



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® Adhesive Slides /

Tissue-Tek® AutoWrite® Adhesive Slides, Ground Edges 90°, Clipped Corner White, 20x72/pcs	9597
Tissue-Tek® AutoWrite® Adhesive Slides, Ground Edges 90°, Clipped Corner Blue, 20x72/pcs	9598
Tissue-Tek® AutoWrite® Adhesive Slides, Ground Edges 90°, Clipped Corner Pink, 20x72/pcs	9599
Tissue-Tek® AutoWrite® Non-Adhesive Slides, Ground Edges 90°, Clipped Corner Yellow, 20x72/pcs	9600
Tissue-Tek® AutoWrite® Non-Adhesive Slides, Ground Edges 90°, Clipped Corner Green, 20x72/pcs	9601
Tissue-Tek® AutoWrite® Non-Adhesive Slides, Ground Edges 90°, Clipped Corner Orange, 20x72/pcs	9602
Tissue-Tek® AutoWrite® Non-Adhesive Slides, Ground Edges 90°, Clipped Corner Purple, 20x72/pcs	9603

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.
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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