



## 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: Grossing, Trimming and dissecting products  
Model name/number: Tissue Tek® Accu- Edge®  
See Annex I for products

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 with the essential requirements of the In Vitro Diagnostics Directive and its amending directives

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

ISO 13485:2003, ISO 14971:2000, EN 980:2003, ISO 15223:2000, ISO 14644-1:1999.

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: 01/04/2011

