RADIOMETER

EC Declaration of Conformity

Radiometer Medical ApS
Åkandevej 21
DK-2700 Brønshøj
Denmark

We hereby declare that the product(s) described below meets the applicable requirements of Directive 98/79/EC of the European Parliament and of the Council of October 27, 1998, on *in vitro* diagnostic medical devices (IVDD) as specified in Annex III.

Class: ☒ General ☐ Annex II/List A ☐ Annex II/List B

Product family: Glass capillaries for blood sampling

Name	Ref. No.	GMDN	First CE-mark
Clinitubes D956-70-35	905-661	34590	2002-02
Clinitubes D956-70-100	905-663	34590	2002-01
Clinitubes D956-70-125	905-664	34590	2002-02
Clinitubes D956-70-210	905-666	34590	2002-01
Clinitubes D957G-70-35	942-869	34590	2002-01
Clinitubes D941G-240-85	942-875	34590	2002-01
Clinitubes D941G-240-85	942-876	34590	2002-01
Clinitubes D957G-70-100	942-878	34590	2002-01
Clinitubes D957G-70-125	942-880	34590	2002-01
Clinitubes D941G-80-140	942-882	34590	2002-01
Clinitubes D957G-70-210	942-884	34590	2002-01
Clinitubes D941G-240-55	942-885	34590	2002-01
Clinitubes D941G-240-55	942-873	34590	2002-01

Issuance:

Name: Gitte Juel Friis
Title: Director Regulatory Affairs

Place: Copenhagen, Denmark

Signature: Gitte Juel Friis

Date: 2013-01-31