

EU REP CERTIFICATE

EUR SRN: LV-AR-000011602

MDR 2017/ 745

Class 1 as Annexure VIII

Conformity Assessment as per Annexure IV

Confirm that as per Medical Device Regulation (MDR 2017/745) we perform all duties and responsibilities as EU Representation of

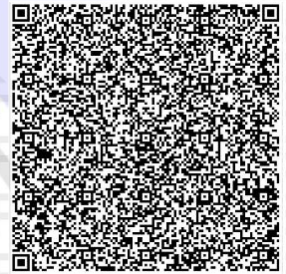
Immane Corporation

Khadim Ali Road, Near City Public School, Kotli Behram
51330 Sialkot - Pakistan

HAS GIVEN A MANDATE TO

CS ACTIVE

Usmas Iela 19 dz 24
1083 Riga LATVIA



Assessment of technical documentation has been completed by CS Active on October 04th 2025. No non-compliance according to the MDR 2017/745 was found during the assessment. Manufacturer declare the Medical Devices listed in attached "ANNEXURE A" as Class-I as per Medical Device Regulatory (MDR 2017/745).

This certificate will be valid until the expiry date or in the case of any change in Product manufacturing process and specification, Quality System and relevant legislation by manufacturer. The end of certificate validity will subject to the surveillance.

Reference No: CS/PK/EC-3102230LV

Date of Initial Registration: 24.05.2021

Certificate Date of Issue: 04.10.2025

Certificate Valid Until: 03.10.2026

Authorized Signatory

CS Active



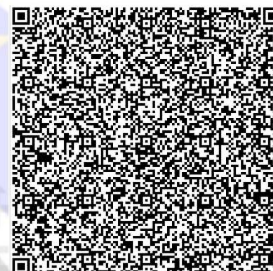
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ANNEX I PAGE 1/1:



List of medical devices under the scope of Agreement

Obstetrical Forceps	Gastrointestinal Instruments
Hollow ware	Urology Instruments
Amalgam Instruments	Rhinology Instruments
Gynecology Instruments	Oral Instruments
Diagnostic Instruments	Tracheotomy Instruments
Mirror Handles	Ring Clamp Forceps
Plaster Instruments	Tooth Extracting Forceps
Impression Trays	Rubber Dam Punches and Clamps
Explorers	Mouth Gags
Sterilizing Instruments	Periodontal Instruments
Orthodontic Pliers	Separating Spatulas
Filling Instruments	Scalers
Cement Spatulas	
Endodontic Instruments	
Ophthalmic Instruments	
Otology Instruments	
Dermatology Instruments	

Authorized Signatory


CS Active



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