

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

COVID-19 Antigen Rapid Test Kit, Other device
DIMDI Registration Number DE/CA22/1311-419.1-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, 09.10.2020




on behalf of MedNet EC-REP GmbH

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