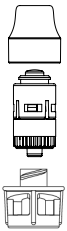
EN

ProSeal™ Injection Site with Priming Cap



REF **422120**

Priming Volume 0.17 ml
External Fitting Male Luer Lock (ISO 80369-7)
Compatible with DMA (Dimethylacetamide)

NOT MADE with NATURAL RUBBER LATEX or DEHP

EC REP

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Indications for Use:
The ProSeal Closed System drug Transfer Device (CSTD) mechanically prohibits environmental contaminants from entering the system and the escape of drug or vapor concentrations from the system, thereby minimizing individual and environmental exposure to drug vapor, aerosols, and spills. The ProSeal system also prevents the introduction of microbial contaminants into the drug or fluid path for up to 168 hours or 7 days.

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Rx Caution: Federal (U.S.A.) law restricts this device to sale by or on the order of a physician.

ProSeal Injection Site is the interface between any standard female luer lock port and ProSeal CSTD component for closed system fluid transfer. When connected to a female luer lock port and engaged with a ProSeal Injector (Syringe Adaptor) Ref 421010, fluid can be transfer to the connecting device in a closed system.

The ProSeal injection site and all its corresponding interface membranes exhibit a dry connection with the communicating surfaces in a fluid transfer. The use of this component and its appropriate ProSeal CSTD connecting component reduces the risk of microbial ingress for up to 168 hours or 7 days.

DIRECTIONS - Use Aseptic Technique

1. Prime the ProSeal Injection Site by connecting to ProSeal Injector and push syringe. Ensure the priming cap is securely attached during priming. The hydrophobic filter on the priming cap stops liquid flow once priming completed.
2. Remove the priming cap and connect the ProSeal Injection Site to the female luer lock port of connecting device.
3. Prior to every access, swab the top of ProSeal Injection Site with 70% isopropyl alcohol (15 seconds) and allow to dry (approximately 30 secs).
Drying time is dependent on temperature, humidity, and ventilation area.
4. Attach ProSeal Injector onto ProSeal Injection Site for fluid transfer to connecting device.
5. Discard used ProSeal Injection Site with connecting device intact in accordance to disposal procedures for biohazardous materials of your facility.

Contraindications
The device is contraindicated whenever:
- The drug to be prepared is contraindicated to Copolyester, TPE, stainless steel, polypropylene, silicone and polyisoprene.

Warnings
- Use accepted IV and pharmacy practice.
- The performance of the self-sealing membrane of the device is reduced after multiple perforations.
- Do not re-use to avoid contamination.

Precautions
- Do not use when caps and /or components are loose.
- The device is sterile unless packaging is damaged. Do not use when packaging is damaged.