EC Certificate Production Quality Assurance System: Certificate GB19/964485

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 V

Restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions

For the following products

Sterile skin barrier film.

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 22 January 2020 until 07 July 2022
and remains valid subject to satisfactory surveillance audits.
Issue 2. Certified since 26 November 2014
and first certified by SGS Belgium NV since 16 December 2019.

Certification is based on reports numbered GB/PC/ 216465

Authorised by

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