EC Certificate

FULL QUALITY ASSURANCE SYSTEM
Directive 93/42/EEC on Medical Devices, Annex II (3)

Certificate Number
41316939-01

Initial Certification Date
May 15, 2009

Certificate Valid from
May 16, 2014

Certificate Expiry Date
May 15, 2019

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.

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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:

Allied Healthcare Products, Inc.
1720 Sublette Avenue, St. Louis, MO 63110, USA

Product Category:

Emergency Transport Ventilators, Demand Valves, Humidifiers, Oxygen Regulators, Adapters, Air Compressors, Bag Mask Resuscitators, Burn Towels, Oxygen Cannulas, Flowmeters, Medical Gas Hoses, Oxygen Masks, Portable Suction Pumps & Aspirators, Thermotic Drainage Pumps, Selector Valves, Selector Valve Kits, Vacuum Regulators, Water Traps, CO_2 Absorbents

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

May 9, 2014
Signed date

Mats Premfors, Certification Authority MDD
Intertek Semko AB, Kista, Sweden