DENTALSTUDIO

MICRO BONE SCRAPER

www.dentalstudio.co.kr

Manufacturer **Dental Studio Co., Ltd.**
B101, B102, B125, B1, Geumjeong Station 2nd SKV1 Tower, 136 LS-ro, Dongan-gu, Anyang-si, Gyeonggi-do, Republic of Korea
Tel: +82 31 8069 7007 / Fax: +82 31 8069 7009

EC REP
CMC Medical devices & Drugs S.L.
C/Horacio Lengo N 18 CP 29006, Málaga-Spain
Málaga-Spain
Tel. +34951214054, Fax. +34952330100

CE	● CE conformity marking
	● Date of manufacture
LOT	● Batch code
	● Non-sterile
	● Manufacturer
	● Caution, consult accompanying documents
Rx Only	●Federal law(USA) restricts this device to sale by or on the order of a licensed dentist
	● Consult instructions for use
EC REP	● Authorised representative in the european community

IFU-DS-I-3500-1(Rev. 1)/ 2024-11-06

[Product name] MICRO BONE SCRAPER
[Model name] 3500-1
[Weight or packaging unit] 1set (2ea in a case)
This product is a single use medical device.

Purpose of use

❑ A manual instrument used to cut or adjust hard tissue such as bone

Instruction for Use & Procedure

1. Preparations before use

1) This product is intended for professional use only by professional training providing medical devices.
2) The practitioner must fully understand the product's usage instructions and precautions before using it.
3) Before the procedure, prepare by selecting a product appropriate for the surgical area and purpose.
4) Before performing the procedure, check the condition of the product, including whether it is damaged, cleaned, and sterilized.
5) Since this is a disposable non-sterilized product, please follow the hospital's own cleaning and sterilization methods, or follow the cleaning and sterilization conditions below. Sterilize and prepare accordingly.
   - Washing method: Before use, disinfect and clean the equipment with distilled water and alcohol, then dry it
   - Sterilization method: After sterilization at 121℃for 30 minutes by high-pressure steam sterilization method, drying for 30 minutes

2. How to use

1) Hold the instrument so that the groove (gap) on the blade faces toward the bone.
2) Place the product's blade on the treatment area.
3) Be careful not to use excessive force and cut hard tissue such as bone by pulling in the direction of the handle. Used for adjustment.
4) After the procedure, the remaining cut bone tissue is stored in the groove (gap) of the blade, which watches the system closure. Turn in the opposite direction and push to remove bone tissue.

3. How to store and manage before use Store in a dry place at room temperature.

Precautions for use

1. This product is a medical device and can only be used by those who have received professional training.
2. Practitioners must be fully familiar with the product's usage instructions and precautions before using it, and only sufficiently trained or skilled practitioners should use it.
3. It must not be used for purposes other than its intended use.
4. Since this product is non-sterilized, it must be sterilized before use.
5. The manufacturer is not responsible for any damage to the product caused by sterilization using a method other than the sterilization conditions suggested by our company or improper handling.

Warnings, side effects and contraindications

1. Contraindication

1) This product is a disposable medical device and can be used only once.
2) Re-sterilization or cleaning after use is absolutely not possible.
3) We do not repair products damaged by the user.

2. Warning

For safe and effective use, we strongly recommend:

1) Do not use the product for purposes other than its intended use or modify the product.
2) It is not used in patients unsuitable for other surgeries.
3) The product must be handled with care to avoid damage or deformation.

3. Side effects

Careless use and storage of the product may cause reactions such as infection or metal allergy.

⚠ Caution

1) Use for purposes other than the intended use is prohibited.
2) Do not use products contaminated by surgeon's mistake during surgery.
3) The manufacturer is not responsible for any damage to the product caused by sterilization or improper handling of the product other than the sterilization conditions provided by the company.
4) Thorough pre-diagnosis and product inspection must be performed on the patient before the procedure.
5) This product is a non-sterilized product and must be used after sterilization.

IFU-DS-I-3500-1 (Rev. 2) : 2026. 08. 26