



HANS RUDOLPH, inc.
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Date of Issue: May 5, 1999

Authorized Representative in EU:



Medical Devices Group
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EC DECLARATION OF CONFORMITY

Council Directive 93/42/EEC on Medical Devices

Manufacture's Name: HANS RUDOLPH, inc.

Manufacture's Address: 8325 Cole Parkway, Shawnee KS 66227 USA

Type of Equipment: Volume Calibration Syringes

Product Series (Model Number): 5520, 5510, 5550, 5540, 5530, 5570 and 4900

Classification: I

Rule(s) used for Classification: 1

**Harmonized Standards to which Conformity is Declared: EN ISO 5356-1:2004,
EN 980:2003, EN 1041:1998**

Hans Rudolph, inc. (HRI) declares that the above mentioned devices are in conformity with the applicable provisions of the EC Council Directive, 93/42/EEC of 14 June 1993, Annex VII and are entitled to bear the CE Mark.

Approved By: 

Date: JAN 4, 2008

Kevin Rudolph
Vice President of Hans Rudolph, inc.

Declaration of Conformity Certificate Number: 691184 Rev F