

RB - Resistance Band

EN BODY/LIMB ELASTIC EXERCISER

DE GANZKÖRPER-BEWEGUNGSTHERAPIEGERÄT,
ELASTISCH

FR APPAREIL D'EXERCICE ÉLASTIQUE
POUR LE CORPS

ES EJERCITADOR ELÁSTICO PARA
CUERPO/EXTREMIDADES

IT ATTREZZO ELASTICO PER ESERCIZI DEL CORPO

NL ELASTISCH LICHAAMSOEFENTOESTEL

DK ELASTISK KROPSTRÆNINGSSAPPARAT

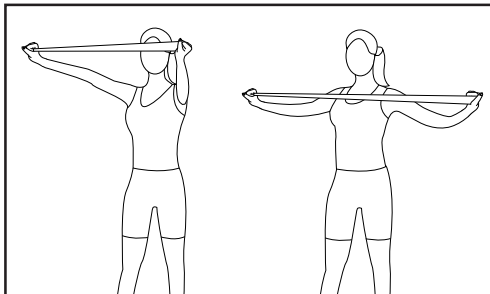
FI KEHON ELASTINEN HARJOITUSVÄLINE

SE KROPPSTRÄNINGSMOTSTÅND, ELASTISKT
MOTSTÅND

NO TRENINGSMOTSTYR, SMIDIGHET

PL PRZYRZĄD DO ĆWICZEŃ CIAŁA WYKORZYSTUJĄCY
OPÓR ELASTYCZNY

GR ΕΛΑΣΤΙΚΗ ΔΙΑΤΑΞΗ ΕΚΓΥΜΝΑΣΗΣ ΣΩΜΑΤΟΣ



ENGLISH

Product Description

Elastic resistance bands made from non-latex material, used to support rehabilitation by gently strengthening muscles, improving mobility, and aiding recovery.

Usage Information

Follow exercise plans provided by your healthcare professional.

Indications for Use

- Injury rehabilitation
- Managing joint pain
- Improving flexibility

Contraindications

- Do not use over open wounds.
- Any known allergy and/or hypersensitivity to any material listed in product composition.

Warnings and Precautions

- We recommend you have been advised to use this device by a suitably qualified healthcare professional.
- Carefully read all instructions and warnings prior to use.
- Follow all instructions to ensure proper performance of the device.
- Do not use if liniments, ointments, gels, creams or any other substances have been applied to the affected area.
- Perform regular skin and circulation checks, especially patients with diabetes, vascular deficiencies and neurological conditions.
- Should any adverse reactions occur, please cease use and contact your healthcare professional or provider of this product.

Washing and Care Instructions

Wipe clean with a damp cloth.

Storage and Transport Conditions

Store in a cool, dry place out of direct sunlight, in the original packaging.

Recycling and Disposal

Packaging and constituent parts should be recycled or safely disposed in accordance with local or national laws.

Serious Incident


Report any serious incident to the manufacturer and the competent authority of the EU Member state, or country in which you reside.

 **MEDICAL DEVICE**

 **SINGLE PATIENT - MULTIPLE USE**

 **CAUTION**



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