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To our customers

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**RoHS 2011/65/EU compliance and announcement of future compliance to the delegated directive (EU) 2015/863**

Tuttlingen, 2020-09-30

Dear Sirs or Madam,

We hereby inform you that Andreas Hettich GmbH & Co. KG offers the products of the product groups

- Medical devices according to Directive 93/42/EEC, as well as their accessories and spare parts from 2014-07-22
- In vitro diagnostic products according to Directive 98/79/EC, and their accessories and spare parts from 2016-07-22

in compliance with EU Directive 2011/65/EU of the European Parliament and of the Council of 2011-06-08 on the restriction of the use of certain hazardous substances in electrical and electronic equipment (RoHS II).

The EC Declarations of Conformity of these products include the 2011/65/EU as a co-existing directive. They are part of the operating instructions of each delivered product.

Furthermore, we would like to inform you that we are currently working on the implementation of the delegated directive (EU) 2015/863 which will be valid for us from the middle of next year. Currently we don't have all information that we need from our suppliers. Therefore, we ask for your understanding that no declaration of conformity can be issued at this time.

Please also understand that we do not process standard questionnaires on this issue.

Best regards,
**Andreas Hettich GmbH & Co. KG**

i. A.

Michael Eberhard
Head of Regulatory & Quality Affairs

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