

Instruction For Use

Absorbent Paper Points

(English Version) 

Sterilized endodontic points

Used to remove moisture from the root canal or to introduce medicaments.

Composition

Paper 100%

Application

1. Check if there is any abnormality in the appearance of the product such as foreign substances or cracks.
2. Check if the product is within the expiration date.
3. After forming the root canal, select the size that fits the file and dry the root canal, and use it with medication after.
4. This product is a gamma ray sterilized disposable medical device.

Intended Purpose

1. Intended Use: Used to remove moisture from the root canal or to introduce medicaments.
2. Indication: Used to remove moisture from the root canal or to introduce medicaments.
3. Contraindication: No contraindication has been reported that can cause health problems.
4. Patient Population: All age group
5. Intended user: Dentist
6. Side Effects/Adverse Effects: There are no side effects or adverse effects reported.
7. Potential risk if re-used: Re-use of used Paper point may cause inflammation.
8. Can be used with medicaments and tools and it shall be used under the control of the dentist according to the clinical indication.
9. Single Use Device

Precaution

1. Do not use other than dental specialist.
2. Use disinfected tools.
3. This product is a gamma-ray sterilized disposable medical device.

Storage

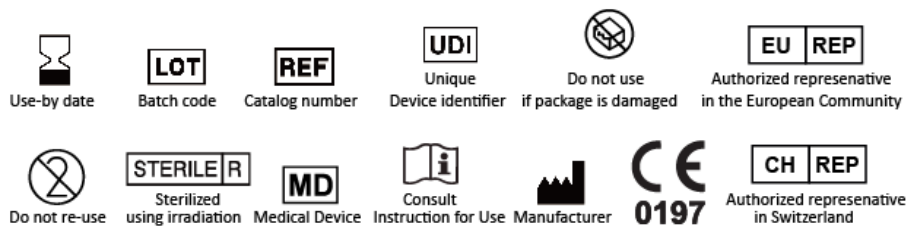
Store at room temperature.

Disposal : Dispose of contents/container according to the related regulations of the country.

Shelf Life : 4 years from the date of manufacture

Nominal Length : 28 mm

Symbols



SURE DENT CORPORATION (Manufacturer)

#809, 52, Sagimakgolro, Jungwon-gu, Seongnam-si, Gyeonggi-do, 13210, Republic of Korea
Tel: +82-31-737-4720 Fax: +82-31-737-4724 <http://www.suredent.com>

AUTHORIZED REPRESENTATIVE

Europe - OBELIS S.A. - Bd. Général Wahis, 53, 1030 Brussels, Belgium

Switzerland - Obelis GmbH - Ruessenstrasse 12, 6340 Baar / ZG, Switzerland

■ In case that any serious incident that has occurred in relation to the device, please report to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

■ A free printed copy of the IFU in the appropriate language is available upon request, within 7 days in the EU. Please contact us at suredent@suredent.com. An electronic version of IFU is available at <https://www.suredent.com/media/InstructionOfProducts>.