

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: VidaQuick Biotech S.L.

Address: No.132, Rosello Street, Barcelona, Barcelona Province, 08036, Spain

Product Name: Norovirus+Rotavirus/Adenovirus+Astrovirus+Enterovirus Combo
Rapid Test

Model: Cassette

Cat. No.: IMVD-655

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

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

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 ISO14971:2019, 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 15223-1:2021

Date of Issue of DOC: in Hangzhou on 29/04/2022

Signature: 

Name: GAO FEI (Position: General Manager)



HANGZHOU ALLTEST BIOTECH CO., LTD.

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