Ref: Vietnam 4160

غرفة التجارة العربية الايرلندية
Arab-Irish Chamber of Commerce

HPRA
An tÚdarás Rialála Táirgí Sláinte
Health Products Regulatory Authority

HPRA CERTIFIED
HPRA Reference: C1a/1753
13th August 2019

CERTIFICATE OF FREE SALE

To Whom It May Concern

The Health Products Regulatory Authority hereby certifies that:

1) Siemens Healthcare Diagnostics Manufacturing Ltd, Chapel Lane, Swords, Co Dublin, Ireland is the authorised representative for the in-vitro diagnostic medical devices specified in the attached schedule. These devices are manufactured by Diatron MI Plc, Tablas Str. 39, Budapest, Hungary H-1097. Manufacturer (as defined in the in-vitro Diagnostic Medical Devices Directive 98/79/EC) is Siemens Healthcare Diagnostics Inc., 511 Benedict Avenue, Tarrytown, NY 10591, United States of America.

2) The in-vitro diagnostic medical devices specified in the attached schedule are CE marked in accordance with the European Communities (In-vitro Diagnostic Medical Device) Regulations, 2001 (which transposed the In-vitro Diagnostic Medical Devices Directive 98/79/EC into Irish law) and may be marketed and sold in Ireland.

3) Exportation of the in-vitro diagnostic medical devices listed in the attached schedule is not prohibited.

4) The granting of this certificate is based on the information available to the Health Products Regulatory Authority on the date of issue of the certificate.

Issued To: Siemens Healthcare Diagnostics Manufacturing Ltd
Chapel Lane
Swords
Co. Dublin
Ireland

Expiry Date: 13th August 2024

Seen for the authentication of signature/seal
of Paulina Nulty
Health Products Regulatory Authority
Signed
Position EO
Date 05/03/2020

This Authentication 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 Authentication does not certify the content of the document for which it was issued. To verify an Authentication issued by the Ministry of Foreign Affairs in Ireland, see www.authentications.dfat.ie

AN ROINN GNÓTHAÍ EACHTRACHA AGUS TRÁDÁLA

Patrick Keating
Compliance Department
Health Products Regulatory Authority

CFS010229

An tÚdarás Rialála Táirgí Sláinte, Teach Kevin O'Malley, Ionad Phort an Iarla, Ardán Phort an Iarla, Baile Átha Cliath 2, Éire
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CHỨNG NHẬN / HỢP PHÁP HÓA LÃNH SỰ
CONSULAR AUTHENTICATION

1. Quốc gia: **Việt Nam**
Country

Giấy tờ, tài liệu này
This public document

2. do Ông (Bà): **Matthew Sewell** ký
has been signed by

3. với chức danh: **Cán bộ lãnh sự**
acting in the capacity of

4. và con dấu của **Bộ Ngoại giao và Thương mại Ai-len**
bears the stamp of

được chứng nhận / hợp pháp hóa lãnh sự
Certified

5. tại **London** 6. Ngày **11/ 03/ 2020**
at *the*

7. Cơ quan cấp: Đại sứ quán Việt Nam tại VQ Anh và CH Ai-len
by

8. Số: **1125**
N°

Ký tên và đóng dấu
Signature and seal/stamp
Bí thư thứ nhất
First Secretary

Nguyễn Diệu Hà

ĐẠI SỨ C.H.X.H.C.N VIỆT NAM TẠI V.Q ANH VÀ C.H AI-LEN

MEDICAL DEVICE SCHEDULE

Item Number/Product Code	Description of Device
SMN 11170841	ADVIA 360 Hematology System
SMN 11170842	ADVIA 560 Hematology System
SMN 11170843	ADVIA Autoloader
SMN 11170845	ADVIA 360/560 Dil
SMN 11170846	ADVIA 560 Lyse
SMN 11170847	ADVIA 560 5P Diff
SMN 11170848	ADVIA 360 Lyse 3P Diff
SMN 11170849	ADVIA 360 Cleaner
SMN 11170850	ADVIA 360/560 Hypoclean
SMN 11170851	ADVIA 360/560 Hypoclean CC