Positioned to Perform

Innovative Medical Products™

Instructions For Use

De Mayo Adapt2Fit®

Modular Knee Positioner

Catalog Number **1020**

Doc Title **IFU-IMP-0018**

Version **1**

Intended Use/Intended Users: The De Mayo Adapt2Fit® Modular Knee Positioner's intended use is to position the patient's leg during a surgical procedure, including but not limited to total knee replacement, partial knee replacement and revision. Orthopedic surgeons are the intended users.

Target Patient Group: Patient selection depends on the judgment of the surgeon. Surgeons must consider the size of the patient. The heel to popliteal region should be in between 13" and 18"

Contraindications: This device is not designed, sold, or intended for use except as indicated.

Warnings/Precautions

- Do not use device in a manner that does not follow these instructions for use.
- Untrained personnel - review and understand the IFU.
- Incorrect set up – follow IFU.
- Safety: Always use IMP Patient Protective Pads
- ***DO NOT PLACE POSITIONER IN A MILK BATH OR LUBRICATE***
- Max number of reuses:
    - Unplanned carriage movement
    - Excessive medial lateral movement of boot when the carriage is locked.
    - Unrepairable wear and/or damage to the baseplate including locking pins.

Risks:

- Device should not come in direct contact with the patient. Patient is protected with Patient Protective Pad
- Sterilized by end user.
- No shelf-life issues.
- Re-usable
- Must be in compliance with the IFU with IMP accessories and components.

Complaints and Adverse Events: For complaints and adverse events, contact IMP and the appropriate regulatory authorities for specific country.

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Tower Street, Swatar, BKR 4013 Malta

Product Identification

Part No	Product Name	UDI-DI
1020-SYS KIT	De Mayo Adapt2Fit®Modular Knee Positioner	00696588006536

Consumables:

- Sterile Patient Protective Pads
- Coflex

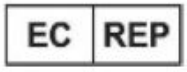
Disposal of unit: If a device is being returned for repair or disposal, please contact Innovative Medical Products at sales@IMPmedical.com. If the device is not being returned, instruments are to be disposed of in accordance with applicable laws, rules, and regulations for the disposal of biohazardous waste. Follow all guidelines for biohazardous waste in accordance with the Centers for Disease Control and Prevention guidelines as well as applicable federal/national, state, and local regulations.

Acceptable Accessories:

Part No	Product Name	GTIN
120	Quad® Clamp	00696588006178
621	De Mayo Universal Distractor® 2.0	00696588007151
713-717	De Mayo Push Button Clamp -Locking Pin	00696588001517
803-ABD	Aluminum Boot with Distractor Block	00696588001463
803-ABDA	Aluminum Boot with Distractor Block 110°	00696588006468
803-CBD	Composite Boot with Distractor Block for De Mayo Knee Positioner®	00696588000794
907*	De Mayo Universal Distractor®	00696588001388

*Denotes discontinued item

Symbol Glossary

Symbol	Title	Description	Standard
	Authorized Representative in the European Community	Indicates the authorized representative in the European Community.	ISO 15223-1:2016
	Batch Number	Indicates the manufacturer's batch code so that the batch or lot can be identified.	ISO 15223-1:2016
	Catalogue number	Indicates the manufacturer's catalogue number so that the medical device can be identified. The manufacturer's catalogue number shall be placed after or below the symbol and adjacent to it.	ISO 15223-1:2016
	Caution	Indicates the need for the user to consult the instructions for use for important cautionary information such as warnings and precautions that cannot be presented on the medical device itself.	ISO 15223-1:2016
	Complies with European Directives.		
	Consult instructions for use.		ISO 15223-1:2016

Symbol	Title	Description	Standard
	Date of manufacture	The date must be presented in the following format: YYYY-MM-DD	FDA 21 CFR 801
	Manufacturer .	This symbol shall be accompanied by the name and address of the manufacturer.	ISO 15223-1:2016
MD	Medical device		ISO 15223-1:2016
LATEX	Not made with natural rubber latex	.	Manufacturer defined
SN	Serial Number	The manufacturer's serial number shall be placed after or below the symbol and adjacent to it.	ISO 15223-1:2016
UDI	Unique device identifier		ISO 15223-1:2016

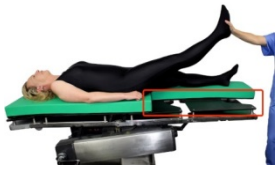
Instructions for use:

1. Replace OR Table Pads with green IMP® Pads.

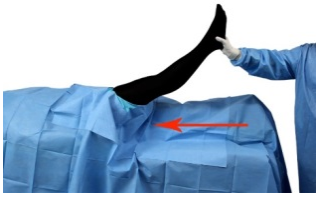
Note: It is not necessary to remove X-Ray cassette plates.

2. Position the patient with the gluteal fold at the separation of the pads.

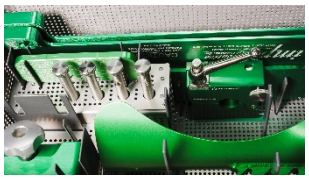
3. During Prep of the surgical leg, remove single OR Table Pad.



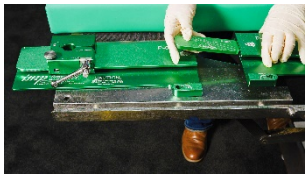
4. During Prep of the surgical leg, remove single OR Table Pad.



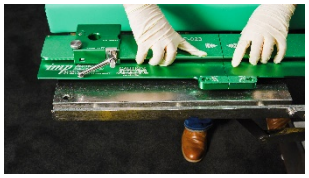
5. Open the sterilization tray and remove the baseplate halves and the baseplate connector.

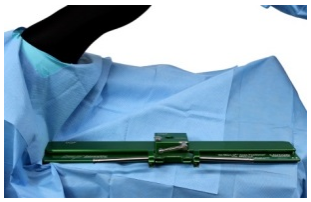
6. Insert the baseplate connector into one half of the baseplate until it stops.



7. Connect the other baseplate by sliding it onto the protruding baseplate connector until the baseplates meet.

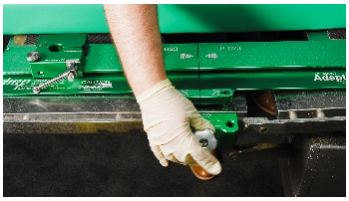



8. Place the De Mayo Adapt2Fit® Modular Knee Positioner in the well to achieve maximum flexion. The Adapt2Fit® is designed to fit perfectly into the well created with no overhang.



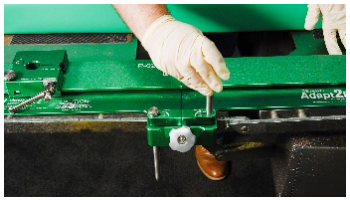
9. Check flexion for final approval of freedom of the drape to allow full range of motion, when the positioner is locked on the drapes.

10. Position clamp directly under the holes in the baseplates in accordance with the clamp IFU



See Individual Clamp IFU

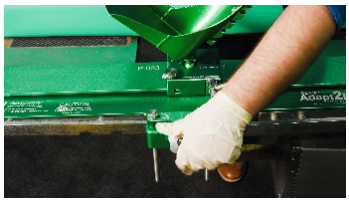
11. Insert a guide pin so that the protruding pin fits into the keyhole and goes into the clamp and turn it ¼ turn following the arrow on the baseplate.



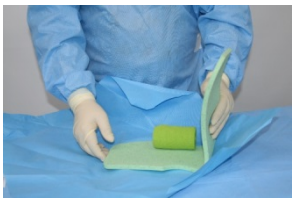
*****Note: Only 2 guide pins are needed to set up the positioner 2 extra guide pins have been provided in case of dropping or missing pins*****

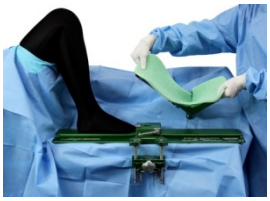
12. Place the other guide pin into the baseplate and awaiting clamp

13. Tighten the knob of the Clamp



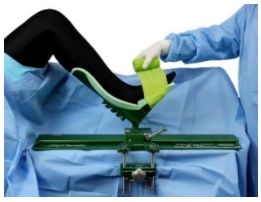
14. Insert IMP® Patient Protective Pad® into the sterile boot.
For patient's safety, always use IMP® Patient Protective Pads®.





15. Wrap cohesive bandage around the foot starting with a minimum of six (6) foot wraps, tear, and finish the wrap above the distractor block around the calf.

Note: Do not wrap cohesive bandage over the distractor block on the back of the boot.

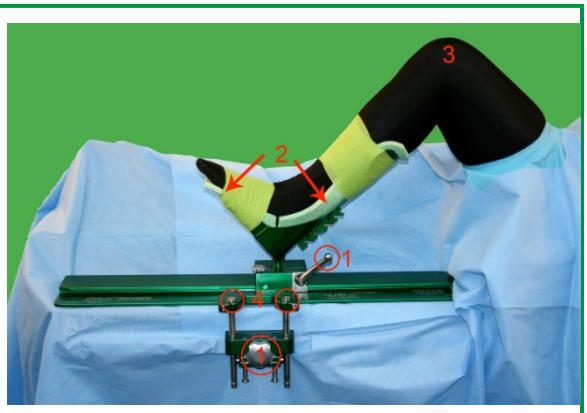


After surgery - recommend spraying the positioner with a pre-treatment prior to resterilization

Safety Test

Tighten clockwise to lock the knob and lever.

1. Check all knobs and levers are locked.
2. Patient is fully protected by IMP® foam and cohesive wrap
3. Patient's leg can reach full flexion.
4. Check that both Guide Pins are fully seated and locked.



Cleaning and Sterilization Procedure

NOTE: ALL SOLUTIONS MUST BE COMPATIBLE WITH ALUMINUM & STAINLESS STEEL

Recommended Washer / Decontaminator Instructions:
• Remove carriage from positioner – Reference De Mayo Adapt2Fit Carriage Instructions for Use
• **Note: All carriage components including the handle, boot and baseplate should be controlled so that the same components all get reassembled together**
• Soak the product in an enzyme solution (example: TRI-POWER ENZYMATIC CLEANER from UNITED BIOTECH www.united-biotech.net or similar). Follow the manufacturer's direction for dilution and soaking time. ***Validated by 3rd party for 15-minute soak time with Tri-Power**
• Put through washer / decontaminator according to manufacturer's instructions **with a detergent up to a pH of 9.0.** ○ **NOTE**: Select cycle that does not include lubrication.
Recommended Hand Cleaning Instructions:
• Pre-Soak the product / components in an enzyme solution. Follow the manufacturer's direction for dilution ratio and soaking time.
• Rinse the product in warm tap water.
• Wash the product with an instrument detergent up to a pH of **9.0** or enzyme product
• Rinse the product in warm tap water
• Soak or wipe the product down with a hospital approved and / or EPA approved germicide according to instructions.
• Rinse the product in warm tap water
• **DO NOT LUBRICATE THE KNEE POSITIONER PRIOR TO STERILIZATION OR IN THE OR.**
• Assemble carriage to baseplate
• Assemble Pin Base to baseplate. Ensure original boot is placed with the assembled baseplate
• Dry thoroughly and wrap

Recommended Sterilization Instructions:
• Ensure that all parts are thoroughly cleaned.
• Make sure that all movable parts are loose and can move freely.
• *DO NOT PLACE POSITIONER IN A MILK BATH OR LUBRICATE*
• If using a sterilization case, follow Instructions For Use for the product
• Double wrap in two disposable wraps. Use 48 inch x 48 inch wraps. ○ (Approximately 122 cm x 122 cm).
• If using sealed container, follow Instructions For Use for the product

• Run normal vacuum cycle for your institution
• **STEAM STERILIZATION ONLY- ALL OTHER STERILIZATION METHODS NOT VALIDATED**

MINIMUM PARAMETERS PRE-VAC STERILIZATION

Product Part Number	**With Sterilization Case Case Part Number**	**Temperature Setting**	**Exposure Time**	**Dry Time**
De Mayo Adapt2Fit Modular Knee Positioner 1 Baseplate: 1020 1 Clamp: 713-717, 120 1 Boot: 803-ABD, 803-ABDA or 803-CBD	706-A2	270°F to 272°F 132°C to 134° C	4 minutes	30 minutes
	Without Sterilization Case			
De Mayo Adapt2Fit Modular Knee Positioner 1 Baseplate: 1020 1 Clamp: 713-717, 120 1 Boot: 803-ABD, 803-ABDA or 803-CBD	Double Wrap or Sealed Container	270°F to 272°F 132°C to 134° C	4 minutes	20 minutes
Accessories				
1 Distractor: 907* or 621	706-M	270°F to 272°F 132°C to 134° C	4 minutes	20 minutes
1 Distractor: 907* or 621	No Case Wrap **or** Sealed Container	270°F to 272°F 132°C to 134° C	4 minutes	20 minutes

*Denotes discontinued item

CAUTION: The positioner must be cool before applying to the patient

Sterilization Parameters Certified by:

- Micro Test Laboratories (now Accuratus Lab Services)
- Accuratus Lab Services
- HIGHPOWER Validation Testing and Lab Services

Scan for additional documentation

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