



RETRAX[®] FS SOLUTION

REF 15-640

15mL

Contents: 15.5% Ferric Sulfate in an aqueous base

Indications for Use:

For the purpose of aiding in gingival retraction and to suppress excessive capillary bleeding associated with dental impression procedures, as well as for tissue management problems in other phases of dentistry. This product is for topical use only. Retrax FS Solution is compatible with all types of retraction materials except those containing epinephrine.

Contraindications:

None known.

Warnings:

For topical intraoral use only. Do not ingest. Keep out of reach of children.

Precautions:

- For professional office use only.
- Do not use on patients with an allergy to ferric sulfate.
- High volume suction is recommended to avoid possible ingestion.
- Temporary tissue discoloration may occur. This discoloration should last no more than 24 – 48 hours.
- Do not use Retrax FS Solution with products containing epinephrine.

Dosage:

Use as needed to achieve and maintain hemostasis.

Directions for Use:

- Soak cotton pellet or retraction cord in dispensed solution.
- Apply pellet or cord to gingival site.
- Wait 2 - 6 minutes.
- If bleeding persists, rinse area and reapply soaked pellet or cord. Wait 2 - 6 minutes.
- Remove cord or pellet.
- For best results, thoroughly rinse and dry application site.
- Take impression as usual.
- Any soaked pellets or cord should be discarded immediately after procedure.

Retrax FS Solution should only be applied using cotton pellets, retraction cord, or cotton tipped applicators. Solution should never be dispensed directly into patient's mouth or directly to human tissue.

For product SDS information please see company website at www.pascaldental.com or contact Pascal directly.

Pascal's Retrax FS Solution is free of gluten, soy, milk, egg, peanut and tree nut products. Retrax FS Solution contains no artificial color or flavor.

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.



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