Certificate number: 2022-IVDR/CE374

Certificate of CE-Registration

This is to certify that, in accordance with the *In Vitro* Diagnostic Medical Device Regulation (EU) 2017/746, **CEpartner4U BV** agrees to perform all duties and responsibilities as the Authorized Representative for

SAFE-TEC Clinical Products LLC

500 Parkway
Broomall, PA 19008
United States

as stipulated and demanded by the aforementioned Regulation.

IVDR devices were registered on the 2nd of June, 2022 under number:

Product Group	**Registration number**
MicroSafe® Tubes	NL-CA002-27916

see appendix

The manufacturer has provided CEpartner4U with all necessary documentation, together with an appropriate Declaration of Conformity that the IVD medical devices fulfil the general safety and performance requirements of Regulation (EU) 2017/746.

Issue date: 2022-08-23

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CEpartner4U BV

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3951 DB Maarn NL
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www.cepartner4u.nl

Appendix

Model Name	Model number	UDI-DI Number
5 Microliters MICROSAFE®, 2000 tubes	1005	0560586450137
10 Microliters MICROSAFE®, 2000 tubes	1010S	0560586450144
15 Microliters MICROSAFE®, 2000 tubes	1015	0560586450151
17 Microliters MICROSAFE®, 2000 tubes	1017	0560586450168
20 Microliters MICROSAFE®, 2000 tubes	1020	0560586450175
25 Microliters MICROSAFE®, 2000 tubes	1025	0560586450182
30 Microliters MICROSAFE®, 2000 tubes	1030	0560586450199
35 Microliters MICROSAFE®, 2000 tubes	1035	0560586450205
40 Microliters MICROSAFE®, 2000 tubes	1040	0560586450212
50 Microliters MICROSAFE®, 2000 tubes	1050	0560586450229
60 Microliters MICROSAFE®, 2000 tubes	1060	0560586450236
75 Microliters MICROSAFE®, 2000 tubes	1075	0560586450243
80 Microliters MICROSAFE®, 2000 tubes	1080	0560586450250