

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



SGS United Kingdom Ltd, Notified Body 0120

202B Worle Parkway, Weston-super-Mare, BS22 6WA UK t +44 (0)1934 522917 f +44 (0)1934 522137 www.sgs.com

SGS CE 14 0215

Page 1 of 1





This document is issued by the Company subject to its General Conditions of Certification Services accessible at www.sgs.com/terms_and_conditions.htm. Attention is drawn to the limitations of liability, indemnification and jurisdictional issues established therein. The authenticity of this document may be verified at http://www.sgs.com/en/Our-Company/Certified-Client-Directories/Certified-Client-Directories.aspx. Any unauthorized alteration, forgery or falsification of the content or appearance of this document is unlawful and offenders may be prosecuted to the fullest extent of the law.