

Certificate of CE-Registration

This is to certify that, in accordance with the *In Vitro* Diagnostic Medical Device Directive 98/79/EC, **CEpartner4U BV** agrees to perform all duties and responsibilities as the Authorized Representative for:

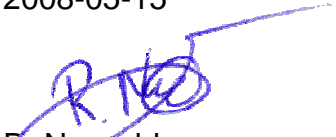
Safe-Tec Clinical Products, LLC.
142 Railroad Drive
Ivyland, PA 18974
United States

as stipulated and demanded by the aforementioned Directive.
The Dutch Competent Authorities have accepted the manufacturer's medical device registration by CEpartner4U as listed below:

Description	Date of registration	EDMS code
MICROSAFE® TUBE	2008-05-15	51 01 01 01

The manufacturer has provided CEpartner4U with all necessary documentation, together with an appropriate Declaration of Conformity that the IVD medical devices fulfil the essential requirements of Directive 98/79/EC.

2008-05-15



R. Nusselder
CEpartner4U BV

C e p a r t n e r 4 U

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