Certificate of Registration

QUALITY MANAGEMENT SYSTEM – ISO 13485:2016

This is to certify that:

NeoMed, Inc.
100 Londonderry Court
Suite 112
Woodstock
Georgia
30188
USA

DUNS Number: 79-977-2079

Holds certificate No: **MDSAP 662076**

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 1 (excluding Part 1.6) - Full Quality Assurance Procedure [if design controls are part of the certification]; Canada - Medical Devices Regulations - Part 1- SOR 98/282; USA - 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 – Subparts A to D

Design, development, manufacture and distribution of sterile feeding tubes, sterile extension sets, sterile urinary drainage catheters, sterile neonatal procedure drapes; sterile and non-sterile oral and enteral syringes, sterile and non-sterile oral and enteral syringe and feeding tube accessories. Provision of servicing activities for customer supplied product.

For and on behalf of BSI:

Stewart Brain, Head of Compliance & Risk – Medical Devices

Original Registration Date: 2018-12-13 Effective Date: 2018-12-13 Expiry date: 2021-11-06

BSI Group America Inc. is an MDSAP authorized auditing organization

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