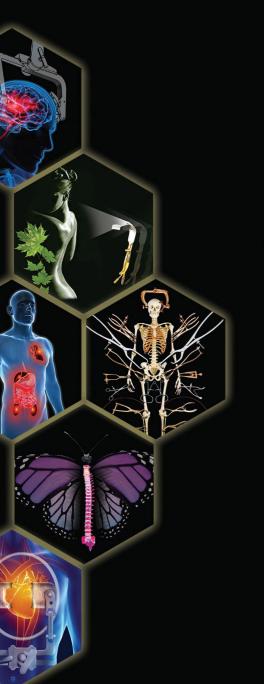
International Instruments

Designers & Manufacturers of Surgical Instruments









Orthopaedic | Hip Surgery | Knee Surgery Shoulder Surgery | Hand Surgery Foot & Ankle Surgery | Arthroscopy Spine Surgery | Neuro Surgery Micro Surgery | General Surgery | Uro - Gastro Thoracic - Cardiovascular Surgery | Gynecology Laparoscopy | Plastic Surgery | E.N.T Surgery

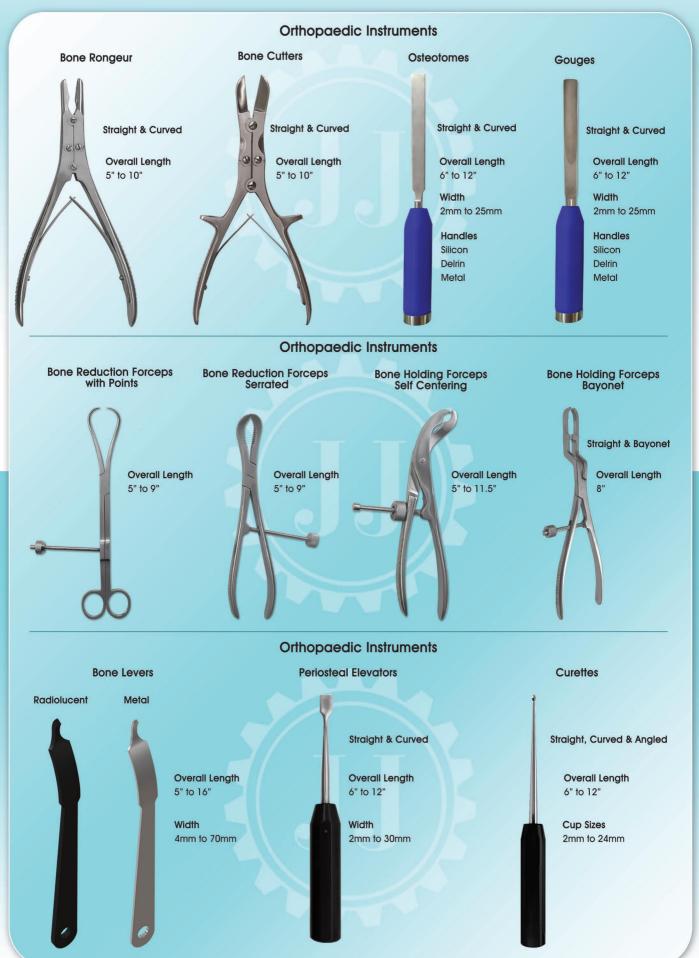
www.myjjonline.com

sales@myjjonline.com | myjjonline@yahoo.com | jjcare@myjjonline.com

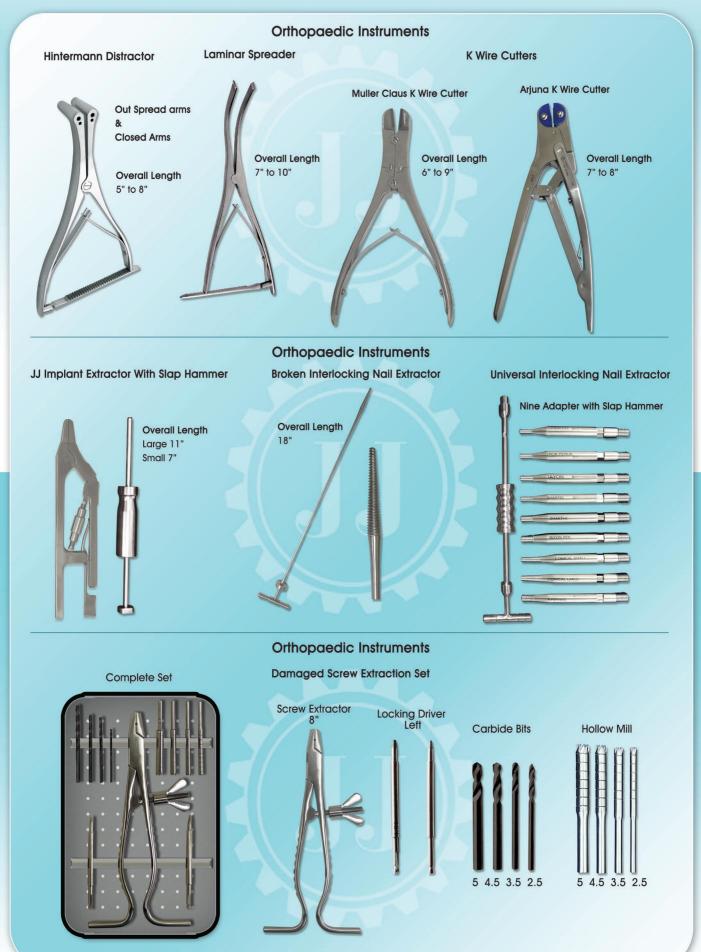
Kinfra Park P.O., Koratty – 680 309, Thrissur District, Kerala, India +91 480 2735959 / 5858 | +91 94970 66333 | +91 94970 77555

INDIA | USA | OMAN | UAE

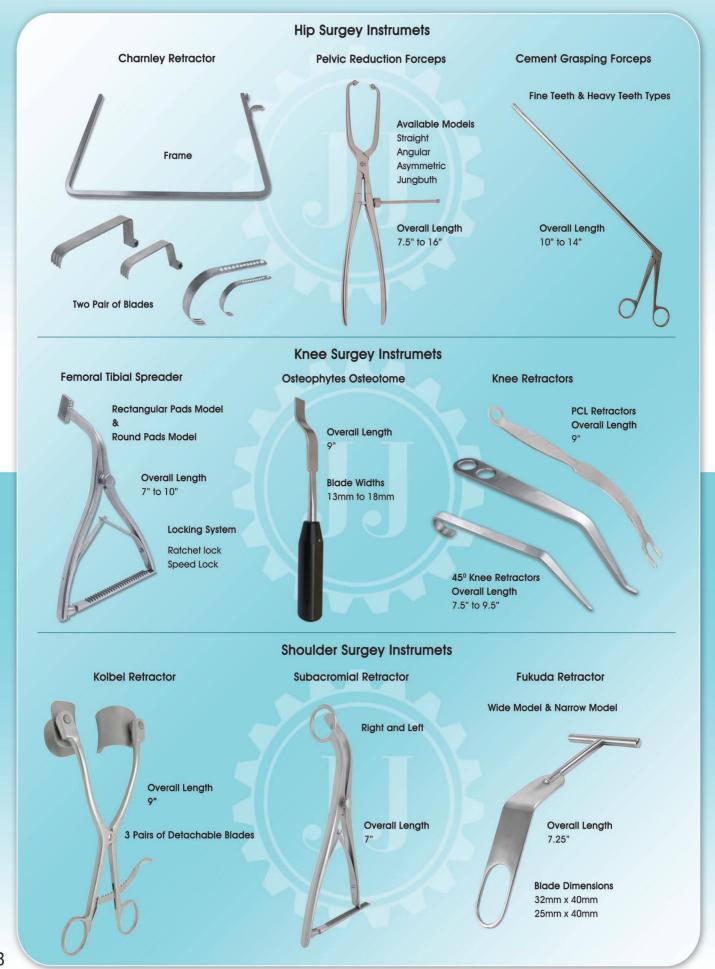




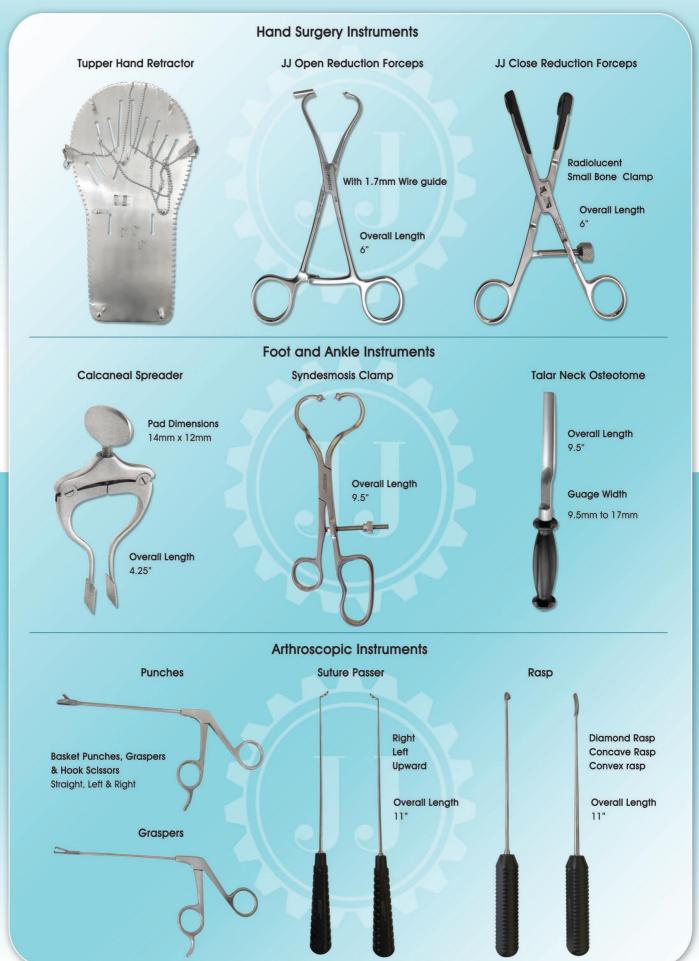




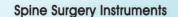












Casper Micro Lumbar Discectomy Retractor



Micro Discectomy Retractor



Hook Blades



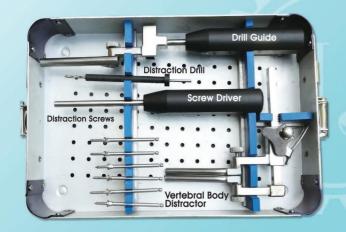
Toothed Blades

Narrow Blades

Spine Surgery Instruments

Caspar Vertebral Body Distraction Set

Cervical Retractor set





Spine Surgery Instruments

Spine Rongeurs

Laminectomy Retractor

Super -Slide Laminectomy Retractor

Thin Foot Plate

Upward cutting 1mm to 5mm Downward Cutting 1mm to 5mm



Finocchietto Retractor With Three Pair of Blades and one Hook



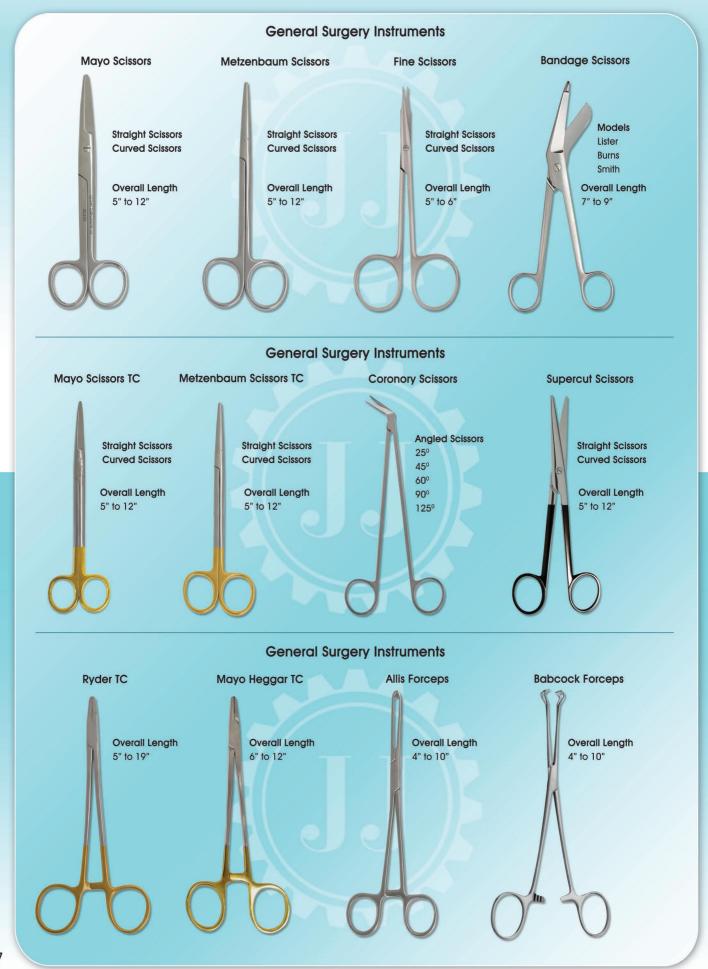
Intervertebral Disc Rongeur

Upward cutting 1mm to 5mm Downward Cutting 1mm to 5mm

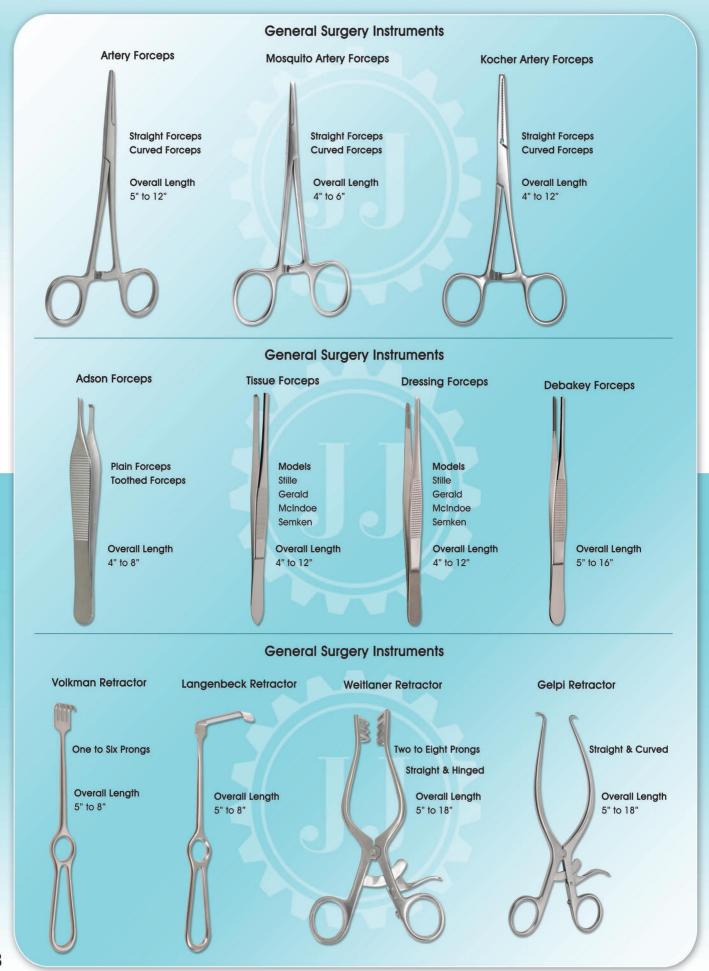


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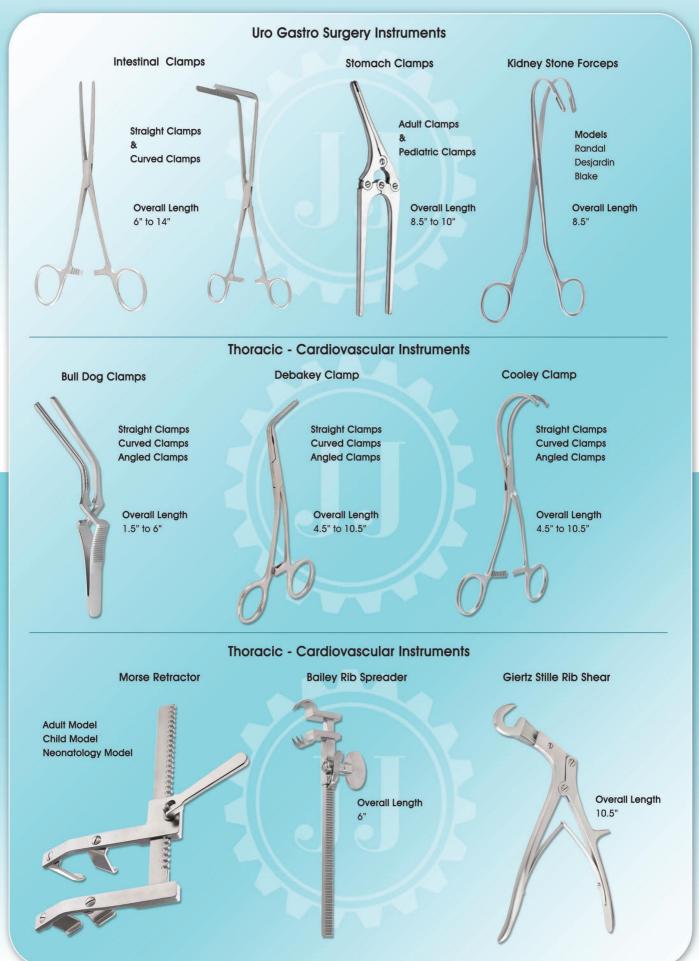




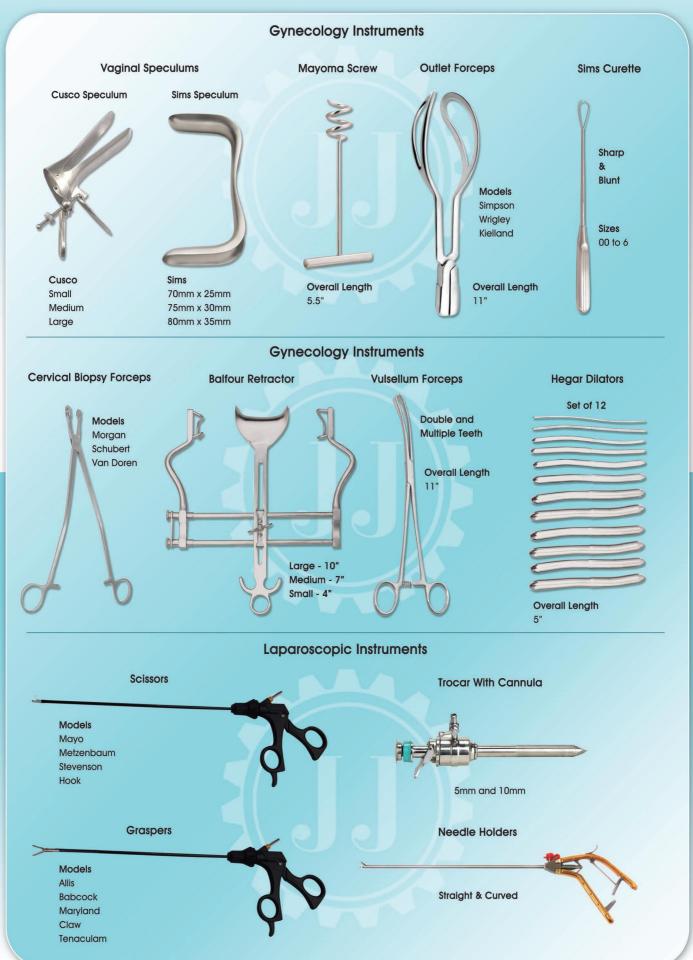




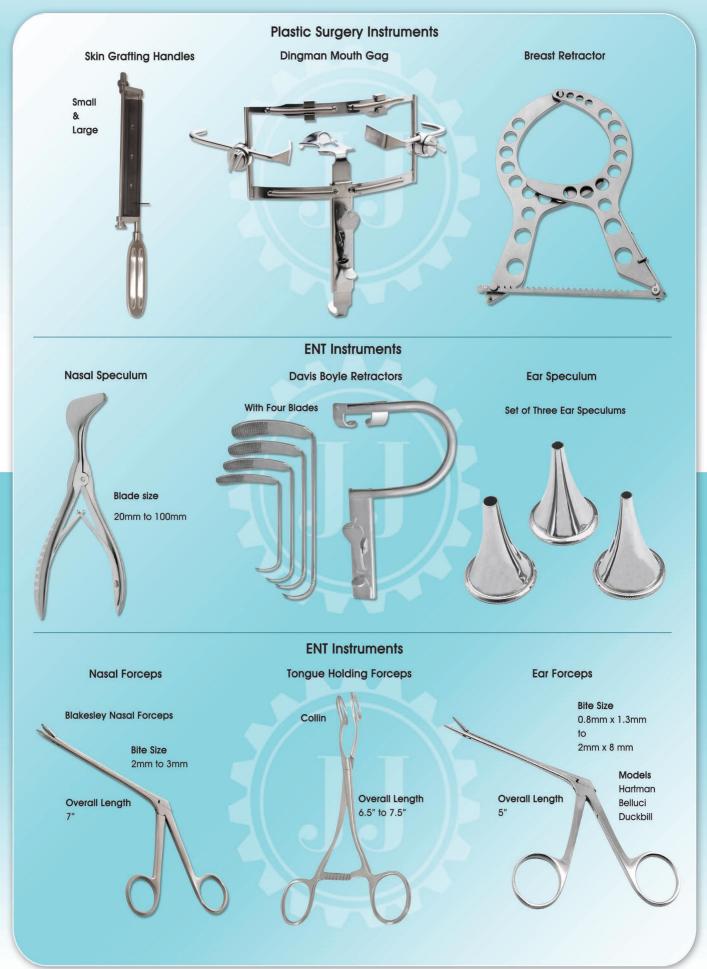












JJ Quality Certificates



Dear Joshi Thomas

This e-mail provides confirmation that the annual registration for the following medical device establishment has been successfully completed for 2021:

Registration Number: 3010077069 Owner Operator Number: 10043535 INTERNATIONAL INSTRUMENTS Kinfra Park PO Koratty Thrissur INDIA 680308

If you do not see a registration number assigned to the establishment and your establishment previously had one, please send an email to reglist@cdrh.fda.gov and include the registration number you believe is assigned to your establishment. We will review and determine if a duplicate registration has been created for your establishment.

Your registration is valid until December 31, 2021. Registration for 2022 will be conducted between October 1 and December 31, 2021.

Please note that registering your device facility and listing your devices does not, in any way, constitute FDA approval of your facility or your devices.

Should you have any questions, please send an e-mail to the CDRH Registration and Listing Helpdesk at reglist@cdrh.fda.gov.

CDRH Registration and Listing Helpdesk Imports & Registration and Listing Team Division 2 Establishment Support Office of Regulatory Programs
Office of Product Evaluation and Quality Center for Devices and Radiological Health
U.S. Food and Drug Administration
Tel: 301-796-7400, Option 1
Email: reglist@cdrh.fda.gov



d'Evaluation Rapport 더 뻐 전 Report Review

(€ Documentation Review No. 2P191227.IIWT46 Holder: International Instruments Plot No. XIX-350, Kinfra Park P.O, Koratty, Thrissur District, Kerala – 680 309, India. Verification of the presence of the Technical File in regards of the Medical Devices Directive 93/42/EEC Annex VII Review goal: Non-active Surgical Instruments (see the following annex I) Model(s): Classification: Class I We aftest that a Technical Re in reference to the Directive 93/42/EEC is in place for the C Marking process. This document has been sixed on voluntary basis and not as NB. Whereast the Manufacturer is responsible and not exempted to carry out all the necessary activities, as required by the Directive, before placing the C Mark on the product. Date of issue 27 December 2019 Expiry date 26 December 2022

Ente Certificazione Macchine
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