CERTIFICATE

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Of Compliance

This is to certify that the technical documentation has been independently assessed and compliant with the requirements of MDD directive 93/42/EEC.

Manufacturer

Name	:	**Company Name**
Address	:	**Address**
Product Details	:	**Xxxx**
Applicable Standard	:	**Medical Device Regulation (EU) 2017/745 and Medical Device Directive 93/42/EEC**

This certificate refers to the information examined and read with manufacturer's declaration of conformity. Further, the product liability rests with the manufacturer or his representative in accordance with the council Regulation (EU) 2017/745 and directive 93/42/EEC. The CE Mark as shown below can be used, under the responsibility of manufacturer, after completion of a CE Declaration of conformity and compliance with the relevant CE Directives.

Certificate No.: xxxxxxxxxxx

Date of Initial Registration:	xx-xx-xxxx
1st Surveillance audit on or before:	xx-xx-xxxx
2nd Surveillance audit on or before:	xx-xx-xxxx
Date of Recertification:	xx-xx-xxxx

Authorised Signatory

SP Certification Limited
130, Old Street, London,
EC1V9BD, U.K.
Email: info@spcertification.co.uk
www.spcertification.co.uk
Company No. 11668116

Verify Certificates

Accredited by Accreditation Forum of International Standards [AFIST (UK) LTD.]
9, Park View Road, Leeds, LS4 2LG, United Kingdom (UK), Email : info@afist.org, Website : www.afist.org

The approval is subject to the company maintaining its system to the required standards, which will be monitored by SPC.
The Certificate remains the property of SPC and must be returned on request.

To check validity of this certificate please visit **www.spcertification.co.uk** and **www.afist.org**