

# **CERTIFICATE OF CONFORMITY**

## No. ECSPK2021.183

This is to certify that the products submitted are described items of Meeting the

**Requirements of:** 

**Medical Devices Directive 93/42/EEC** 

**Manufactured By:** 

### **ULTRA TRUSTED IMPLANTS & INSTRUMENTS**

H.O. 4F-9/4, Nazimabad # 4, Karachi-Pakistan

The Manufacturer technical file documentation of the undermentioned product(s) has been reviewed and found to comply with requirements of above regulation for article 52 (MDR).

Scope: "Manufacturing & Supply of Face mask, Orthopedic Implants, Neuro Implants & Instruments"

#### **Limitations:**

For Placing the devices on the market, an EC Design-Examination Certificate will be required. This certificate does not imply assessment of the series production of the product. The holder must inform ECS, of any substantial changes occurred in the product or process in order to examine whether this certificate remains valid.

On Behalf of ECS:
Technical Manage

CE

Active Certification: 07/April/2022 to 06/April/2023

This certificate is the property of European Certification Services and remains valid subject to satisfactory annual assessment.

#### **IONTEX OÜ**

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