

EC Declaration of Conformity

Manufacturer:

Name: HANGZHOU ALLTEST BIOTECH CO., LTD.

Address: #550, Yin Hai 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 (Group) Name: FOB Rapid Test

Cat. No.: See Attachment 1

Analyte: See Attachment 1

Model: Attachment 1

Classification: non-listed Device of IVDD 98/79/EC

Conformity Assessment Route: IVDD 98/79/EC Annex III (excluding point 6)

EDMA Code: 12 70 03 21 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:

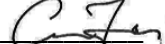
DIRECTIVE 98/79/EC OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 27 October 1998 on in vitro diagnostic medical devices

Standard Applied: EN ISO13485:2016, EN ISO14971:2012, EN 13975:2003, EN ISO 18113-1:2011, EN ISO 18113-2:2011, EN 13612:2002/AC:2002, EN ISO 17511:2003, EN ISO23640:2015, EN 13641:2002, EN ISO 15223-1:2016

Date of First Issue of DOC: in Hangzhou on 14/06/2016

Date of Issue of DOC on 29/04/2022

Date of DOC Corrigendum on 26/08/2024

Signature: 

Name: GAO FEI (Position: General Manager)



Attachment 1

Catalog No.	Product Name	Analyte	Model
TFO-602	FOB Rapid Test Cassette	Feces Occult Blood (FOB) in Feces	Cassette
TFO-601	FOB Rapid Test Dipstick	Feces Occult Blood (FOB) in Feces	Dipstick