EC Certificate Full Quality Assurance System: Certificate GB19/964486

The management system of

Avery Dennison Medical Limited

IDA Business Park, N39 DX73 Longford, Ireland

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

Sterile:
Polyurethane foam wound dressings
Gel Hydrogel wound dressings
Silfoam and Silfoam Border wound dressings
Silfoam V and Silfoam V Border Silicone adhesive sterile wound dressing-double sided lamination
FinSap Superabsorbent dressing for management of wounds that have penetrated the epidermis.
Sterile Manuka Honey Tulle dressing
InterSil Silicone wound contact layer wound dressings

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 18 May 2021 until 24 May 2024
and remains valid subject to satisfactory surveillance audits.
Issue 4. Certified since 07 July 2008

Certification is based on reports numbered GB/PC 216465

Authorised by

Global Medical Devices Head of Notified Body

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