ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-245.10.07

Product Service

EC Certificate

EC Design-Examination Certificate
Directive 98/79/EC on In Vitro Diagnostic Medical Devices (IVDD), Annex III (6)
(Devices for self-testing)

No. V9 090019 0003 Rev. 04

Manufacturer: **MP Biomedicals Germany GmbH**
Thüringer Straße 15
37269 Eschwege
GERMANY

Product: **In Vitro diagnostic devices for self testing**

Model(s): **Rapid SARS-CoV-2 Antigen Test Card**

Parameters:

Model Name:	Model Nº:
Rapid SARS-CoV-2 Antigen Test Card	REF 07AG6001BS
Rapid SARS-CoV-2 Antigen Test Card	REF 07AG6005BS
Rapid SARS-CoV-2 Antigen Test Card	REF 07AG6007BS
Rapid SARS-CoV-2 Antigen Test Card	REF 07AG6008BS
Rapid SARS-CoV-2 Antigen Test Card	REF 07AG6020BS
Rapid SARS-CoV-2 Antigen Test Card	REF 07AG6001BSNL
Rapid SARS-CoV-2 Antigen Test Card	REF 07AG6005BSNL

The Certification Body of TÜV SÜD Product Service GmbH declares that a design examination has been carried out on the respective devices in accordance with IVDD Annex III (6). The design of the devices conforms to the requirements of this Directive. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:V9 090019 0003 Rev. 04

Report No.: 713237599 / 713239369-02

Valid from: 2022-04-29
Valid until: 2024-05-26

Date, 2022-04-29

Christoph Dicks
Head of Certification/Notified Body

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany