



Commonwealth of Virginia
The Honorable Kay Coles James - Secretary of the Commonwealth
Document revised January 24, 2022

APOSTILLE

(Convention de La Haye du 5 octobre 1961)

1. Country: **United States of America**
Pays / País:
- This public document
Le présent acte public / El presente documento público
2. has been signed by **Sami A. Swadek**
a été signé par
ha sido firmado por
3. acting in the capacity of **Notary Public**
agissant en qualité de
quien actúa en calidad de
4. bears the seal / stamp of **Sami A. Swadek – Commonwealth of Virginia**
est revêtu du sceau / timbre de
y está revestido del sello / timbre de

Certified
Attesté / Certificado

5. at **Richmond, VA**
à / en
6. the **1st day of April 2022**
le / el día
7. by **The Secretary of the Commonwealth**
par / por
8. N° **2974378-11**
sous n°
bajo el número
9. Seal / stamp:
Sceau / timbre :
Sello / timbre:
10. Signature:
Signature :
Firma:



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By Royal Charter

Certificate of Registration

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This is to certify that:

Becton, Dickinson and Company
Belliver Industrial Estate
Belliver Way
Roborough
Plymouth
PL6 7BP
United Kingdom

City/County of Alexandria
Commonwealth of Virginia

I certify this to be a complete, exact and true
copy of the original document. Certified this

24th day of March, 2022

Notary Public

SAMI A. SWADEK

Reg # 132353

Commission Exp. 4/30/2023

Facility ID Number: F000080

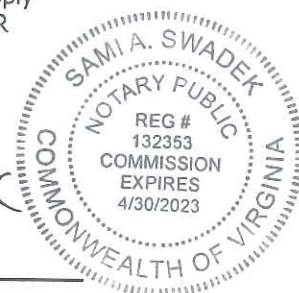
Holds Certificate No:

MDSAP 662394

Statement of Conformity: The company listed on this certificate has been audited to and found to conform with the following criteria: ISO 13485:2016 and Australia - Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 4 - Production Quality Assurance Procedure; Brasil - RDC ANVISA n. 16/2013, RDC ANVISA n. 23/2012, RDC ANVISA n. 67/2009; Canada - Medical Devices Regulations - Part 1 - SOR 98/282; Japan - MHLW Ministerial Ordinance 169, Article 4 to Article 68, PMD Act; USA - 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 - Subparts A to D

The manufacture of sterile blood collection tubes, urine tubes, urine specimen receptacles, syringes, blood collection needles, wingsets and lancets, non-sterile needle holders and supply and outsourced servicing of in vitro diagnostic instruments used in the determination of ESR for professional use.

Gary E Slack



For and on behalf of BSI:

Gary E Slack, Senior Vice President - Medical Devices

Original Registration Date: 2017-02-07

Effective Date: 2022-01-08

Expiry Date: 2025-01-07



BSI Group America Inc. is an MDSAP authorized auditing organization

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To be read in conjunction with the scope above or the attached appendix.

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