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**REFLEXX S.p.A.**

**DECLARATION OF CONFORMITY CE rev. 01 del 16 marzo 2017**

Product: reflexx 51 art. R51/ XS - art. R51/ S – art. R51/ M – art. R51/ L– art. R51/ XL

*Emessa da G.Isetti – Amm.re Unico*

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## **DISPOSABLE LATEX GLOVES TRADEMARK REFLEXX 51**

The company REFLEXX SpA, located in Via Passeri, 2 in Viadana (MN) as Manufacturer declares that the product meets the requirements of the following European Community Directives:

- ❑ As a PPE category I risk according to IT Legislative Decree 475/92 implementing Dir 89/686/EEC . The product is certified according to the EN 420 and EN 374.
- ❑ As Medical Device Class I fall in between devices by invasive examination also intended for temporary use in accordance with Directive 93/42/EEC and subsequent Legislative Decree 37/2010 on the implementation Dir 2007/47/EEC. The product is in accordance with the EN 455 1-2 and 3. – CND T010201 LATEX GLOVES mis. XS/1522253 S/1524280 M/1526319 L/1526320 XL/1526321

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