

Certificate FR99/51155

MICROVAL S.A.

Z.A. de Champ de Berre,
F43240 Saint-Just-Malmont, France

Device Identification:
GRF Sterile Surgical Glue

Medical Purpose of Device:

Biological glue for use in the surgical repair of acute aortic dissection.

has been assessed and certified as meeting the requirements of

EC Directive 93/42/EEC

on Medical Devices Annex II, section 4

It is certified that the manufacturer's design dossier (and product, where applicable) for the above device has been examined and, based on the evidence submitted, it is considered that the device conforms to the relevant Essential Requirements of EC Directive 93/42/EEC.

This certificate is issued in conjunction with a certificate covering the full quality assurance system to Annex II, which must be subject to regular compliance visits.

This certificate is valid from 10 December 2009 until 9 December 2014
Issue 3

Certification is based on report number FR/MD 202023 dated 5th January 2010

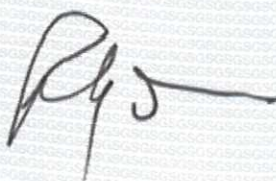
Addenda to that report have been issued on the following dates:

Addendum Date

Reason for Addendum

Notified Body Number 0120

Authorised by



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