NOTARIAL CERTIFICATE

TO ALL TO WHOM THESE PRESENTS SHALL COME,

I, **CHU SIU LUN IVAN**, Notary Public, duly admitted, authorized and sworn, practising in Hong Kong Special Administrative Region of the People's Republic of China DO HEREBY CERTIFY that the following document hereunto annexed is a copy document provided by ABON BIOPHARM (HANGZHOU) CO., LTD., namely:-

1. GMED Additional Document n° 38757 rev. 0 dated 16 May 2022.

For the authenticity and the contents of the annexed document, I assume no responsibility.

This Apostille only certifies the authenticity of the signature and the capacity of the person who has signed the public document, and, where appropriate, the identity of the seal or stamp which the public document bears. This Apostille does not certify the content of the document for which it was issued. To verify the issuance of this Apostille, see "https://www.judiciary.hk/en/court_services_facilities/apostille_verification.html"

此項文件加簽僅就公共文件上簽署的真確性、簽署人的身分及，如適用的話，文件上的蓋章/蓋印予以證明。此項文件加簽並不就文件的內容作出證明。就發出此文件加簽之查證，見 "https://www.judiciary.hk/zh/court_services_facilities/apostille_verification.html"

APOSTILLE
(Convention de La Haye du 5 octobre 1961)

1. Country: 國家/地區	Hong Kong, China 中國香港		
This public document 此公共文件			
2. has been signed by 簽署人為	CHU Siu Lun Ivan 朱兆麟		
3. acting in the capacity of 其行事的身分為	Notary Public 公證人		
4. bears the seal / stamp of 蓋有的蓋章/蓋印	CHU Siu Lun Ivan 朱兆麟		
Certified 加簽證明			
5. at 在	High Court 高等法院	6. the 於	02 JUN 2022 2022 年 06 月 02 日
7. by 由	Andy HO Registrar, High Court 何志賢 高等法院司法常務官		
8. No 編號	34150 / 2022		
9. Seal / stamp: 蓋章/蓋印		10. Signature: 簽署	Andy Ho

Reference Code 參考編號: 2D071EDF

IN TESTIMONY whereof I have hereunto subscribed my name and affixed my Seal of Office this 31st day of May in the Year Two Thousand and Twenty-two.

CHU SIU LUN IVAN
Notary Public,
Hong Kong
Special Administrative Region of
the People's Republic of China

GROUPE LNE

Document complémentaire GMED n° 38757 rev. 0 **page 1/1**
GMED additional document n° 38757 rev. 0
Dossier(s) / *File(s)* N° P600210 – P605735
Délivré à Paris le 16/05/2022
Issued in Paris on 05/16/2022

Ce document complémentaire GMED n° 38757 rev. 0 atteste de la validité du certificat CE n° 19820 rev. 4 au regard des informations listées ci-dessous.

This GMED additional document n° 38757 rev. 0 attests to the validity of CE certificate n° 19820 rev. 4 with regard to the information listed below.

Fabricant / *Manufacturer:* **ABON BIOPHARM (HANGZHOU) CO., Ltd**
#198 12th, Street East, Hangzhou Economic & Technological Development Area
ZJ 310018 Hangzhou
CHINA

Identification des dispositifs / *Identification of devices*

Désignation du dispositif / Accessoires marqués CE *Device designation / CE marked accessories*	**Nom commercial du dispositif ou code article** *Device commercial name or article code*	**Classe du DM** *MD class*
Test rapide pour la détection des anticorps anti-VIH1, anti-VIH2 et sous-type O *Rapid test for the detection of antibodies to Human Immunodeficiency Virus (HIV) type 1, type 2 and subtype O*	**ABON HIV 1/2/0 Tri-Line Human Immunodeficiency Virus Rapid Test Réf. IHI-T402 (Code GMDN 48454)** *ABON HIV 1/2/0 Tri-Line Human Immunodeficiency Virus Rapid Test ref. IHI-T402 (GMDN Code 48454)*	**Annexe II Liste A** *Annex II List A*

Sites couverts et Activités / *Locations and Activities*

Site / *Location*	**Activités** / *Activities*
ABON BIOPHARM (HANGZHOU) CO., Ltd #198 12th, Street East, Hangzhou Economic & Technological Development Area, ZJ 310018 Hangzhou CHINA	**Conception - Fabrication - Contrôle Final** *Design - Manufacture - Final control*

DocuSigned by:
Beatrice Lys
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GMED SAS GMED SAS GMED GROUPE LNE

GMED 0459

GMED - 38757 rev. 0

On behalf of the President
Béatrice LYS
Technical Director

GROUPE LNE

ATTESTATION / *CERTIFICATE* **N° 19820 rev. 4**
Délivrée à Paris le 16 mai 2022
Issued in Paris on May 16th, 2022

ATTESTATION CE / *EC CERTIFICATE*

Approbation du Système Complet d'Assurance Qualité */ Approval full Quality Assurance System*
Annexe IV excluant les points 4 et 6 Directive 98/79/CE relative aux dispositifs médicaux de diagnostic in vitro
Annex IV excluding sections 4 & 6 Directive 98/79/EC concerning in vitro diagnostic medical devices
Pour les dispositifs de la liste A IVD, un certificat CE de la conception est requis
For list A IVD devices, a EC design certificate is required

Fabricant */ Manufacturer*

ABON BIOPHARM (HANGZHOU) CO., LTD
#198 12th Street East Hangzhou Economic & Technological Development Area
ZJ 310018 HANGZHOU CHINA

Catégorie du(des) dispositif(s) */ Device(s) category*

Dispositifs médicaux de diagnostic in vitro couverts par l'Annexe II liste A :
Test rapide pour la détection des anticorps anti-VIH1, anti-VIH2 et sous-type O.

In-vitro diagnostic medical devices covered by Annex II list A :
Rapid test for the detection of antibodies to Human Immunodeficiency Virus (HIV) type 1, type 2 and subtype O.

Voir document complémentaire GMED / See GMED additional document
n° 38757

GMED atteste qu'à l'examen des résultats figurant dans le rapport référencé P600210 - P605735, le système d'assurance qualité - pour la conception, la production et le contrôle final - des dispositifs médicaux énumérés ci-dessus est conforme aux exigences de l'annexe IV excluant les points 4 et 6 de la Directive 98/79/CE.

GMED certifies that, on the basis of the results contained in the file referenced P600210 - P605735, the quality system - for design, manufacturing, and final inspection - of medical devices listed here aboved complies with the requirements of the Directive 98/79/EC, annex IV excluding sections 4 & 6.

La validité du présent certificat est soumise à une vérification périodique ou imprévue
The validity of the certificate is subject to periodic or unexpected verification

Début de validité */ Effective date :* **May 16th, 2022 (included)**
Valable jusqu'au */ Expiry date :* **May 26th, 2025 (included)**

DocuSigned by:
Beatrice Lys
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On behalf of the President
Béatrice LYS
Technical Director

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GMED 19820 rev. 4
Renouvelle le certificat 19820-3