

USE INSTRUCTIONS



Figure 1. CMH Slim



Figure 2. CMH Standard



Figure 3. CMH Crestal Level

ATTENTION: The figures are merely illustrative. They do not represent the real dimensions.

This device is intended to a specialized procedure, which shall be made by professionals qualified in Dental Implants. To achieve optimized results, use the product knowing the proper techniques. Always apply them in proper conditions in a surgical environment.

DESCRIPTIVE INFORMATION

INDICATION OF USE

The DSP Implant System is intended to be surgically installed on the human upper and lower jawbone, serving as support for prosthetic devices, such as artificial teeth, with the purpose of restoring the masticatory function and the esthetics of the patient. The DSP Implant System may be used in one- or two-stage procedures, for unit or multiple restorations, enabling performance of immediate loading, when achieving a good primary stability and the proper occlusal loading.

DESCRIPTION OF THE DEVICE

The Morse Hexagonal Cone Implants (CMH) are dental implants made of commercially pure titanium (Grade 4), their external surfaces are treated with mechanical attack and chemical attack. The implants may be installed using a surgical motor or a torque wrench (manual).

They are available according to the table below:

IMPLANT	DIAMETER (mm)	HEIGHTS (mm)
CMH SLIM	3.3	8.5, 10, 11.5, 13, 15, 17
CMH Standard	3.75; 4.0; 5.0	8.5, 10, 11.5, 13, 15, 17
CMH Crestal Level	3.8; 4.0; 4.3; 5.0	7; 8.5; 10; 11.5; 13; 15; 17

WARNING

The non-recognition of the real lengths in relation to the radiographic measures may result in permanent injury to the nerves and other vital structures. The drilling beyond the depth intended to the surgery of the lower jaw may result in permanent numbness on the lower lip and chin or lead to bleeding on the lower part of the mouth.

Follow the mandatory procedures of any surgery, such as: asepsis during the bone drilling, avoid damages in blood vessels and nerves, using the pre-surgery anatomical and radiographic knowledge.

CONTRAINDICATIONS

This product is contraindicated for patients that present signs of allergy or hypersensitivity to the composition of the material: titanium.

In the presence of acute inflammatory or infectious processes, inadequate bone volume or quality, serious clinical problems, such as: disorders of the bone metabolism, disorders of blood coagulation, inadequate capacity of regeneration, insufficient oral hygiene, incomplete growth of the jaw, non-collaborative and non-motivated patient, undue use of drugs or alcohol, psychosis, extended functional disorders that resist to any treatment with medicines, xerostomia, weakened immune system, diseases requiring the use of steroids, endocrine diseases, pregnancy.

RISKS AND BENEFITS

As any surgery, there is no total guarantee of operation, because achieving a good performance involves several factors, being them usability, clinical conditions of the patient, and the product itself. Non-observance of the indicated limitations of use and work stages may result in failure.

The non-recognition of the real lengths of the risks in relation to the radiographic measures may result in permanent injury to the nerves and other vital structures. The drilling beyond the depth intended to the surgery of the lower jaw may result in permanent numbness on the lower lip and chin or lead to bleeding on the lower part of the mouth.

APPLICATION

The CMH implants are indicated for intraoral installation through surgical procedures in bones with density I, II, III, or IV, according to Lekholm & Zarb's jawbone quality classification (1985), they are used as temporary or definitive support, for unit or multiple restorations, including conventional protocols with immediate loading, provided that the required primary stability is achieved. The CMH implants may be installed immediately after the dental extraction.

IMPLANT	BONE DENSITY
CMH SLIM / CMH	I, II, III, IV
CMH CRESTAL LEVEL	III, IV

WARNINGS AND PRECAUTIONS

- Do not use the product if the package is violated.
- Do not use the product if the validity is expired.
- The material to be used during the procedure shall be sterile.
- This product shall be used immediately after the opening of the package, at the surgery time. If it is not used, dispose of it.
- This product is of single use and may not be re-sterilized.
- Reprocessing is Forbidden.
- The reuse of this product may cause adverse biological effects due to microorganisms and/or substances resulting of previous uses and/or reprocessing, being able to generate changes in the physical, mechanical, and chemical properties of the products, macro- and micro-structural, which may put the function desired at risk.
- The reuse of this product does not guarantee its safety and efficacy and exempts any guarantee of the product.
- Observe the conditions of the intraoral tissue, the bone quality, and the quantity of the bone bed, through radiographic exams and/or tomography. The absence of the pre-surgery assessment may compromise the success of the procedure.
- The inadequate surgical and/or prosthetic planning may compromise the performance of the implant/prosthesis set, resulting in failure in the system, such as loss or fracture of the implant, loosening, or fracture of prosthetic components and/or screws.
- The maximum installation torque suggested is 55 N.cm. The insertion torque higher than the recommended one may make the system inoperative.
- Before each procedure, certify that the parts are duly laid down.
- Certify that the parts are not swallowed or aspirated by the patient.
- Check the passivity and make the occlusal and interproximal adjustments after the installation of the prosthesis, avoiding the impairment of the implant/prosthesis set.
- Before each procedure, check the conditions of DSP Biomedical surgical instruments, always respecting their service life. Replace the instruments if there are damages, marks removed, sharpening compromised, deformation, or wear and tear.
- Always use the sequence of DSP Biomedical products, the use of prosthetic components and/or instruments of other manufacturers does not guarantee the perfect function of the DSP Implant System, and exempts any guarantee of the product.
- It is the professional's responsibility to use the DSP Biomedical products according to the instructions of use.

IMPORTANCE OF THE NEED OF ADHESION TO A CARE REGIME

The products shall be protected against aspiration when handled through intraoral way. The aspiration of products may lead to infection or physical injury not planned. If you want to protect it, use a rubber barrier. If an implant or an instrument is swallowed or aspirated, immediately call a doctor. In addition to the mandatory precautions to each surgery, such as asepsis, during the drilling in the jawbone, it shall be avoided damages to the inferior alveolar nerve and to the facial, deep facial, superior and inferior lip blood vessels. The anatomic knowledge and the pre-surgery medical images (for example, radiographies) shall be referred to.

The non-recognition of the real duration in relation to the radiographic measures may result in permanent injury of the nerves and other vital structures. The drilling beyond the depth intended to the surgery of the lower jaw may potentially result in permanent numbness on the inferior lip and chin or lead to bleeding on the floor of the mouth.

The inadequate use of the products lead to a work poorly performed, and increase of the risk. In particular, the users of manual tools shall take care of gently using them and with attention. The user shall always avoid touching on the instruments and pieces with no protection (sterile protective gloves and aprons shall be used). The thermal bone damages caused by rotary and oscillating tools shall always be avoided (user's training, work at low speed and with sufficient cooling. During the intraoral application, it shall pay attention to the fact that the products are protected against aspiration or dropping on the floor. The rotary instruments need to be fixed as further as possible with their speed set before the application. Do not exceed the recommended drilling speeds, since it may cause bone necrosis or fracture of components of the system. The inadequate cleaning and sterilization of the instruments may

result in the patient's infection with harmful bacterias. To avoid damaging the instruments, they shall be individually taken out of the blister package.

Do not use the device if the primary package had been damaged or previously opened.

Do not use damaged or forceful instruments for drilling. The broken land lips of the instruments cause vibrations and high pressure forces, which, on their turn, leads to broken preparation corners and rough surfaces. Instruments that are folded and/or do not work shall be immediately disposed of. Damaged, corroded, or worn devices shall not contact intact instruments to avoid contact corrosion.

OPERATION INSTRUCTIONS

DRILLING

Under abundant irrigation, make the drilling using drills in good cutting conditions and with proper rotation speed, as indicated in table:

IMPLANT	DRILLING ROTATION (rpm)
CMH SLIM / CMH STANDARD/ CMH CRESTAL LEVEL	600 - 1200

Select the sequence of drills according to the model of implant to be installed, according to the indications in the tables below:

SEQUENCE OF CMH SLIM DRILLS BONE TYPE I AND II		
IMPLANT DIAMETER (mm)	LANCE DRILL 2.0mm	CYLINDRICAL DRILL 2.8mm
3.3	●	●

● Indicated

SEQUENCE OF CMH SLIM DRILLS BONE TYPE III AND IV		
IMPLANT DIAMETER (mm)	LANCE DRILL 2.0mm	CYLINDRICAL DRILL 2.5mm
3.3	●	●

● Indicated

SEQUENCE OF CMH STANDARD DRILLS – BONE TYPE I, II, III, IV									
IMPLANT DIAMETER	Initial Drill Ø2.0	Pilot Drill Ø2.0/3.0	Cylindrical Drill Ø3.0	Pilot Drill Ø3.0/3.8	Cylindrical Drill Ø3.3	Cylindrical Drill Ø3.5	Cylindrical Drill Ø3.8	Cylindrical Drill Ø4.3	Countersink Drill 5.0
3.75	●	●	●		▲				
4.0	●	●	●		●	▲			
5.0	●	●	●	●	●		●	●	▲

● Indicated

▲ 3.5 mm perforation at corticalized height

SEQUENCE OF CHM CRESTAL LEVEL DRILLS – BONE TYPE III AND IV							
IMPLANT DIAMETER	INITIAL DRILL 2.0mm	DRILL 3.5mm	DRILL 3.8mm	DRILL 4.3 mm	DRILL 5.0mm	COUNTERSINK 4.1 mm	COUNTERSINK 5.0 mm
3.8	●	●				●	
4.0	●		●	●		●	
4.3	●		●	●			
5.0	●		●		●		●

● Indicated

The drilling depth of the drills, as well as their size, shall be in compliance with the model of Implant selected during the planning, considering measures of the Implant, installation level, three-dimension spacing, among other factors (short, regular, or long drills).

SEQUENCE OF IMPLANT HANDLING

1. The box of the Implant shall be manually opened, with no sterile gloves.
 2. Break the seal of the box and remove the Blister. Open the Blister and put the Pendriver on the sterile surgical field.
 3. Using sterile surgical gloves, hold the tube with the non-dominant hand, and remove the Badock (cap of the tube) with the dominant hand.
 4. For installation using a surgical motor, capture the implant with the proper connection driver. Take the implant to the surgical cavity. In the surgical motor, use maximum torque of 35 Ncm and insertion rotation of 20-30 rpm.
 5. Complete the installation of the implant with the torque wrench. The maximum installation torque suggested is 55 N.cm.
- The indication of application of loads in relation to the torque is described in the table as follows:

LOAD APPLICATION	MIN. TORQUE (N.cm)	MAX. TORQUE (N.cm)
Delayed Loading*	10	55
Immediate Loading	35	55

*Associated to the use of Healer.

When the installation torque obtained is lower than 10 N.cm, it is recommended the use of the Cover Screw.

PROSTHETIC

To use the CMI SLIM in two-stage procedures, the prior preparation of the soft tissues may be made using a compatible Protection Cylinder.

For the procedure of modeling the head in a pillar form of the implant.

1. Fit the corresponding transfer, assure the proper fitting, and make the molding with proper materials.
2. Prepare the cast model.
3. Prepare the prosthesis using the corresponding coping. (Temporary metal coping, calcinable coping, definitive coping). It may be cemented or screwed, or use Abutment Oring, according to the proper laboratory techniques.
4. The tests shall be made on the passivity and the adjustment of the structure of the prosthesis.
5. Cement or screw the final prosthesis on the implant head, use its indexer, and check the perfect fitting between prosthesis and implant.

TRACEABILITY LABEL

This product is accompanied by three labels that allow its traceability and shall be attached to the following documents: • patient's records; • prosthetic records; • Document to be delivered to the patient.

The identification and traceability are carried out through the REF and LOT number codes.

PRESENTATION AND STERILIZATION

This product is indicated for single use and is provided sterile by gamma radiation, packed unit by unit in packages that offer quadruple protection: clear tube, capsule, blister, and box.

MAGNETIC RESSONANCE (MR) - SAFETY INFORMATION

The DSP Implant System was not assessed for safety and compatibility in the MR environment. Tests were not carried out regarding factors of heating, migration, or image artifact in the MR environment.

The safety of the DSP Implant System in the MR environment is unknown, submit a patient that has this device to magnetic resonance may result in injury to the patient.

STORAGE INSTRUCTIONS

This product shall be stored in its original package, in clean and ventilated place, at maximum temperature of 45°C, and protected against direct sunlight.

FURTHER INFORMATION

Instruct the Patient regarding the need of professional medical follow-up after the surgery, and follow the guidelines relative to the precautions, hygiene, and prescription of medicines. Such guidelines are responsibility of the professional in charge.

DISPOSAL OF MATERIALS

Every product and consumable used during the surgery for installation of dental implants may put at risk the health of those who handle them, after the use. Before disposing of them in the environment, it is recommended to observe the effective legislation and adhere thereto.

SERVICE LIFE

This product is of single use; it may not be reused.

EXPIRATION DATE

See package.

ADVERSE EFFECTS

The installation of dental implants, as well as any other surgical procedure, may cause a slight discomfort and localized edema. More persistent symptoms may occur, such as: chronic pain related to the dental implant, permanent paresthesia, dysesthesia, maxillary/mandibular bone reabsorption, localized systemic infection, oroantral or oronasal fistula, adjacent teeth unfavorably affected, irreversible damages to the adjacent teeth, fracture of the implant, jaw, bone, or prosthesis, aesthetic problems, injury to the nerves, exfoliation, hyperplasia.

Failure in the osseointegration and loss of the prosthesis during the treatment may be caused by:

Inadequate osteotomy, infections, diseases, or systemic problems, low quality or insufficient volume of bone, absence or failure of irrigation, use of instruments and/or non-specific instruments with no power of cutting, poor oral hygiene, occlusal trauma, lack of prosthetic passivity, and lack of specific training.

PRODUCT GUARANTEE



























DSP Biomedical assures to the owner of this product guarantee against any material or manufacturing defect, the presence of any defect shall be immediately informed to the manufacturer, respecting the legal term. The guarantee of the products manufactured by DSP Biomedical is strictly connected to following the information described in the instruction of use. The inadequate use of the product disregarding the indications releases the manufacturer and/or vendor of any responsibility.

Note: the guarantee does not cover wear and tear from using the product.

INFORMATION OF TECHNICAL ASSISTANCE

If there is need of further information, or the product presents any adverse effect, with potential of risk to the patient, which generates or has potential of injury or threat to public health, or any dissatisfaction of the client, contact DSP through the phone numbers 0800 600 88 66, or send an e-mail to sac@dspbiomedical.com.br.

SYMBOLS

SYMBOL	DESCRIPTION	SYMBOL	DESCRIPTION
	Batch number		Consult instructions for use or consult electronic instructions for use
	Date of manufacture		Do not resterilize
	Manufactured by		Keep dry
	Sterilized using irradiation		Keep away from sunlight
	Product Code		Single sterile barrier system with protective packaging inside
	Model Number		Used by-date
	Do not reuse		Unique Device Identifier
	Limit of temperature		Country of manufacturer
	Authorized representative in the European Community		Do not use if package is damaged and consult instructions for use
	Caution		Humidity limitation
	Importer		Medical device
	Fragile, handle with care		Mandatory medical prescription Notification required by FDA for United States market
	CE Mark		CE marking with number of Notified body; SIQ, number 1304

The SSCP is available in the European database on medical devices (Eudamed), where it is linked to the Basic UDI. Please see Eudamed public website: <https://ec.europa.eu/tools/eudamed>. BASIC UDI: 7908467800460CMHIMPLANTV2

REF : Products

Device Name(s)	Code	Device Name(s)	Code
CMH SLIM IMPLANT Ø 3,3 X 7,0	60.3307	CMH IMPLANT Ø 3,8 X 7,0	60.3807C
CMH SLIM IMPLANT Ø 3,3 X 8,5	60.3308	CMH IMPLANT Ø 3,8 X 8,5	60.3808C
CMH SLIM IMPLANT Ø 3,3 X 10,0	60.3310	CMH IMPLANT Ø 3,8 X 10,0	60.3810C
CMH SLIM IMPLANT Ø 3,3 X 11,5	60.3311	CMH IMPLANT Ø 3,8 X 11,5	60.3811C
CMH SLIM IMPLANT Ø 3,3 X 13,0	60.3313	CMH IMPLANT Ø 3,8 X 13,0	60.3813C
CMH SLIM IMPLANT Ø 3,3 X 15,0	60.3315	CMH IMPLANT Ø 3,8 X 15,0	60.3815C

CMH SLIM IMPLANT Ø 3,3 X 17,0	60.3317	CMH IMPLANT Ø 3,8 X 17,0	60.3817C
CMH IMPLANT Ø 3,75 X 7,0	60.3707	CMH IMPLANT Ø 4,0 X 7,0	60.4007C
CMH IMPLANT Ø 3,75 X 8,5	60.3708	CMH IMPLANT Ø 4,0 X 8,5	60.4008C
CMH IMPLANT Ø 3,75 X 10,0	60.3710	CMH IMPLANT Ø 4,0 X 10,0	60.4010C
CMH IMPLANT Ø 3,75 X 11,5	60.3711	CMH IMPLANT Ø 4,0 X 11,5	60.4011C
CMH IMPLANT Ø 3,75 X 13,0	60.3713	CMH IMPLANT Ø 4,0 X 13,0	60.4013C
CMH IMPLANT Ø 3,75 X 17,0	60.3717	CMH IMPLANT Ø 4,0 X 15,0	60.4015C
CMH IMPLANT Ø 4,0 X 7,0	60.4007	CMH IMPLANT Ø 4,0 X 17,0	60.4017C
CMH IMPLANT Ø 4,0 X 8,5	60.4008	CMH IMPLANT Ø 4,3 X 7,0	60.4307C
CMH IMPLANT Ø 4,0 X 10,0	60.4010	CMH IMPLANT Ø 4,3 X 8,5	60.4308C
CMH IMPLANT Ø 4,0 X 11,5	60.4011	CMH IMPLANT Ø 4,3 X 10,0	60.4310C
CMH IMPLANT Ø 4,0 X 13,0	60.4013	CMH IMPLANT Ø 4,3 X 11,5	60.4311C
CMH IMPLANT Ø 4,0 X 15,0	60.4015	CMH IMPLANT Ø 4,3 X 13,0	60.4313C
CMH IMPLANT Ø 4,0 X 17,0	60.4017	CMH IMPLANT Ø 4,3 X 15,0	60.4315C
CMH IMPLANT Ø 4,3 X 7,0	60.4307	CMH IMPLANT Ø 4,3 X 17,0	60.4317C
CMH IMPLANT Ø 4,3 X 8,5	60.4308	CMH IMPLANT Ø 5,0 X 7,0	60.5007C
CMH IMPLANT Ø 4,3 X 10	60.4310	CMH IMPLANT Ø 5,0 X 8,5	60.5008C
CMH IMPLANT Ø 4,3 X 11,5	60.4311	CMH IMPLANT Ø 5,0 X 10,0	60.5010C
CMH IMPLANT Ø 4,3 X 13	60.4313	CMH IMPLANT Ø 5,0 X 11,5	60.5011C
CMH IMPLANT Ø 4,3 X 15	60.4315	CMH IMPLANT Ø 5,0 X 13,0	60.5013C
CMH IMPLANT Ø 4,3 X 17	60.4317	CMH IMPLANT Ø 5,0 X 15,0	60.5015C
CMH IMPLANT Ø 5,0 X 7,0	60.5007	CMH IMPLANT Ø 5,0 X 17,0	60.5017C
CMH IMPLANT Ø 5,0 X 8,5	60.5008		
CMH IMPLANT Ø 5,0 X 10,0	60.5010		
CMH IMPLANT Ø 5,0 X 11,5	60.5011		
CMH IMPLANT Ø 5,0 X 13,0	60.5013		
CMH IMPLANT Ø 5,0 X 15,0	60.5015		
CMH IMPLANT Ø 5,0 X 17,0	60.5017		

MANUFACTURED BY

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Reference

1-Lekholm U, Zarb G. Patient selection and preparations. Branemark, PI, Zarb,G & Albrektsson, T, eds Tissue-Integrated Prostheses: Osseointegration in Clinical Dentistry. Chigado: Quintessence; 1985. p. 233-40.