Rauland-Borg Corporation

TO WHOM IT MAY CONCERN

QUALITY ASSURANCE

The Responder 4000 communications system is listed by Underwriter's Laboratories under UL Standard 1069 - 6th Edition (published March, 2001).

The Responder 4000 communications system is compliant with European Union Medical Device Directive 93/42/EEC (MDD).

The Responder 4000 communications system is compliant with CE Marking for Conformity of Medical Devices, in compliance with Article 17 of Medical Device Directive 93/42/EEC (MDD), and EU Council Directive 93/68/EEC.

The Responder 4000 communications system is supplied by an FDA-registered medical device manufacturer, and is list their device in accordance with the provisions set forth in 21CFR Part 890.3710 or 890.3725, Class II Medical Device.

The Responder 4000 communications system is in compliance with European Union's DIRECTIVE 2002/95/EC OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 27 January 2003, commonly known as the RoHS Directive.

Sincerely,

Jonathan Wacks 4/19/10
Director, Regulatory Affairs

RAULAND-BORG CORPORATION