

# EC Certificate - Full Quality Assurance System

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

**No.**

**CE 607662**

Issued To:

**Medeco B.V.  
Alexander Flemingstraat 2  
Oud-Beijerland  
3261 MA  
The Netherlands**

In respect of:

**The design, manufacture and final inspection of sterile pen needles, sterile enteral pump feeding sets including connectors and adapters, non-sterile gauzes and superabsorbent dressings, sterile non-medicated wound dressings comprising alginate, foam, silicone contact layer, superabsorbent, gauze, non-woven, hydrocolloid and petroleum tulle gras technologies**

**Those aspects of Annex II concerned with securing and maintaining sterility of wound dressings, plasters, urine collection bags including connecting and fixation materials, syringes and enteral gravity feeding sets including connectors and adapters.**

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 2797):



Gary E Slack, Senior Vice President Medical Devices

First Issued: **2016-06-24**

Date: **2020-04-17**

Expiry Date: **2024-05-26**

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Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

This certificate was issued electronically and is bound by the conditions of the contract.