GLOVE AIRFLOW MANUAL SLINGS

Instructions For Use

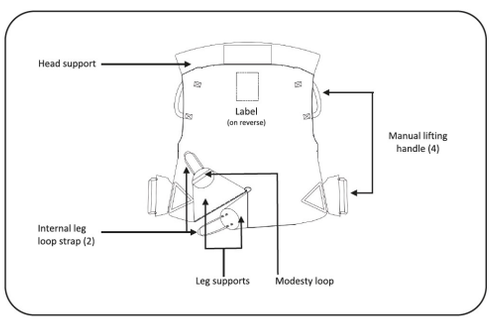
Please read these instructions befor

Care Independence
A GBUK COMPANY

INSTRUCTIONS FOR USE
Please read these instructions before use.

PRODUCT CODES & DESCRIPTIONS
All Glove Slings. Product codes prefixed with:
- GAIRM

DEVICE DESCRIPTION
The Glove Airflow Manual Sling are non-sterile medical devices, complaint with BS EN 10535. Each sling has a sewn woven label containing device Serial Number and Safe Working Load (kg), and a printed label containing all pertinent information.



For further information, support and product training requests visit: www.gbukgroup.com

PATIENT INFORMATON
The following information is provided for use by trained healthcare professionals, and carers, and those with the appropriate training or knowledge, however as the learned intermediary between the company and the patient, the trained healthcare professionals and carers, and those with the appropriate training or knowledge must convey the aspects they consider relevant to the individual patient. The patient must be informed of the potential adverse effects & residual risks contained within this document.

PATIENT TARGET GROUP(S)
Intended for patients who require lifting and/or repositioning following a suitable risk assessment.

INTENDED USE
Intended to be used to transfer patients following a risk assessment. The sling should be used in conjunction with a mechanical lifting device (either mobile or ceiling hoist) to ensure safe lifting and repositioning of a patient.

INTENDED USER(S)
The device is intended to be used by trained healthcare professionals and carers, and those with the appropriate training
or knowledge. Refer to the Royal College of Nursing Guidelines.

INDICATION
The sling is indicated for use on patients with limited mobility requiring lifting and/or repositioning.

SAFE WORKING LOAD (SWL)
The Glove Airflow Manual Sling has a SWL of 200 kg. The SWL is also provided on the label(s) directly attached to the device. The SWL of a sling labelled as patient matched is detailed on the label(s) directly attached to the device. Refer to the device label(s) for applicable SWL of the patient matched sling.

CONTRAINDICATIONS
Do not use on patients whose weight exceeds the safe working load limit of the sling.

WARNINGS & PRECAUTIONS
- A risk assessment must be carried out by a suitably qualified individual before any sling is selected and used, to ensure safety for the patient and handler. See 'RISK ASSESSMENT' section.
- Always carry out a visual check of the sling prior to every use for any damage, rips, tears, wearing, loose stitching, material degradation, or any other general signs of wear and tear. The sling should be disposed of in accordance with the disposal instructions, if any of these are present, and/or in the instance of failed LOLER inspection, and a new sling selected.
- Do not use with hoists.
- Follow local manual handling policy from your organization, National Back Exchange or HSE guidelines.
- All users must be trained or have the appropriate training or knowledge before using the Sling.
- Inspect the device, packaging and labelling prior to use and do not use if damaged.
- Do not exceed SWL of the sling.
- The Sling is subject to Lifting Operations and Lifting Equipment Regulations (LOLER) Examinations. A LOLER examination must be performed every 6 months, by trained and qualified persons.
- Wash the sling in accordance with the instructions provided in the 'CLEANING' section.
- Do not bleach, iron or dry clean the sling. Do not use stain removers.
- Do not exceed maximum washing and drying temperatures.
- Do not wash below minimum washing temperature.
- Do not place the sling on hot pipes or heaters. Keep away from flames.
- Where the sling has hooks and loops, ensure any hooks and loops are closed prior to laundering.
- Repeated washing may lead to sling deterioration and/or discolouration to some materials or fading of labels.
- All devices labelled as a Patient matched device must not be used with any patient other than the single patient who has been risk assessed and fitted with the specific patient matched device. Patient matched devices are single patient use only. Devices not labelled as patient matched are reusable.

EXPECTED LIFETIME
The expected performance and safety lifetime of the device is 5 years, however slings may last longer. All slings must be LOLER inspected and the inspection must be within date prior to use. Carry out a risk assessment prior to each use of the sling - Refer
to 'WARNINGS & PRECAUTIONS' & 'RISK ASSESSMENT'.

PERFORMANCE CHARACTERISTICS
- SWL as stated on label(s)
- ISO 21856 and ISO 10535 compliant

DIRECTIONS FOR USE
Refer to 'Warnings & Precautions' and 'Risk Assessment' Sections prior to use of the Sling.
1. With the user on their side, fold sling in half lengthways with the label on the inside.
2. Position the sling with the top of the aperture aligned with the user's coccyx and the centre of the sling in alignment with the user's spine.
3. Fold the uppermost section over the user like a blanket.
4. Tuck the remaining sling material under the user.
5. Roll the user onto their opposite side.
6. Unfurl the sling beneath the user.
7. Reposition user onto their back with sling now spread out fully underneath them.
8. Check the alignment of the sling and reposition if required.
9. Feed the leg supports carefully between the users thighs.
10. Feed the internal leg strap through the modesty loop.
11. Feed the manual lifting handles through the left and right internal leg loop straps.
12. Use the four handles to manually lift the user. Depending on the risk assessment, this may be performed by either two or four carers.

POTENTIAL ADVERSE EFFECTS & RESIDUAL RISKS
As with any procedure involving manual handling, use of manual handling aids, and hoisting equipment the typical risks associated apply. Failure of the device during use may expose the patient and user to risks such as falling, and potential subsequent associated injury, and/or user exposure to sudden mechanical stress and potential associated pain and/or musculoskeletal injuries.

RISK ASSESSMENT
A risk assessment and visual inspection must be performed by competent person(s) before using the sling. This should consider the following factors:
- The physical and medical status of the patient, including skin integrity,
- The status of the sling, including the presence of any damage, rips, tears, soil, loose stitching, or any other general signs of wear and tear,
- The compatibility of the hoist and sling, including if the correct size and type of the sling is being used
- The capability of the users, using the equipment; and,
- The situation in which it is being used

WARRANTY
Glove Slings and Patient Matched Glove Slings are covered by a lifetime warranty. Any manufacturing defect will be rectified free of charge. Product damage through wear and tear, neglect or accidental damage is not covered.

CLEANING
Routinely clean the sling, in accordance with these instructions and your facility's infection control protocols. Always follow your organisation's infection control guidelines to determine the appropriate cleaning schedule. Provide a copy of these instructions to the person(s) laundering the sling.
1. Where present, close hook & loop fastenings before washing to prevent damage to the fabric and webbing.
2. Wash gently at a minimum of 40°C and a maximum of 95°C. Use a mild soap solution or a non-biological detergent without bleaching additives.
3. Do not add bleach or stain removers.
4. Air dry, cabinet dry or tumble dry on a cool cycle (below 60°C). Do not tumble dry on a hot setting.

STORAGE CONDITIONS
Standard storage conditions. Keep off the floor. Keep out of direct sunlight. Keep dry.

DISPOSAL
No specific disposal requirements, handle contaminated items as clinical waste. Dispose of device in line with local and/or hospital policy.

COMPLAINTS
Any user who has a complaint or experiences dissatisfaction in the product quality, identity, durability, reliability, safety, usability, effectiveness, and/or performance, should notify the manufacturer and distributor immediately.

In the event of a serious incident involving the device, the user should report the event to the manufacturer (GBUK Group) and the competent authority of the member state in which the user and/or patient is established, immediately, by telephone, email (complaints@gbukgroup.com) or written correspondence.
When filing a complaint, provide the component(s) REF and SN number(s), your name and contact details, and the nature of the complaint.

In the event of a suspected device fault the user should report the event to GBUK Group with the above information and return the device to GBUK Group for examination.

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