EC Certificate Production Quality Assurance System: Certificate MY00/51710.02

The management system of

Biopro (M) Sdn. Bhd.

Lot 14, PT 4204 Lingkaran Sultan Hishamuddin
North Port Industrial Estate, 42000 Port Klang
Selangor Darul Ehsan
MALAYSIA

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

Sterile Natural (Latex) and Synthetic Latex (Nitrile) Surgical Gloves.

Where the above scope includes class IIb or class III medical device(s), a valid EC Type Examination Certificate according to Annex III is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 19 February 2016 until 19 February 2021 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 19 December 2018
Issue 5. Certified since 04 August 2004

Certification is based on reports numbered MY/KUL MY00370

Multiple certificates have been issued for this scope
The main certificate is numbered MY00/51710.00

Authorised by

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