



## DECLARATION OF CONFORMITY

Regarding In Vitro Diagnostic Directive (98/79/EC)

**Manufacturer:** GUANGZHOU BALLYA BIO-MED CO LTD  
**Address:** Room 308, Building 1, No. 1, Yangkan 1st Road, Beicun, Baiyun District,  
Guangzhou, Guangdong, China

**EC Representative:** SUNGO Europe B.V.  
**Address:** Olympisch Stadion 24, 1076DE Amsterdam, Netherlands

**Product Name:** COVID 19 Neutralizing Antibody Rapid Test  
**Model:** CNA-19 / CNA-20 / CNA-21 / CNA-22 / CNA-23  
**Classification:** Others (IVDD)  
**Conformity Assessment Procedure:** Annex III of In Vitro Diagnostic Directive (98/79/EC)

We here with declare that the above-mentioned products meet the requirements of In Vitro Diagnostic Directive (98/79/EC) and the following harmonized standards.

EN ISO 14971:2019      EN ISO 18113-1:2011      EN ISO 18113-2:2011  
EN 13612:2002+AC:2002      EN ISO 23640:2015      EN 13641:2002

Signature: Liang Zhan Hong

Name/ Position: / GM Liang Zhan Hong

Date: 2021.3.26

Place: Guangzhou / China

