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REFLEXX S.p.A.

DECLARATION OF CONFORMITY CE rev 01 del 31 gennaio 2018

Product: *reflexx Care N350* art. N350/S - art. N350/M - art. N350/L

Emessa da G.Isetti - Amm.re Unico

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DISPOSABLE NITRILE GLOVES TRADEMARK REFLEXX care N350

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 1-2-3 and 4. CND T01020204 NON SURGICAL NITRILE GLOVES **mis. S/ 1636541 M/ 1636542 L/ 1636543.**

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