

ATTESTATION CE / EC CERTIFICATE

Examen de type / Type Examination

ANNEXE III de la directive 93/42/CEE relative aux dispositifs médicaux

ANNEX III DIRECTIVE 93/42/EEC concerning medical devices

Fabricant (nom et adresse) / Manufacturer (name and address)

MEDEFIL, INC.

250 Windy Point Drive, GLENDALE HEIGHTS

ILLINOIS 60139 UNITED STATES

Catégorie du(des) dispositif(s) / Device(s) category

solution saline 0,9% NaCl de rinçage des cathéters

Saline solution 0,9% NaCl for rinsing of catheters

Identification du(des) dispositif(s) / Identification of device(s)

Normal Saline I.V. Flush Solution

Normal Saline I.V. Flush Solution

Le LNE/G-MED atteste qu'à l'examen des résultats figurant dans le(s) rapport(s) référencé(s) L110066-3, un échantillon représentatif de la production est conforme aux exigences de l'annexe I de la Directive 93/42/CEE

LNE/G-MED certifies that, on the basis of the results contained in the file(s) referenced L110066-3, a representative sample of the production complies with the requirements of the Directive 93/42/EEC, annex I

Début de validité / Effective date : **December 26th, 2011 (included)**

Valable jusqu'au / Expiry date : **December 25th, 2016 (included)**



LNE - 22625 rev. 0

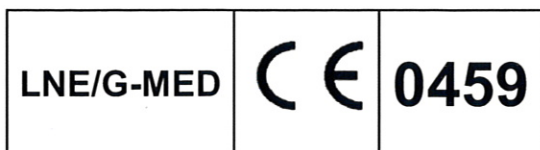
On behalf of the Deputy Director
Thierry THOMAS
G-MED Certification Division Manager

Identification des dispositifs / Identification of devices

**MEDEFIL Inc.
Normal Saline I.V. Flush Solution**

Commercial reference	Syringe volume	Solution volume
MIS-1121	6 ml	1 ml
MIS-1122	6 ml	2 ml
MIS-1123	6 ml	3 ml
MIS-1125	6 ml	5 ml
MIS-1130	12 ml	10 ml
MIS-1133	12 ml	3 ml
MIS-1135	12 ml	5 ml
MIS-1152	6 ml	2.5 ml

8 alinéas / indented lines.




**On behalf of the Deputy Director
Thierry THOMAS
G-MED Certification Division Manager**