

Surgical Manual

Sinus KIT

Introduction

Welcome,

and thank you for choosing Osstem Implant products. This catalogue is designed to support dental professionals with product information, clinical workflows, and practical guidance for daily use. It is important to inform patients about the option of dental implant treatment and the potential benefits it may provide. For further information, please contact your local Osstem representative.

Important Notice

This document is provided for **informational and educational purposes only** and does not replace the applicable product label, the current product-specific Instructions for Use (IFU), formal clinical training, or independent professional judgment. All product information, specifications, and protocols are subject to change without notice. Not all products may be approved, cleared, released, licensed, or available in all markets. Product illustrations are not shown to scale. Despite careful preparation of this catalogue, typographical, editorial, translation, or printing errors may occur. **All critical information must be verified against the current product-specific IFU and product label before use.**

Electronic IFU (per (EU) 2021/2226)

- Surgical Drill & KIT System is eligible for provision of electronic instructions for use (e-IFU) under Regulation (EU) 2021/2226 for professional users.
- e-IFUs are available at [website URL: ifu.osstem.com] in the official languages required by the Member State(s) where the device is placed on the market.
- The e-IFU content is consistent with the paper version; all updates are promptly reflected in both versions.
- If requested, a paper copy of the IFU will be supplied free of charge, within 7 calendar days.
- The e-IFU website maintains historical versions for traceability of all previously applicable instructions.
- Labeling on the product/package indicates the provision of e-IFU and how to access it online.

Surgical Manual | English Edition

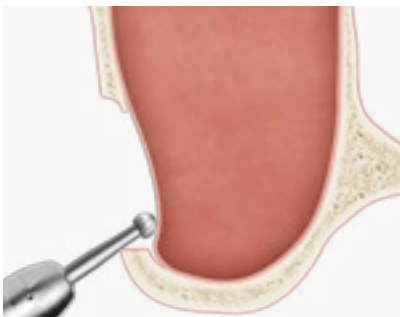
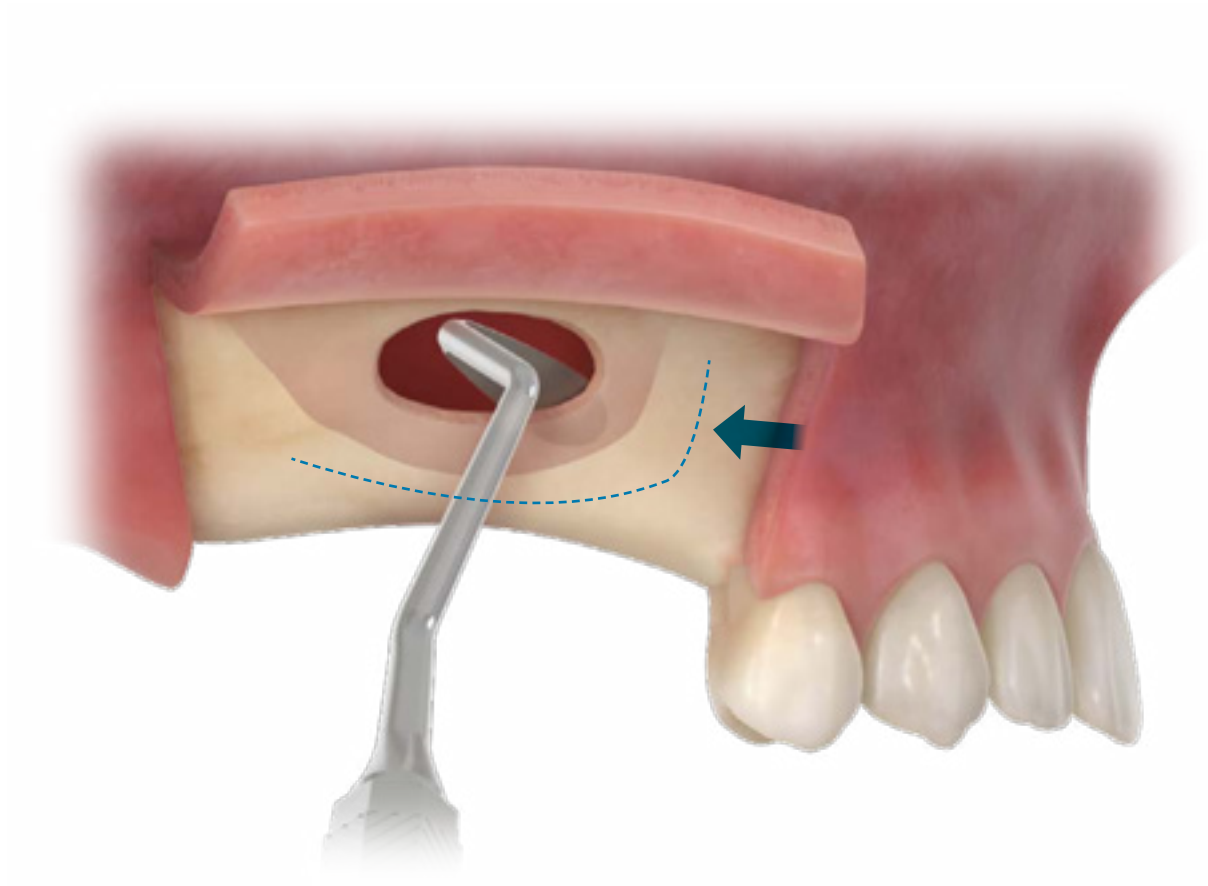
Publication date: April 2026

Publisher: Osstem Europe s.r.o.
Radlická 740/113c
158 00 Prague, Czech Republic

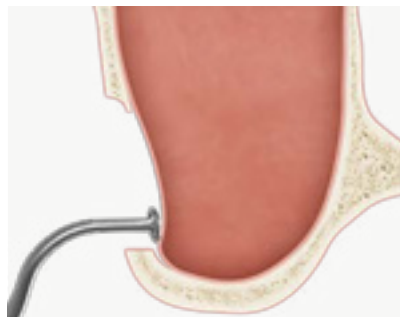
Note: *This brochure is based on the global 2021 Osstem Surgical Manual and has been visually revised and adapted for the European market. Product availability and specifications may vary by country and are subject to change without notice. Images are for illustrative purposes only. For professional use only.*

Sinus KIT

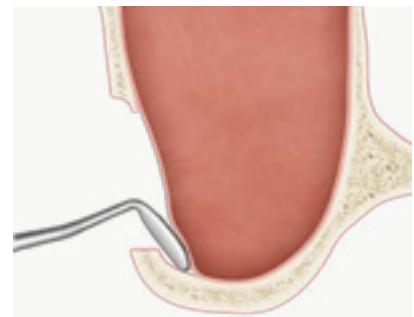
KIT with various configurations used for sinus procedures



Remove bony window



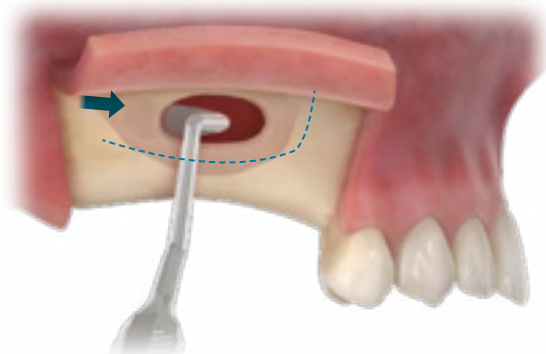
Peel maxillary sinus membrane



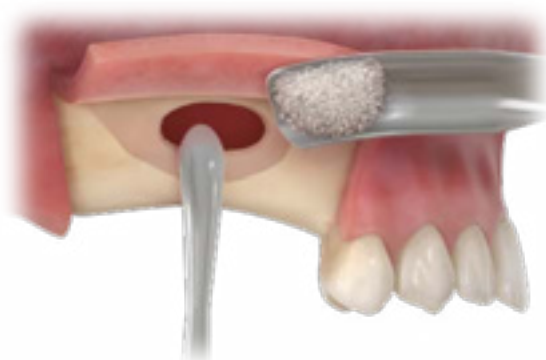
Lift maxillary sinus membrane

1 Indication

- A When lifting the maxillary sinus membrane after creating a bony window using the lateral approach

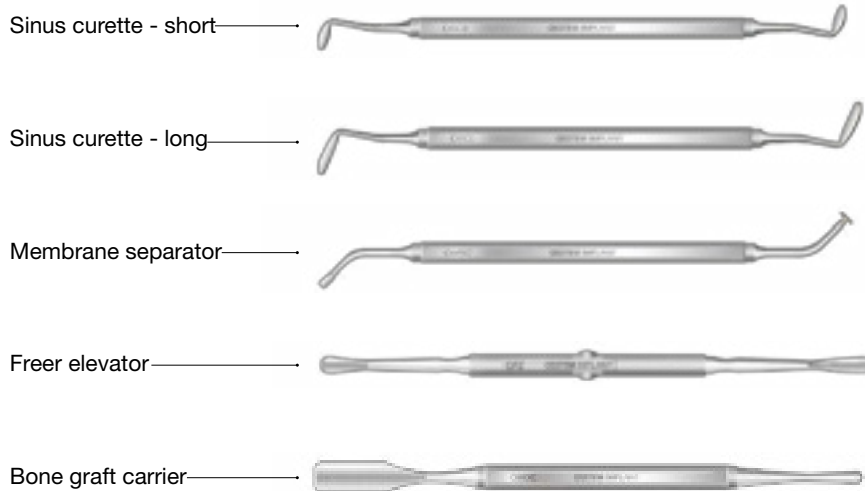


- B When bone graft after creating a bony window using the lateral approach



2 Feature

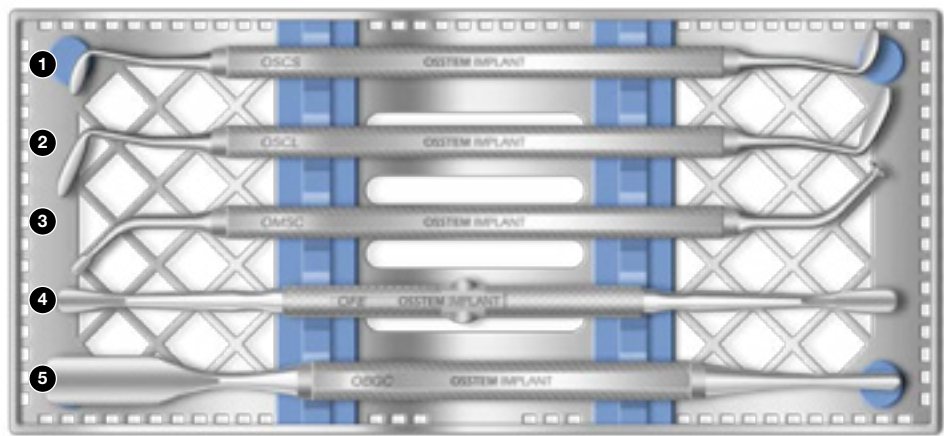
KIT used for maxillary sinus lifting and bone graft at the same time



3 KIT (included components)

Sinus KIT

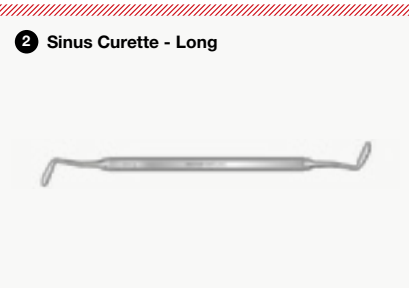
/// Sinus surgery tool
(1~5)



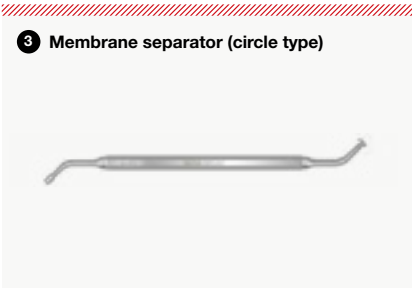
1 Sinus Curette - Short



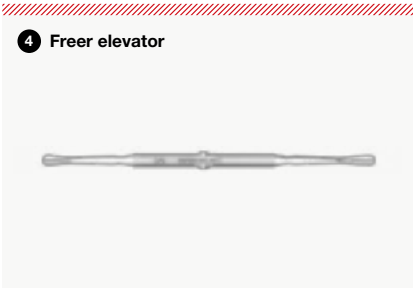
2 Sinus Curette - Long



3 Membrane separator (circle type)



4 Freer elevator



5 Bone graft carrier



4 KIT user instructions

Sinus surgery tool

Includes tools used for sinus procedure and bone graft filling.



1 Sinus Curette - Short



2 Sinus Curette - Long



3 Membrane separator (circle type)



4 Freer elevator



5 Bone graft carrier

1 Sinus curette-short

Use for maxillary sinus lifting.

User instructions

- Lift the sinus membrane in the distal, floor, mesial, and medial directions with a sinus curette (short).



Distal



Floor



Mesial



Medial

2 Sinus curette-long

Use for maxillary sinus lifting.

User instructions

- Lift the sinus membrane in the distal, floor, mesial, and medial directions with a sinus curette (long).



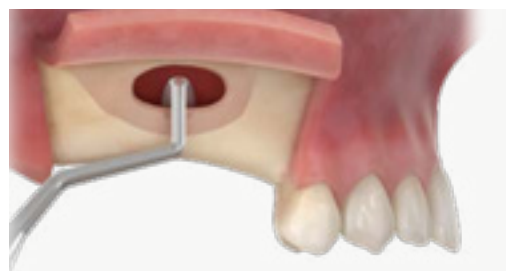
Distal



Floor



Mesial



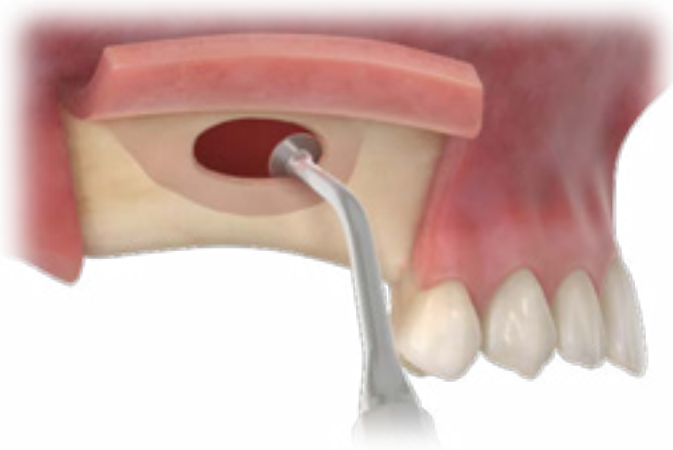
Medial

3 Membrane separator (circle type)

Use for initial detachment of the maxillary sinus membrane.
Use for filling bone grafts in the maxillary sinus.

User instructions

- The rounded tip is used in the initial stage of dissection of the maxillary sinus membrane after bone spur removal.
- The plugger tip is used when filling the bone graft in the maxillary sinus.



4 Freer elevator

Use to remove bone spurs in the maxillary sinuses.
Also use to raise the mucous membrane of the maxillary sinuses,
and to fill bone grafts in the maxillary sinuses.

User instructions

- Case 1: Use to remove the maxillary sinus bone.
- Case 2: Use to lift the maxillary sinus mucosal membrane.
- Case 3: Use to filling bone grafts in the maxillary sinuses.



Case 1



Case 2



Case 3

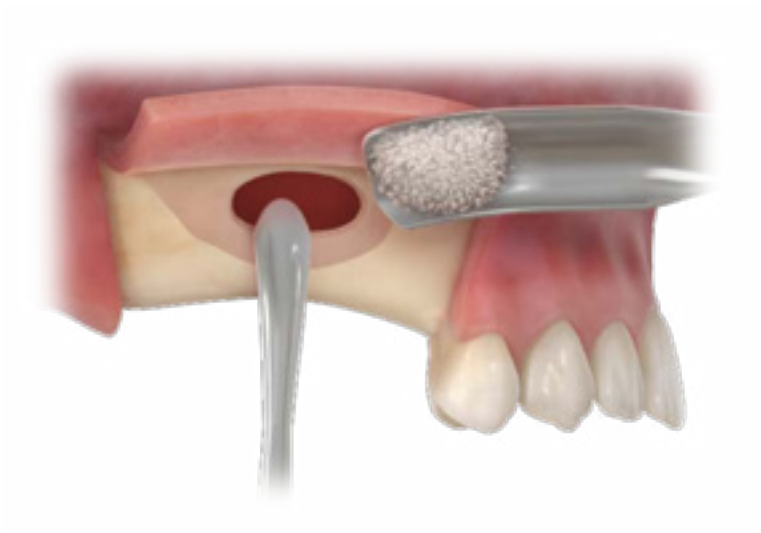


5 Bone graft carrier

A device used to transport autogenous bone or bone material through the entrance to the maxillary sinus.

User instructions

- Perform bone grafting in the maxillary sinus using a bone graft carrier and freer elevator.
- Fill forward and downward to prevent the bone graft from being pushed backwards.



How to take care of the KITS

1



Soak (saline/distilled water)

- Soak the surgical instruments in saline or distilled water

2



Drying (remove moisture)

- Completely dry all drills, drivers, tools, etc by using a towel or fan.

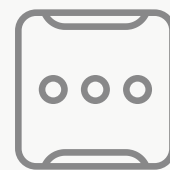
3



First wash

- After surgery, immediately separate and wash all the used instruments.

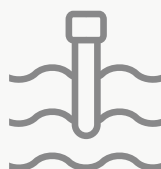
4



Organize instruments in the KIT

- Place the completely dried instruments in the KIT
- Make sure they are properly placed in the correct location
- Refer to the color coding for reference

5



Second wash

- Thoroughly wash with distilled water or running water to avoid remnants of blood or foreign debris.

6



Sterilization and storage at room temperature

- Wrap clean kit in a sterilization wrap or pouch and place into sterilizer.
- Sterilize temperature - 121°C to 132°C, time duration 15 - 30 minutes, dried and stored at room temperature.
- KIT re-sterilization is recommended immediately before surgery.
- Before and after sterilization, thoroughly dry (the drills will corrode if not fully dried after sterilization)

Important Information and Legal Notices 2026.03 ver.1.1

1. IMPORTANT NOTICE

This catalogue is intended solely as an informational and educational guide for trained dental professionals. It does not replace the applicable Instructions for Use (IFU), product labelling, formal clinical training, treatment planning, or independent professional judgment.

All clinical protocols, drilling sequences, cleaning instructions, sterilization requirements, torque recommendations, indications, contraindications, warnings, and procedural steps must be verified against the current product-specific IFU and the applicable product label for the relevant REF/product code prior to use.

In the event of any discrepancy between this catalogue and the applicable IFU, product labelling, or other official Osstem documentation, the IFU, labelling, and official product documentation shall prevail.

2. PRODUCT INFORMATION, CHANGES, AND AVAILABILITY

All products, specifications, protocols, recommendations, illustrations, and other information contained in this catalogue are subject to change without prior notice.

Not all products may be approved, cleared, released, licensed, or otherwise available in all markets. Product availability, indications, and regulatory status may vary by country. For information on the current product portfolio, approved indications, and local availability, please contact your local Osstem representative or Customer Service and consult the current official Osstem documentation.

3. PROFESSIONAL USE ONLY

Osstem Implant products are intended for use by appropriately trained dental professionals only. Dental implant treatment involves complex professional procedures and requires appropriate education, clinical training, patient selection, treatment planning, and radiographic as well as clinical evaluation.

The suitability of any procedure must be assessed individually for each patient, taking into account anatomy, bone quality and quantity, occlusion, systemic conditions, oral hygiene, compliance, and any other relevant clinical factors.

4. PRODUCT DESCRIPTION AND COMPATIBILITY

Osstem Implant offers implant fixtures, prosthetic components, surgical instruments, and related materials for dental implant treatment. Product codes, specifications, lot numbers, dates of manufacture, and expiration dates, where applicable, must be checked on the product label before use.

Unless expressly stated otherwise in the applicable product documentation, Osstem Implant abutments, prosthetic components, instruments, and related accessories are intended to be used only with compatible Osstem Implant fixtures and components. Use in combination with components or instruments from other manufacturers may result in improper fit, incomplete locking, loosening, fracture, reduced performance, or other clinical complications.

5. STERILITY, CLEANING, REPROCESSING, AND STORAGE

Sterile products supplied in sterile packaging must be used only if the packaging is intact and the expiration date has not passed. If sterile packaging has been opened, damaged, or has expired, the product must not be used.

Single-use products must not be reused, reprocessed, or resterilized.

Reusable instruments must be cleaned, disinfected, inspected, maintained, and sterilized strictly in accordance with the applicable Osstem IFU before reuse.

Products must be stored in accordance with the applicable labelled

storage conditions and protected from moisture, contamination, direct sunlight, and other adverse environmental conditions.

6. CLINICAL PROTOCOLS AND PROCEDURAL GUIDANCE

Any surgical, prosthetic, drilling, insertion, loading, cleaning, maintenance, or other procedural guidance shown in this catalogue is provided for general informational purposes only and must be adapted to the individual patient, the specific product, and the current approved IFU.

Clinicians remain solely responsible for selecting the appropriate treatment protocol and for determining whether the intended procedure, component selection, loading protocol, and clinical application are appropriate for the individual case and within the approved indications for the relevant product.

7. WARNINGS, CONTRAINDICATIONS, AND POSSIBLE COMPLICATIONS

Improper patient selection, inadequate treatment planning, non-compliance with the applicable IFU, improper use, off-label use, product modification, poor oral hygiene, infection, insufficient bone quality or quantity, excessive occlusal loading, or other unfavorable clinical conditions may result in complications or treatment failure.

Possible complications and adverse events may include, without limitation, implant instability or failure, loosening, fracture, bone loss, infection, soft- or hard-tissue complications, prosthetic complications, delayed healing, or the need for revision or removal.

Contraindications and precautions must always be assessed in accordance with the applicable Osstem product documentation and accepted professional standards of care.

8. INTENDED PURPOSE

The products are tools and instruments for surgical placement of Osstem implant fixtures. The drill is used to make implant sites. The cortical drill and tap removes cortical bones or forms threads on bone for the purpose of preventing excessive torque generated when implanting a fixture on hard bone. The drivers are for the placement of the fixture, and the prosthesis is used for setting. In addition, other instruments and tools will be used as aids in the Implant procedure.

The applicable product-specific IFU must always be consulted to confirm the intended purpose, indications, limitations, and approved clinical applications of the relevant product.

9. ACCURACY OF INFORMATION

Although reasonable care has been taken in preparing this catalogue, typographical, editorial, translation, printing, and formatting errors may occur. Information may also become outdated as a result of product updates, regulatory changes, technical revisions, or clinical developments.

No representation is made that this catalogue is complete, current, or error-free in every respect. Users must verify all critical information against the current IFU, product labels, and other official Osstem documentation before clinical use.

10. ILLUSTRATIONS AND EXAMPLES

Product illustrations, diagrams, radiographic examples, case images, and step-by-step demonstrations are for illustrative purposes only. Unless expressly stated otherwise, they are not shown to scale and do not guarantee any clinical outcome.

Example cases do not constitute a promise or representation of treatment success in any individual case.

11. TRADEMARKS AND COMPANY NAMES

All trademarks, trade names, product names, brand names, and company names are the property of their respective owners.

Important Contact Information



International HQ

Osstem Implant Co., Ltd.
3, Magokjungang 12-ro, Gangseo-gu
Seoul 07801, Republic of Korea

☎ +82-2-2016-7000

☎ +82-2-2016-7001

🌐 www.osstem.com



European HQ

Osstem Implant s.r.o.
Radlická 740/113c
158 00 Prague, Czech Republic

☎ +420-296-238-800

✉ info@osstem.eu

🌐 www.osstem.eu



Hiossen HQ

Hiossen Implant Inc.
85 Ben Fairless Drive, Fairless Hills
PA 19030, USA

☎ +1-888-678-0001

✉ master@hiossen.com

🌐 www.hiossen.com



Osstem Pharma HQ

Osstem Pharma Co., Ltd.
123, Gasan digital 2-ro, Geumcheon-gu
Seoul 08505, Republic of Korea

☎ +82-70-5118-0330

☎ +82-70-4369-1930

🌐 www.osstempharma.com



Osstem Orthodontic HQ

OSSTEM Orthodontics Inc.
13, Ojeongongeop-gil, Uiwang-si
Gyeonggi-do 16072, Republic of Korea

☎ +82-31-477-9902

☎ +82-31-477-9904

🌐 www.osstemortho.com

WE ARE GLOBAL

Global Leader in Implant Dentistry

Driven by advanced technology and exceptional product quality, we are exporting to more than 90 countries.



We are on Social Media



/osstem
/osstem-europe
/hiossen



/osstem
/osstem.eu
/hiossenimplants



/osstem
/osstem_europe
/hiossen



/@osstemglobal
/@osstem_europe
/@hiossenimplants



© Osstem Implant Co., LTD. All rights reserved.

Osstem Implant® and/or other trademarks and logos of Osstem Implant® mentioned herein are trademarks or registered trademarks of Osstem Implant Co., Ltd. and/or its affiliates. For dental professionals only. Informations in this document are provided for general information purposes and do not constitute medical advice. Specifications, content and availability are subject to change without notice. Errors and omissions excepted.