

Certificate US06/68558

RTI Biologics, Inc.

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

Device identification:

**Sterling® Suture Anchor, Sterling® Interference Screw ST, Sterling®
Interference Screw HT.**

Medical Purpose of Device:

**Fixation of hard tissue and soft tissue in orthopaedic surgical
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

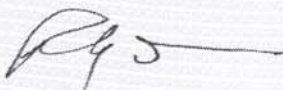
Certification is based on report number(s) PC DDE 214682 dated 23 May 2006

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

Addendum Date	Reason for Addendum
30 May 2006	Addition of 10, 11, and 12 mm diameter ST and HT Interference Screws
25 June 2007	Amend medical purpose.
26 September 2007	Change in packaging configuration.
31 March 2008	Change of Compan Name to RTI Biologics, Inc.

Notified Body Number 0120

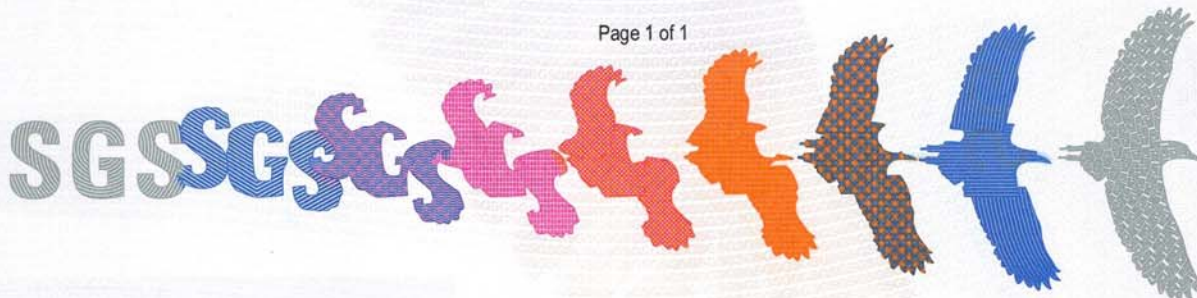
Authorised by



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