





Certificate

No. Q5 003706 0001 Rev. 01

Holder of Certificate: ANHUI DEEPBLUE MEDICAL

TECHNOLOGY CO.,LTD.

4th Floor, D-1# Zone Pearl Industrial Park

106 Innovation Avenue, High-Tech Development Zone

230088 Hefei, Anhui

PEOPLE'S REPUBLIC OF CHINA

Certification Mark:



Scope of Certificate: Design and Development, Production and Distribution of In Vitro Diagnostic Reagents by Colloidal Gold and Enzyme

Chemical Reaction Method, Medical Ultrasonic Couplant, Acetowhite Solution, Epithelial Tissue Staining Solution, Rapid Test for Vaginitis(Polyamines) and Cell Preservation

Solution

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). 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:Q5 003706 0001 Rev. 01

Report No.: SH21130301

 Valid from:
 2021-06-22

 Valid until:
 2024-06-21

Date. 2021-06-16 Christoph Dicks

Head of Certification/Notified Body



Certificate

No. Q5 003706 0001 Rev. 01

Applied Standard(s): EN ISO 13485:2016

Medical devices - Quality management systems -

Requirements for regulatory purposes

(ISO 13485:2016) DIN EN ISO 13485:2016

Facility(ies): ANHUI DEEPBLUE MEDICAL TECHNOLOGY CO.,LTD.

4th Floor, D-1# Zone, Pearl Industrial Park, 106 Innovation Avenue, High-Tech Development Zone, 230088 Hefei, Anhui,

PEOPLE'S REPUBLIC OF CHINA

See Scope of Certificate

TÜV®



www.dbluemedical.com

DECLARATION OF CONFORMITY

MANUFACTURER: ANHUI DEEPBLUE MEDICAL TECHNOLOGY CO.,LTD.

4th Floor, D-1# Zone, Pearl Industrial Park, 106 Innovation Avenue,

High-Tech Development Zone, 230088 Hefei, Anhui, People's

Republic of China

EUROPEAN Luxus Lebenswelt GmbH

REPRESENTATIVE: Kochstr. 1, 47877, Willich, Germany

PRODUCT: COVID-19 (SARS-CoV-2) Antigen Test Kit (Colloidal Gold) -Saliva

Models: SEE ATTACHMENT

REF: SEE ATTACHMENT

CLASSIFICATION: SELF-TESTING

EDMA CODE: 15 70 90 90 00

CONFORMITY ASSESSMENT ROUTE: Following the procedure relating to the EC Declaration of Conformity set out in Annex III Section 6 of Directive 98/79/EC.

WE HEREWITH DECLARE THAT THE ABOVE MENTIONED PRODUCTS MEET THE PROVISIONS OF THE COUNCIL DIRECTIVE 98/79/EC. ALL SUPPORTING DOCUMENTATION IS RETAINED UNDER THE PREMISES OF THE MANUFACTURER.

THE MANUFACTURER IS EXCLUSIVELY RESPONSIBLE FOR THE DECLARATION OF CONFORMITY.

STANDARDS APPLIED: EN ISO 13485:2016

EN ISO 18113-1:2011, EN ISO 18113-4:2011, EN 13612: 2002/AC:2002, EN ISO 23640:2015, EN 13641: 2002, EN ISO 15223-1: 2016, EN 13975:2003, EN 13532:2002, EN ISO

14971:2012.

NOTIFIED BODY: Polish Center for Testing and Certification

469 Puławska Street,02-844 Warsaw,Poland

(EN) CERTIFICATE(S): 1434-IVDD-484/2021

START OF CE-MARKING: 2021-11-10

SIGNATURE:

PLACE, DATE OF ISSUE: HEFEI, 2021-11-12

,

CHEN FENGLING

GENERAL MANAGER

DOC-COVID-19 Ag(M/0)



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DECLARATION OF CONFORMITY ATTACHMENT

DEEPBLUE, ShenLanTest, Highbluetest, NewBluetest, Quicklyblue

Specification	REF
1 piece per box	SL030101SST-1
2 pieces per box	SL030101SST-2
3 pieces per box	SL030101SST-3
4 pieces per box	SL030101SST-4
5 pieces per box	SL030101SST-5
6 pieces per box	SL030101SST-6
7 pieces per box	SL030101SST-7
8 pieces per box	SL030101SST-8
9 pieces per box	SL030101SST-9
10 pieces per box	SL030101SST-10
11 pieces per box	SL030101SST-11
12 pieces per box	SL030101SST-12
13 pieces per box	SL030101SST-13
14 pieces per box	SL030101SST-14
15 pieces per box	SL030101SST-15
16 pieces per box	SL030101SST-16
17 pieces per box	SL030101SST-17
18 pieces per box	SL030101SST-18
19 pieces per box	SL030101SST-19
20 pieces per box	SL030101SST-20
21 pieces per box	SL030101SST-21
22 pieces per box	SL030101SST-22
23 pieces per box	SL030101SST-23
24 pieces per box	SL030101SST-24
25 pieces per box	SL030101SST-25



EC Certificate No. 1434-IVDD-484/2021

EC Design-examination
Directive 98/79/EC concerning
in vitro diagnostic medical devices

Polish Centre for Testing and Certification certifies that manufactured by:

Anhui Deepblue Medical Technology Co., Ltd. 4th Floor, D-1#Zone, Pearl Industrial Park, 106 Innovation Avenue, High-Tech Development Zone, 230088 Hefei, Anhui, China

in vitro diagnostic medical devices for self-testing

COVID-19 (SARS-CoV-2) Antigen Test Kit

(Colloidal Gold) - Saliva

The list of medical devices covered by this certificate is provided in the annex 1

in terms of design documentation, comply with requirements of Annex III (Section 6) to Directive 98/79/EC (as amended) implemented into Polish law, as evidenced by the audit conducted by the PCBC

Validity of the Certificate: from 10.11.2021 to 27.05.2024

The date of issue of the Certificate: 10.11.2021

The date of the first issue of the Certificate: 10.11.2021

C E 1434

Issued under the Contract No. MD-113/2021 Application No: 230/2021 Certificate bears the qualified signature. Warsaw, 10.11.2021 Module A1 Anna Elektron podpisal Anna Małgorzata Wyroba Data: 20

Elektronicznie podpisany przez Anna Małgorzata Wyroba Data: 2021.11.10 16:15:24 +01'00'

Vice-President



ANNEX TO THE CERTIFICATE

VALID ONLY WITH CERTIFICATE No 1434-IVDD-484/2021

List of medical devices covered by the certificate:

DEEPBLUE, ShenLanTest, Highbluetest, NewBluetest, Quicklyblue

REF: SL030101SST-1, SL030101SST-2, SL030101SST-3, SL030101SST-4, SL030101SST-5, SL030101SST-6, SL030101SST-7, SL030101SST-8, SL030101SST-9, SL030101SST-10, SL030101SST-11, SL030101SST-12, SL030101SST-13, SL030101SST-14, SL030101SST-15, SL030101SST-16, SL030101SST-17, SL030101SST-18, SL030101SST-19, SL030101SST-20, SL030101SST-21, SL030101SST-22, SL030101SST-23, SL030101SST-24, SL030101SST-25



Issued under the Contract No. MD-113/2021 Application No: 230/2021 Certificate bears the qualified signature. Warsaw, 10/11/2021 Anna Elektronicznie podpisany przez Anna Małgorzata Wyroba Data: 2021.11.10 16:16:26 +01'00'