

Instructions For Use

Contraindications:

PosiSep® X should not be used on patients who have known allergies to shellfish.

Intended Use:

PosiSep X Hemostat Dressing/Intranasal Splint is indicated for use in patients undergoing nasal/sinus surgery as a space occupying splint and hemostat to:

- Separate tissue or structures compromised by surgical trauma;
- Separate and prevent adhesions between mucosal surfaces during mesothelial cell regeneration in the nasal cavity;
- · Help control minimal bleeding following surgery or trauma;
- Help control minimal bleeding following surgery or nasal trauma by tamponade effect, blood absorption and platelet aggregation; and
- Act as an adjunct to aid in the natural healing process.

PosiSep X is indicated for use as a nasal hemostat to treat epistaxis.

PosiSep X is intended for use under the direction of a licensed healthcare provider.

Description:

Hemostasis, LLC offers two different platforms of hemostat dressings/intranasal splints. Both PosiSep and PosiSep X are patient-comfortable sponges manufactured from naturally occurring Chitosan polymers. Both PosiSep and PosiSep X expand when hydrated providing separation between the sinus tissues. However, if extra gelling properties are desired, PosiSep will form a smooth gel surface when exposed to liquids. If extra expansion properties are desired, PosiSep X will expand more than PosiSep to fill in spaces.

Warnings and Precautions:

- PosiSep X is not bacteriostatic toward pre-existing infections, nor does it prevent the occurrence of new infections.
 Appropriate treatment to be instituted in case of infection.
- When utilized as an intranasal splint, if the PosiSep X becomes completely coated with blood prior to hydration then the function of the device as a hemostat may subsequently diminish the expansion of the device via hydration.
- Sterility not guaranteed if package is damaged or opened. Discard damaged or open packages.
- Single Use Only.
- To prevent product contamination prior to application, always follow aseptic techniques.
- Clinician must inform patient that the device contains material derived from shellfish.
- In rare instances, the physiochemical condition associated with nasal surgery, both with and without nasal packing, may present a risk of toxic shock syndrome (TSS).
- Appropriate treatment to be instituted in case of infection.
- In the case of prolonged bleeding seek medical attention.
- · Do Not Ingest, do not inhale.
- To avoid risk of aspiration, do not place in the posterior nasal cavity.
- Do not use after the end of the year and month shown.
- When used as a topical dressing, PosiSep X should be kept dry prior to use.
- This product shall be disposed of in compliance with pertinent government regulations regarding medical devices.
- PosiSep X is intended for topical use only.
- PosiSep X has not been tested for use with other devices or medicines. Use only according to labeled instructions.

Instructions for Topical Dressing:

- 1. Peel open PosiSep X pouch and remove dressing using sterile technique to maintain product integrity.
- Blot excess blood from the wound with gauze.
- Apply PosiSep X dressing to cover wound.
- As necessary, apply pressure and hold in place until hemostasis is achieved, which will generally take
 minutes depending upon the nature of the bleeding site and specifics of the patient.
- If hemostasis has not been achieved, repeat steps 2-4 or use an alternate method of hemostasis treatment.
- 6. Once hemostasis is achieved, remove PosiSep X from the wound site using gentle irrigation with saline or water and aspirate as needed. If decided to leave the PosiSep X in place, PosiSep X will be naturally eliminated between 5 to 10 days via mucociliary action, assisted by daily irrigation with fluid which helps with fragmentation of the device.
- 7. Discard any unused product after opening.

Instructions for Intranasal Splint:

- 1. Peel open PosiSep X pouch and remove dressing using sterile technique to maintain product integrity.
- 2. PosiSep X may be cut or bent to a size appropriate for the application.
- 3. Using forceps, insert PosiSep X longitudinally and apply to area of treatment.
- 4. If PosiSep X is broken during insertion, assess if size is appropriate for application. If not, dispose and reapply PosiSep X.
- 5. After placement, hydrate the PosiSep X using a sterile fluid.
- Discard any unused product after opening.
- PosiSep X will be naturally eliminated between 5 to 10 days via mucociliary action, assisted by daily irrigation with fluid which helps with fragmentation of the device

How Supplied:

See pouch and carton labels for package quantity.

PosiSep X is terminally sterilized by gamma irradiation.

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









Store at room temperature.



5000 Township Parkway St. Paul, MN 55110 USA 866-612-2568

www.hemostasisllc.com
Made in USA. Patent Pending.

Symbol	Description
RONLY	CAUTION: Federal (U.S.) law restricts this device to sale by or on the order of a physician
#	Quantity Symbol: numeral in symbol (in place of #) indicates the quantity of units in package.
\triangle	Caution: See Instructions for Use
STERILE R	Sterilized by Irradiation
2	Single Use Only. Do not reuse.
STEPREZE	Do Not Re-sterilize
	Do Not Use if Package is Damaged
- 	Keep Dry
	Manufacturer
REF	Catalog Number
LOT	Lot Number
	Expiration Date (Use By). This device should not be used after the end of the year and month shown.

©2017 Hemostasis, LLC. PosiSep is a registered trademark of Hemostasis, LLC. 50059 Rev. L 03/2018