Declaration of Conformity

ACON Biotech (Hangzhou) Co., Ltd.
No.210 Zhenzhong Road, West Lake District,
Hangzhou, P.R. China, 310030

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

Flowflex SARS-CoV-2 Antigen Rapid Test

classified as Others according to the Annex II of the directive 98/79/EC, 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 Annex III (excluding Section 6) of the Directive.

Authorized Representative:
MedNet GmbH
Borkstrasse 10
48163 Muenster, Germany

This Declaration of Conformity is valid until 25 May, 2022.

Signed this 28 day of 9, 2020
in Hangzhou, China

Junny You
International Regulatory Affairs Senior Director
ACON Biotech (Hangzhou) Co., Ltd.

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