

## Female Luer Lock Cap

The sterile, red Female Luer Lock Cap is for use on medical devices to cover male Luer fittings to reduce the risk of touch contamination and medication leakage. Luer lock fittings are securely joined by means of a tabbed hub on the female fitting which screws into threads in a sleeve on the male fitting.

### Specifications:

- Packaged Individually
- Sterile and non-pyrogenic in unopened, undamaged package
- Non-vented
- Single use only
- Not made with natural rubber latex
- Contains no di(2-ethylhexyl) phthalate additives
- Made in the USA

### Material:

Verify compatibility with the drug manufacturer.

- High Density Polyethylene

### Directions:

Use aseptic technique. Peel open package to expose Female Luer Lock Cap. Hold the cap without contaminating the Luer end. Align the cap with the Luer fitting. Attach cap to the Luer fitting with simultaneous push and twist, while applying force.

### Indications for Use:

The Female Luer Lock Cap is indicated for use as a cap for male Luer fittings on medical devices such as manifolds, stopcocks or IV sets.













### Warnings and Contraindications:

#### WARNINGS:

- Female Luer Lock Cap terminates the fluid path.
- Female Luer Lock Caps are a choking hazard. Keep out of the reach of children.

**CONTRAINDICATIONS:** Not intended for blood administration sets.

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Symbol	Title of Symbol/Definition/Reference
	Do not re-use / Indicates device is intended for one use or for use on a single patient during a single procedure / Ref. 5.4.2 ISO 15223-1:2016
	Consult instructions for use / Indicates the need for the user to consult the instructions for use / Ref. 5.4.3 ISO 15223-1:2016
	Use-by-date / Indicates the date after which the device is not to be used / Ref. 5.1.4 ISO 15223-1:2016
	Sterilized using irradiation / Indicates device has been sterilized using irradiation / Ref. 5.2.4 ISO 15223-1:2016
	CAUTION! Federal (US) law restricts this device to sale, distribution, and use by or on order of a physician. / Ref. 21 CFR § 801.109
	Do not use if package is damaged / Indicates device should not be used if package is damaged or opened / Ref. 5.2.8 ISO 15223-1:2016
	Manufacturer / Indicates the medical device manufacturer name & address / Ref. 5.1.1 ISO 15223-1:2016
	Batch code / Indicates the manufacturer's batch code / Ref. 5.1.5 ISO 15223-1:2016
	Catalogue Number / Indicates the manufacturer's catalogue number / Ref. 5.1.6 ISO 15223-1:2016
	CAUTION! Indicates important cautionary information. / Ref. 5.4.4 ISO 15223-1:2016