Declaration of Conformity

Meditrade GmbH
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We, the manufacturer

Meditrade GmbH
Medipark 1
83088 Kiefersfelden
Germany

herewith declare in our own responsibility, that the following medical products of **class Is (classified following rule 5 acc. Appendix IX of directive 93/42/EEC)**

Sterile latex examination gloves, powderfree
Gentle Skin® steril (REF 8021RW / 8022)
Sterile nitrile examination gloves, powderfree
Nitril® steril (REF 8023)
Sterile copolymere examination gloves, powderfree
Copolymed® (REF 8091 / 8092)

comply with the essential requirements of Medical Device Directive 93/42/EEC (last amended by Directive 2007/47/EC).

NBOG-Code:	MDS 7006 + MD 0101
Conformity Assessment Procedure:	Appendix VII of MDD 93/42/EEC in connection with Procedure acc. to Appendix V of MDD 93/42/EEC
Notified Body:	DEKRA Certification GmbH, Handwerkstr. 15, 70565 Stuttgart ID-Number: 0124

Kiefersfelden, 17.05.2016

CE 0124

Martin Unterberg
Product Manager
Meditrade GmbH