

QUIKLITTER XL™

REF #50-0449

Multi-Use Disposable Litter

Rated to 1000 lbs (453 kg)

Size: 80" x 40" Weight: 2 lbs 5 oz



RESCUE ESSENTIALS

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LOT



QuikLitter XL™ Instructions

The QuikLitter XL™ is rated to transport up to 1000 pounds (453 kg). It is recommended to use at least four people to ensure safe transport and transfer, minimizing the chance of accident or injury. Not recommended when spinal stability is an issue. Improper handling or use of damaged QuikLitter XL™ can result in death or serious injury. Please follow all instructions for best results.

CONTRAINDICATIONS AND WARNINGS

- Do not use QuikLitter XL™ if unit is excessively worn or punctured.
- Avoid contact with sharp objects.
- Avoid dragging QuikLitter XL™ over rough surfaces.
- Do not store QuikLitter XL™ in contact with heat sources greater than 190°F.
- Not to be used in conjunction with any type of mechanical lifts for either vertical or horizontal movement.

PATIENT HANDLING

- Do not exceed load capacity of 1000 pounds.
- Use a minimum of 4 people to lift patient.
- Ensure proper grasping technique before lifting patient.
- Ensure retention straps are properly buckled before lifting patient.

CLEANING

- Soiled QuikLitter XL™ can be cleaned using a damp cloth with soap, detergent, or a mild disinfectant.
- Do not machine wash.
- If the entire device is contaminated, place it in a red biohazard bag and dispose according to your hazardous waste protocol.
- Avoid use of harsh detergents or disinfectants.

INTENDED USE

The QuikLitter XL™ is intended for use by healthcare professionals for general patient stabilization, transport, and transfer. Due to the lightweight and compact construction of the device, it is especially useful in austere environments where rapid casualty evacuations are needed.

 MD	Medical Device	 EC REP	Authorized European Representative	 LOT	Lot Number	 PK	Country Code
 LATEX	Not Made with Natural Rubber Latex		Manufacturer		Consult Instructions For Use		Date of Manufacture

Made in Pakistan

Notice: Any serious incident that has occurred in relation to this device should be reported to Tri-Tech Forensics and the Competent Authority of the Member State in which the user and/or patient is established.