



**CE-Declaration of Conformity**  
**in accordance with Council Directive 93/42 / EEC of 14 June 1993**  
**concerning medical devices**

Medical Device	<b>Surgical latex gloves, powdered</b> <ul style="list-style-type: none"><li>- Reference OP / REF 9321</li></ul> <b>Surgical latex gloves, powderfree</b> <ul style="list-style-type: none"><li>- Gentle Skin Premium OP (REF 9021W)</li><li>- Gentle Skin Smooth OP (REF 9026)</li><li>- Gentle Skin Micro OP (REF 9031)</li><li>- Gentle Skin Securitex OP (REF 9051)</li></ul> <b>Surgical polyisoprene gloves, powderfree</b> <ul style="list-style-type: none"><li>- Gentle Skin Isopretex OP (REF 9522)</li></ul> <b>Surgical polychloroprene gloves, powderfree</b> <ul style="list-style-type: none"><li>- Neopretex OP (REF 9521)</li></ul>
Classification	Class IIa according directive 93/42/EEG Appendix IX; Rule 7
NBOG-Code:	MDS 7006 + MD 0101
Manufacturer	Meditrade GmbH Medipark 1 83088 Kiefersfelden

Herewith we declare in our own responsibility, that the above mentioned medical devices meet the essential requirements of 93/42/EEC.

Chosen conformity procedures	According the Directive 93/42/EEC Appendix V
Notified Body	TÜV SÜD Produkt Service GmbH Ridlerstraße 65 80330 München ID-Nummer: 0123
EC-Certificate	Nr. G2 011655 0014 Rev.02 valid until 2024-05-26
Validity of the Declaration of Conformity	2024-05-25

Kiefersfelden, 20.04.2021

Martin Unterberg  
Safety Officer of MD