EC Declaration of Conformity
In accordance with EN ISO/IEC 17050-1:2004

We, **Sakura Finetek Europe B.V., Flemingweg 10A, 2408 AV, Alphen aan den Rijn, The Netherlands**

as Legal Manufacturer declare that:

Product: Slides
Product name/number: Tissue-Tek® AutoWrite® Non-Adhesive Slides /

Tissue-Tek® AutoWrite® Non-Adhesive Slides, Ground Edges 90º, Clipped Corner White, 50x50/pcs	9562
Tissue-Tek® AutoWrite® Non-Adhesive Slides, Ground Edges 90º, Clipped Corner Blue, 50x50/pcs	9563
Tissue-Tek® AutoWrite® Non-Adhesive Slides, Ground Edges 90º, Clipped Corner Pink, 50x50/pcs	9564
Tissue-Tek® AutoWrite® Non-Adhesive Slides, Ground Edges 90º, Clipped Corner Yellow, 50x50/pcs	9565
Tissue-Tek® AutoWrite® Non-Adhesive Slides, Ground Edges 90º, Clipped Corner Green, 50x50/pcs	9566
Tissue-Tek® AutoWrite® Non-Adhesive Slides, Ground Edges 90º, Clipped Corner Orange, 50x50/pcs	9567
Tissue-Tek® AutoWrite® Non-Adhesive Slides, Ground Edges 90º, Clipped Corner Purple, 50x50/pcs	9568

are manufactured in accordance with the following Directive:

98/79/EC	Conforms with the essential requirements of the In Vitro Diagnostics Directive and its amending directives. Classification: Other (General). Conformity Assessment route: Annex III applied.

In addition the following internal standard applies:

ISO 9001:2015 Quality Management System requirements.

I hereby declare that the products named above have been tested and found to comply with the relevant sections of the above referenced specifications. The products comply with all essential requirements of the Directive.

Signed:

C.Koeman
General Manager

Alphen aan den Rijn, 15 October 2018