

Certificate US06/68524

RTI Biologics, Inc.

11621 Research Circle, PO Box 2650,
Alachua, FL, 32616-2650, United States

Device identification:

**Sterling® Impacted Cortical Wedge, Sterling® Impacted Cortical Ring,
Sterling® SR, Sterling® Machined Dowel.**

Medical Purpose of Device:

Load-bearing spinal fusion applications.

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, and Directive 2003/32/EC on medical devices manufactured utilising tissues of animal origin.

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 3 April 2008 until 30 May 2011
Issue 4

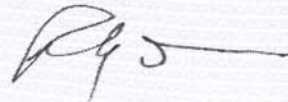
Certification is based on report number(s) PC DDE 208856 dated 23 May 2006

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

<u>Addendum Date</u>	<u>Reason for Addendum</u>
25 June 2007	Removed Sterling® HTO Wedge; Amended medical purpose.
26 September 2007	Packaging configuration change.
31 March 2008	Changed Company Name to RTI Biologics, Inc.

Notified Body Number 0120

Authorised by

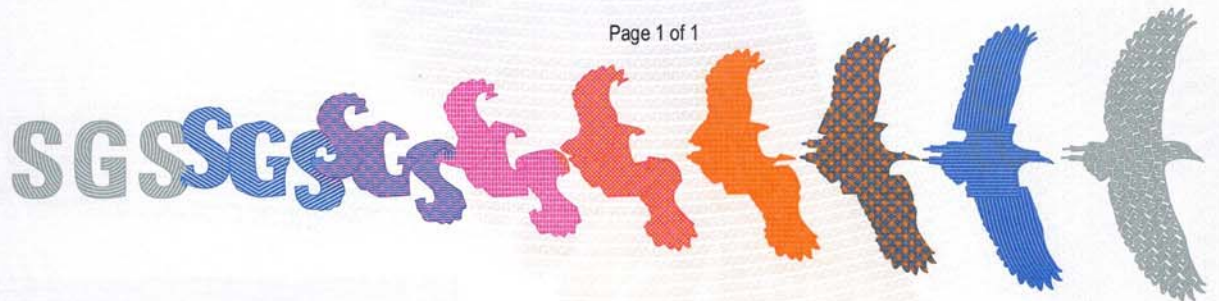


SGS United Kingdom Ltd Systems & Services Certification
202B Worle Parkway, Weston-super-Mare, BS22 6WA, UK
t +44 (0)1934 522917 f +44 (0)1934 522137 www.sgs.com

CE 0120

SGS EC 04 1007

Page 1 of 1



COPY