

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

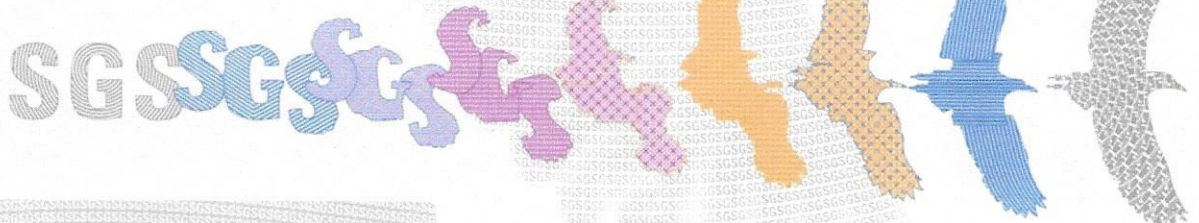
Global Medical Devices Head of Notified Body

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LPMD5007 - Certificate CE1639 Annex II-4_EN rev. 02

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