GLIDE AND LOCK – DISPOSABLE / SINGLE PATIENT USE

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

The **Glide and Lock – Disposable** for those who struggle to sit up in bed without sliding down or have trouble sitting upright in a chair/wheelchair or bed

USER INSTRUCTIONS

+1 +2 6-3 FIM ASSESSED

UNDERSTANDING THE GLIDE AND LOCK – DISPOSABLE

ANTI-SLIP PANEL TO STOP IT SLIPPING AGAINST THE CHAIR SURFACE

INTERNAL LOCKING STRIPS ENSURE SMOOTH GLIDE IN ONE DIRECTION ONLY AND LOCK IN THE OTHER

1

Open up your **Glide and Lock – Disposable** and hold it with your hand through the loop, ensuring the anti-slip panel is at the back and at the top.

The directional label will be on the left hand side of the product as you look at it.

Place the **Glide and Lock – Disposable** on the chair prior to seating.

Check that the direction of the locking mechanism is correct (see directional arrows on the label).

The normal positioning is for the anti-slip panel to be face down against the chair surface.

If tested, the product will slide away from you.

2

3

Ask the patient/client to sit on the **Glide and Lock – Disposable**, making sure that their bottom does not overshoot the product.

By using their hands and feet to push back and their nose over their toes, the user will be able to glide to the back of the chair and rest in a more comfortable and seated position. The locking strips inside the **Glide and Lock – Disposable** will now lock them into place and prevent them from slipping forwards.

4

MATERIAL CONTENT

Top: 100% Cotton
Filling: 100% Polyester
Back: 100% Polyester
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.
- Inspect product, before use for signs of damage or tears.

CLEANING INSTRUCTIONS

- Wipeclean, if needed for 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.

Single Patient Use

Keep dry

Keep away from sunlight

Caution

Do not use if package is damaged

40°c
-5°c
Storage Temperature Limit

LATEX

Patient Information website

MD
Medical Device

CE

EC REP

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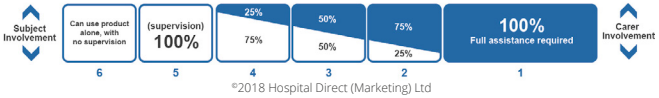
HOSPITAL DIRECT (HD) ADDITIONAL SYMBOLS

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.