SURGICAL INSTRUMENTS

Fasteners, Keys, Rods, Analog, Pullers, Hammer and Curettes



The Instruments are intended for specialized procedures, which must be performed by qualified professionals. The way to use the product and surgical techniques are inherent to the professional's training. The use of the product must be performed in a surgical environment and under adequate conditions for the patient's health and safety.



PRODUCT DESCRIPTION

Instruments are indicated to aid in the placement of implants, components, plates and other surgical procedures following the appropriate dental techniques.

INDICATIONS OF USE

The instruments are recommended to aid in the placement of implants, components and other surgical procedures according to appropriate dental techniques.

Fixer and Driver: They assist in the installation or removal of components and implants.

Rounder: Used for easy handling of the components and implants in the surgical procedure.

Folding tool: They aid in folding plates.

Analog: It assists in the polishing procedure of the component.

Mucosa Extractor: An instrument used to guide the placement of the implant in the technique of guided surgery.

Prosthesis Extractor: Instrument used for the removal of prosthesis on implants.

Hammer: Used with surgical hammer in surgeries with osteotomes.

Roinder 02 and 03: Angled sinus membrane strainer. Used for sinus membrane displacement.

Maxillary sinus curette: It has sinus membrane displacement function.

OPERATION PRINCIPLE

All dental instruments are based on mechanical principles of power transmission, for the activation of secondary elements (such as driver).

HOW TO USE

- Open the original packaging and remove the Instruments;
- Prior to first use or after having been used in surgical procedures, perform the cleaning and sterilization procedures contained in these Use Instructions.
- After sterilization, the instruments are ready for use.



ATTENTION

The instruments are intended for specialized procedures, which must be performed by qualified professionals in implant dentistry. The use of the product must be performed in a surgical environment and under adequate conditions for the patient's health and safety.

PRECAUTIONS

The product should only be used by qualified dentistry professionals who already have all the scientific information necessary for the correct use of the product.

RECOMMENDATIONS

It is recommended for use by dental professionals only.

CONTRAINDICATIONS

The Instruments do not present contraindications once they follow their recommendations correctly. They are used by specialized professionals.



SIDE EFFECTS

Not Applicable. Adverse effects will occur only if the choice of instruments is inadequate.

WARNING

Do not use the instrument if you notice cracks, wear or oxidation/corrosion points. This may cause instrument performance problems. All items may appear a natural wear due use and they should be replaced whenever the professional identifies loss of fitting capacity or accuracy of these products, as they may interfere with final work results.

TRACEABILITY

All S.I.N. products - Implant System products have sequential batches that allow traceability, which promotes greater safety for the professional qualified to the procedure. Through this batch number, it is possible to know the entire history of the product from the manufacturing process to the distribution time.

STORAGE

Instruments should be stored in a dry, fresh and ventilated, place away from direct sunlight;

HANDLING

Once sterilized, instruments should be handled only in a sterile environment by properly trained professionals and in appropriate suits at the time of the surgical procedure.

DISPOSAL OF MATERIALS

The disposal of materials should comply with local hospital regulations and applicable local laws.

TRANSPORTATION

Instruments should be transported at room temperature away from direct sunlight, avoiding locations where greater variations in temperature and humidity occur.

The transportation must be carried out properly, to avoid falls and it must be carried out in its original package.

COMPLEMENTARY INFORMATION

Multiple use product. Exclusive of Odontological use. Passible of reprocessing. See cleaning and sterilization conditions contained in this use instruction. In case of any incident caused by the product, the professional must inform the manufacturer immediately. If you need the printed version of this instruction for use, without any cost, please request by e-mail to sin@sinimplante.com.br or call to 0800 770 8290 will receive until 7 days calendar.

CLEANING INSTRUCTIONS

- Disassemble the product (if applicable). For the torque wrench, disassembly it completely, remove all the internal organic matter using tap water and follow to the next step only after performing such procedures.
- Prepare the enzymatic detergent according to the manufacturer's recommendation.
- Immerse all parts of the product into the prepared detergent solution and keep in contact for at least 5 minutes, then using soft bristle brush, scrub the parts to remove organic matter from the products.
- Remove parts from detergent solution and rinse with tap water for 1 minute, repeat the rinse for two more times, a total of three rinses of 1 minute each.
- 5. Visual inspection of each part for cleaning process residue or organic waste from product use.
- If residue is detected in the product, repeat the cleaning process until the residue is completely removed.
- Dry with a soft, clean, dry cloth or disposable paper.
- 8. Follow to sterilization process.

RECOMMENDATIONS

- a. Use the proper PPEs (gloves, masks, goggles, caps, etc.).
- b. Start the cleaning right after the surgical use.
- Never let the instruments dry with organic waste after the surgical use.
- d. Never let the instrument dry naturally after cleaning.



- e. Never use saline solutions, include sodium hypochlorite, disinfectant, hydrogen peroxide or alcohol to cleaning or rinsing the surgical instruments and Kits.
- Never use steel wool and abrasive products, so that the instruments are not damaged.
- g. Do not stack the instruments in lots to avoid the deformation of smaller and delicate pieces.

STERILIZATION

Reusable product and provided non-sterile. It must be clean and sterilized in autoclave before use.

- Dry all instruments before the steam sterilization cycle.
- 2. The product must be enclosed in a steam sterilizable wrap.
- Steam sterilize in cycles of 121°C at 1 ATM pressure for 30 minutes or of 134°C at 2 ATM pressure for 20 minutes. Drying time 30 minutes.
- 4. Always accommodate the case in autoclave over a plane surface and away of device walls.
- 5. Never stack objects or other cases.

RECOMMENDATIONS

- Sterilize the products in the same day or one day earlier the procedure.
- b. The chemical sterilization is not recommended, once some products may cause the discoloration and damages to the case.
- Do not use temperature higher than 60°C to drying process.
- d. Do not use dry heat stoves for sterilization of the instruments and kits from S.I.N. Implant System.

LIFE TIME

The instruments should be used as follows depending on proper handling, cleaning and sterilization.

Fixators, Hammer, Curettes, Keys and Pullers: Use up to 250 times;

Rods: Estimate 20 perforations;



NON STERRLE	NÃO ESTÉRIL	NON-ESTERILE	NO ESTÉRIL
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CE	MARCAÇÃO CE	CE MARK	MARCA CE
Ť	MANTENHA SECO	KEEP DRY	MANTÉNGALO SECO
类	MANTENHA AO ABRIGO DO SOL	KEEP AWAY FROM SUNLIGHT	MANTÉNGALO LEJOS DE LA LUZ SOLAR
®	NÃO UTILIZAR SE A EMBALAGEM ESTIVER VIOLADA	DO NOT USE IF PACKAGE IS DAMAGED	NO LO UTILICE SI EL ENVOLTORIO ESTÁ DAÑADO
lack	ATENÇÃO	CAUTION	PRECAUCIÓN
EC REP	REPRESENTANTE AUTORIZADO NA COMUNIDADE EUROPEIA	AUTHORIZED REPRESENTATIVE IN THE EUROPEAN COMMUNITY	REPRESENTANTE AUTORIZADO EN LA COMUNIDAD EUROPEA
- 35°C	LIMITE SUPERIOR DE TEMPERATURA	UPPER LIMIT OF TEMPERATURE	LÍMITE SUPERIOR DE TEMPERATURA
Rx only	ATENÇÃO: A LEI FEDERAL (EUA) LIMITA A VENDA DESTE DISPOSITIVO POR OU POR ORDEM DE UM PROFISSIONAL DE SAÚDE LICENCIADO.	(USA) RESTRICTS THIS DEVICE TO SALE BY OR ON	PRECAUCIÓN: LAS LEYES FEDERALES (USA) RESTRINGEN LA VENTA DE ESTE DISPOSITIVO POR O EN EL ORDEN DE UN PROFESIONAL DE LA SALUD LICENCIADO.
	FABRICANTE	MANUFACTURE	FABRICANTE
	DATA DE FABRICAÇÃO	DATE OF MANUFACTURE	FECHA DE FABRICACIÓN
REF	CÓDIGO DE REFERÊNCIA	REFERENCE CODE	CÓDIGO DE REFERENCIA
MD	DISPOSITIVO MÉDICO	MEDICAL DEVICE	DISPOSITIVO MEDICO
UDI	IDENTIFICADOR ÚNICO DO DISPOSITIVO	UNIQUE DEVICE IDENTIFIER	IDENTIFICADOR DE DISPOSITIVO ÚNICO
	IMPORTADOR	IMPORTER	IMPORTADOR
	DISTRIBUIDOR	DISTRIBUTOR	DISTRIBUIDOR
BRA	PAÍS DE FABRICAÇÃO	COUNTRY OF MANUFACTURE	PAÍS DE FABRICACIÓN
LOT	LOTE	BATCH CODE	LOTE
Δ	EMBALAGEM RECICLÁVEL	RECYCLABLE PACKAGING	EMBALAJE RECICLABLE



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