



# 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. 01**

<b>Manufacturer:</b>	<b>Xiamen Boson Biotech Co., Ltd.</b>	
	90-94 Tianfeng Road Jimei North Industrial Park 361021 Xiamen, Fujian PEOPLE'S REPUBLIC OF CHINA	
<b>Product:</b>	<b>In Vitro diagnostic devices for self testing</b>	
<b>Model(s):</b>	<b>Rapid SARS-CoV-2 Antigen Test Card</b>	
<b>Parameters:</b>	Model Name:	Model No.:
	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
	Rapid SARS-CoV-2 Antigen Test Card	REF 1N40C5-7
	Rapid SARS-CoV-2 Antigen Test Card	REF 1N40C5-8
	Rapid SARS-CoV-2 Antigen Test Card	REF 1N40C5-9
	Rapid SARS-CoV-2 Antigen Test Card	REF 1N40C5-10

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. 01](http://www.tuvsud.com/ps-cert?q=cert:V9_061317_0006_Rev_01)

**Report No.:** SH21075CN02

**Valid from:** 2021-11-12

**Valid until:** 2024-05-26

**Date,** 2021-11-12

Christoph Dicks  
Head of Certification/Notified Body