EC Declaration of Conformity

In accordance with EN ISO/IEC 17050-1:2004

We, as Authorized representative:

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

declare that:

Equipment:	Cassettes
Model name/number:	Tissue-Tek® Uni/ Biopsy/Embedding, 4117-4120, 4135, 4153-4158, 4170, 4073-4077, 4045, 4086-4090, 4172, 4174, 4125-4129, 4179, 4182-4187, 4191, 4199, 9229, 4173, 4073-4077, 4306-4309, 4312-4314.

Manufactured for:

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

Is in accordance with the following Directive:

98/79/EC	Conforms to the essential requirements of the In Vitro Diagnostics Directive and its amending directives

has been designed and manufactured to the relevant parts of the following standards:

EN 14254 (2001)

I hereby declare that the equipment named above has been tested and found to comply with the relevant sections of the above referenced specifications. The unit complies with all essential requirements of the Directives.

Signed by:

Name: W. Buijteweg
Position: President

Done at: Sakura Finetek Europe B.V.
On: 01/04/2010

Digitally signed by Şarco Alexandru
Date: 2021.12.21 16:41:55 EET
Reason: MoldSign Signature
Location: Moldova

CE