# PTFE Tape

**Manufacturer:**

Smart Denture Conversions, LLC
1800 N. Salem St.
Suite 104
Apex NC, 27523
855-550-0707
www.SmartOnX.com




**Training:**

It is strongly recommended that clinicians, new as well as experienced implant and abutment users, always go through special training before undertaking a new treatment method. Smart Denture Conversions offers a wide range of courses for various levels of knowledge and experience. For more information, please visit www.SmartOnX.com.

**Product Description:**

PTFE Tape is polytetrafluoroethylene tape that is placed over bite registration during a pickup procedure. The table below summarizes information on the device:

| Name | Width | Material | Use |
|---|---|---|---|
| PTFE Tape | 2 in. | Polytetrafluoroethylene | Single Use |

**Intended Use/Indications for Use:**

PTFE Tape is intended to act as a barrier to prevent bite registration material from drying onto any components or oral tissue or bone. This helps the bite registration material properly capture the location of prosthetic components. PTFE Tape is intended for single use. Re-use of single use devices creates a potential risk of patient or user infection and misfitting components.

PTFE Tape is indicated for use during intraoral pickup procedures using the Smart Denture Conversions technique. For more specific information on process steps, please refer to the Technique manual located on the website www.SmartOnX.com.

**Contraindications:**

It is contraindicated to using Smart Denture Conversions' Teflon Tape in:

- Patients who are medically unfit for an oral surgical procedure.
- Patients in whom adequate sizes, numbers or desirable positions of implants are not reachable to achieve safe support of functional or eventually parafunctional loads.
- Patients who show signs of allergy or hypersensitivity to the chemical components of the materials listed in the chart above.

**Warning:**

- Components are to be used by dental health care professionals and are to be used in patients subject to dental implant treatment.
- If the indication or the nature of use is not clear, do not use until all points have been clarified.
- Do not use if package is damaged.
- Even if the product is used according to the instructions for use, the clinical outcome of dental treatment is influenced by multiple variables and may lead to any or all of the following complications: anaphylaxis (severe allergic reaction); aspiration or swallowing of components; pain; local infection; inflammation; local irritation; loss of product function; follow-up treatment.
- Always inspect components before use. Do not use damaged, deformed, corroded, or discolored components.
- Ensure products are secured against aspiration when handled intraorally. Aspiration of products may lead to infection or unplanned physical injury.

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- Smart Denture Conversions is not liable for damage resulting from use outside the intended use of the product.
- Failure to follow the procedures outlined in these instructions may lead to any or all of the following complications: Aspiration or swallowing of components, follow-up treatment, incorrect impression resulting in incompatible restorations.

**Cautions/Precautions:**

The following precautions are required before or during treatment:

- Do not use Smart Denture Conversion components after the expiration date on packaging (if applicable).
- All products intended for single use must not be reused. Re-use of single use devices creates a potential risk of patient or user infection and misfitting components.
- Before every procedure make sure that all required components, instruments, and auxiliary equipment are complete, in operating order and available in the required quantity.
- If, due to unfavorable anatomical conditions, instruments do not fit or cannot be used for other reasons, the course of treatment planned with them must not be continued and alternatives must be sought.
- Always wear suitable personal protective equipment for your own safety.
- Position the patient such that the danger of aspiration of components is minimized.

**Magnetic Resonance Imaging (MRI) Safety Information:**

Smart Denture Conversion LLC PTFE Tape is a tool used in the operatory with transient patient contact, and does not remain in the body. This product has not been evaluated for safety and compatibility in the MR environment. It has not been tested for heating, migration, or image artifact in the MR environment. The safety of the Smart Denture Conversion LLC PTFE Tape in the MR environment is unknown.

**Disinfection Instructions:**

Components are delivered non-sterile by Smart Denture Conversions and are intended for single use. Devices should be disinfected prior to use on a patient to ensure safety and prevent the risk of infection. An intermediate disinfectant, such as Chlorhexidine gluconate (0.12%), should be used. Follow the manufacturer's instructions for the correct application, ensuring adequate contact time to achieve effective disinfection. Ensure that the device is clean and free from any contaminants before applying the disinfectant to maximize efficacy.

**Note:** This device should not be steam sterilized, as exposure to high temperatures and moisture may cause the part to deform and compromise its functionality.

**Storage, Handling and Transportation:**

The devices must be stored in a dry place in their original packaging at room temperature and protected from direct sunlight. Improper storage may compromise essential material and design characteristics, leading to device failure.

**Disposal:**

Safely discard potentially contaminated or no longer usable medical devices as healthcare (clinical) waste in accordance with local healthcare guidelines, country and government legislation or policy. Separation, recycling or disposal of packaging material shall follow local country and government legislation on packaging and packaging waste, where applicable. If there is no current legislation, pack them in a perforating waste/sharps disposal container and dispose of them in hospital waste.

**Validity:**

Upon publication of these instructions for use, all previous versions are superseded.

**Availability:**

Some items of Smart Denture Conversions are not available in all countries.

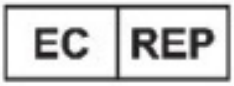
**Warranty:**

Please visit www.SmartOnX.com for the most up to date warranty information.

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**Symbols Glossary:**

The following Symbols may be present on the device labeling or in information accompanying the device. Refer to the device labeling or accompanying information for the applicable symbols.

| | | | |
|---|---|---|---|
|  | Manufacturer |  | Date of Manufacture |
|  | Use By Date |  | Serial Number |
|  | Batch Code |  | Catalog Number |
|  | Unique Device Identifier |  | Medical Device |
|  | CE Mark |  | UK Representative |
|  | United Kingdom Conformity Assessment Mark |  | United Kingdom Conformity Assessment Mark with Approved Body Number |
|  | European Representative |  | Non-Sterile |
|  | Do Not Use if Packaging is Damaged |  | Consult Instructions for Use |
|  | Keep Dry |  | Keep Away from Sunlight |
| RX Only | For Prescription Use Only |  | Caution, Consult Accompanying Documents |