HD EEZI-TRANSFER™ – OVAL BOARD – LARGE

This guide covers product HDOVLLRG (supplied as a pair).

The **Eezi-transfer™ Oval Boards** are strong, flexible and very easy to use. They are most commonly used to assist with sling fitting for patients in chairs but also for patients lying flat in bed.

To insert a sling you will need two **HD Eezi-Transfer™ Oval Boards – Large**.

USER INSTRUCTIONS

+1 +2 4-1 FIM ASSESSED

1

Slide the first board horizontally behind the sitting or lying patient. (if they can, ask them to lean forwards slightly).

Once the board is central to the person, rotate it to a vertical position, so that it mirrors the person's spine.

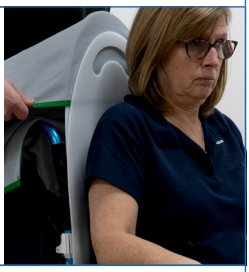
Insert the second board BEHIND the first board using the method outlined above.

Insert a sling BETWEEN the two boards so it is not in direct contact with the patient's clothing or skin.

Make sure the centre line of the sling matches the person's spine.

Slide the sling down between the two boards and pull through down to the buttocks.

2



3

Gently remove both boards one at a time,

Continue to pull the leg straps of the sling through and under the person's legs, or....

If you have the **HD Eezi-Transfer™ Oval Boards – Small** too, use them to help you pull the leg straps of the sling through and under the person's legs.

Just slide one gently under each leg and then pull the leg straps of the sling through and under each small board.

Remove the **HD Eezi-Transfer™ Oval Boards – Small** gently once sling strap is pulled through.

4

PRODUCT FEATURES

- Supplied as a pair but an individual board can be purchased if needed as a replacement.
- All edges tapered for user comfort and ease of insertion and removal.
- Comfortable carry handle.

MATERIAL CONTENT

100% HDPE

STORAGE AND HANDLING

- Store in a clean dry environment that is not subject to excessive heat.
- Keep away from sunlight.
- Store flat and avoid placing heavy objects on top of the product.

WARNINGS AND PRECAUTIONS

- Do not leave on the floor when not being used.
- Inspect before use, including the non-slip panels, for signs of wear or damage.

CLEANING INSTRUCTIONS

- Wipeclean only.
- Wipeclean with natural water and detergent or wipes that don't contain alcohol, solvents, bleaching or abrasive agents.
- Cleaning materials used should be patient safe and biodegradable.

DISPOSAL INSTRUCTIONS

- Easily recycled.
- If it has been used by an infectious person, thoroughly disinfect before recycling.
- Otherwise dispose of through normal recycling waste management.


Keep dry


Keep away from sunlight


Caution


Do not use if package is damaged

Storage Temperature Limit


Patient Information website

Medical Device

MANUFACTURER
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Quality UK Manufacturing Since 1997

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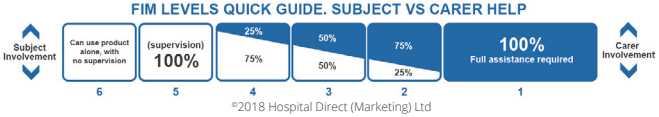
HOSPITAL DIRECT (HD) ADDITIONAL SYMBOLS

At HD we understand that not everyone is familiar with all equipment, so to help therapists and healthcare professionals in assessing a product's suitability for a client, we have included two assessment tools, Functional Independence Measure (FIM) and Easy Guide Symbols. These will enable the healthcare professional to assess their client and decide whether or not a product will help the person based on their level of dependence and functional ability.

FUNCTIONAL INDEPENDENCE MEASURE (FIM) ASSESSMENT

Our FIM assessment guide for each product will help you decide the suitability of this product for the person's ability and need. Based on the standard criteria from Level 6 where the person can use the product unaided and unsupervised to Level 1 where all the assistance is provided by the carer and the client can do nothing, this guide is easy, quick and simple to use to check suitability against ability and circumstance.



EASY GUIDE SYMBOLS

We use four symbols to indicate where a product is suitable and safe for the person to use unassisted, with a single carer or with multiple carers. These symbols indicate the minimum recommendation.

HD has a policy of continuous development and, as such, reserves the right to alter specifications (including measurements, materials and colours) without prior notice. If any serious incident has occurred in relation to the device it should be reported to the manufacturer and the competent authority in which the user and/or patient is established.