RoHS Recast
Declaration of Conformity

We, Auritec Medizindiagnostische Systeme GmbH, Dernauer Straße 12, 22047 Hamburg, Germany, hereby declare under our sole responsibility that all aour medical equipment the provisions of Directive 2011/65/EU of the European Parliament and of the Council of 8th June 2011 to meet restriction of use of certain hazardous substances in electrical and electronic equipment.

Auritec is in compliance with the following European Union Directives and is actively restricting the use of designate substances.

- 2011/65/EU RoHS Recast Directive (RoHS-II)

Restriction of Hazardous Substances	Content of Compliance	
Lead – Pb	<0,1%	(1000ppm)
Mercury – Hg	<0,1%	(1000ppm)
Cadmium – Cd	<0,01%	(100ppm)
Hexavalent Chrome – (Cr+6)	<0,1%	(1000ppm)
Polybrominated Biphenyls – PBB	<0,1%	(1000ppm)
Polybrominated Diphenyl Etehrs - PBDE	<0,1%	(1000ppm)

This declaration only applies medical device with original accessories.

Hamburg, 27.03.2014

Michael Knuhr
Quality Manager