



- ✓ Accurato
- ✓ Facile da usare
- ✓ Risultato rapido

Self Test *H. Pylori*

Cassetta Per Test Rapido Antigenico
Per la rilevazione
dell'*Helicobacter Pylori* (Feci)

contiene 1 test

RISULTATO IN 10 MINUTI



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

Full Quality Assurance System

Directive 98/79/EC on In Vitro Diagnostic Medical Devices (IVDD), Annex IV excluding (4, 6)
(List A and B and devices for self-testing)

No. V1 095123 0008 Rev. 04

Manufacturer:

Hangzhou AllTest Biotech Co., Ltd.

550#, Yinhai Street
Hangzhou Economic and Technological Development Area
310018 Hangzhou
PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies): Products for determination of infection markers
tumor markers and products for self testing**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device families in accordance with IVDD Annex IV. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of List A devices an additional Annex IV (4) certificate is mandatory. 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:V1 095123 0008 Rev. 04

Report no.:

SH221064A02

Valid from:

2022-04-05

Valid until:

2025-05-26

Date,

2022-04-05

Christoph Dicks
Head of Certification/Notified Body



EC Certificate

Full Quality Assurance System

Directive 98/79/EC on In Vitro Diagnostic Medical Devices (IVDD), Annex IV excluding (4, 6)
(List A and B and devices for self-testing)

No. V1 095123 0008 Rev. 04

Model(s):

Toxo IgG/IgM Rapid Test,
Rubella IgM Rapid Test,
CMV IgM Rapid Test,
ToRCH IgM Combo Rapid Test,
PSA Rapid Test,
PSA Qualitative Rapid Test,
Chlamydia Rapid Test,
Sperm Concentration Rapid Test,
SP-10 Male Fertility Rapid Test,
hCG Rapid Test,
Digital hCG Pregnancy Test
LH Rapid Test,
FSH Rapid Test,
Vaginal pH Rapid Test,
Ferritin Rapid Test,
TSH Rapid Test,
H.pylori Rapid Test,
Urinary Tract Infections Test,
FOB Rapid Test,
Vitamin D Rapid Test

Facility(ies):

Hangzhou AllTest Biotech Co., Ltd.
550#, Yin Hai Street, Hangzhou Economic and Technological
Development Area, 310018 Hangzhou, PEOPLE'S REPUBLIC OF
CHINA



EC Declaration of Conformity

Manufacturer:

Name: HANGZHOU ALLTEST BIOTECH CO., LTD.

Address: #550, Yinhai Street, Hangzhou Economic & Technological Development Area, Hangzhou -310018, P.R. China

European Representative:

Name: MedNet EC-REP GmbH

Address: Borkstrasse 10, 48163 Muenster, Germany

Product Name: H. pylori Antigen Rapid Test (Feces)

Cat. No.: IHP-602H

Analyte: *H.pylori* antigens in human Feces

Model: Cassette

Classification: Self-testing of IVDD 98/79/EC

Conformity Assessment Route: IVDD 98/79/EC Annex IV

EDMA Code: 15 70 01 02 00

We, HANGZHOU ALLTEST BIOTECH CO., LTD., herewith declare that we are exclusively responsible for this declaration of conformity. We herewith declare that the above mentioned products meet the transposition into national law, the provisions of the following EC Council Directives and Standards. All supporting documentations are retained under the premises of the manufacturer.

DIRECTIVES

General applicable directives:

DIRECTIVE 98/79/EC OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 27

October 1998 on in vitro diagnostic medical devices

Standard Applied: Standard Applied: EN ISO 13485:2016, EN ISO 14971:2012, EN 13975:2003, EN ISO 18113-1:2011, EN ISO 18113-4:2011, EN 13612:2002/AC:2002, EN ISO 17511:2003, EN ISO 23640:2015, EN 13641:2002, EN 13532:2002, EN ISO 15223-1:2016

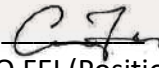
Notified body: TUV SUD Product service GmbH, Ridlerstrasse 65, 80339 Munich, Germany (0123)

(EC) Certificate(s): V1 095123 0008 Rev.04

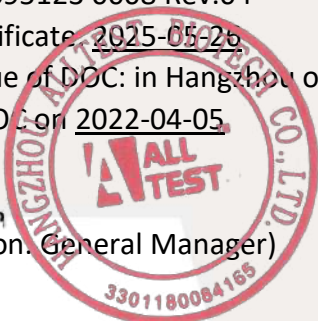
Expire date of the Certificate: 2025-05-26

Place, Date of First issue of DOC: in Hangzhou on 2017-09-12

The Date of Issue of DOC on 2022-04-05

Signature: 

Name: GAO FEI (Position: General Manager)



杭州奥泰生物技术股份有限公司 Hangzhou AllTest Biotech Co.,Ltd	文件号 Document No.: ZTC-QC-005-R-003
传染病、心肌、肿瘤类 COA The Infectious Disease、Cardiology、Tumor COA	生效日期 Effective Date: 2018 年 07 月 02 日

Certificate of Analysis

Product Name: H.pylori Antigen Rapid Test Cassette (Feces)

Catalog No.: IHP-602H

Batch No.: HPG25020028

Quantity:1000PCS

Expiry Date:2027-01

Date of Sampling:2025-03-01

Date of Analysis: 2025-03-01

Other information:

Buffer Lot: 25025023, Exp:2027-01

QC Item		QC Criterion	QC Result	Conclusion
Physical	Appearance	Good	Good	Pass
Functional Performance	Low Positive	Positive	100% Positive	Pass
	Middle Positive	Positive	100% Positive	Pass
	Negative Sample	Negative	100% Negative	Pass

Others	/
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Final QC Conclusion:	This batch of product met the QC Criteria.
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QC supervisor: Freeman.zheng

Date: 2025.03.01

