

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

**SGS Belgium NV, Notified Body 1639**

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

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