

# EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

**No.****CE 515162**

## Issued To:

**Zimmer Inc.  
1800 W. Center Street  
Warsaw  
Indiana  
46580  
USA**

In respect of:

**Design, development and manufacture of joint replacement implants, fracture fixation devices, bone cement accessories, trial prostheses and instruments for use with active devices. Those aspects related to securing and maintaining sterility in the manufacture of orthopaedic instruments, fracture fixation devices and bone cement instruments. Those aspects related to the accuracy of metrology in the manufacture of orthopaedic instruments and fracture fixation devices.**

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 2797):



Gary E Slack, Senior Vice President Medical Devices

First Issued: **2007-08-13**Date: **2021-01-25**Expiry Date: **2024-05-26**

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Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. 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.

This certificate was issued electronically and is bound by the conditions of the contract.