IBC CERTIFICATE OF REGISTRATION

CE

CERTIFICATE # IBC 02105201

This is to certify that

Frix Surgical Instruments

P.O Box # 950, New Miana Pura Roras Road, Sialkot-51310-Pakistan

Has been assessed for applicable requirements of Directive 93/42/EEC as updated 2007/47/EC for Class I – Reusable Medical devices (Non-Active/Non-Sterilized/Non-Measuring) and found to meet the requirements of Medical Device Directive satisfactorily and is registered with Sweden Medical Products Agency (Läkemedelsverket) with reference number 6.6.1-2021-42038.

Frix Surgical Instruments comply with the mentioned standards and the status of Sweden`s Competent Authority remains legible until compliance and successful surveillance. Manufacturer of a medical device is responsible to meet the applicable requirements in the Medical Devices Act (1993:584) and the Swedish Medicine Agency`s Regulations, LVFS 2003:11 or LVFS 2001:7. This assessment exercise was carried with all due care though conformance verification practices and all the regulatory requirement is the sole responsibility of manufacturer.

EU Authorized Representative
IBC - Sweden
Norsborg, Stockholm-Sweden
info@ibcsweden.eu

Active Certification:	20-05-2022 to 18-05-2023
First Surveillance:	19-05-2022
Second Surveillance:	18-05-2023

Certification Cycle

Issue Date: 20-05-2021 Expiry Date: 17-05-2024

Remains valid subject to satisfactorily surveillance audit

881014-6707

Bolagsverket LÄKEMEDELSVERKET MEDICAL PRODUCTS AGENCY