HANS RUDOLPH, inc.
8325 COLE PARKWAY
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Authorized Representative in EU: EC REP

Medical Devices Group
Marlborough House and Riding Street
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EC DECLARATION OF CONFORMITY
Council Directive 93/42/EEC Annex VII
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, ISO 15223-1:2012 , EN 1041:2008

Hans Rudolph, inc. (HRI) declares that the above mentioned devices are in conformity with the applicable provisions of the Medical Device Directive: Council Directive 93/42/EEC, as amended by Directive 2007/47/EC, Annex VII and are entitled to bear the CE Mark.

Approved By: _____ **Date:** MAY 5, 2014

Kevin Rudolph
CEO of Hans Rudolph, inc.