



## CERTIFICATE OF NOTIFICATION

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

**Applicant:** Xiamen AmonMed Biotechnology Co.,Ltd.  
**Address:** Unit 503, 120 Xinyuan Road, Haicang District, Xiamen, Fujian, China.

The Manufacturer has provided SUNGO 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):** COVID-19 IgM/IgG test kit (Rare earth nano fluorescence immunochromatography); COVID-19/Influenza A virus/Influenza B virus IgM combo test kit (Rare earth nano fluorescence immunochromatography)

**Type(s):** See Annex

**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.

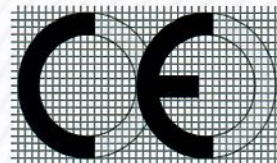
The notification of aforementioned device has been completed by the European Representative in Netherlands. The Netherlands Competent Authority is notified of the manufacturer's medical devices and has allocated registration. NOTIS number is CIBG- 20200640.

Executive Director

**Issued: Mar. 23 2020**

**Cert. No.: EU213518**

**Expiration Date: Mar. 22 2025**



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

SUNGO CONTACT INFORMATION

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Annex to Cert. No.: EU213518

Product Name	Type
COVID-19 IgM/IgG test kit (Rare earth nano fluorescence immunochromatography)	25 tests/kit,50 tests/kit
COVID-19/Influenza A virus/Influenza B virus IgM combo test kit (Rare earth nano fluorescence immunochromatography)	25 tests/kit,50tests/kit

END OF THE ANNEX

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