



E SEI IN MANI SICURE!

**REFLEXX S.p.A.**

**DECLARATION OF CONFORMITY CE rev. 07 del 20 dicembre 2017**

Product: reflexx 40 art. R40/ XS - art. R40/ S – art. R40/ M – art. R40/ L– art. R40/ XL

*Emessa da G.Isetti – Amm.re Unico*

Pag.: 1/1

## **DISPOSABLE LATEX GLOVES TRADEMARK REFLEXX 40**

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 III risk according to IT Legislative Decree 475/92 implementing Dir 89/686/EEC (CIMAC, authorized Ministerial: 0465). The product is certified according to the EN 420:2003 + A1 2009, EN 374-1:2016 TYPE B, EN 374-2:2015, EN 374-4:2013 e EN 374-5:2016.
  
- ❑ 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/1557586 S/1557587 M/1557588 L/1557589 XL/1557590

**REFLEXX S.p.A.**  
Via Passeri, 2 - 46019 VIADANA (MN)  
P.I. e C.F. 02085450209  
Tel.: 0375/633164 - Fax: 0375/464504  
e-mail: info@reflexx.com