Instructions For Use RHINO KNEE IMMOBILIZER

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DESCRIPTION

The Rhino Knee Immobilizer is a medical device intended for use with pediatric patients receiving post operative or non-operative treatment for medical conditions that impact the knee, such as ligament deficiencies or repair/reconstruction of the extensor mechanism and as instructed by a healthcare provider. A consistent examination scheduled with the clinical provider is needed to ensure proper fit, use, and instruction for caregivers and/or users. Compliance with instructions for use and clinical care treatment plan is important to achieve clinical and therapeutic outcomes.

MATERIALS

The Rhino Knee Immobilizer is made from threaded navy blue, blue foam, double fold navy bias, lacteal polymer, stabilizers, carbon black, formaldehyde, polypropylene, aluminum, and nylon.

INDICATIONS FOR USE

The Rhino Knee Immobilizer is intended for use with pediatric patients receiving post operative or non-operative treatment for medical conditions that impact the knee, such as ligament deficiencies or repair/reconstruction of the extensor mechanism and as instructed by a healthcare provider.

WARNINGS AND PRECAUTIONS

- Follow the qualified healthcare provider and physician instructions or treatment plan.
- Follow instructions to ensure straps are tightened properly, to reduce risk of irritation, blisters, and pressure
 ulcers.
- Consult your healthcare provider immediately if the child is experiencing pain, irritation, blisters, swelling, pressure ulcers, or if the brace is not fitting properly.
- · Protective barriers (i.e. clothing) should be worn when wearing the Rhino Knee Immobilizer.
- Do not apply the brace directly to the skin. Use protective barrier/clothes between stirrups and skin (e.g., long stretchy pants, lightweight pajamas, etc.).
- The Rhino Knee Immobilizer should be used without additional components or accessories, unless instructed by the healthcare provider.
- Do not use a Rhino Knee Immobilizer that is damaged, broken or not functioning properly.
- Modification is not recommended for the Rhino Knee Immobilizer.

MRI SAFETY INFORMATION

The Rhino Knee Immobilizer brace is MR safe.

CLEANING INSTRUCTIONS

Wash on delicate cycle or hand wash with mild detergent. Rinse thoroughly.

STORAGE AND HANDLING

- The Rhino Knee Immobilizer requires no special handling during transport or storage.
- The Rhino Knee Immobilizer is a nonsterile device.

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APPLYING THE BRACE



- 1. Carefully place the immobilizer behind the injured leg so that it flaps can wrap around the front of the leg.
- 2. Fold the flaps over the front of the leg.



- Thread the straps through the loops above and below the knee.
 - Note: There should be no straps across the knee.
- 4. Once the strap is through the loop fold it back and Velcro the strap onto itself.

REUSE LIFE

All parts were designed to be used by a single patient under normal use conditions and as instructed by a healthcare provider. Normal use includes the daily application of the Rhino Knee Immobilizer as healthcare provider instructed.

SAFE DISPOSAL

The Rhino Knee Immobilizer can be disposed of in the regular trash. None of the components of the device are recyclable.

IMPORTANT STATEMENT

- MD Orthopaedics, Inc. does not provide medical treatment, advice or recommendations about the risks and benefits of medical treatment, including treatment that involves the use of MD Orthopaedics, Inc. products. This information should be provided solely by the physician or other qualified health care provider treating your child. If you have questions about your child's treatment, it is important for you to discuss those questions with the appropriate health care provider.
- The manufacturer and distributor are not liable for cases of material damage or personal injury caused by
 incorrect handling or non-compliance with instructions. Normal use is defined as a single user following intended
 use
- Instructions for Use (IFU) are available in English at https://opsb.com/ifu. To obtain a copy of the paper IFU, please call Customer Service at MD Orthopaedics, Inc. at (877) 766-7384.
- If a user and/or patient experiences any serious incident that has occurred in relation to the device, it should be reported to MD Orthopaedics, Inc. and the competent authority of the EU and EEA Member State in which the user and/or patient is established.

SYMBOL LEGEND

Symbol	Meaning	Symbol	Meaning
	Manufacturer	3	Date of manufacture
REF	Catalog number	LOT	Batch code
(1m)	Single Patient, Multiple Use	NON STERILE	Non-sterile
i	Consult Instructions for Use or Consult Electronic Instructions for Use	MR	MR Safe