



## 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



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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