By Royal Charter

# 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):

Digitally signed by Kalcev Ghennadi
Date: 2022.05.27 10:38:30 EEST
Reason: MoldSign Signature
Location: Moldova

Gary E Slack, Senior Vice President Medical Devices

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