



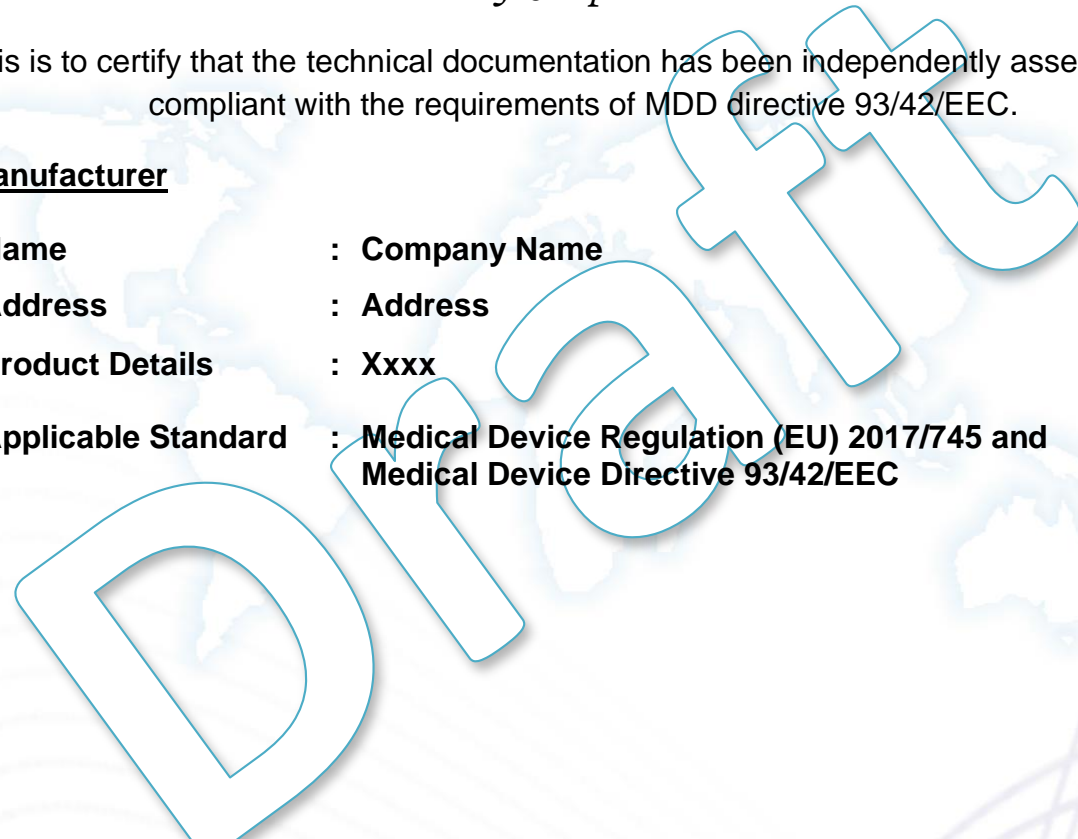
CERTIFICATE

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 : Xxx
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.: xxxxxxxxxxxx **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

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www.spcertification.co.uk
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Verify Certificates

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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.
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