TECHNIMOUNT
EMS®

BRACKET PR SERIE® 60 – FL

USER MANUAL

AIRPLANES/ROTOCRAFTS FAA 14 CFR TECHNIMOUNT ENGINEERING

SAFETY AND FLEXIBILITY
WHERE IT MATTERS MOST

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For any issues with your Technimount product, its components, or for any technical questions during the installation, operation, or maintenance, please contact Technical Support at techsupport@technimount.com.

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1. General Mentions and Considerations

This user manual includes detailed product information, standards and guidelines to assist the administrator/ manager/supervisor and biomedical technician (or equivalent) with the unpacking, assembling (when indicated), maintenance, training and skills assessment of the EMS and clinical personnel, as well as specific user-related information to safely and effectively operate the Bracket Pro Serie 60 – FL (hereinafter called BP60–FL).

Please read the user manual thoroughly to fully assess, comprehend, then relay its content to EMS and clinical personnel during training, to warn them of any potential danger of its abuse, how to safely use the product and provide a safe environment for patients as well as themselves. Your established internal protocols should be updated to include the Technimount product(s) standards, guidelines, requirements and safety recommendations included within this documentation. The user manual should remain available to users when needed and relayed if the product is subsequently sold.

NOTE : Technimount continually seeks advancements in product design and quality. While the user manual contains the most updated product information available at the time of printing, it may contain minor differences from the current version, including image references. For more information, please contact Technical Support at techsupport@technimount.com.

NOTE : Technimount reserves the right to change part numbers and products without notice. Please contact Customer Service at customerservice@technimount.com to ensure product options and availability.

1.1. Intended Use

The BP60–FL is a mounting bracket designed to aid trained EMS and clinical personnel secure and transport the Hamilton-T1 Transport Ventilator during air emergency medical services and critical care transport.

1.2. User Competency

To safely operate the BP60–FL, personnel must have the required skill level to comply with their function and level of interaction with the mounting bracket. Training should be given to EMS and clinical personnel prior to them using the BP60–FL. Refer to the « Skills Assessment of the EMS and Clinical Personnel » section on page 29 to evaluate their competency.

Indicates who the content is intended for and the level of competency required. The definitions of the three (3) levels of competency are specified below.

- **Competent (trained EMS and clinical personnel):** Has received the required training, is sufficiently knowledgeable to safely operate the product and have passed the skills assessment (refer to the « Skills Assessment of the EMS and Clinical Personnel » section on page 29).

  NOTE : Any member of the EMS and clinical personnel who has not received the required training and lacks the knowledge needed to safely operate the mounting bracket must not use the product.

- **Proficient (administrator/manager/supervisor):** Has in-depth knowledge and product comprehension, and is familiar with standards and guidelines. Skilled to train the EMS and clinical personnel on how to safely

use the product.

- **Advanced (biomedical technician or equivalent):** Has extensive mechanical experience. Skilled to perform the unpacking, assembly, safety checks and condition-based maintenance procedures as detailed herein, or the basic troubleshooting, upgrade and/or replacement procedures if applicable.

1.3. Warranties

1.3.1. Warranty Policy

This statement constitutes Technimount's entire warranty policy with regards to Technimount products. Technimount makes no other warranty or representation, neither expressed nor implied, except as stated herein. There is no warranty of merchantability or warranty of fitness for any particular purpose. Under no circumstances will Technimount be held liable hereunder for incidental or consequential damages, arising from or in any manner, related to sales or use of any such product.

Technimount E.M.S. Holding Inc. guarantees to the original "Purchaser" of the "Product" with which this "Limited warranty" is included, that the product will be free from "Defects" in workmanship and materials under normal use for a "Warranty period" of one (1) year from the product purchase date by the purchaser. During the warranty period, the product will be repaired or replaced according to the "Limited warranty" without charge to the purchaser for parts or labor. The parts and product may be repaired or replaced with new or refurbished parts or products. Herein this Limited Warranty, "Refurbished" means parts and products which have been returned to the factory, specifically. If the product is repaired or replaced within the warranty period, the greater of the remaining warranty period will apply, or three (3) months from the date of repair or replacement. If the product is repaired or replaced after the warranty period has expired, the warranty period for the repair or replacement will expire three (3) months after the repair or replacement date.

1.3.2. Limited Warranty

Technimount products are intended to secure in place medical devices only in the case of a single emergency landing, and must thereafter be immediately replaced. If the end user uses a Technimount product following an emergency landing, it is at the end user's own risk, and Technimount will not be held liable.

The limited warranty does not apply to normal wear that could result from normal use. It does not apply when the product or any of its components have been disassembled or repaired by someone not authorized by "Technimount". It does not cover repair or the replacement of any product or part thereof damaged by neglect, misuse, moisture, liquids, exposure to heat, accidents, abuse, and non-compliance with the instructions for installation and use provided with the product. It does not cover physical damage to the surface of the product. The decision to repair, replace or refuse the coverage is final and at the sole discretion of Technimount, without any compensation or obligation from Technimount. The product, defined as a "mounting bracket" or "bracket", is specifically designed to secure and transport the Hamilton-T1 Transport Ventilator during air emergency medical services and critical care transport, and should only be used to fulfill this requirement. Any other use will void the warranty and Technimount shall not be held liable on any claim if the product has been modified or adapted for use.

1.3.3. International Warranty Clause

This warranty abides by the Canadian domestic policy. Warranty outside Canada may vary by country. Please contact Customer Service at customerservice@technimount.com for more information.

1.3.4. User Liability

The purchaser and administrator are responsible to validate regulations and standards for safety in their region, to comply with applicable safety regulations. Technimount is not responsible to inform the purchaser or the administrator of any applicable legislation for safety in their area.

The administrator is responsible for providing proper training to any personnel who will install, operate and perform maintenance on Technimount products.

1.4. Claims

1.4.1. Damaged or Defective Merchandise

ICC Regulations require that claims for damaged merchandise must be made with the carrier within fifteen (15) days of the reception date. **Do not accept** damaged merchandise unless such damage is noted on the delivery receipt at the time of reception. Upon prompt notification, Technimount will file a freight claim with the appropriate carrier for damages incurred. Claims are limited in amount to the actual replacement cost. If the claim has not been received by Technimount within the fifteen (15) day period following the date of delivery, or the damage was not noted on the delivery receipt at the time of receipt, the customer will be responsible for the full payment of the original invoice.

Claims for any short or broken merchandise must be made within thirty (30) days of invoicing. For details, refer to the « Claim Process » section on page 8 or contact Customer Service at customerservice@technimount.com.

1.4.2. Return Policy

Technimount products may be returned up to sixty (60) days from the reception date, if:

- The received product does not match what was originally ordered.
- The product does not meet the Technimount technical sheet specifications.
- The product is not compatible with the system on which it was intended to be installed on.

To return a Technimount product,

- A Return of Merchandise Authorized (RMA) must be requested and approved by Technimount prior to returning the product.
- Products must be returned undamaged and in its original packaging, appropriately identified with the approved RMA number. Returns will not be approved on a modified or damaged item.
- Charges may apply if the package received is damaged or items are missing.
- Purchaser is responsible for a restocking fee (refer to « Table 1: Restocking fees » table on page 8).

Table 1: Restocking fees

RESTOCKING FEES	
Prior to thirty (30) days	10%
Prior to forty-five (45) days	25%
Prior to sixty (60) days	30%

For any manufacturing defect, refer to the conditions within the warranty policy or contact Customer Service at customerservice@technimount.com for additional information.

1.4.3. Return of Material Authorization (RMA)

The Technimount Customer Service department is responsible for all merchandise returns and will provide a Return of Merchandise Authorization (RMA) number, upon approval. The RMA must be printed and placed on the returned merchandise. Technimount reserves the right to charge shipping and restocking fees (refer to « Table 1: Restocking fees ») for the returned items. Special, modified, or discontinued items are not subject to returns.

1.4.4. Claim Process

Upon reception of the returned merchandise, a thorough inspection will be performed. If the merchandise is compliant with the return policy or it is found that the product is defective, Technimount will take corrective actions and close the claim. If, however, it is found that the product is not defective, but rather misused or abused, the product will not be covered by the warranty. Details of our findings and conclusions will be provided shortly thereafter. To submit a claim, contact Customer Service at customerservice@technimount.com to obtain a Return of Material Authorization (RMA) form and return instructions.

2. General Safety Guidelines

The content in this section is intended for personnel who have a competent, proficient or advanced of competency. Refer to the « User Competency » section on page 5 if needed.

Pictograms, safety symbols and labels are used to alert the user to a potentially hazardous situation, which, if not avoided, may endanger the patients or the EMS and clinical personnel or clinical personnel, and/or damage the product. This includes the special care necessary for the safe and effective use of the Technimount product to avoid damage that may occur from use or misuse.

The terms "Warning" and "Caution" herein carry special meaning and should be carefully reviewed before reading the « Safety Measures » section on page 12.

WARNING – Indicates a hazardous situation that, if not avoided, could result in death or serious injury, and/or damage the product.

CAUTION – Indicates a hazardous situation that, if not avoided, could result in minor or moderate injury, and/or damage the product.

2.1. Symbols and Definitions

WARNING – Risk of Injury

Indicates when a misuse of the Technimount product could result in injuries to the patients or EMS and clinical personnel, and/or damage to the product.

CAUTION – Safe Practice

Alerts the reader to pay special attention to the recommendations and methods outlining how to safely operate the product to minimize risks to the patients, EMS and clinical personnel, and/or to the product.

CAUTION – Safe Handling and Operation

Alerts the reader to pay special attention to the recommendations for safe use of the product, and of potentially hazardous situations that could result in minor injuries to the patients or EMS and clinical personnel. This includes the special care necessary for the safe and effective use of the product to avoid damage that may occur from use or misuse.

CAUTION – Safe Working Load (SWL)/Load Balance

Indicates the maximum charge for a safe use of the product.

CAUTION – Follow Instructions for Use

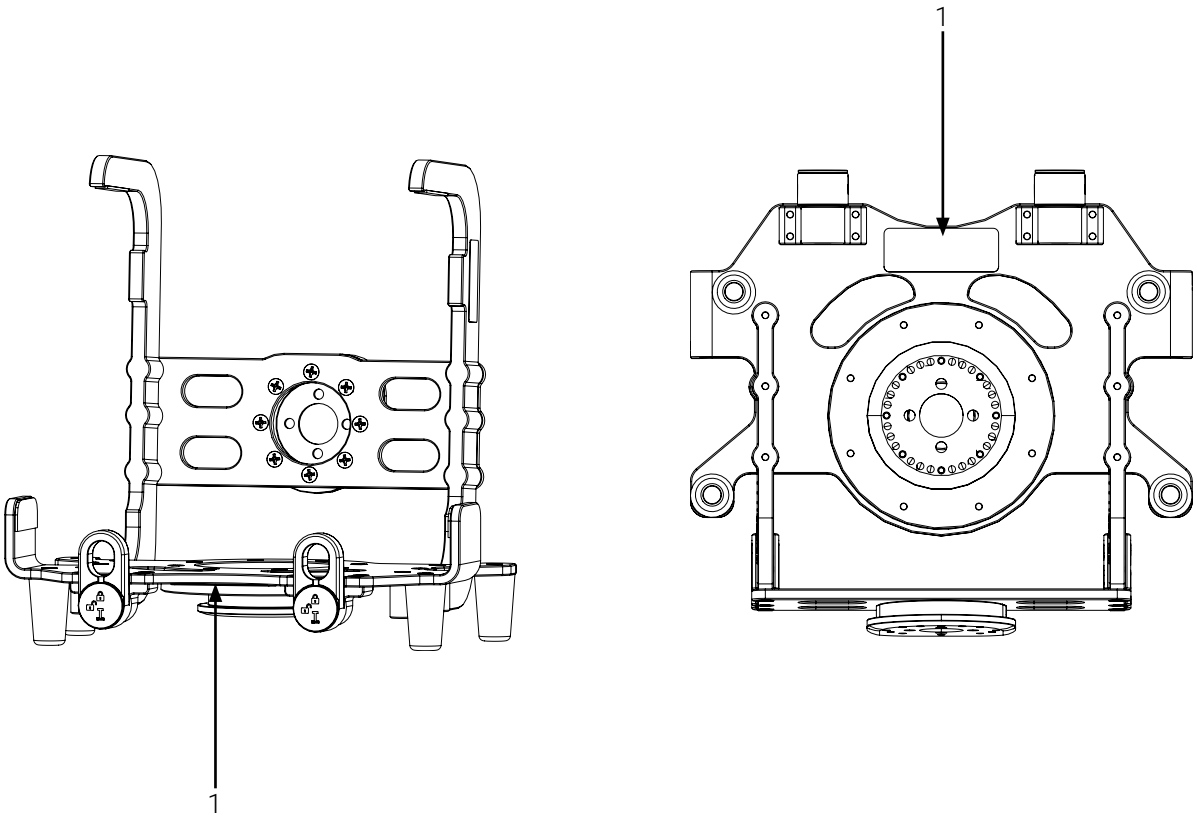
Call for action. Reminds the reader to consult the user manual for information.

2.2. Labels

Labelling on the surface of the Technimount product, quickly identify potential risks and provides information to the user. A manufacturing label, including the serial number and the safe working load specifications (Figure 1), as well as identification labels (Figure 2), can be seen on the Technimount product.

NOTE : The illustrations herein are for user-comprehension purposes and may differ from your actual configuration. The labelling specifications below apply to any compatible mounting bracket. Refer to their user documentation for the safety guidelines and safe use, if needed.

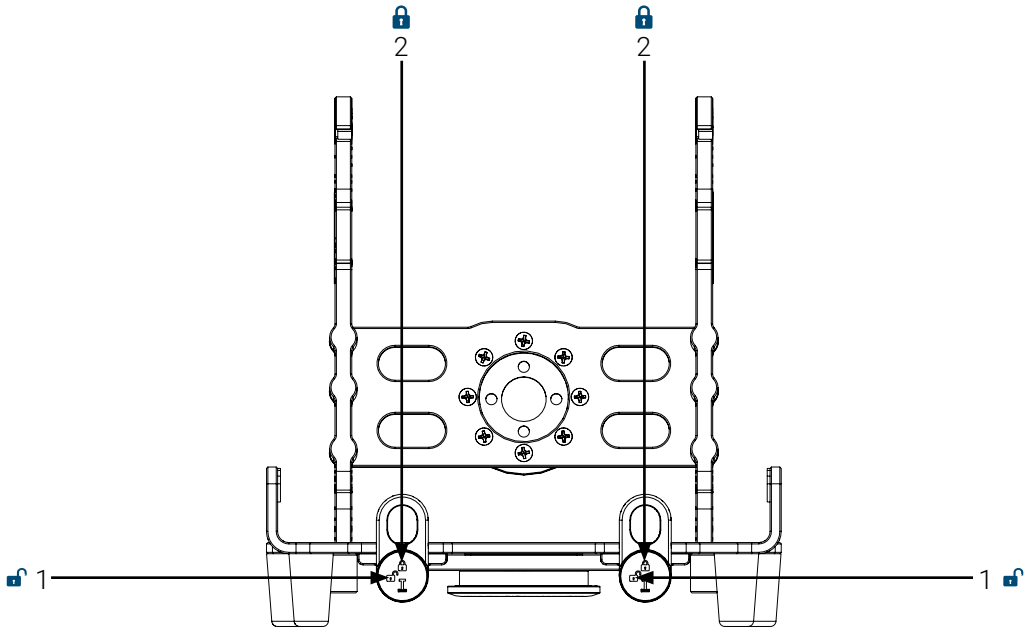
2.2.1. Manufacturing Label



1. Manufacturing label

Figure 1: Location of the manufacturing label

2.2.2. Identification Labels



1. "Unlocked" position label (2X; 1 per rotary locking handle)	2. "Locked" position label (2X; 1 per rotary locking handle)

Figure 2: Location of the identification labels

2.3. Safety Measures

Carefully read all the safety measures herein before operating the Technimount product, and always abide by all the safety guidelines identified within this document.

Specific safety measures relating to the safety checks and the conditioned-based maintenance, intended for personnel who have an advanced level of competency, can be found in the « Maintenance » section on page 32.

WARNING – Unauthorized Systems

- **Under no circumstances** should the BP60–FL be installed on, connected to, or used in combination with any third party/competitor mounting system, mounting bracket, medical device or similar equipment, unless Technimount has provided a formal written confirmation of compatibility for the specific configuration.
- The BP60–FL was specifically engineered to be used with the Hamilton-T1 Transport Ventilator. Using any other medical device may cause the equipment to fall, operate unpredictably, or fail, resulting in serious injury to the patients or to the EMS and clinical personnel.
- The BP60–FL was engineered and tested exclusively for use with specific authorized and compatible Technimount components and systems. Using any unauthorized and/or incompatible system may cause the equipment to fall, operate unpredictably, or fail, resulting in serious injury to the patients or to the EMS and clinical personnel.
- Technimount assumes no responsibility for any damage, malfunction, or injury arising from use of unauthorized or incompatible systems. Using any unauthorized and/or incompatible system may result in non-compliance with the applicable safety standards and will immediately void all warranties and liability coverage.
- Technimount products are intended for use only within their defined applications, including but not limited to, emergency medical services, critical care transport, and approved healthcare environments. Refer to the « Technical Specifications » section on page 14.
- Regulations and standards for safety are the sole responsibility of the end user. Ensure that your established internal protocols are updated and meet the technical specifications requirements herein (refer to the « Technical Specifications » section on page 14), as well as the local and regional compliance requirements before use.

WARNING – Risk of Injury

- **Do not use** the BP60–FL if there are any loose or missing screws, to avoid risks that may cause damage or the equipment to fall or operate unpredictably, resulting in serious injury to the patients or to the EMS and clinical personnel.
- Immediately stop using the BP60–FL if any serious incident occurs and inform authorized personnel, to contact the Technical Support at technicalsupport@technimount.com for a remedial action plan and report the incident to the applicable regulatory agency.

CAUTION – Safe Handling and Operation

- Always wait until the transport vehicle is immobilized before manipulating installing/removing the medical device in/from the BP60–FL.
- Always ensure that the medical device is secured on the BP60–FL before it is used, to avoid risks that may cause damage or the equipment to fall or operate unpredictably, resulting in serious injury to the patients or to the EMS and clinical personnel.
- Always pay close attention **not to wedge** the power cords or tubing during the installation/removal of the medical device in/from the BP60–FL and/or the BP60–FL on/from the mounting system.

CAUTION – Safe Practice

- Always pay close attention to the safety mechanisms of the BP60–FL, to avoid risks that may cause damage or the equipment to fall or operate unpredictably, resulting in serious injury to the patients or to the EMS and clinical personnel Follow the recommended maintenance plan and its guidelines, as described in this user manual.
- Always practice safely operating the BP60–FL until the manipulations have been perfected, before use with patients. Improper use of the mounting bracket may cause damage or the equipment to fall or operate unpredictably, resulting in serious injury to the patients or to the EMS and clinical personnel.

CAUTION – Safe Working Load (SWL)/Load Balance

- **Do not overload or exceed** the total Safe Working Load (SWL) of the BP60–FL, to avoid tipping incidents or risks of collapsing. Refer to the « Technical Specifications » section on page 14 for the SWL specifications.
- The Safe Working Load (SWL) of the BP60–FL **does not override** the maximum load capacity of another Technimount product or system, neither those of the medical devices, accessories and/or medical equipment accessories that could be placed in bags and drawers for example. To comply with the SWL specification, the entire configuration should be taken into account. Refer to the user documentation of each product applicable to your specific configuration for specifications.

CAUTION – Follow the Instruction for Use

Always refer to your established internal protocols, as well as read and abide by the safety guidelines and instructions provided in the user documentation of the BP60–FL, the mounting system and the Hamilton-T1 Transport Ventilator.

3. Technical Specifications

The content in this section is intended for personnel who have a competent, proficient or advanced of competency. Refer to the « User Competency » section on page 5 if needed.

Product Name	Bracket Pro Serie 60 – FL
Description	Mounting bracket specifically designed to secure and transport the Hamilton-T1 Transport Ventilator during air emergency medical services and critical care transport
Product Code	- 700-20-HM–FL (standard bottom disc) - 700-21-HM2D–FL (standard bottom disc and anti-rotation micro disc) - 700-23-HM-FL-MCD (micro disc and anti-rotation micro disc)
Operating Environment	EMS/CCT (air)
Compliance	Tested in compliance with 14 CFR 23.561, 14 CFR 25.561, 14 CFR 27.561 and 14 CFR 29.561
Expected Service Life	5 years
Compatible Mounting System	- Standard Surface Base (compatible with standard bottom disc) - Micro Base (compatible with micro disc and anti-rotation micro disc) - Q-Protrak (compatible with micro disc and anti-rotation micro disc)
Compatible Medical Devices/ Accessories	Hamilton-T1 Transport Ventilator
Dimensions (W X D X H)	- 700-20-HM–FL – 292.1 mm (11.5 in.) W X 254 mm (10 in.) D X 300.6 mm (11.84 in.) H - 700-21-HM2D–FL – 292.1 mm (11.5 in.) W X 269.6 mm (10.62 in.) D X 300.6 mm (11.84 in.) H - 700-23-HM-FL-MCD – 292.1 mm (11.5 in.) W X 269.6 mm (10.62 in.) D X 300.6 mm (11.84 in.) H
Weight	- 700-20-HM–FL – 2.54 kg (5.6 lb) - 700-21-HM2D–FL – 2.67 kg (5.9 lb) - 700-23-HM-FL-MCD – 2.58 kg (5.69 lb)
Composition	Aluminum, stainless steel and plastic
Total Safe Working Load (SWL)	7.48 kg (16.5 lb)
Operating Temperature	- 35° C to 45° C (- 31° F to 113° F)
Tested and Approved Cleaning Solutions	- Oxivir, 5% Hydrogen Peroxide with Peracetic Acid (AHP) - Lavo 12, 10 000 ppm Sodium Hypochlorite - TNT-100, 5% Quaternary Ammonium Compound - Spectro-Sept, 5% Ethyl Alcohol - Spectrol, 5% EDTA salt
Options	N/A

4. Orientation Illustrations

The content in this section is intended for personnel who have a competent, proficient or advanced of competency. Refer to the « User Competency » section on page 5 if needed.

NOTE : The orientations referenced herein are from the EMS and clinical personnel standpoint, when facing the BP60–FL (Figure 3, Figure 4 and Figure 5).

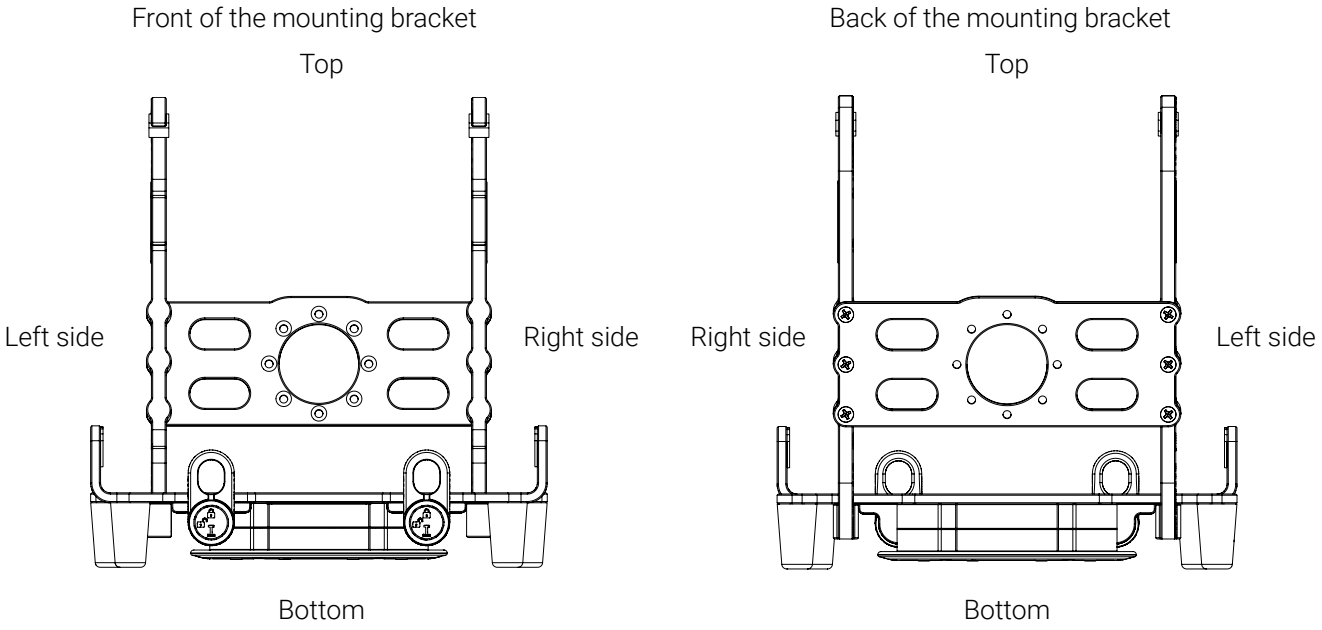


Figure 3: BP60–FL orientation illustration – standard bottom disc

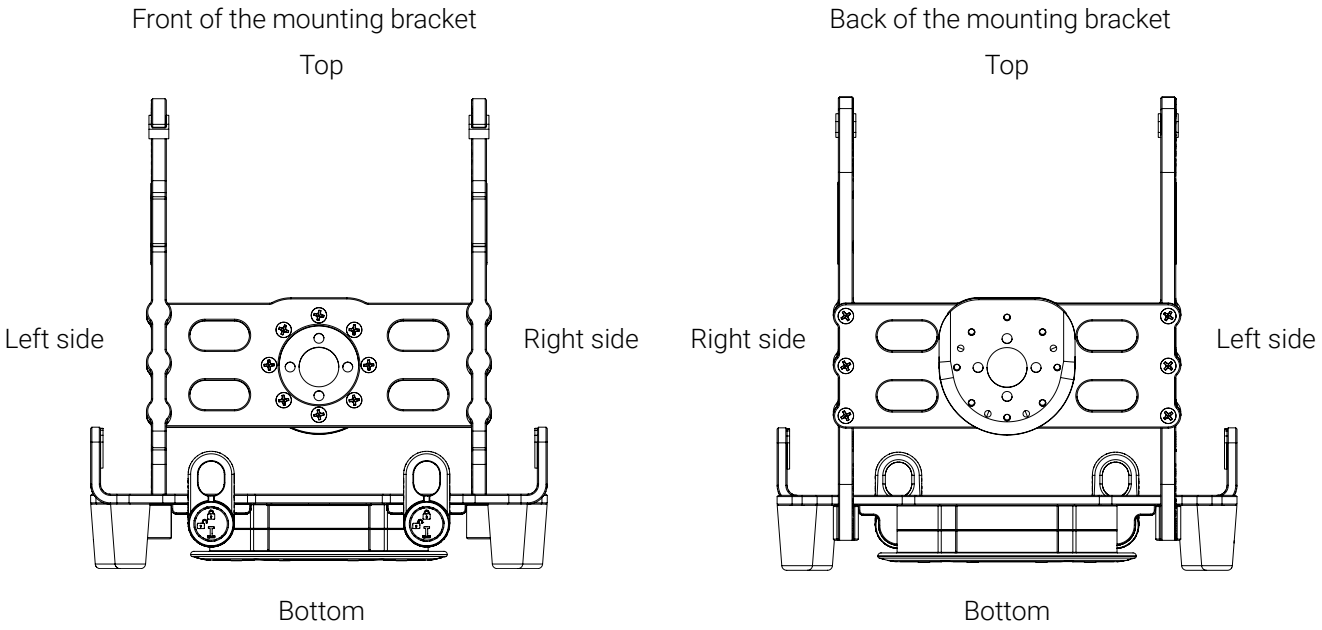


Figure 4: BP60–FL orientation illustration – standard bottom disc and anti-rotation micro disc

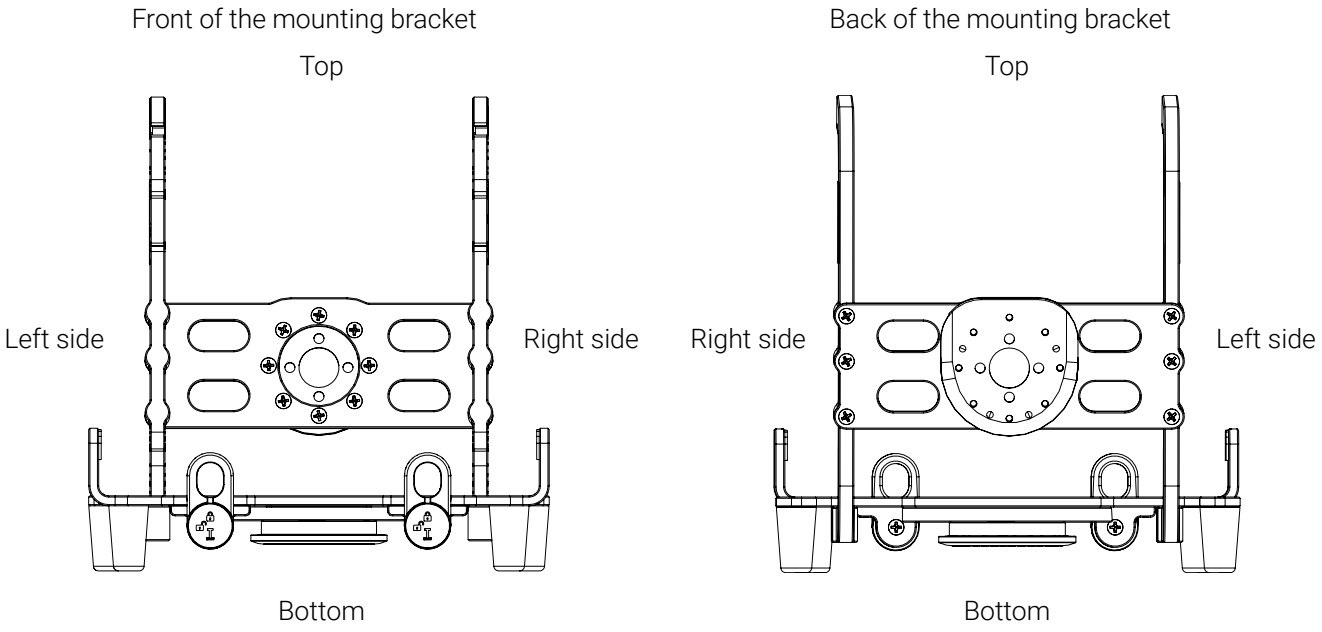


Figure 5: BP60–FL orientation illustration – micro disc and anti-rotation micro disc

5. Illustrated Parts

The content in this section is intended for personnel who have a competent, proficient or advanced of competency. Refer to the « User Competency » section on page 5 if needed.

Figure 6, Figure 7 and Figure 8 illustrates the components of the BP60–FL.

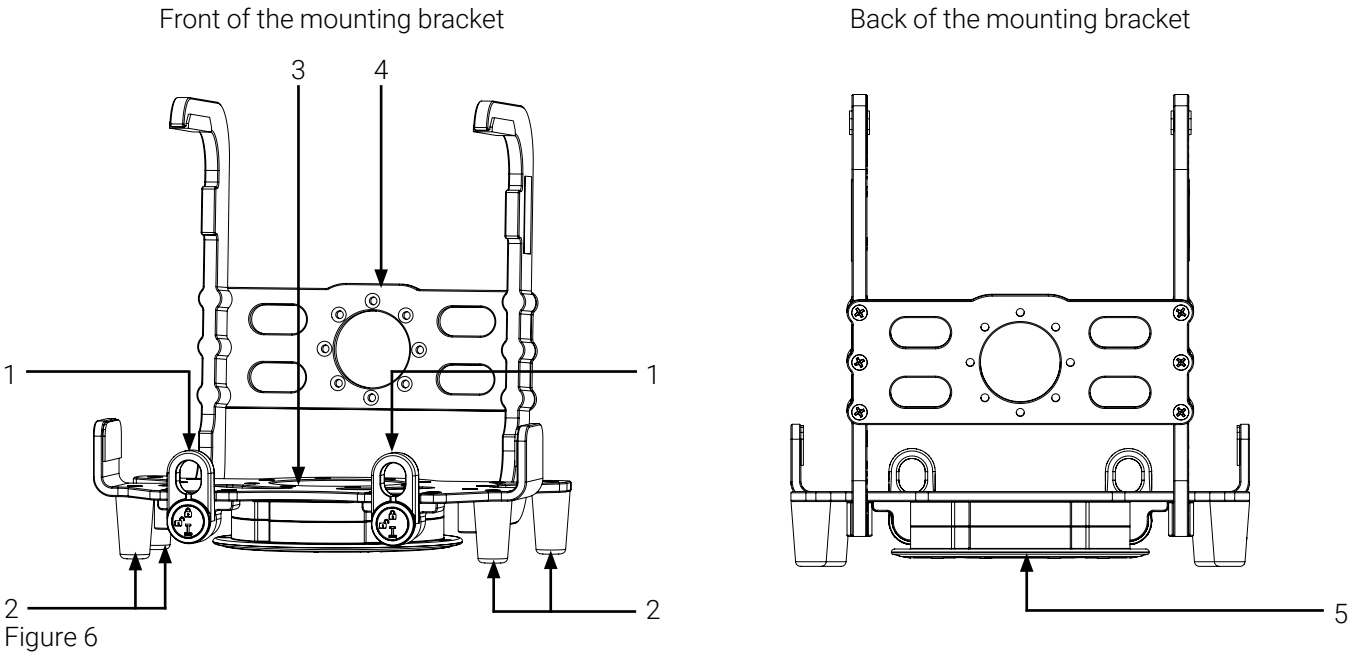
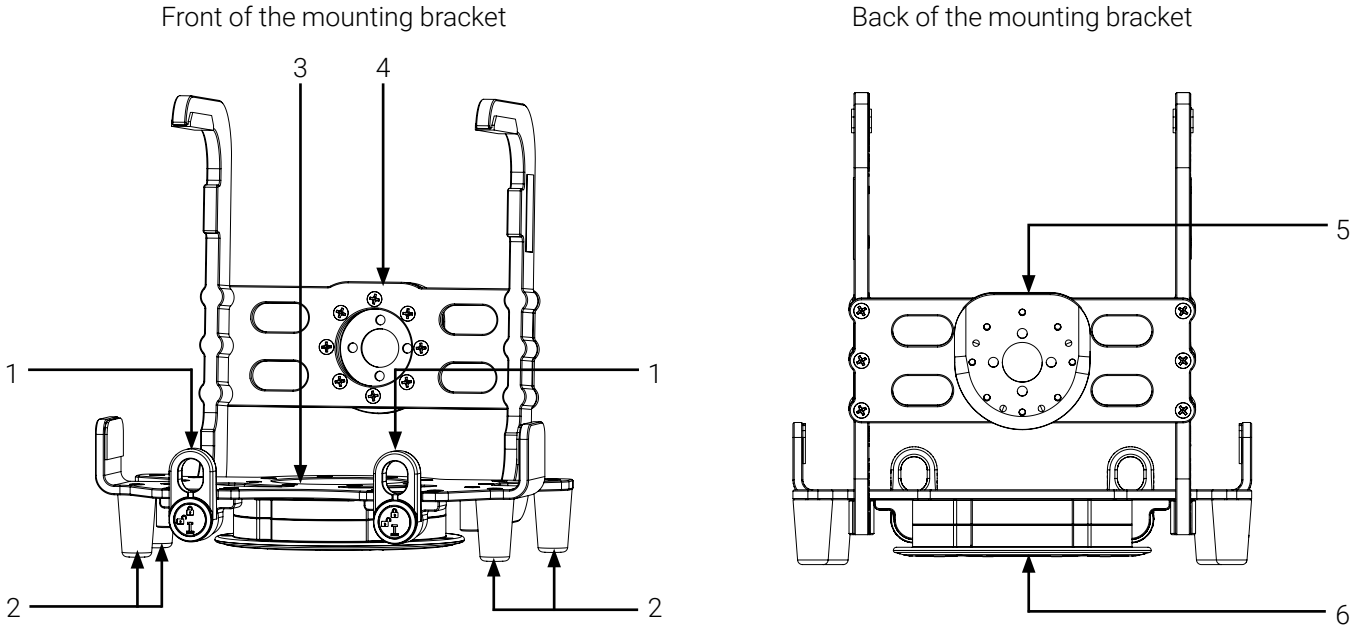


Figure 6

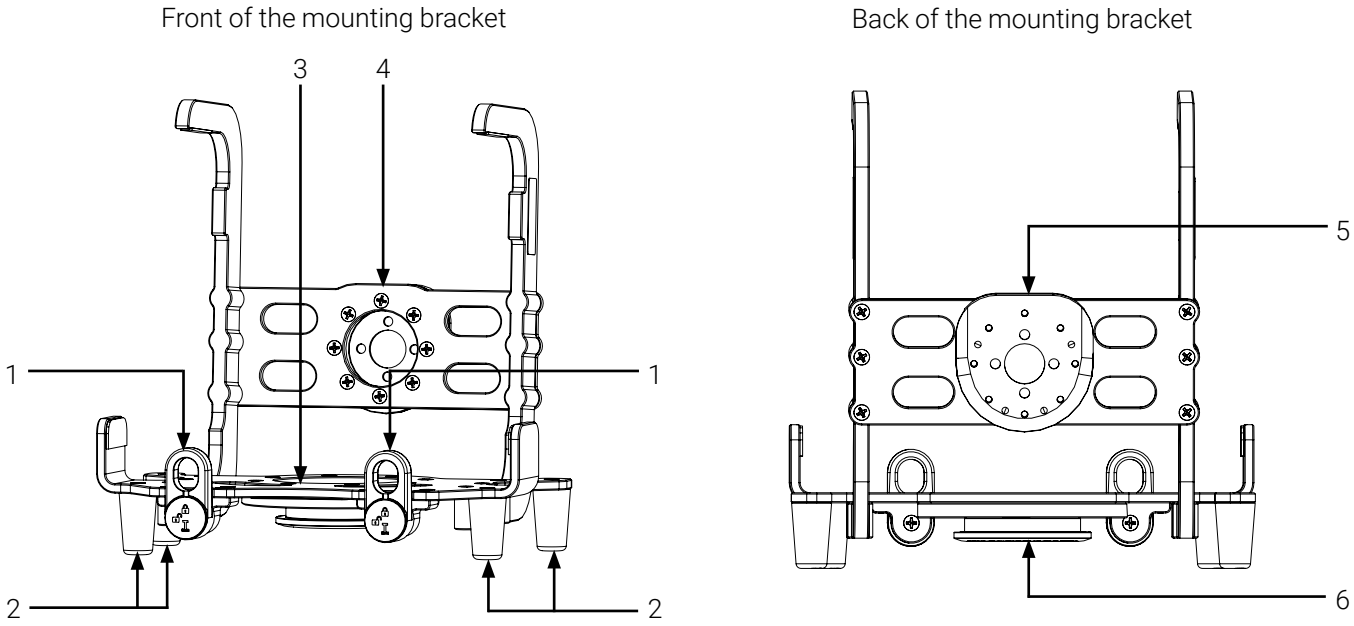
1. Rotary lock (2X)	4. Back plate
2. Foot (4X)	5. Standard bottom disc
3. Bottom plate	

Figure 6: BP60–FL components – standard bottom disc



1. Rotary lock (2X)	4. Back plate
2. Foot (4X)	5. Anti-rotation micro disc
3. Bottom plate	6. Standard bottom disc

Figure 7: BP60–FL components – standard bottom disc and anti-rotation micro disc



1. Rotary lock (2X)	4. Back plate
2. Foot (4X)	5. Anti-rotation micro disc
3. Bottom plate	6. Micro disc

Figure 8: BP60–FL components – micro disc and anti-rotation micro disc

6. Illustrated Dimensions

The content in this section is intended for personnel who have a competent, proficient or advanced of competency. Refer to the « User Competency » section on page 5 if needed.

Figure 9, Figure 10 and Figure 11 illustrates the dimensions of the BP60–FL.

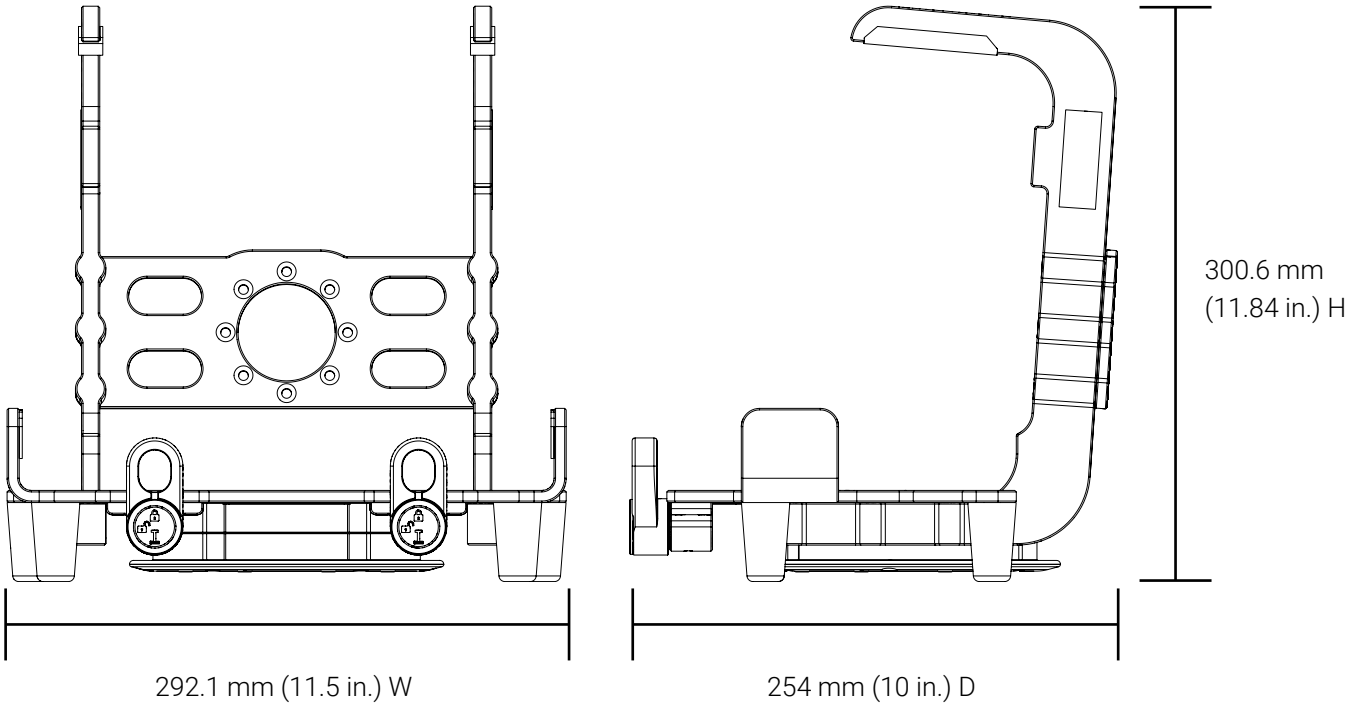


Figure 9: BP60–FL dimensions – standard bottom disc

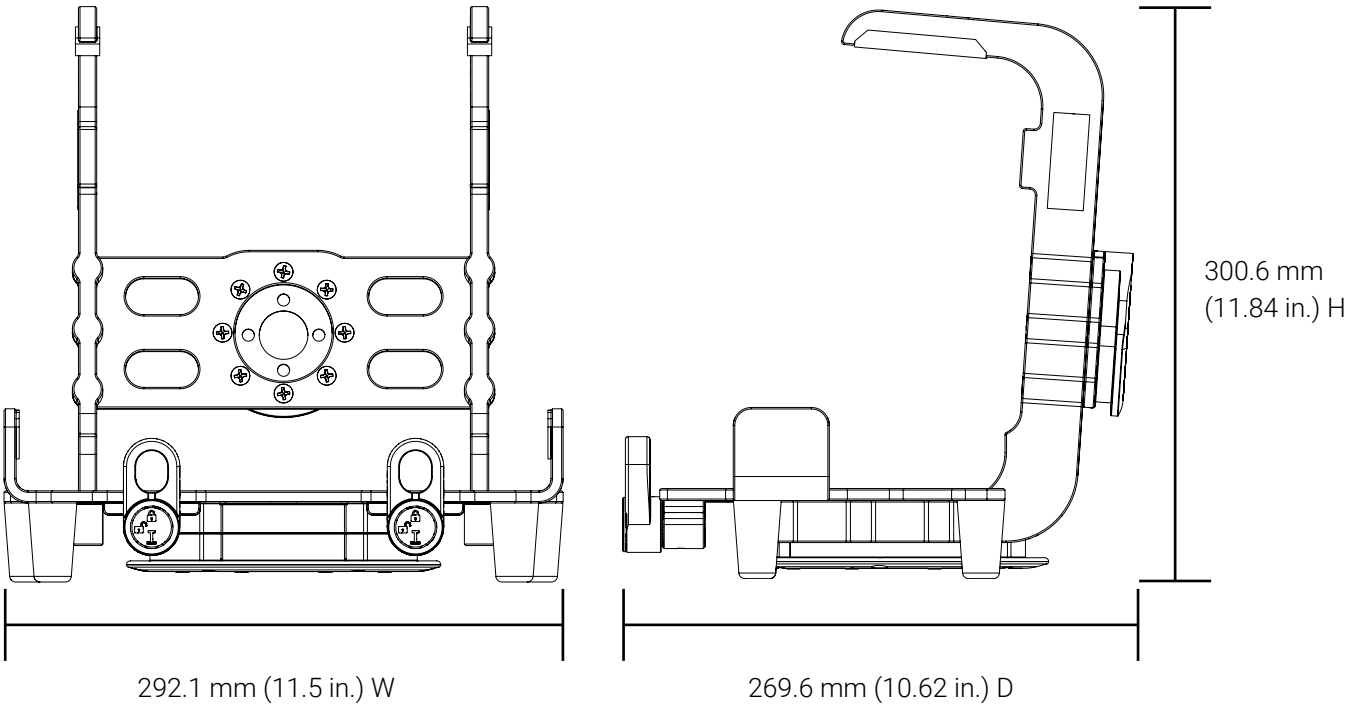


Figure 10: BP60–FL dimensions – standard bottom disc and anti-rotation micro disc

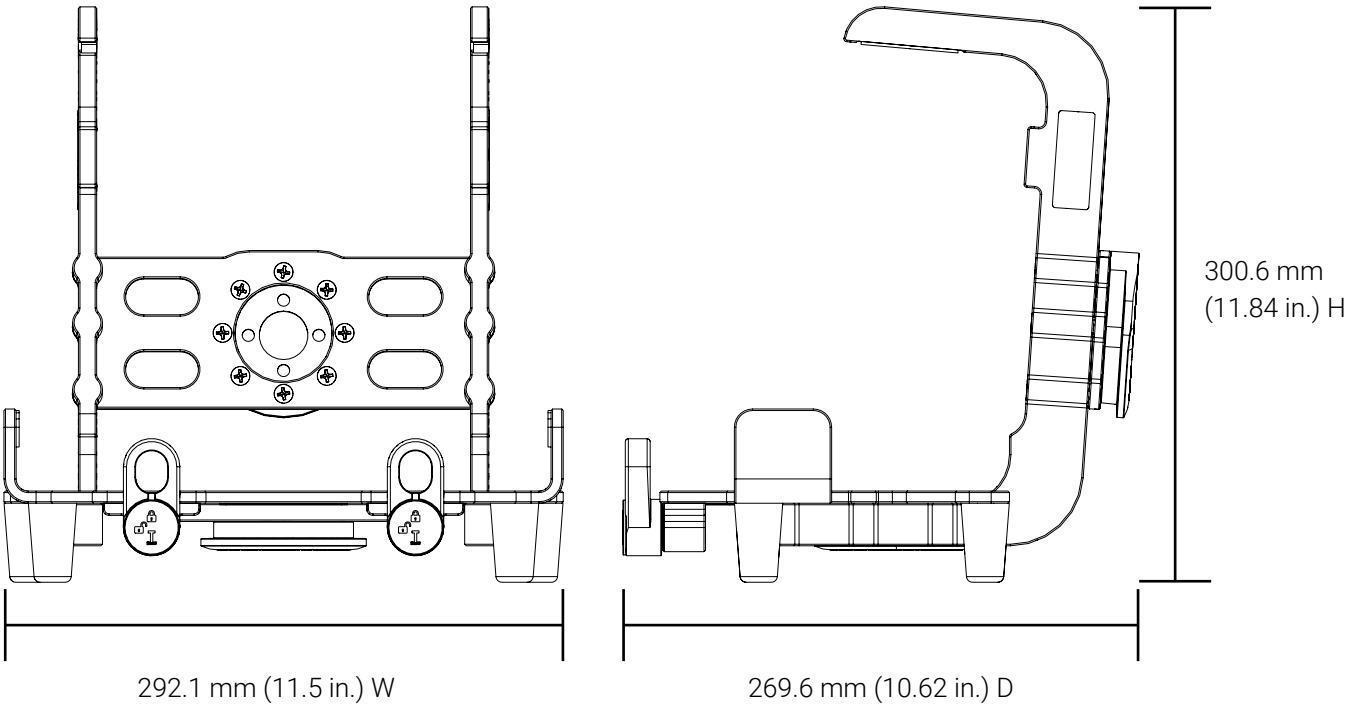


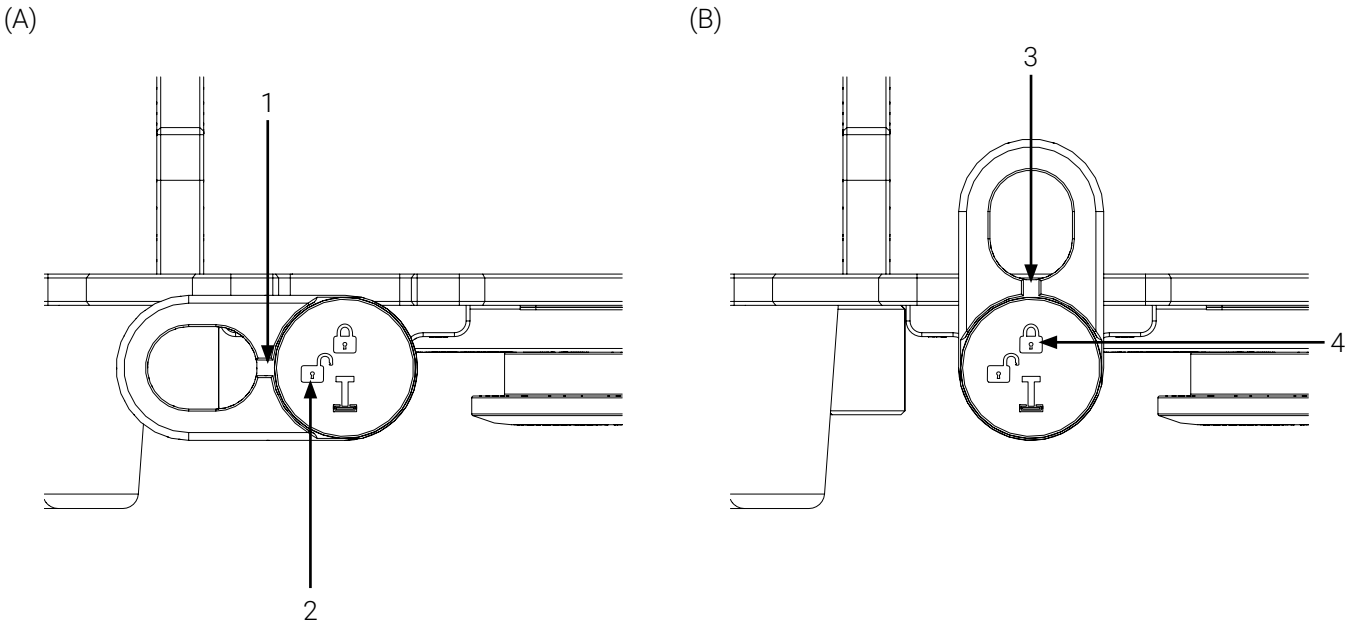
Figure 11: BP60–FL dimensions – micro disc and anti-rotation micro disc

7. Safety Mechanism

The content in this section is intended for personnel who have a competent, proficient or advanced of competency. Refer to the « User Competency » section on page 5 if needed.

NOTE : The illustrations herein are for user-comprehension purposes and may differ from your actual configuration. The instructions apply to any compatible mounting bracket and mounting system. Refer to their user documentation for the safety guidelines and safe use, if needed.

The medical device is secured in the BP60–FL by two (2) rotary locks. When the "Unlocked" icons of the both rotary locks are aligned with the indicators of the mounting bracket, the medical device is secured in the bracket (Figure 12A). When the "Locked" icons of the both rotary locks are aligned with the indicators of the mounting bracket, the medical device can be removed from the bracket (Figure 12B). Refer to the « Illustrated Parts » section on page 17 if needed.



1. Indicator aligned with the "Unlocked" icon
2. "Unlocked" icon
3. Indicator aligned with the "Locked" icon
4. "Locked" icon

Figure 12: Rotary locks

7.1. Locking the BP60–FL

Rotate both rotary locks a quarter turn clockwise, until the "Lock" icons are aligned with the indicators (Figure 13).

(A) (B)

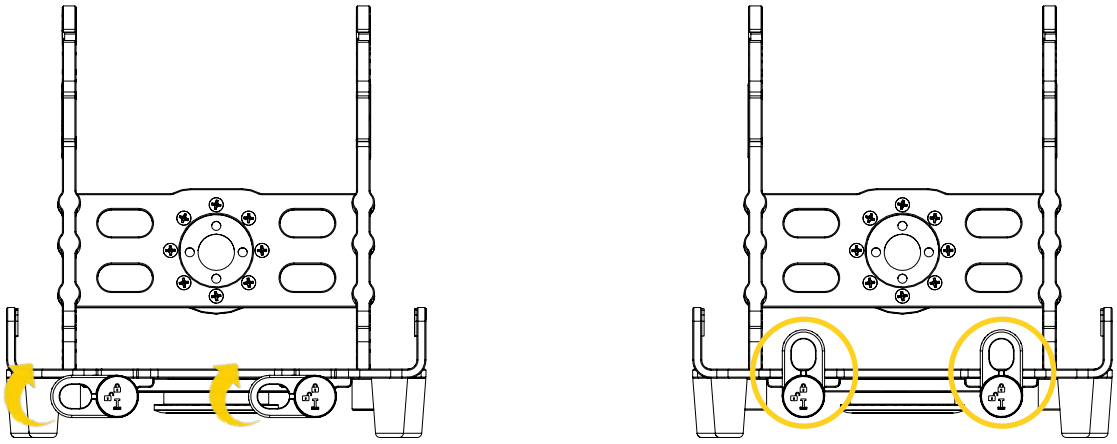


Figure 13: Locking the BP60–FL

7.2. Unlocking the BP60–FL

Rotate both rotary locks a quarter turn counterclockwise, until the "Unlock" icons are aligned with the indicators (Figure 14).

(A) (B)

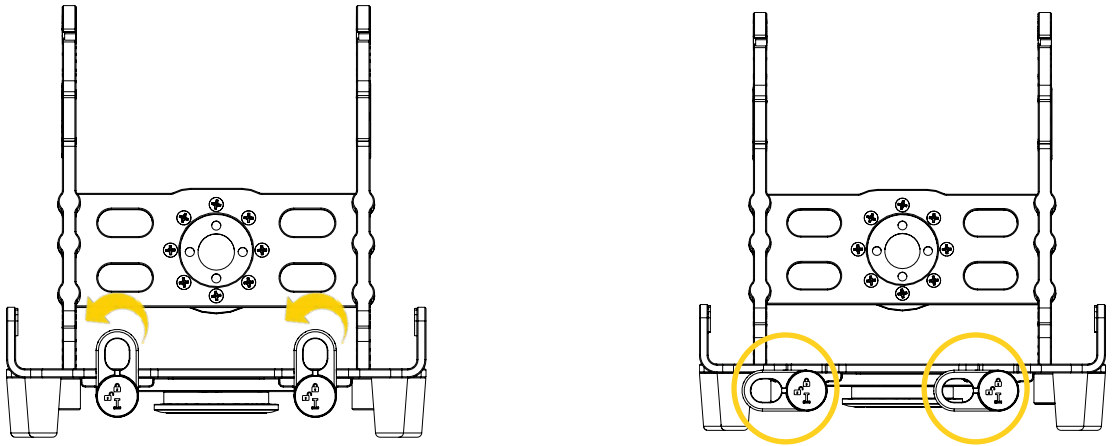


Figure 14: Unlocking the BP60–FL

8. Operate the BP60–FL

The content in this section is intended for personnel who have a competent, proficient or advanced of competency. Refer to the « User Competency » section on page 5 if needed.

NOTE : The illustrations herein are for user-comprehension purposes and may differ from your actual configuration. The instructions apply to any compatible mounting bracket and mounting system. Refer to their user documentation for the safety guidelines and safe use, if needed.

8.1. Install the Hamilton-T1 Transport Ventilator in the BP60–FL

1. Unlock the BP60–FL using the rotary locks. Refer to the « Unlocking the BP60–FL » section on page 23 if needed.
2. Align and insert the medical device in the bracket horizontally until it reaches the back of the bracket (Figure 15A).
3. Lock the bracket using the rotary locks (Figure 15B). Refer to the « Locking the BP60–FL » section on page 23 if needed.

(A) (B)

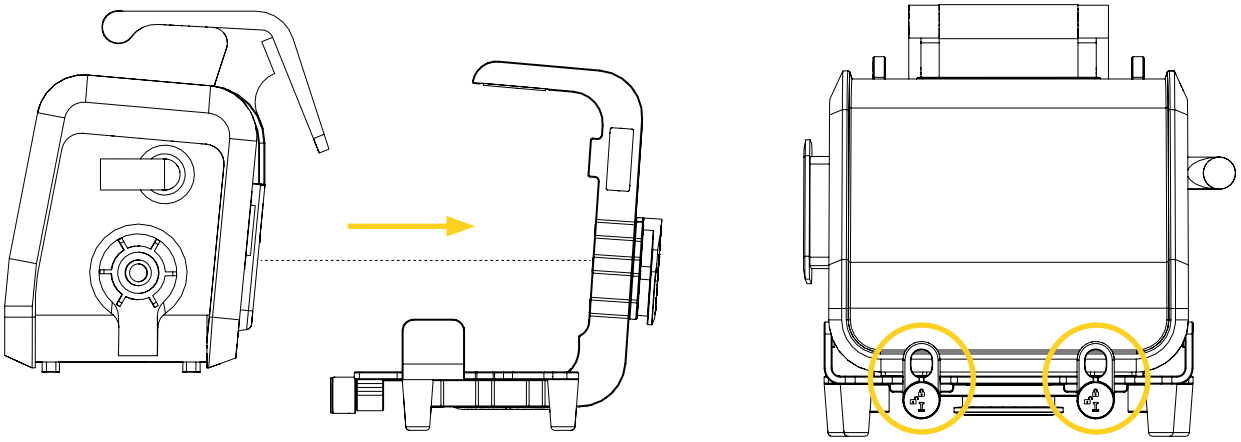


Figure 15: Installing the Hamilton-T1 Transport Ventilator in the BP60–FL

4. Move the medical device back and forth a few times to ensure it is locked and secured in the bracket. If it stays in the bracket after the verification, it is locked and secured.

The installation of the Hamilton-T1 Transport Ventilator in the BP60–FL is complete.

8.2. Remove the Hamilton-T1 Transport Ventilator from the BP60–FL

1. Unlock the BP60–FL using the rotary locks (Figure 16A). Refer to the « Unlocking the BP60–FL » section on page 23 if needed.
2. Pull the medical device out of the mounting bracket horizontally (Figure 16B).

(A) (B)

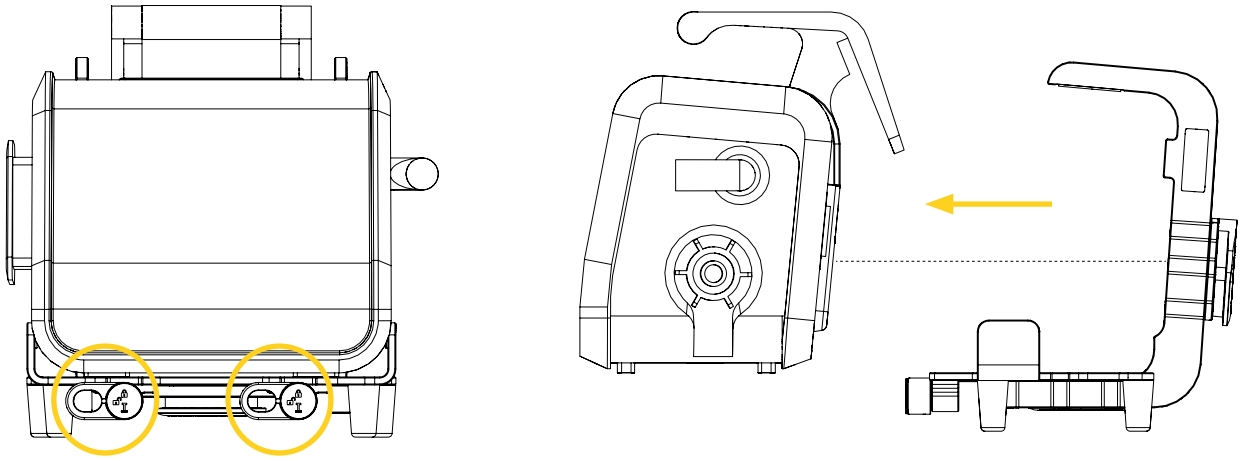


Figure 16: Removing the Hamilton-T1 Transport Ventilator from the Bracket Pro Serie 60 – FL or BP60–FL

3. Set the bracket aside on a clean surface, or place it in its dedicated storage space. Refer to your established internal protocols if needed.

The removal of the Hamilton-T1 Transport Ventilator from the BP60–FL is complete.

8.3. Install the BP60–FL in a Horizontal Mounting System

1. Align and insert the standard bottom disc and/or micro disc located under the BP60–FL in the mounting system horizontally until it locks (Figure 17).

(A) (B)

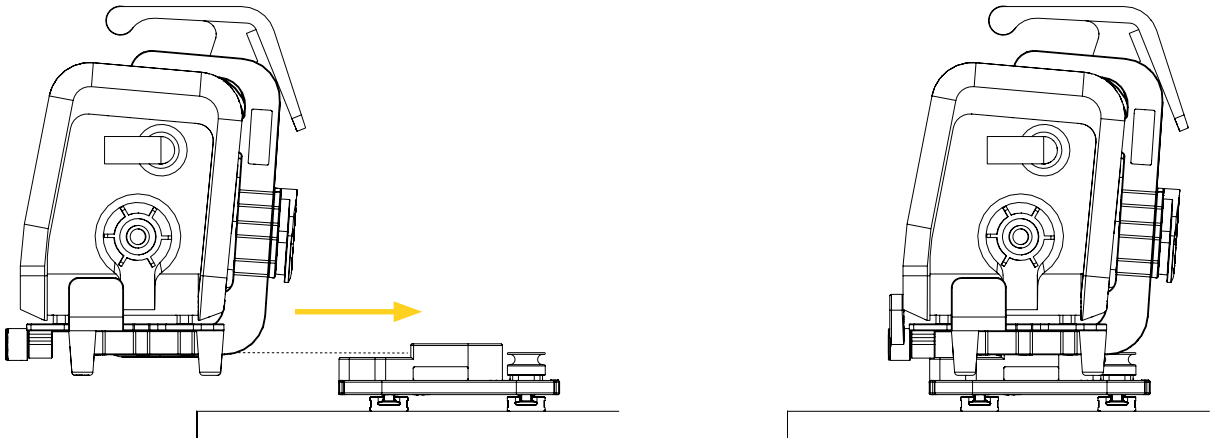


Figure 17: Installing the BP60–FL in a horizontal mounting system

2. Move the mounting bracket back and forth a few times to ensure it is locked and secured in the base. If the disc stays in the base after the verification, it is locked and secured.
3. Turn the bracket up to 360° clockwise or counterclockwise to the desired position (Figure 18) using the medical device's top handle.

(A) (B) (C)

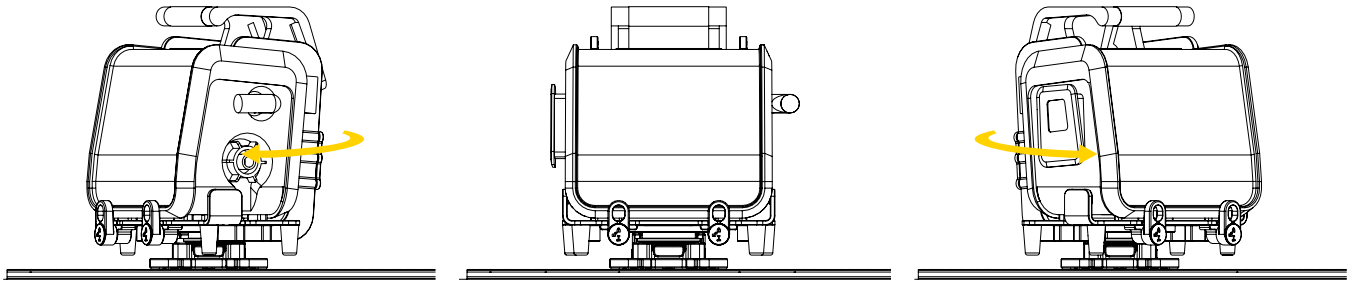


Figure 18: Rotating the Bracket Pro Serie 60 – FL or BP60–FL

The installation of the BP60–FL in a horizontal mounting system is complete.

8.4. Remove the BP60–FL from a Horizontal Mounting System

1. Press and hold the quick-release button of the mounting system (Figure 19A), then slide the standard bottom disc and/or micro disc located under the BP60–FL out of the base horizontally (Figure 19B).

(A) (B)

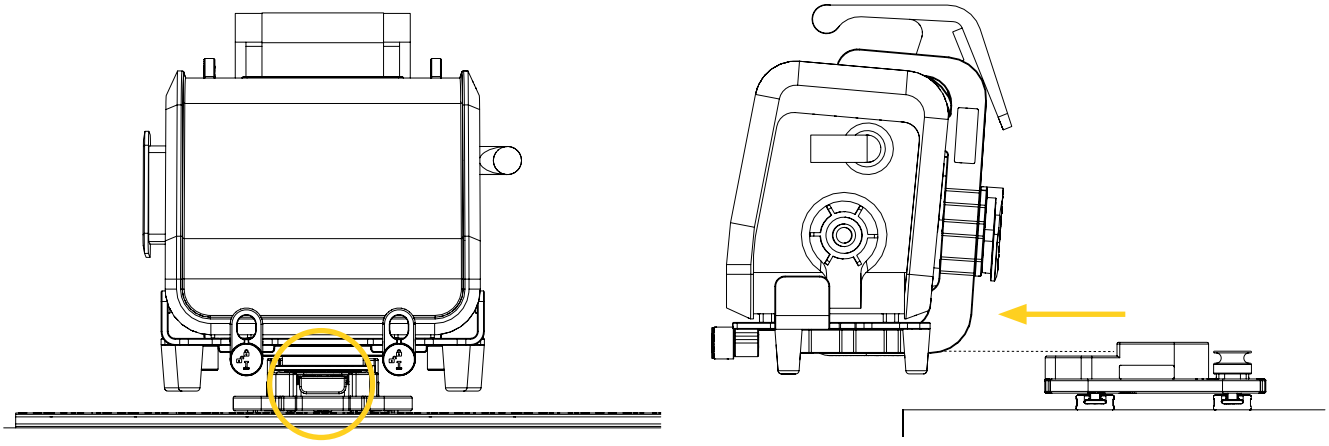


Figure 19: Removing the BP60–FL from a horizontal mounting system

2. Set the mounting system aside on a clean surface, or place it in its dedicated storage space. Refer to your established internal protocols if needed.

The removal of the BP60–FL from a horizontal mounting system is complete.

8.5. Install the BP60–FL in a Vertical Mounting System

1. Align and insert the anti-rotation micro disc located at the back of the BP60–FL in the mounting system vertically until it locks (Figure 20).

(A) (B)

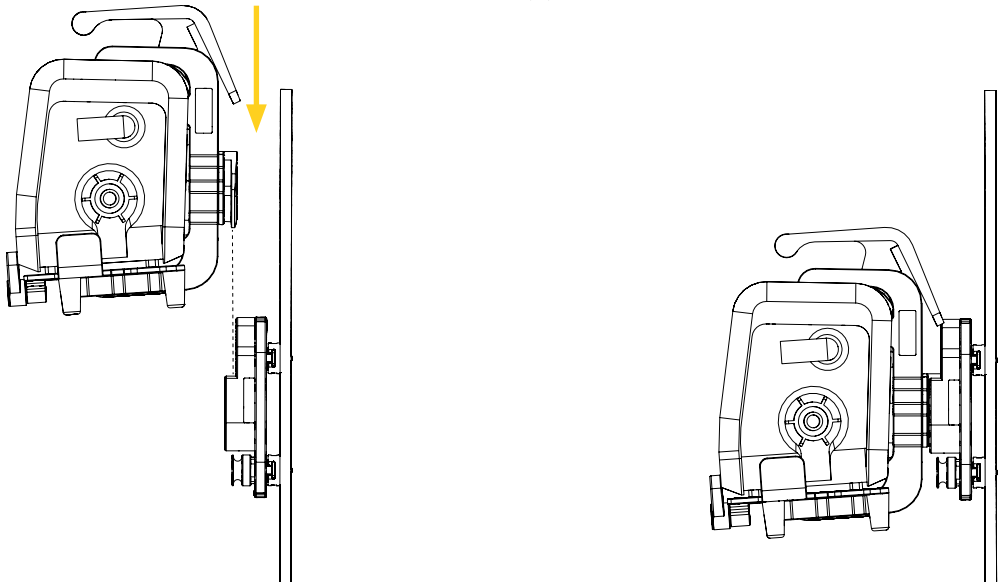


Figure 20: Installing the BP60–FL in a vertical mounting system

2. Move the mounting bracket up and down a few times to ensure it is locked and secured in the base. If the disc stays in the base after the verification, it is locked and secured.

The installation of the BP60–FL in a vertical mounting system is complete.

8.6. Remove the BP60–FL from a Vertical Mounting System

1. Press and hold the quick-release button of the mounting system (Figure 21A), then slide the anti-rotation micro disc located at the back of the BP60–FL out of the base vertically (Figure 21B).

(A) (B)

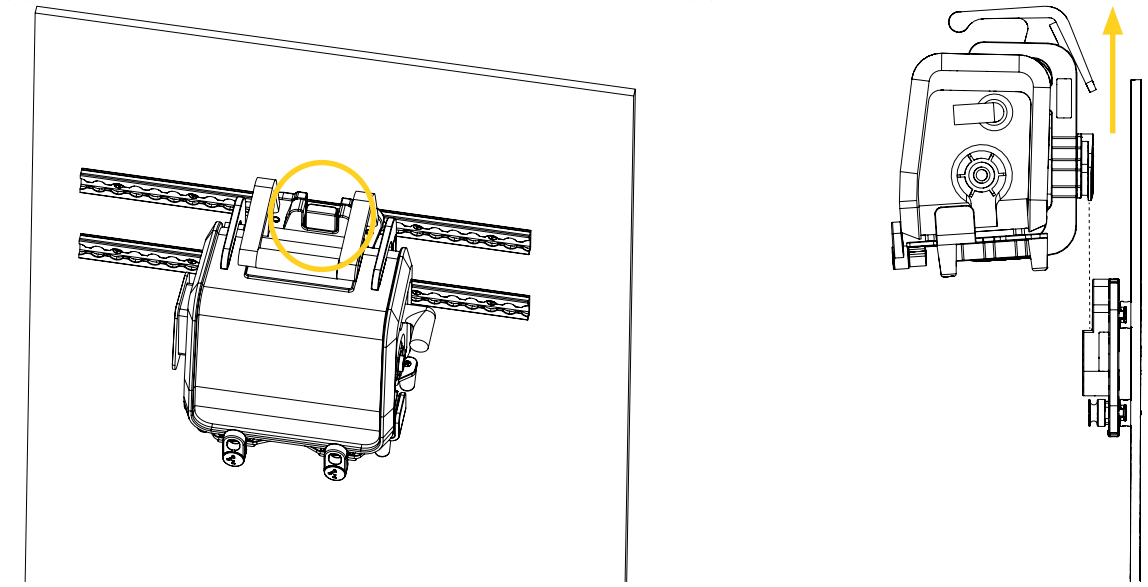


Figure 21: Removing the BP60–FL from a vertical mounting system

2. Set the mounting system aside on a clean surface, or place it in its dedicated storage space. Refer to your established internal protocols if needed.

The removal of the BP60–FL from a vertical mounting system is complete.

Annex I Skills Assessment of the EMS and Clinical Personnel

The content in this section is intended for personnel who have a proficient level of competency. Refer to the « User Competency » section on page 5 if needed.

Following training, a skills assessment should be given to each member of the EMS and clinical personnel to ensure they have fully comprehended the labelling, warnings and cautions, potential risks, safe practices and proper operating procedures needed to safely and effectively use the BP60–FL. Consider adding the following to your internal training protocols.

Trainee name: _____ Unit: _____

Assessor name: _____ Date: _____

SKILLS ASSESSMENT		
SKILL CRITERIA	PASSED	FAILED
Labelling		
- Able to identify meaning and potential risks associated with the different safety labels:		
- Manufacturing label	☐	☐
- Safe Working Load (SWL)	☐	☐
- "Locked" 🔒/"Unlocked" 🔓 identification labels (2X)	☐	☐
Safety Measures		
- Knows that **under no circumstances** should the BP60–FL be installed on, connected to, or used in combination with any third party/competitor mounting system, mounting bracket, medical device or similar equipment.	☐	☐
- Knows that the BP60–FL was specifically engineered to be used with the Hamilton-T1 Transport Ventilator and that using any other medical device may cause the equipment to fall, operate unpredictably, or fail.	☐	☐
- Knows **not to use** the BP60–FL if there are any loose or missing screws.	☐	☐
- Knows to immediately stop using the BP60–FL if any serious incident occurs and to inform authorized personnel.	☐	☐
- Knows to always wait until the transport vehicle is immobilized before manipulating installing/removing the medical device in/from the BP60–FL.	☐	☐
- Knows to always ensure that the medical device is secured on the BP60–FL before it is used.	☐	☐

SKILLS ASSESSMENT		
SKILL CRITERIA	PASSED	FAILED
- Knows to always pay close attention **not to wedge** the power cords or tubing during the installation/removal of the medical device in/from the BP60–FL and/or the BP60–FL on/from the mounting system.	☐	☐
- Knows to always pay close attention to the safety mechanisms of the BP60–FL.	☐	☐
- Knows to always practice safely operating the BP60–FL until the manipulations have been perfected, before use with patients and that improper use of the mounting bracket may cause damage or the equipment to fall or operate unpredictably.	☐	☐
- **Knows not to overload or exceed** the total Safe Working Load (SWL) of the BP60–FL.	☐	☐
- Knows that the Safe Working Load (SWL) of the BP60–FL **does not override** the maximum load capacity of another Technimount product or system, neither those of the medical devices, accessories.	☐	☐
- Knows to always refer to their established internal protocols, as well as read and abide by the safety guidelines and instructions provided in the user documentation of the BP60–FL, the mounting system and the Hamilton-T1 Transport Ventilator.	☐	☐
Operation		
- Able to install/remove the Hamilton-T1 Transport Ventilator in/from the BP60–FL.	☐	☐
- Able to install/remove the BP60–FL in/from the mounting system.	☐	☐
- Able to rotate the Hamilton-T1 Transport Ventilator on a horizontal mounting system.	☐	☐

Annex II Unpack the BP60–FL

The content in this section is intended for personnel who have an advanced level of competency. Refer to the « User Competency » section on page 5 if needed.

1. Inspect the shipping box(es) for signs of damage before accepting shipment. Take pictures and report them promptly if applicable.
2. Move the shipping box(es) to the location of the installation.
3. Open the shipping box(es).
4. Unpack the box(es) and ensure that all shipping and packaging materials have been properly removed, prior to installation.

   NOTE : Keep all packaging material for future use.
5. Identify the item(s) included in the shipping box(es), then set them aside.
6. Inspect the item(s) for signs of damage. Take pictures and report them promptly if applicable.

Annex III Maintenance

The content in this section is intended for personnel who have an advanced level of competency. Refer to the « User Competency » section on page 5 if needed.

Factors such as weather, environment, geographical location and individual usage will necessitate different needs. For the maintenance of the BP60–FL, follow the guidelines listed herein and in accordance with your service's current maintenance practices and established internal protocols. Please contact Technical Support at techsupport@technimount.com for replacement parts or repair related issues, if needed.

WARNING – General Warning

- **Do not perform** the safety checks or the condition-based maintenance before having read the entire « Safety Measures » section on page 12 and the maintenance specific safety measures in this section, having read the entire content of this user manual, having gained in-depth knowledge and product comprehension, as well as having familiarized yourself with the standards and guidelines.
- Safety checks and a condition-based maintenance plan are required and should be established for all Technimount products.
- Perform the safety checks and maintenance operations as described herein. Failing to follow the recommended maintenance plan or its guidelines could cause premature damage to the product.
- Use only Technimount parts, maintenance procedures, cleaning solutions and lubricants (if applicable), as described herein. Using unapproved modified parts or procedures for the maintenance of the Technimount product may cause the system to be unstable and could cause injury to the patients or EMS and clinical personnel and void the product warranty.
- Replace damaged or worn-out parts if past their expected service life or when damaged (refer to the « Replacement Parts/Kits » section on page 39). Recycle damaged parts or dispose according to the environmental laws that apply to your jurisdiction and consult the Safety Data Sheets (SDS).

CAUTION – Safe Handling and Operation

- **Do not use** unauthorized, untested or unapproved cleaning products and disinfectants to perform condition-based maintenance, to avoid damaging the surface of your Technimount product and void the warranty. Technimount will not be held liable for damages resulting from the use of an unauthorized, untested or unapproved cleaning product.
- **Do not use** powered tools to screw the hardware during installation, as there is a risk of damage to the threads.
- **Do not steam clean or use ultrasonic cleaners** on the system or any of its components.
- **Do not immerse** the metal parts/components in water.
- To spot clean, the maximum water temperature should not exceed 82° C (180° F). The maximum water pressure should not exceed 1500 psi/103.5 BAR. If using a high pressure washer, the pressure nozzle must be kept a minimum of 609.6 mm (24 in.) from the product.
- When cleaning, always use the appropriate Personal Protection Equipment (PPE) based on your established internal protocols (e.g., gloves, eyewear, etc.).

CAUTION – Corrosion

- Always rinse and dry the BP60–FL components properly after using cleaning products. Certain types of cleaners may leave a corrosive residue on the surface of the product and could cause the premature corrosion of critical components. Refer to the product Safety Data Sheets (SDS) for chemical information or handling, storage and emergency measures in case of accident.
- Dispose of corrosive wastes according to the environmental laws that apply to your jurisdiction and consult the Safety Data Sheets (SDS).

CAUTION – Follow Instructions for Use

Always read and abide by all the safety guidelines identified, as well as follow all of the instructions provided by the manufacturer of the cleaning product.

Maintenance Frequency

- Safety checks and the condition-based maintenance should be performed minimally every month or as frequently needed, to prolong the longevity of the BP60–FL in optimal conditions.
- Decontaminate the BP60–FL as recommended in your established internal protocols, as well as the regulations and standards in virtue of the infection prevention and control procedures.

Customer-Provided Tools

- Clean and dry cloth
- Soft brush
- Pressure washer
- Cleaning solutions
- Phillips-head screwdriver #2 and #3
- Medium strength thread lock adhesive ( )

Tested and Approved Cleaning Solutions

- Oxivir, 5% Hydrogen Peroxide with Peracetic Acid (AHP)
- Lavo 12, 10 000 ppm Sodium Hypochlorite
- TNT-100, 5% Quaternary Ammonium Compound
- Spectro-Sept, 5% Ethyl Alcohol
- Spectrol, 5% EDTA salt

Maintenance Plan

1. Check the boxes upon completion of the « Safety Checks » section on page 34 and the « Condition-Based Maintenance » section on page 35, then add your comments and/or observations in the « Maintenance Log » section on page 36 if needed. In case of a non-conformity, immediately stop using the BP60–FL and contact Technical Support at techsupport@technimount.com for a remedial action plan.
2. Keep records of your maintenance activities.

SAFETY CHECKS	CHECKED
BP60–FL (Figure 22, Figure 23 and Figure 24)	
- Visually inspect all the components of the BP60–FL to ensure that the hardware is in good condition and there are no loose screws. • *If the hardware is not in good condition, replace it. Contact Technical Support if needed.* • *If the screws are loose, there is a non-conformity.*	☐
- Visually inspect all the components of the BP60–FL to ensure there is no chemical attack. • *If there are traces of chemical attack, clean the mounting bracket. Refer to the « Condition-Based Maintenance » section on page 35 if needed.*	☐
- Visually inspect feet of the BP60–FL to ensure there is wear and/or damage. • *If there are signs of wear and/or damage, replace the feet. Refer to the « Replacement Parts/Kits » section on page 39 if needed.*	☐
- Rotate the rotary locks of the BP60–FL clockwise and counterclockwise a few times to ensure proper functioning of the safety mechanism of the mounting bracket. • *If the rotary locks do not easily rotate and/or you feel resistance when doing so, there is a non-conformity.*	☐
- Install and remove the medical device in/from the BP60–FL a few times to ensure proper functioning of the safety mechanism of the mounting bracket. • *If the medical device does not lock and/or unlock when using the rotary locks of the mounting bracket, replace the rotary lock(s). Refer to the « Replacement Parts/Kits » section on page 39 if needed.*	☐
- Insert and remove the disc of the mounting bracket in/from the base(s) horizontally and/or vertically a few times to ensure it is secured in the mounting system and does not move after hearing the click sound. • *If the mounting bracket moves after correctly having been installed in the base(s), there is a non-conformity.*	☐
- Insert and remove the disc(s) of the BP60–FL in/from the mounting system(s) horizontally and/or vertically a few times to ensure it is secured in the base and does not move after hearing the click sound. • *If the mounting bracket moves after correctly having been installed in the base(s), there is a non-conformity.*	☐

SAFETY CHECKS	CHECKED
- Insert the standard bottom disc/micro disc in the mounting system horizontally and turn it clockwise and counterclockwise a few times to ensure proper functioning of the mounting bracket's mechanism. • *If the mounting bracket does not easily rotate from left to right and/or you feel resistance when doing so, there is a non-conformity.*	☐
- Look at the manufacturing label to ensure that specifications are still visible and that the BP60–FL has not passed its expected service life. Refer to the « Labels » section on page 10 and the « Technical Specifications » section on page 14 if needed. • *If the estimated period of safe and reliable use has passed, there is a non-conformity.*	☐
- Look at the safety labels and identification labels to ensure that they are still visible and that they are all accounted for. Refer to the « Labels » section on page 10 and the « Technical Specifications » section on page 14 if needed. • *If the labels are no longer visible and/or missing, there is a non-conformity.*	☐

CONDITION-BASED MAINTENANCE	CHECKED
Clean the BP60–FL, after you have completed the safety checks: 1. Remove the excess dirt using a clean cloth, if needed. 2. Remove the contaminants using a pressure washer or as recommended in your established internal protocols and control procedures. 3. Clean using a cloth and cleaning solution. 4. Spot clean stains by applying the solution directly on the stain and let sit on the surface, if needed. **NOTE :** Avoid over saturation and ensure that the product **does not sit** on the surface of the BP60–FL components longer than recommended by the cleaner's manufacturer. 5. Thoroughly rinse the solution with a clean cloth dampened with lukewarm water, then dry all the components using a clean cloth before returning to service.	☐
Clean the mounting system, after you have completed the safety checks of the BP60–FL. Refer to the mounting system' user documentation if needed.	☐

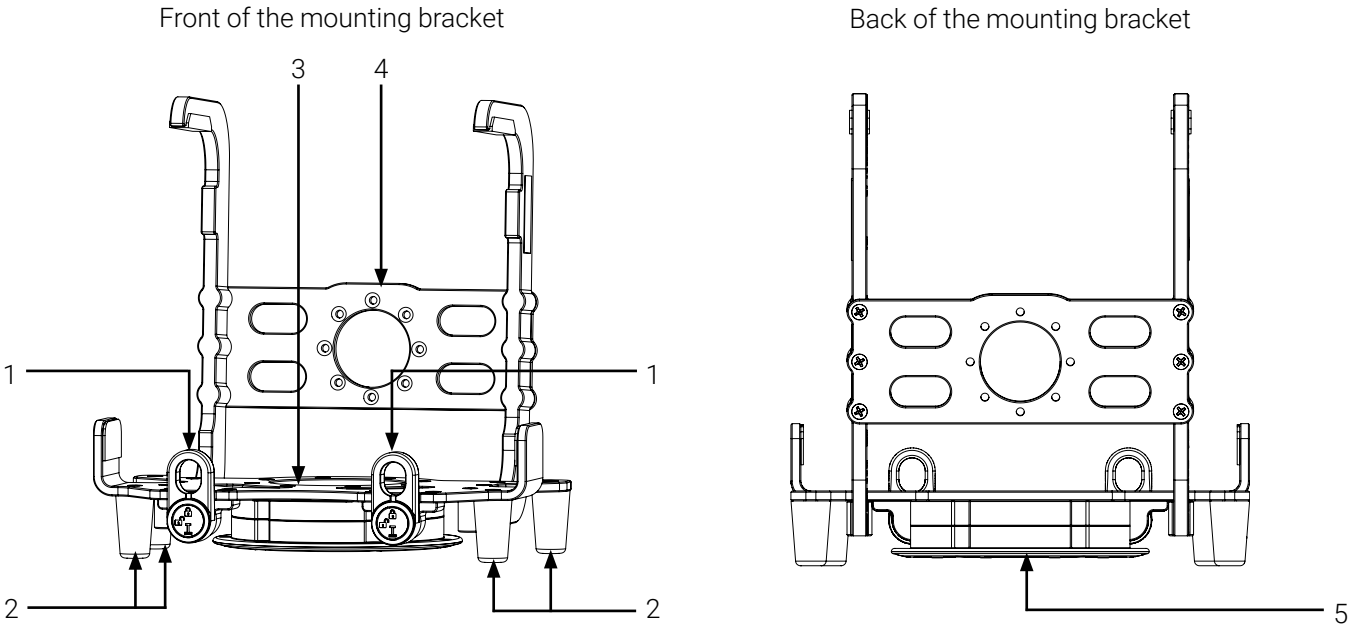
MAINTENANCE LOG

Add your comments and/or observations, after you have completed the safety checks and condition-based maintenance, if needed:

Maintenance plan completed on (dd/mm/yyyy): _____

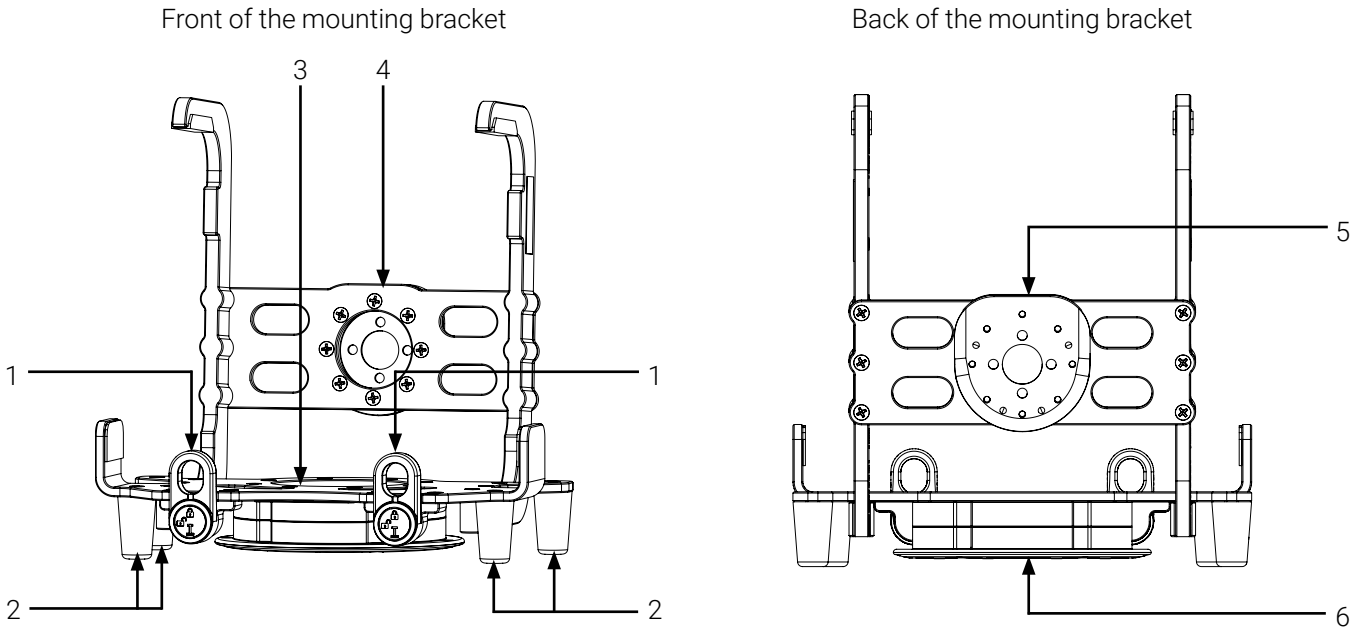
Maintenance plan completed by: _____

Illustrated Inspection Points



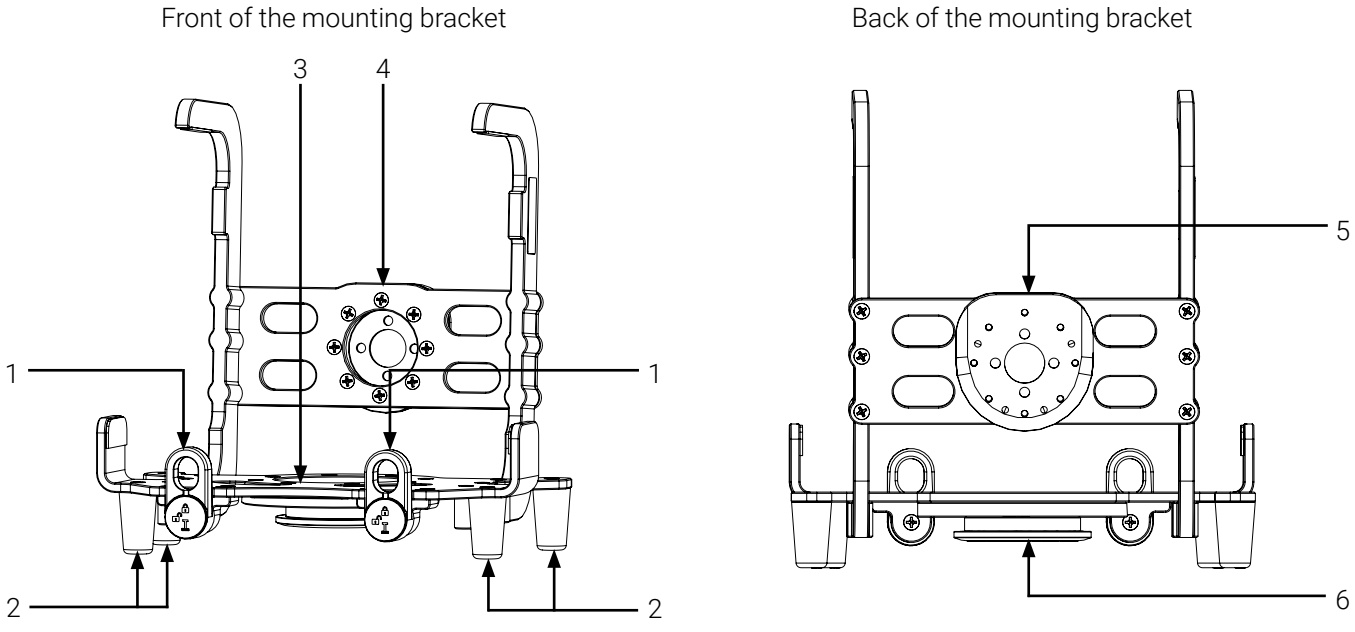
1. Rotary lock (2X)
2. Foot (4X)
3. Bottom plate
4. Back plate
5. Standard bottom disc

Figure 22: Illustrated inspection points – standard bottom disc



1. Rotary lock (2X)	4. Back plate
2. Foot (4X)	5. Anti-rotation micro disc
3. Bottom plate	6. Standard bottom disc

Figure 23: Illustrated inspection points – standard bottom disc and anti-rotation micro disc



1. Rotary lock (2X)	4. Back plate
2. Foot (4X)	5. Anti-rotation micro disc
3. Bottom plate	6. Micro disc

Figure 24: Illustrated inspection points – micro disc and anti-rotation micro disc

Annex IV Replacement Parts/Kits

The content in this section is intended for personnel who have a proficient or advanced level of competency. Refer to the « User Competency » section on page 5 if needed.

Technimount reserves the right to change part numbers and products without notice. Please contact Customer Service at customerservice@technimount.com to ensure product options and availability, or Technical Support at techsupport@technimount.com for replacement parts/kits or repair related issues.

REFERENCE #	PART/KIT #	PART/KIT DESCRIPTION	REPLACEMENT PROCEDURE
Figure 22, #1 Figure 23, #1 Figure 24, #1	9001-10-UN	Rotary Lock kit	Refer to the IN-RPRC-RL-202607EN-01 replacement procedure
Figure 22, #2 Figure 23, #2 Figure 24, #2	923-00-1625-INS	1-1⁄8 in. acetal foot	Refer to the IN-RPRC-FEET-202605EN-04 replacement procedure

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