EC Certificate Production Quality Assurance System: Certificate BE19/819943317

The management system of

BEDAL NV

Agoralaan A bis
3590 Diepenbeek, Belgium

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V
Restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions

For the following products

Sterile, single use patches for stabilization of catheters, tubing & cables.

Where the above scope includes class IIb or class III medical device(s), a valid EC Type Examination Certificate according to Annex III is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 12 November 2020 until 03 January 2024
and remains valid subject to satisfactory surveillance audits.
Issue 3. Certified since 13 February 2019.

Certification is based on reports numbered BE/AMD-19/1065.CMD

Authorised by

SGS Belgium NV, Notified Body 1639

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