Declaration of Conformity

VivaChek Biotech (Hangzhou) Co., Ltd.
Level 2, Block 2, 146 East Chaofeng Rd., Yuhang Economy Development Zone, Hangzhou,Zhejiang,
311100, P.R.China

We declare under our sole responsibility that the in vitro diagnostic device:

Product Name	Model
VivaDiag™ Pro SARS-CoV-2 Ag Rapid Test	VCD16-10-013
	VCD16-10-015
	VCD16-10-014
	VCD16-10-011
Verino® Pro SARS-CoV-2 Ag Rapid Test	VCD16-10-043
	VCD16-10-045
	VCD16-10-044
	VCD16-10-041

meets all the provisions of the directive 98/79/EC on *in vitro* diagnostic medical devices which apply to it

This declaration is according to Conformity Assessment Route: Annex III , section 6, and the Classification/Qualification of medical device is for self-testing

The declaration is base on the approval by the notified body

Polskie Centrum Badań i Certyfikacji S.A.ul. Puławska 469 02-844 Warszawa（PCBC）,notified under No.1434 to the EC commission.
The EC certificate No.is 1434-IVDD-478/2021

European Representative:

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Julie zhou
R.A Director
Date: Nov 10, 2021