



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: Coverslips
Product name/number: Tissue-Tek® Glas™ Coverslips /

Tissue-Tek® Glas™ Coverslips, 24x55 mm, 10x100/pcs	9579
Tissue-Tek® Glas™ Coverslips, 24x40 mm, 10x100/pcs	9580
Tissue-Tek® Glas™ Coverslips, 24x50 mm, 10x100/pcs	9581
Tissue-Tek® Glas™ Coverslips, 24x60 mm, 10x100/pcs	9582

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