











HANDY STRAP – WASHA

This guide covers all products with the prefix HS in the product code.

The Handy Strap is an ideal back support helping to sit forward in bed or in a chair. It is perfect for people who can't sit forward without assistance from carers.

USER INSTRUCTIONS









Open out the **Handy Strap**, as shown, so you can see the anti-slip panel. Now close it so that the red handles are on the top.

With the red handles on the top, wrap the Handy Strap around your hand and slide it under the patient's pillow towards your colleague on the other side of the bed.







Make sure that it is lying flat and then working together, hold the red straps with your inner hands and unpeel the top blue layer towards the headboard to expose the anti-slip panel. This will help to grip the persons clothing for extra security.



Now turn to face the head of the bed. Holding the red handles with your outer hand and the green or yellow handles (to suit) with your inner hand adopt a safe posture and after co-ordinated commands, help the patient to sit up/sit forward.



UNDERSTANDING THE HANDY STRAP



UNDERSIDE YELLOW HANDLES

BACK SLIPPY LAYER



ANTI-SLIP PANEL TO HELP WITH GRIP. STANDING AND SUPPORT LEGS ETC

MATERIAL CONTENT

Outer: 100% Polyester Handles: Polypropylene

Anti-slip Panel: PVC coated polyester, phthalate free

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 or drop on the floor; it is a slip hazard.
- Test for slippiness before use. If over-laundered and doesn't slide please discard or replace.
- Inspect product (including handles) before use for signs of damage or tears.

CLEANING INSTRUCTIONS

- · Wash as per laundry instructions.
- Can be wiped clean, between use with the same patient, 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

- Non-recyclable
- Dispose of as clinical waste if it has been used by an infectious person.
- · Otherwise dispose of through normal waste management.





Keep away from sunlight







LOW









Wash at 30-74°C. Tumble dry on low heat. Do not iron. Do not use bleach. Do not use fabric conditioner



wehsite









MANUFACTURER

Hospital Direct (Marketing) Ltd, Units 2, 3 & 4, The Green, Clun, Shropshire SY7 8LG, United Kingdom

EU REPRESENTATIVE: Obelis S.A., Bd Général Wahis 53, 1030 Brussels, Belgium



www.hospitaldirect.co.uk

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.

FIM LEVELS QUICK GUIDE. SUBJECT VS CARER HELP











Four Carers

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.