





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 061317 0006 Rev. 00

Manufacturer: Xiamen Boson Biotech Co., Ltd.

> 90-94 Tianfeng Road Jimei North Industrial Park 361021 Xiamen, Fujian

PEOPLE'S REPUBLIC OF CHINA

Product: In Vitro diagnostic devices for self testing

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

Model Name: Model No .: Parameters:

> Rapid SARS-CoV-2 Antigen Test Card REF 1N40C5-2 Rapid SARS-CoV-2 Antigen Test Card REF 1N40C5-4 Rapid SARS-CoV-2 Antigen Test Card REF 1N40C5-6

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 061317 0006 Rev. 00

713210321 Report No.:

Valid from: 2021-04-01 Valid until: 2022-05-26

2021-04-01 Date.

Christoph Dicks

Head of Certification/Notified Body

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