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 (Self-Testing)
（L031-118M5: 1 Test; L031-118P5: 5 Test; L031-118R5: 25 Test）

classified as self-testing 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.6 of the Directive and thus is based on approval by the notified body TÜV SÜD Product Service GmbH, Ridlerstraße 65 80339 MÜNCHEN, Germany, notified under No. 0123 to the EC Commission.

Authorized Representative:
MedNet GmbH
Borkstrasse 10
48163 Muenster, Germany

This declaration is valid until expiration of EC certificate
No. V9 042074 0032 Rev.00
Expiration Date: 2024-05-26

Signed this 31 day of May, 2021
in Hangzhou, China

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