic of Germany at the Physikalisch-Technische Bundesanstalt (PTB). Competent authority code DE/CA90, Thüringer Landesamt für Lebensmittelsicherheit und Verbraucherschutz, Abteilung 2, Dezernat 24 (Pharmazie). This declaration is made under the sole responsibility of the producer.

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Nordhausen, 2010-02-17

This certificate was sent online and is valid without signature.

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