



Certificate

Full Quality Assurance

No. CE 596074



Issued to:

Tyber Medical
89 Headquarters Plaza, 1464
Morristown
New Jersey
07960
USA

In respect of:

Design and manufacture of sterile and non-sterile interbody implants and associated instrumentation

on the basis of our examination under the requirements of Council Directive 93/42/EEC, Annex II, Section 3.2.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 0086):

Gary Fenton, Global Assurance Director

First Issued: **29 Aug 2013**

Date: **29 Aug 2013**

Expiration Date: **28 Aug 2018**

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Conditions of Approval

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

...making excellence a habit.™