Eon Biotechnology Limited
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EC Declaration of Conformity

According to Directive 98/79/EC on in-vitro-diagnostic devices, Annex III

Product Name	**EONBT COVIDAG 2019-nCOV-RBD Rapid Antigen Detection Kit**
Name and address of the Manufacturer	Eon Biotechnology Limited 110, Hilmanton Road, Lower Earley, Reading, RG6 4HJ United Kingdom
Authorised Representative	Diadeni sro ICO 02489902 Velešinska 659 - 19900 Praha, Czech Republic Contact details : Mr Bashkim Binaku, Tel: +420733677257 eMail: diadenicz@seznam.cz
Conformity Assessment Route	Directive 98/79/EC Annex III

We herewith declare on our sole responsibility that all batches of above In-vitro-diagnostic device is conform with the Essential Requirements Annex I of the directive 98/79/EC of the European Parliament and of the Council of 27 October 1998 on in vitro diagnostic medical devices. The product is suitable for the intended application (only professional users). Relevant standards and guidelines are applied.

Valid from 15 February 2021

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Suraz Kottakki, Director
Eon Biotechnology Limited
London