

Multi-Use Disposable Litter

Red Handles Rated to 500 lbs (226 kg) Size: 72" x 27" Weight: 17 oz

RESCUE





www.rescue-essentials.com

info@rescue-essentials.com | 910-830-0286

MD

Tri-Tech Forensics, Inc. dba Rescue Essentials 3811 International Blvd NE, Ste 100, Leland, NC 28451 USA



MedEnvoy Global B.V. Prinses Margrietplantsoen 33, Suite 123 2595AM The Hague The Netherlands





BALINTHING

QuikLitter[™] Instructions

The QuikLitter[™] is rated to transport up to 500 pounds (226 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[™] can result in death or serious injury. Please follow all instructions for best results.

CONTRAINDICATIONS AND WARNINGS

- Do not use the QuikLitter[™] if unit is excessively worn or punctured.
- Avoid contact with sharp objects.
- Avoid dragging the QuikLitter[™] over rough surfaces.
- Do not store the QuikLitter[™] 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 500 pounds.
- Use a minimum of 4 people to lift patient.
- Ensure proper grasping technique before lifting patient.

CLEANING

- Soiled the QuikLitter[™] 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[™] 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.

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.

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

Made in Pakistan

