greiner bio-one

EC DECLARATION OF CONFORMITY

according to annex III of the Directive 98/79/EC of the European Parliament and of the Council of 27. Oct. 1998 on in vitro diagnostic medical devices

Manufacturer:	Greiner Bio-One GmbH
Address:	Bad Haller Str. 32 A-4550 Kremsmünster AUSTRIA
Production Location:	Greiner Bio-One GmbH Bad Haller Str. 32 A-4550 Kremsmünster AUSTRIA
Product	**VACUETTE® Transport Box (VTB)**
Classification	Directive 98/79/EC; Other device (all devices except Annex II and self-testing devices)

We herewith declare under our sole responsibility that the above mentioned products meet the provisions of the following EC Council Directives and applicable Standards. All supporting documentations are retained under the premises of the manufacturer.

This product has been CE marked since 29 Oct 2003.

DIRECTIVES

Directive 98/79/EC of the European Parliament and of the Council of 27. Oct. 1998 on in vitro diagnostic medical devices

STANDARDS

Standards applicable are:
ISO 9001, EN ISO 13485

Place, Date: Kremsmünster, 02. FEB. 2010

Signature: Gabriele Rose

Gabriele Rose
QM, Regulatory Affairs