EC Declaration of Conformity

for In Vitro Diagnostic Medical Devices
according to Annex III of the 98/79/EC IVD Directive

BioMaxima S.A., Vetterów 5, 20-277 Lublin, Poland hereby declare under our sole responsibility that the products:

2019-nCoV IgG/IgM Rapid Test Cassette (Cat. No 1-360-K025)

classified as "Other IVD Medical Devices (all except listed in List A or List B and devices for self-testing)" according to Article 9 rules, conform to the relevant provisions of the EC Council Directive 98/79/EC and are in accordance with Annex III of the IVDD, as implemented by the European Union's Medical Devices Regulations.

Place and date of issue:
Lublin, 11.03.2020

Signed on behalf of BioMaxima S.A.:

Name: Łukasz Urban
Function: President

Name: Henryk Lewczuk
Function: Vice President