Notification of a Body in the framework of a technical harmonization directive

From : RAPPRESENTAZIONE
PERMANENTE D'ITALIA
PRESSO L'UNIONE EUROPEA
1040 BRUXELLES

To : **European Commission**
GROWTH Directorate-General
200 Rue de la Loi,
B-1049 Brussels.

Other Member States

Reference : Legislation : 93/42/EEC Medical devices

Body name, address, telephone, fax, email, website :

MTIC InterCert S.r.l.
Sede Legale: Via G.Leopardi, 14 - Sede Operativa: Via Moscova, 11
20123 - Milano (MI) - 20017 - Rho (MI)
Italy
Phone : +39 02 97071800
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Email : info@mtic-group.org
Website : www.mtic-group.org

Body : **NB 0068**

The body is formally accredited against :

EN ISO/IEC 17020 - Inspection

EN ISO/IEC 17021 - Certification of management systems

EN ISO/IEC 17065 - Product certification

Name of National Accreditation Body (NAB) :

The accreditation covers the product categories and conformity assessment procedures concerned by this notification : No

Tasks performed by the Body :

Last approval date :

Product family, product /Intended use/Product range	Procedure/Modules	Annexes or articles of the directives	Limitations
Dental instruments (non sterile) equipment	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	
Equipment for aerosol therapy, humidification and nebulisation	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	
Equipment for anaesthesia, assisted ventilation and support of vital functions			
- surgical equipment and ancillary surgical equipment	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	
Equipment for electrosurgery	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	
Equipment for infusion, transfusion, extracorporeal circulation, plasmapheresis and blood collection	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	
Equipment for measuring and/or monitoring and/or recording and/or analysing physiological parameters and vital functions	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	
Equipment for nerve, heart or muscle stimulation or electrotherapy	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	
Equipment for peritoneal dialysis and hemodialysis			
- Equipment for endoscopy	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	
Equipment for rehabilitation	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	
Equipment for the production, visualisation and processing of images using non-ionising radiation or ultrasound	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	
Equipment for treatment using non-ionising radiation or ultrasound	Full quality assurance system EC type-examination EC verification Production quality assurance	Annex II Annex III Annex IV Annex V	

Product family, product /Intended use/Product range	Procedure/Modules	Annexes or articles of the directives	Limitations
	Product quality assurance	Annex VI	
Incubators and equipment for neonatal pathology	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	
Non-active devices for anaesthesia and assisted ventilation	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	
Non-active devices for measuring physiological parameters	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	
Non-active devices for surgery and endoscopy	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	