EC Certificate Full Quality Assurance System: Certificate CH12/0881

The management system of

pfm medical cpp SA

Allée du Quartz 9,
2300 La Chaux-de-Fonds, Neuchâtel, Switzerland
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

Sterile Huber needles, Sterile perfusion catheters for oncology applications, Sterile explantation cannulae. Sterile implantable vascular access port for administration of systemic chemotherapy, long term parental nutrition and long term medication.

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 16 June 2017 until 16 June 2022
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 4 June 2020
Issue 3. Certified since 25 June 1996

Certification is based on reports numbered CH/GE 3302042.1

Authorised by

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