RoHS Adherence

Tuttlingen, den 03. April 2018

Dear Sir or Madam,

We hereby inform you that the product groups

- Medical devices as per Directive 93/42/EEC and their accessories and replacement parts beginning 22 July 2014
- In vitro diagnostics products as per Directive 98/79/EC and their accessories and replacement parts beginning 22 July 2016


The EC conformity declarations of the products are expanded to include the provisions of the co-valid Directive 2011/65/EU. They belong to the operating instructions of every delivered product.

We ask that you understand that we will not fill out any standard questionnaires regarding this matter.

Mit freundlichen Grüßen / Best regards,
Andreas Hettich GmbH & Co. KG

i.A.

Christian von der Grün
Head of Regulatory- & Quality Affairs