M.S.D.N: 0312593757 - C.T.C.P
CÔNG TY CỔ PHẦN THIẾT BỊ Y TẾ VIỆT GIA
Q. PHÚ NHUẬN - T.P HỒ CHÍ MINH

NSAI

Certificate of Registration of Quality Management System to ISO 13485:2016

Australia - Therapeutic Goods (Medical Devices) Regulations, 2002,
☒ Schedule 3 Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure
Brazil - 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 (as applicable)
United States - 21 CFR 803, 21 CFR 806, 21 CFR 807 – Subparts A to D,
☒ 21 CFR 820 – Quality System Regulation

The National Standards Authority of Ireland is an MDSAP Recognized Auditing Organization and certifies that:

The Seaberg Company, Inc.
dba SAM Medical Products
12200 SW Tualatin Road, Suite 200
Tualatin, OR 97062
USA
D-U-N-S: 131084188

has been assessed and deemed to comply with the requirements of the above standard and regulations in respect of the scope of operations given below:

The Design, Manufacture and Distribution of Orthopaedic Appliances, Supplies and Wound Care Devices

Approved by:
Geraldine Larkin
Chief Executive Officer

Approved by:
Caroline Dore Geraghty
Director of Medical Devices /
Head of Notified Body

Certificate Number: MP19.4360 / Rev 1
Certification Granted: 2019/01/01
Effective Date: 2019/07/01
Expiry Date: 2022/06/30

MDSAP
MEDICAL DEVICE SINGLE AUDIT PROGRAM

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National Standards Authority of Ireland, 20 Trafalgar Square, Nashua, New Hampshire, NH 03063, USA T +1 603 882 4412
All valid certifications are listed on NSAI's website – www.nsaiinc.com The continued validity of this certificate may be verified under "Approved Client Listing