Declaration of Conformity

Date: 03 Sep 2015

The Products specified below are defined as in vitro diagnostic medical devices as specified in Article 1 of EC Directive 98/79/EC on In Vitro Diagnostic Medical Devices and conform with the Essential Requirements listed in Annex I of EC Directive 98/79/EC together with Statutory Instrument 2002 No. 618 The UK Medical Devices Regulations 2002.

125 Series -	**Polystyrene Container, 60ml**
128 Series -	**Universal Polystyrene Container, 30ml**
129 Series -	**Polystyrene Containers 7ml Bijou**
131 Series -	**Polystyrene Urine Tube**
165 Series -	**Polystyrene Container, 150ml**
185 Series -	**Polystyrene Container, 100ml**
190 Series -	**Polystyrene Container, 250ml**

The intended use of these products is as specimen receptacles and as detailed in Article 9 of EC Directive 98/79/EC the products are classified as General Devices. Conformity is achieved following the procedure detailed in Annex III of EC Directive 98/79/EC.

Signed on behalf of Thermo Fisher Scientific:

Robert Tomsa
Operations Director & Site Leader

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