OPTHALMIC GLIDE AND LOCK

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

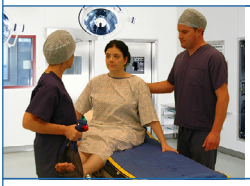
Developed with an opthalmic surgeon to help position patients for eye surgery, the **Ophthalmic Glide and Lock** prevents slippage during eye surgery and allows patients to sit up safely post surgery without the risk of sliding down.

USER INSTRUCTIONS

+1 +2 6-3 FIM ASSESSED

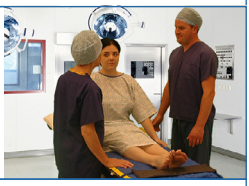
1 Place the **Ophthalmic Glide and Lock** on the day surgery trolley/ operating table with the glide arrow indicating towards the head end and on the LEFTHAND SIDE. Attach the disposable cover.

Ask the patient to sit on the edge of the surgery trolley/operating table as shown, so that when they turn and lie down their head will be on the pillow. 2



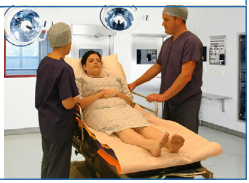
3 For those with difficulties raising legs, a **Leg Lifter** can be used to support legs whilst turning. A small slide sheet or **Rota Cushion** can also be useful here to rotate the hips.

Once the patient is safely in position on the surgery trolley/operating table, ask them or help them to lie down. 4



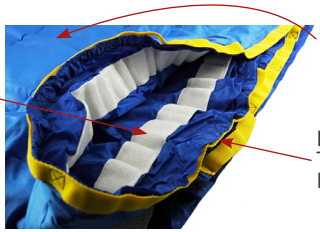
5 The **Ophthalmic Glide and Lock** can now be slid further up the surgery trolley/operating table until the patient's head is exactly where the surgeon requires it. Once in position the patient will not slip during surgery.

Post surgery, the patient can sit up in Recovery and, due to the internal locking strips, will not slip down. 6



MATERIAL CONTENT

Top: 100% Nylon
Filling: 100% Polyester
Back: 100% Polyester
Handles: 100% Polypropylene



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.

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

MD
Medical Device

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

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

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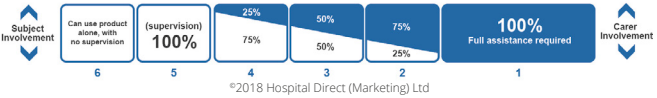
HOSPITAL DIRECT (HD) ADDITIONAL SYMBOLS

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.