C € Documentation Review

No. 0H200315.GJIUD99



Holder:

Guizhou Junjiang Industry Co., Ltd.

South 7th Road, Hongxing Pharmaceutical Industrial Park, Maling Town, Xingyi City, Guizhou

Province China.

Review goal:

Verification of the presence of the Technical File in regards of the Medical Devices Directive 93/42/EEC Annex VII

Product: Model(s): Medical Rubber Gloves (no sterile) Inspection Gloves, Surgical Gloves

Classification:

Class I (no sterile) (accordingly to the Manufacturer's declaration)

Review output:

We attest that a Technical File in reference to the Directive 93/42/EEC is in place for the CE Marking process. Test Report identified with the no. XMT0201901559S/MDD.

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 basis of the regulation on ECM Voluntary Mark for the certification of products. RG01_ECM rev.3 available at: www.entecerma.it

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> Reviewer **Technical expert** Amanda Payne

Approver ECM Service Director Luca Bedonni