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REFLEXX S.p.A.

DECLARATION OF CONFORMITY CE

rev 00 del 03 gennaio 2018

Product: N70plus art. N70P/S – art. N70P/M – art. N70P/L– art. N70P/XL– art. N70P/XXL

Emessa da G.Isetti - Amm.re Unico

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DISPOSABLE NITRILE GLOVES POWDER FREE TRADEMARK REFLEXX N70PLUS

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 1-2-3 and 4. CND T01020204 – NITRILE GLOVES - mis S/1526796 M/1526797 L/1526798 XL/ 1526799 XXL/1526800.

REFLEXX S.p.A.

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