



E SEI IN MANI SICURE!

REFLEXX S.p.A.

DECLARATION OF CONFORMITY CE rev 03 del 29 gennaio 2018

Product: reflexx 77 art. R77/XS - art. R77/ S – art. R77/ M – art. R77/ L– art. R77/ XL

Emessa da G.Isetti - Amm.re Unico

Pag.: 1/1

DISPOSABLE NITRILE GLOVES TRADEMARK REFLEXX 77

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 TIPE 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 1-2-3 and 4. CND T01020204 – NITRILE GLOVES
–

**Progressivo di Sistema Attribuito al DM: mis. XS/1528651 S/1528652
M/1528653 L/1528654 XL/1528655.**

REFLEXX S.p.A.

Via Passeri, 2 - 46019 VIADANA (MN)

P.I. e C.F. 02085450209

Tel.: 0375/833164 - Fax: 0375/464504

e-mail: info@reflexx.com