Form QAT_10-M04, version 00, effective since March 6th, 2020

CEDocumentation Review



No. 00374463EIC

Holder: Export. Import. Crespo SL - VAT: ES B94061645

Review goal:

Verification of the presence of Technical Documentation compatible with the Medical Devices Directive 93/42/EEC

Rapport d'Evaluation

Annex VII

Product(s): Model(s):

Classification:

Medpride Medical Exam. Blue Nitrile Latex Free & Powder Free. https://disposable-wholesale.com Class I (accordingly to the Manufacturer's declaration)

Review output:

This document has been issued on a voluntary basis and upon request of the manufacturer. It is our opinion that the Technical Documentation shared with us by the manufacturer is compatible with the European Standard for Medical Devices.

The manufacturer is responsible for the CE Marking process, and not exempted to carry out all necessary compliance activities. This document has been issued on the certification of products. RG01_ECM rev.3 available at: www.entecerma.eu

The certificate can be checked, insert the Certificate number at www.entecerma.eu

Date of issue March 4, 2020

Approver ECM Service Director Luca Bedonni Expiry date March 5, 2025



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