EC Certificate Full Quality Assurance System: Certificate KR08/01099

The management system of

Neobiotech, Co., LTD

#103, 104-1, 104-2, 105, 106, 205, 212, 312, 509, 510, 511 & 10F, 36, Digital-ro 27gil, Guro-gu, Seoul, 152-789

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

Dental sterile single use implant with sterile and non sterile abutment for restoration of tooth function;
Sterile single use and non sterile Dental surgical instrument for dental surgery;
Dental sterile single use titanium membrane for guided bone restoration of dental surgery.

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 27 April 2015 until 28 December 2018 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 2 December 2016
Issue 11. Certified since 30 July 2008

Certification is based on reports numbered KR/SEL Y-PC/08180

Authorised by

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