

We hereby declare that the distributed CE marked products, specified below, meet the relevant provisions of the Council Directive 93/42/EEC of 14 June 1993, concerning medical devices.

This declaration is based on the application of the Quality System approved for the manufacture and final inspection of the products concerned, in accordance with Annex V of the EC-Directive and is described in the CE Marking of Conformity Certificate, reference number 2119704CE01 issued for the first time on 20 October 2008 and delivered by KEMA Quality BV, Arnhem, The Netherlands, Notified Body Identification Number 0344.

In addition we ensure and declare that the distributed CE marked products, as mentioned and falling in class III, meet the relevant provisions as specified in Annex I of Directive 2003/32/EC of 23 April 2003 concerning medical devices manufactured utilizing tissue of animal origin.

This declaration covers the following products:

Product	Classification	Followed Annexes	EC Type Examination Certificate
SurfaSoft	IIA	V and VII	n.a.
HoneySoft	IIB	V and III	2119704TE01
Novacol	III	V and III	2119704TE02

Art. code	Description	Size	First batch
2033S	SurfaSoft®	20 x 33 cm	2033081201
HS0506	HoneySoft®	5 x 6 cm	0506090119
HS1010	HoneySoft®	10 x 10 cm	1010090120
HS1025	HoneySoft®	10 x 25 cm	1025090120
2011	Novacol®	1 gram	2011081103
3001	Novacol®	2,5 x 5 cm	3001081104
3002	Novacol®	5 x 8 cm	3002081105
3003	Novacol®	8 x 10 cm	3003090218

This declaration is valid for all products described here above, bearing the CE marking and manufactured by:

TAUREON
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Dick van Kalker
Managing Director
Rijswijk, 20 maart 2009