Compliance Statement for Directive 2002/95/EC (RoHS)

bluepoint MEDICAL GmbH&Co.KG hereby declares on behalf of itself that the substances listed in Directive 2002/95/EC of the European Parliament and of the Council on the Restriction of the Use of Certain Hazardous Substances in Electrical and Electronic Equipment ("RoHS") are not intentionally added to any of bluepoint MEDICAL GmbH&Co. KG's SpiroTrue®H, SpiroTrue®D, SpiroTrue®PED SpiroTrue®HD, SpiroTrue®A and SpiroTrue®APC products.

This statement is based on bluepoint MEDICAL GmbH&Co. KG's current level of knowledge. Since conditions of use by customers and others are outside bluepoint MEDICAL GmbH&Co. KG 's control, bluepoint MEDICAL GmbH&Co.KG makes no warranties, express or implied, and assumes no liability in connection with any use of this information.

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Place, Date: Selmsdorf, 02 July 2014

Legally binding signature:

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Bernd Lindner
(Managing Director)

EU Directive 2002/95/EC (RoHS), restricts the use of the following substances in the manufacture of Electronic and Electrical Equipment imported into the EU after June 30, 2006,

- Lead
  Mercury
  Hexavalent Chromium (Chrome 6)
  Flame retardants
  - Polybrominated Biphenyls (PBB)
  - Polybrominated Diphenyl Ethers (PBDE)

  The maximum allowable concentration is expected to be 0.1% (1000 ppm) by weight in homogeneous material. Homogeneous material means a unit that cannot be mechanically disjointed into single materials.

- Cadmium
  The maximum allowable concentration is expected to be 0.01% (100 ppm) by weight in homogeneous material.