Intended Use

The BOA® Constricting IV Band provides circumferential pressure on either of the upper extremities for the purpose of obtaining peripheral venous access.

Warning

The BOA® should be used only by trained medical professionals. Improper use could result in injury to casualty. Reuse of this device will degrade efficacy, resulting in adverse casualty reaction, including potential death. Continually monitor casualty to ensure BOA® is functioning per medical protocols. In the event of a BOA® malfunction, follow local protocols and report any serious incident to North American Rescue or authorized representative and the competent authority of the Member State.

Contraindications

Within the European Union (EU), the BOA® Constricting IV Band is not indicated for pediatrics and pregnant women due to EU regulatory requirements.

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NORTH AMERICAN RESCUE®

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WARRANTY: The product or components contained in this package may constitute a medical device for which specific training is required for proper use. North American Rescue, LLC. (NAR), warrants that the product is merchantable and fit for its specified purpose. NAR expressly disclaims all other express or implied warranties relating to the product; any use beyond the product's specified purpose; or use by any party who is not trained or legally authorized to use such product. Use only as directed by your EMS authority or under the supervision of a physician.

DISPOSAL: The BOA® is a single use device and designed for disposal after use. Do not attempt to clean or reuse the device, as it may increase the possibility of cross contamination. Dispose of the device in a manner ensuring the isolation of potential substances in accordance with universal protocols. Contact NAR for more information as needed.

BOA®-Constricting IV Band

REF 30-0009 NSN: 6515-01-537-2611

REF 30-0071 BOA XL

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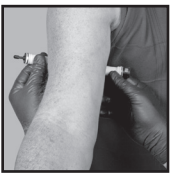
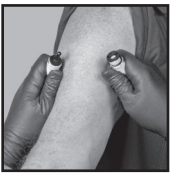
CE EC REP MDI Europa GmbH Langenhagener Str. 71 30855 Hannover-Langenhagen Germany MD

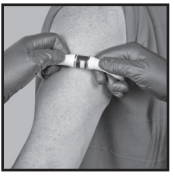
Instructions for Use



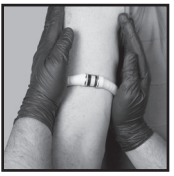
1. To find the correct size, place the BOA® (in its relaxed state) gently around the arm. It should measure 1/2 to 2/3 around the extremity.



2. Stretch BOA® straight out and place as high as possible around upper arm.

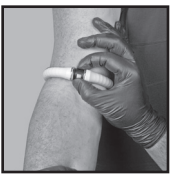


3. Keep fingers underneath the connectors as you secure in place to prevent pinching of the skin.



4. Press the "Quick Release" button and pull to remove the BOA®.

5. Monitor casualties with compromising clinical conditions. Evacuate casualty to secondary treatment facility, advising follow on care to monitor casualty for allergic reactions or infections.



6. In the event of device issues refer to medical protocols.

Harmonized Standard Symbols:

- LOT — Lot number
- Single use
- MD — Medical Device
- Consult instructions for use
- Manufacturer
- EC REP — Authorized Representative
- REF — Device part number
- Not for use in MRI
- Date of manufacture
- Expiration Date
- Keep Away from Sunlight
- Keep Dry