



EXPERTS IN PATIENT HANDLING SOLUTIONS



# EASIDRESS – WASHABLE

This guide covers all products with the prefix DP01, DP02, DP03 and DP04.

The **EasiDress – Washable** sleeves make getting dressed easier, both for independent users and those who need support from a carer.

## USER INSTRUCTIONS



1

Slide the **EasiDress** arm sleeve gently over the patient's arm and up towards the shoulder.

Next, slide the pyjama top, blouse or jumper sleeve over the **EasiDress** arm sleeve, as far up as you can..

Once you have the garment in place, remove the **EasiDress** arm sleeve carefully.  
Repeat with the second arm and then remove the remaining arm sleeve.

2



3

Place the **EasiDress** leg sleeve(s) over the feet and pull up as far as you can. Now, gently slide pyjamas, trousers, jeans, or shorts over the **EasiDress** leg sleeve(s).



### TO REMOVE

From the **EasiDress** foot end, gently pull and remove, the leg sleeve, and repeat for the remaining leg.

4



## PACK INCLUDES

- 1 x EasiDress – Washable Leg Sleeve
- 1 x EasiDress – Washable Arm Sleeve

## MATERIAL CONTENT

100% Polyester

## 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

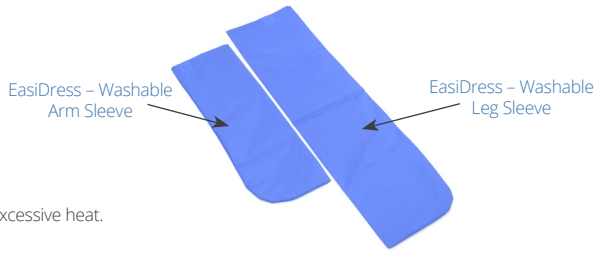
- Do not leave or drop on the floor; it is a slip hazard.
- Test for slippiness before use. If over-laundered and doesn't slide please discard or replace.
- Inspect product before use for signs of damage or tears.

## 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 dry



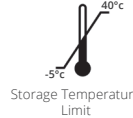
Keep away from sunlight



Caution



Do not use if package is damaged



Storage Temperature Limit



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



Patient Information website



Medical Device



## 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, B-1030 Brussels, Belgium



[www.hospitaldirect.co.uk](http://www.hospitaldirect.co.uk)

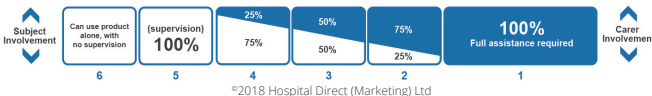
## HOSPITAL DIRECT (HD) ADDITIONAL SYMBOLS

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.

#### FIM LEVELS QUICK GUIDE. SUBJECT VS CARER HELP



### 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.