EC Certificate Full Quality Assurance System: Certificate CH19/1015

The management system of

Unimed SA

Rue du Grand-Pré
CH-1007 Lausanne

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

Medical needles for injection and biopsy

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 2020 until 11 December 2023
and remains valid subject to satisfactory surveillance audits.
Issue 2. Certified since 12 December 1997
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered CH/GE 3302680.1

Authorised by

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