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:	Mounting Media
Product name/number:	Tissue-Tek® Glas™ Mounting Medium, 500 mL / 1408N
	Tissue-Tek® Glas™ Pertex, 500 mL / 1410
	Tissue-Tek® Glas™ Tissue-Mount™, 500 mL / 1467N

is 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 product(s) 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:

Alphen aan den Rijn, 15 October 2018

Digitally signed by Bolocan Vladimir
Date: 2019.04.11 12:28:47 EEST
Reason: MoldSign Signature
Location: Moldova