

# CERTIFICATE

Number: 2119704TE02



## EC TYPE-EXAMINATION MEDICAL DEVICES AND ANIMAL TISSUE DIRECTIVE

Issued to:

**TAUREON**Laan van Zuid Hoorn 61  
2289 DC Rijswijk Zh  
The Netherlands

For the product category:

**Novacol resorbable collagen fibrillar and pad material for haemostasis**

Documents, that form the basis of this certificate:

**Certification Notice 2119704CN, initially dated October 20, 2008**

KEMA hereby declares that the above mentioned manufacturer fulfils the relevant provisions of 'Besluit Medische Hulpmiddelen', the Dutch transposition of the Directive 93/42/EEC of 14 June 1993 concerning medical devices, including all subsequent amendments, and that for the above mentioned product category the Conformity Assessment Procedure Annex III for Class III products is executed by the Manufacturer in accordance with the provisions of the Council **Directive 93/42/EEC of 14 June 1993**. Additionally, KEMA hereby declares that the manufacturer fulfils the relevant provisions as specified in Annex I of the **Directive 2003/32/EC of 23 April 2003** concerning medical devices manufactured utilising tissue of animal origin. The necessary information and the reference to the relevant documentation, of the products concerned and the assessments performed, are stated in the Certification Notice which forms an integrative part of this certificate.

This certificate is valid until: August 1, 2012  
Issued for the first time: October 20, 2008  
Reissued: July 28, 2009

KEMA Quality B.V.

drs. G.J. Zoetbrood  
Managing DirectorM. McCann  
Certification Manager

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