CERTIFICATE OF REGISTRATION

MedNet 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 into the German DIMDI data base

see attached list

on behalf of

ACON Biotech (Hangzhou) Co., Ltd.
No. 210 Zhenzhong Road
West Lake District
310030 Hangzhou
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, October 30th, 2020

i.A.
David Thaler
MedNet GmbH

Network to Market
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List of registered devices

Registration Number	Classification	Product Name
DE/CA22/419-1813.2-IVD	Other device	SARS-CoV-2 IgG/IgM Rapid Test
DE/CA22/419-1826-IVD	Other device	SARS-CoV-2 IgG EIA Test Kit
DE/CA22/419-1827-IVD	Other device	SARS-CoV-2 IgM EIA Test Kit
DE/CA22/419-1830-IVD	Other device	NES-32 Nucleic Acid Extraction System
DE/CA22/419-1831-IVD	Other device	Nucleic Acid(DNA) Extraction Kit
DE/CA22/419-1832-IVD	Other device	Nucleic Acid(RNA) Extraction Kit
DE/CA22/419-1833-IVD	Other device	Whole Blood Genomic DNA Extraction Kit
DE/CA22/419-1836-IVD	Other device	SARS-CoV-2 RT-PCR Test Kit
DE/CA22/419-1842-IVD	Other device	Viral DNA/RNA Isolation Kit
DE/CA22/419-1856-IVD	Other device	SARS-CoV-2 Antigen Rapid Test

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