Ferrosan
Medical Devices

EC - Declaration of Conformity

We, **Ferrosan Medical Devices A/S,**

Sydmarken 5, 2860 Søborg, Denmark
being the manufacturer
within the European Economic Community of the following class III products:

Device name:	*Product code:*
SPONGOSTAN™ Absorbable Haemostatic Gelatin Sponge, Film (20 unit)	MS0001
SPONGOSTAN™ Absorbable Haemostatic Gelatin Sponge, Standard (20 unit)	MS0002
SPONGOSTAN™ Absorbable Haemostatic Gelatin Sponge, Special (20 unit)	MS0003
SPONGOSTAN™ Absorbable Haemostatic Gelatin Sponge, Anal (20 unit)	MS0004
SPONGOSTAN™ Absorbable Haemostatic Gelatin Sponge, Dental (24 unit)	MS0005
SPONGOSTAN™ Absorbable Haemostatic Gelatin Sponge, Standard (2 unit)	MS0006
SPONGOSTAN™ Absorbable Haemostatic Gelatin Sponge, Anal (5 unit)	MS0007
SPONGOSTAN™ Absorbable Haemostatic Gelatin Powder	MS0008
SURGIFLO™ Haemostatic Matrix	MS0010
SURGIFLO™ Haemostatic Matrix Kit with Thrombin	MS0012

Device GMDN code and term:

Device GMDN code:	Term:
48170	Gelatin haemostatic agent

declare that the above is in conformity with the provisions of the Council Directive
93/42/EEC as amended Concerning Medical Devices
and has been subject to the following conformity procedures laid down in
Annex II
under the supervision of the Notified Body, DNV Product Assurance AS, Veritasveien 3, 1363 Høvik, Norway, and carrying the Notified Body Number
2460

Søborg, date 07.06.2021

C. Hudtloff

Camilla Hudtloff
VP, QM & RA