CERTIFICATE OF REGISTRATION

MedNet EC-REP GmbH
Borkstraße 10
48163 Münster
Germany

in its function of the European Authorized Representative, in accordance with the In Vitro Diagnostic Directive 98/79/EC, hereby confirms the registration of the following in vitro diagnostic medical device(s) into the German DIMDI data base

2019-New Coronavirus IgM Rapid Test Cassette (WB/S/P)
DIMDI Registration Number DE/CA22/1311-14.1-IVD

2019-New Coronavirus IgM/IgG Rapid Test Cassette (WB/S/P)
DIMDI Registration Number DE/CA22/1311-15.1-IVD

on behalf of

Beijing Beier Bioengineering Co., Ltd.
NO.99, ChuangXin road LuCheng Industrial development zone, HuangCun Town, Daxing district, Beijing, China

according to the directive 98/79/EC of the European Parliament and of the Council of the European Union relating to *in vitro* diagnostic medical devices.

Münster, 07 May 2020 MEDNET EC-REP

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