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 Medical Device Directive 93/42/EEC.

Manufacturer

Name	:	**NISHTHA MEDIQUIP (INDIA)**
Address	:	**NR. APMC MARKET, 4TH FLOOR, 423, CITY GATE, B/H VISHALA HOTEL, SHAHVADI, AHMEDABAD, GUJARAT – 382405, INDIA**
Product Details	:	**ORTHOPAEDIC IMPLANTS & INSTRUMENTS – EQUIPMENT SUPPLIER**
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.: SPC24C2788

Date of Initial Registration:	12-01-2024
1st Surveillance audit on or before:	11-01-2025
2nd Surveillance audit on or before:	11-01-2026
Date of Recertification:	12-01-2027

Authorised Signatory

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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**