

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

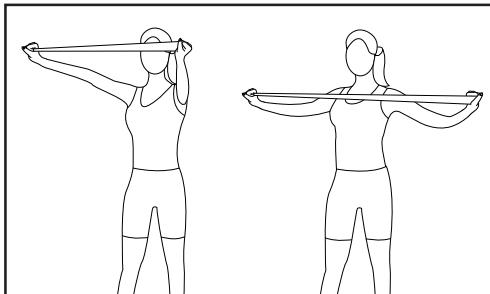
FI KEHON ELASTINEN HARJOITUSVÄLINE

SE KROPPSTRÄNINGSMOTSTÅND, ELASTISKT
MOTSTÅND

NO TRENINGSSUTSTYR, 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

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.
- Improper use of resistance bands can cause serious injury if not used correctly. Please take extra care with exercises that involve stretching the resistance bands near the head to avoid injury to the face or eyes of yourself or others.
- Consider wearing protective eyewear such as safety goggles or protective glasses to reduce the risk of eye injury.
- If you are anchoring the band to an object, ensure the band is securely attached to a stable object before use.
- This product is not a toy and should only be used for its intended purpose.
- 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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