EC Certificate Production Quality Assurance System: Certificate EG08/75891

The management system of

Ultra For Medical Products Co. (ULTRAMED)

Industrial Area, Arab El-Awamer, Abnoub, Assiut, Egypt

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

Sterile intravenous infusion set with needle and without needle, Sterile blood transfusion set; sterile IV Cannula; sterile surgical gloves; sterile guedal airway, sterile Mucus extractor; sterile burette set, sterile Nelaton Catheter; sterile suction tubing, suction units; sterile Rhyle's tubing, sterile infant feeding tubing; sterile 3 way stop cock; sterile nasel Oxygen Catheter; sterile Fistula needle; sterile flow regulator; sterile IV tubing, sterile endotracheal tube with and without cuffs, sterile extension tubes with and without 3 way stop cock. sterile umbilical cord clamp; sterile urine bag; sterile Oxygen face mask

For placing on the market of Class IIb or Class III devices covered by this certificate, an EC Type Examination Certificate according to Annex III is required.

This certificate is valid from 16 February 2015 until 28 July 2019 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 20 July 2017
Issue 1. Certified since 29 October 2008

Authorised by

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