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:	Uni-Cassette® System
Product name/number:	Tissue-Tek® Uni-Cassette® 4154F, 4155F, 4156F, 4157F, 4158F, 4170, 8154, 8155, 8156, 8157, 8158 and 8170 Tissue-Tek® Uni-Cassette® Biopsy Cassette System 4086, 4087, 4088, 4089, 4090, 4174, 8086, 8087, 8088, 8089, 8090 and 8174 Tissue-Tek® Uni-Cassette® Processing/Embedding Cassette 4179, 4182, 4183, 4184, 4185, 4187, 4199, 8130, 8179, 8182, 8183, 8184, 8185, 8187

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:

Digitally signed by Şarco Alexandru
Date: 2020.12.17 12:48:54 EET
Reason: MoldSign Signature
Location: Moldova

Alphen aan den Rijn, 10 September 2020