CERTIFICATE

This is to certify that the product listed below conforms to the requirements of the

Directives Name **Medical Device Directive 93/42/EEC**

Certificate No.: **IN9763U-2024**

Date of Issue: **07 June 2024**

Surveillance audit on or before: 06 June 2027

Date of this Certificate: **07 June 2024**

Date of Expiry: 06 June 2027

Manufacturer :

SPLINT MEDITEC PRIVATE LIMITED

PLOT NO. 1/2, 2 & 3, SURVEY NO. 605, JD INDUSTRIAL AREA -1, RAVKI - MAKHAVAD ROAD, RAVKI, RAJKOT, GUJARAT, INDIA - 360004

Description of Products : **MANUFACTURING, TRADING, EXPORT AND IMPORT DESIGNING OF ORTHOPEDIC IMPLANTS & INSTRUMENT, MAXILLOFACIAL IMPLANTS & INSTRUMENT AND RELATED PRODUCTS**

Equipment Identification : As Shown of Above

Other Certification : **NA**

Standards Applied : **AS per required**

Report Reference: **SMPL/CE/IN9763U-2024**

This Report has been Issued by Certiva Limited according to the provision of the **Medical Device Directive 93/42/EEC.** This Report is issued following the assessment of the documentation and implementation of the Quality System in accordance with the provisions of the quoted Conformity Assessment Module of the above directives. The CE Mark may be affixed to the press **Medical Device Directive 93/42/EEC** the Scope of approvals as described above once the 'EU declaration of conformity' has been signed by the responsible person.

CE

Director

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