

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



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



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