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REFLEXX S.p.A.	DECLARATION OF CONFORMITY CE	rev 00 del 03 gennaio 2018
Product: reflexx N79plus	art. N79P/S – art. N79P/M– art. N79P/L– art. N79P/XL	
Emessa da G.Isetti - Amm.re Unico		Pag.: 1/1

DISPOSABLE NITRILE GLOVES POWDER FREE
TRADEMARK REFLEXX N79PLUS

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/1555786 M/1555785 L/1555788 XL/1555789.

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