EC Certificate Production Quality Assurance System: Certificate BE20/819944096

The management system of

Instrumentmakerij Medeja B.V.

Dorpsstraat 644
1566 EM Assendelft, The Netherlands

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

Low-pressure gas probes for use with medical gases.
Low and high vacuum venturi devices intended to remove substances from the body.

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 26 October 2020 until 24 May 2024
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 26 October 2020.

Certification is based on reports numbered BE/AMD/2/1199.QMD

Authorised by

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