

Quality Disposable Dental Products

Dental Matrix Strips Instructions for Use

DEVICE PURPOSE / INTENDED USE

Matrix Strips are used for restorations in which the proximal wall of an anterior tooth is missing. The matrix strip is placed interproximally before the etching and priming of a tooth. This protects adjacent teeth from these materials.

Materials:

Polyester film

INDICATIONS FOR USE

The matrix strips are indicated for temporary use in dental restorations in which the proximal wall of an anterior tooth is missing.

CONTRAINDICATIONS

The following contraindications apply Matrix Strips:

Patients with known allergy the materials of construction.

WARNINGS

 If patient experiences allergic reaction, immediately remove the Matrix Strip and advise patient to seek medical care.

CAUTIONS

- For single use only
- Federal (USA) law restricts the sale of this device by, or on the order of, a licensed dental professional.

METHOD OF STERILIZATION

Raydenco disposable dental products are delivered nonsterile to the end user.

METHOD OF DISINFECTION

Raydenco disposable dental products do not require disinfection prior to use.

CONDITIONS OF STORAGE

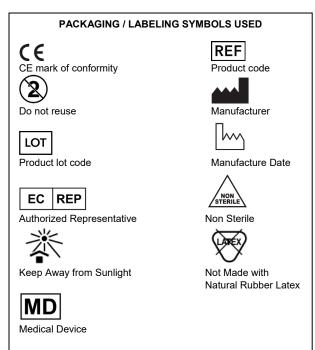
The product should be stored in at ambient temperature and protected from bio-contamination.

NOTICE

Report any serious incident that has occurred in relation to this device to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

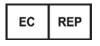
OPERATION AND USE

- For restorations in which the proximal wall of an anterior tooth is missing, place the Matrix Strip interproximal before the etching and priming of a tooth to protect adjacent teeth.
- 2. After placement of composite material pull the Matrix Strip tightly around the tooth to help reconstruct its natural contour.
- 3. The clear plastic Matrix Strip allows the curing light to penetrate the strip and complete the curing process.
- 4. When restoration is completed remove the Matrix Strip and dispose.
- Dispose of used Matrix Strips according to facility procedures
- 6. This product is a non-sterile single-use medical device.





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Revision	CR#	Description of Change
00	CR-2025 003	Initial Release