

Instructions For Use

De Mayo Adapt2Fit® Modular Knee Positioner

Catalog Number 1020

Doc Title IFU-IMP-0018

Version 03



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 doesn't follow these instructions for use.
- Patient is not to be in contact with device for over 24 hours.
- Untrained personnel must review and understand the IFU.
- Follow the IFU to correctly use the product.
- Safety: Always use IMP Patient Protective Pads
- DO NOT LUBRICATE POSITIONER OR PLACE IN A MILK BATH
- Max number of reuses:
 - The maximum number of reuses is not a fixed value. It depends on several factors that influence product wear and safety. The device must be inspected before each use, and reuse should be discontinued if any of the following conditions are observed:
 - Unplanned Carriage Movement: If the carriage does not remain securely in place during
 use, indicating a loss of stability or locking integrity.
 - Excessive Medial-Lateral Movement of the Boot When the Carriage is Locked: If
 noticeable side-to-side movement is present when the carriage is in the locked position,
 suggesting wear or improper function.
 - Irreparable Wear and/or Damage to the Baseplate: If any part of the baseplate, shows significant wear, deformation, or damage that cannot be repaired, compromising the device's safety and function.

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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Product Identification

Part No	Product Name	UDI-DI
1020-BP	De Mayo Adapt2Fit® Modular Knee Positioner 32" Baseplate	00696588006529

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
398-ABG	Armboard Pad Set 26" x 6" (2 per set)	00696588000763
398-ABG	Armboard Pad Set 26" x 6" (2 per set)	00696588000763
398-LG	OR Table Pads Green (3 pc set) 398AG (1 each) and 398CG (2 each)	00696588004112
398-LG	OR Table Pads Green (3 pc set) 398AG (1 each) and 398CG (2 each)	00696588004112
621	De Mayo Universal Distractor® 2.0	00696588007151
706-A2	De Mayo Adapt2Fit® Modular Knee Positioner Sterilization Case - 20" x 10" x 8"	00696588006482
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
803-GP-10	Sterile Patient Protective Pad for IMP® Knee Positioners & Cohesive Wrap (10 / Case)	00696588002880
907*	De Mayo Universal Distractor®	00696588001388
907-GP-10	Sterile Pressure Protector Pad® for De Mayo Universal Distractor® (10/case)	00696588002897
923-A2	De Mayo Adapt2Fit® SteriPod System- Filtered Container & Insert	00696588007618
1020-CT	De Mayo Adapt2Fit® Baseplate Connector	00696588006680
1020-PCT	De Mayo Adapt2Fit® Baseplate Plastic Connector	00696588008066

^{*}Denotes discontinued item



Symbol Glossary

Symbol	Title	Description	Standard
EC REP	Authorized Representative in the European Community	Indicates the authorized representative in the European Community.	ISO 15223-1:2021
LOT	Batch Number	Indicates the manufacturer's batch code so that the batch or lot can be identified.	ISO 15223-1:2021
REF	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 15223-1:2021
\triangle	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 15223-1:2021
C€	Complies with European Directives.		
(ii	Consult instructions for use.		ISO 15223-1:2021
<u>~</u>	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 15223-1:2021
MD	Medical device		ISO 15223-1:2021
DATES	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 15223-1:2021
UDI	Unique device identifier	Indicates a carrier that contains unique device identifier information	ISO 15223-1:2021



Instructions for use:

 Replace OR Table Pads with green IMP® Table 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. Create a 3"-5" fold in the final drape under the buttocks to reduce the possibility of tearing the drape.



5. Open the sterilization tray. Remove the inner sterile basket with the contents of the positioner.





- 6. Inserting the Connector Link
 - a. (1020-PCT) Insert the Plastic Baseplate Connector into one half of the Baseplate until the Connector is fully inserted. When the connector link is fully insterted, there should be an audible click.





b. (1020-CT) Insert the Aluminum Baseplate Connector into one half of the Baseplate until the Connector is fully inserted.







- 7. Connecting the Baseplates
 - a. (1020-PCT) Connect the other half of the Baseplate by sliding it onto the protruding Plastic Baseplate Connector until the Baseplates meet. There should be an audible click when connected.





 (1020-CT) Connect the other half of the Baseplate by sliding it onto the protruding Aluminum Baseplate Connector until the Baseplates meet.





8. Reference De Mayo Adapt2Fit Carriage Removal and Replacement Sheet within this IFU. Place the De Mayo Adapt2Fit® Modular Knee Positioner in the well to achieve maximum flexion. Adapt2Fit is designed to fit perfectly into the well with no overhang.



9. Check flexion for final approval of freedom of the drape to allow full range of motion.



10. Position Clamp directly underneath the holes in the baseplates in accordance with the Clamp IFU.



See Individual Clamp IFU

11. Insert the guide pins through the Baseplate and the Clamp. Once the pins are fully through, turn the pins according to the arrows on the Baseplate.





Note: Only 2 guide pins are needed to set up the positioner. 2 extra guide pins have been provided in case pins are dropped or are missing 12. Tighten the knob of the Clamp.





13. Insert IMP® Patient Protective Pad® into the sterile boot.





For patient's safety, always use IMP® Patient Protective Pads®. CAT #803-GP-10 and/or 907-GP-10.

14. Place the patient's foot in the boot. Wrap the cohesive bandage around the foot at least six (6) times, tear the wrap, and use the rest of the cohesive bandage to wrap around the calf above the distractor block.







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

After Surgery Recommendation - Spray 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
- 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 <u>www.united-biotech.net</u> 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.
 - o **NOTE**: Select cycle that does not include lubrication.
 - NOTE: Use of ultrasonic cleaners are not validated

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

^{*}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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Carriage Removal and Replacement Adapt2Fit® Knee Positioner®

ADAPT2FIT COMPONENTS

e (Z)	uo_	du
Base Plate (2)	Teflon Pad	Clamp
No. of the last		RO.
E-Brake	Boot	Drop Pins (4)
Handle	Carriage	Connector Link
Thumbscrew	Threaded Rod with L-Plate	Sterilization Tray

CARRIAGE REMOVAL

Opt 1: End Remova

with L-Plate is too tight to remove with fingers, put the handle back on and turn counter-clockwise to loosen. and remove the handle. B) Unscrew entire threaded A) Unscrew the Thumbscrew with attached spring rod with L-Plate using your fingers. If threaded rod





Slide the carriage toward the "Loading End" (right side) curved end of the top plate as indicated in the photo. of the Base Plate and slide the carriage around the



Unscrew the E-Break from the carriage. Do not remove Teflon pad.





Opt 2: Middle Removal

A) With the 2 baseplates split apart, B) pull the carriage from the base plate.

Unit must be safety tested before final processing.

SAFETY TEST

the Boot Ball securely in the Check that the handle locks

12 - 3 o'clock range.

Check that the Boot can be removed and carriage can

plate when the handle is slide freely on the base

unlocked.





Remove the handle and E-Break for cleaning. Do not remove the Teflon pad.

CARRIAGE REPLACEMENT -

Opt 1: End Replacement

spring-loaded handle out and turn either clockwise or

With the boot locked into the carriage, pull the

counter-clockwise to make any final adjustments.

Screw the E-Brake into the carriage.

carriage, slide the carriage from the "Loading End" A) While holding the Teflon Pad underneath the onto the base plate. B) Then slide the carriage around the curved end of the Base Plate.





catch the inside of the carriage. B) Insert boot ball into A) Using your fingers, screw in the threaded rod with L-Plate into the carriage until the threads of the rod carriage and continue to tighten until finger tight. 7

A) While holding the Teflon Pad against the carriage,

Opt 2: Middle Replacement

B) slide the carriage onto the left-hand Baseplate.







A) Replace the handle in the 11 - 1 o'clock range and B) screw in the Thumbscrew with attached spring until tight. C) The handle now becomes a spring-loaded nandle.









ensure that all Adapt2Fit components are present.



This guide is for the Adapt2Fit Knee Positioner only. No tools required.



documentation



Catalog Number 1020