EC Certificate Full Quality Assurance System: Certificate CN19/41144

The management system of

NINGBO YINZHOU XIANFENG ELECTRONICS INSTRUMENT FACTORY

No.328 East Wuxiang Road, Bao zhuang, Wuxiang Town, Yinzhou District, Ningbo City, Zhejiang Province, 315112 , P.R. China

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 Oxygen Regulator used in Healthcare Facilities
Model:YR-86, YR-87, YR-88
Oxygen Flowmeter with Humidification used for Administration of Pure Oxygen
Model: LYX AC

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 December 2019 until 07 June 2022
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 15 November 2014
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered CN/NGB 5767

Authorised by

SGS Belgium NV, Notified Body 1639

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