EC Certificate Full Quality Assurance System: Certificate CH12/2218

The management system of

Intensiv SA

Via al Molino 107
CH - 6926, Montagnola

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

Rotating diamond coated instruments for conservative and restorative dentistry
Oscillating diamond coated instruments for conservative, restorative and orthodontic dentistry
Rotating cutting instruments for conservative, restorative and orthodontic dentistry
Instruments to polish dental surfaces in conservative and restorative dentistry
Instruments for finishing of preparations for conservative and restorative dentistry

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 11 February 2018 until 10 February 2023
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 20 November 2020
Issue 5. Certified since 11 March 1998

Certification is based on reports numbered CH/GE 3301461

Authorised by

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