ZERTIFIKAT ◆ CERTIFICATE ◆ ZERTIFIKAT ◆ 認證證書 ◆ CERTIFICATE ◆ ZERTIFIKAT

CERTIFICATE OF NOTIFICATION

This is to certify that, according to the council directive 98/79/EC, SUNGO Europe B.V. performed all notification duties and responsibilities as the European authorized representative of：

Applicant:	Innovita (Tangshan) Biological Technology Co., Ltd.
Address:	No. 699 Juxin Street, High-tech Industrial Development Zone, Qian'an, 064400, Hebei, China.

The Manufacturer has provided SUNGO Europe B.V. with all the appropriate declarations according to the 98/79/EC Directive requirements including the EC Declaration of Conformity confirming that his In vitro diagnostic medical device, as stipulated here below, is fulfilling the applicable requirements of the European Council Directive 98/79/EC.

Product(s):	2019-nCoV Ab Test (Colloidal Gold)
Type(s):	(IgM/IgG Whole Blood/Serum/Plasma Combo)
Product Classification:	IVDD Other

Where the manufacturer affix's the CE marking to the product listed they must ensure that all the requirements of the appropriate EU directive(s) have and continue to be met.

The notification of aforementioned device has been completed by the European Representative in Netherlands. The Netherlands Competent Authority is notified of the manufacturer's medical devices and has allocated registration.

SUNGO
Executive Director
Sungo global service
CERTIFIED

Issued: Mar. 20 2020

Cert. No.: EU208518

Expiration Date: Mar. 19 2025

CE

This is not a CE mark and is only provided as a template for informational purpose.

SUNGO CONTACT INFORMATION
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