

EC CERTIFICATION

FULL QUALITY ASSURANCE SYSTEM

Directive 93/42/EEC on Medical Devices, Annex II excluding (4)

We hereby declare that an examination of the under mentioned full quality assurance system has been carried out following the requirements of the Swedish national legislation LVFS 2003:11 to which the undersigned is subjected, transposing Annex II (with the exemption of section 4) of the Directive 93/42/EEC on medical devices. We certify that the full quality assurance system conforms with the relevant provisions of the aforementioned directive, and the result entitles the organization to use the CE 0413 marking on those products listed below.

Organization:

MedXL Inc.

Main Site: 285, rue Labrosse, Pointe-Claire, QC H9R 1A3, Canada

Product Category:

- Prefilled Syringe for maintaining patency of Vascular Access
Devices (VADs)

For further identification of the products covered, see the MDD product list/product schedule.

Certificate Number:

41316788-03

Initial Certification Date:

25 June 2009

Certificate Valid from:

26 June 2019

Certificate Expiry Date:

27 May 2024



Accred. no. 1003
Certification of
Management
Systems
ISO/IEC 17021-1


Peter Nermander

Certification Authority MDD
Intertek Semko AB, Kista, Sweden

17 June 2019

Signed Date

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The certification is subject to the organization maintaining their system in compliance with the regulations stated in this certificate, allowing regular assessments and following the contracted requirements of the Notified Body.

Intertek Semko AB is a Notified Body according to Directive 93/42/EEC on medical devices, with identification number 0413.

