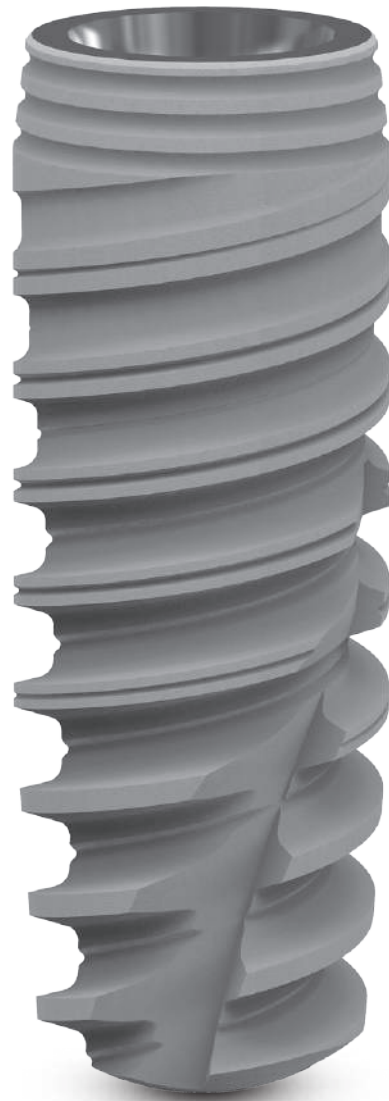


GALAXY

Conical connection implants



GALAXY

Conical connection implants



Important information

Please read carefully before using Ziacom® products

General information

This document contains basic information on the use of original Ziacom® dental implant systems, hereafter referred to as Ziacom® dental implants or simply Ziacom® products. This document has been created as quick guide for clinicians responsible for treatment, hereafter the "user", and, therefore, is neither an alternative nor a substitute for specialized training or professional clinical experience.

Ziacom® products must be used according to a suitable treatment plan and adhering strictly to the surgical and prosthetic protocols established by the manufacturer. Read the product-specific surgical and prosthetic protocols as well as the instructions for use and maintenance before using each Ziacom® product. You can find this information on our website, www.ziacom.com, or request it from your nearest authorised Ziacom® distributor.

Liability, safety and guarantee.

The instructions for the use and handling of Ziacom® products are based on internationally published literature, current clinical standards and our clinical experience, so they should be understood as general guiding information. The handling and use of Ziacom® products is the sole responsibility of the user as it is outside the control of Ziacom Medical SL. Ziacom Medical SL, their affiliates and/or their authorised distributors disclaim all responsibility, whether explicit or implicit, total or partial, for possible damage or injury caused by poor handling of the product or any other situation not considered in their protocols and manuals for the correct use of their products.

The user must ensure that the Ziacom® product is appropriate for the intended procedure and end purpose. Neither these instructions for use nor the work or handling protocols for the products release the user from this obligation. Ziacom® products must be used, handled and applied by professionals with the appropriate training and qualifications required according to current legislation in each country.

The total or partial use, handling and/or application of Ziacom® products at any stage of their implementation by personnel who are unqualified or lack the necessary training will automatically void any type of warranty and may cause severe damage to the patient's health.

Ziacom® products are part of their own system, with their own design characteristics and work protocols, including dental implants, abutments or prosthetic components and surgical or prosthetic instruments. The use of Ziacom® products in combination with elements or components from other manufacturers could result in treatment failure, damage to tissues or bone structures, inadequate aesthetic outcomes and severe damage to the patient's health. Therefore, only original Ziacom® products should be used.

The clinician in charge of the treatment is solely responsible for ensuring the use of original Ziacom® products and that they are used according to the corresponding instructions for use and handling protocols throughout the implant procedure. The use of any other non-original Ziacom® components, instruments or products, whether alone or in combination with any original Ziacom® products, will immediately void the warranty of the original Ziacom® products.

See the Ziacom Medical SL Warranty Programme (available on the website or by contacting Ziacom Medical SL, their affiliates or authorised distributors).

Warning. Not all Ziacom® products are available in all countries. Check availability in your country.

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Ziacom Medical S.L. reserves the right to modify, change, remove or update any of the products, prices or technical specifications referenced on this website or in any of its documents without prior notification. All rights reserved. The reproduction of this document, whole or in part and in any medium or format, without the corresponding written authorisation from Ziacom Medical SL is prohibited.





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The Company

Together for health

Ziacom® has been working for more than 20 years to improve the **oral health** and well-being of patients around the world by **designing and manufacturing innovative**, high-quality dental implant, prosthetic component, surgical instrument and biomaterial solutions.

The company was founded in 2004 with **100% Spanish capital** and began its activity as a manufacturer of dental implants and attachments for several European companies before launching its own **brand of implant systems** in 2006.

In 2015, Ziacom® introduced its **diversification strategy** with the development of **new business lines** and new product lines and the launch of a **new portfolio**, which helped the company achieve a **15% share of the Spanish market** in 2016 with the sale of more than 230.000 implants.

In 2022, the company started up on an **ambitious growth plan** with new goals of **international expansion**, broadening and **diversification** of its portfolio of **products and services** and a Corporate Identity restyle.

Ziacom® quality

Commitment to **quality and innovation** has been part of the values and the essence of Ziacom® since the beginning.

The reason why we used state-of-the-art technology in **every stage of our products' production cycle**, from **design and manufacture to quality assurance, cleaning and packaging**. All of our products are also manufactured using only **high-quality raw materials** after applying **strict controls to select** our main suppliers.

Ziacom Medical SL is a **licensed manufacturer of medical devices** and an AEMPS (Spanish Agency for Medicines and Medical Devices) 6425-PS **marketing authorisation holder**. Our **quality management**

system is certified in accordance with the requirements of ISO standards 9001:2015 and 13485:2018, and is also GMP 21 CFR 820 compliant.



Thanks to our ceaseless endeavours to offer our clients an unsurpassable quality, all our implants have a **lifetime guarantee**.

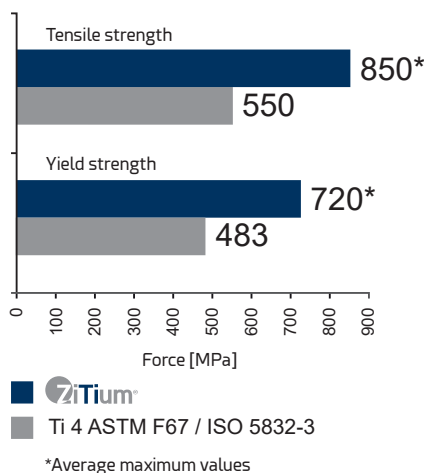
See the General Conditions for Accessing the Guarantee for Ziacom® products.

Zitium® titanium

Ziacom® **Galaxy** implants are manufactured using **extra-high-strength grade 4 Zitium® titanium** which gives them **considerably improved yield strength and mechanical properties**.

Thanks to **Zitium® titanium**, our implants meet the requirements of ASTM F67 and ISO 5832-3 and are certified in accordance with Council Directive 93/42/EEC and its amendment Directive 2007/47/EC by notified body 0051.

Properties of Zitium® titanium



FDA Approved*

*See approved models

Ziacom® implants are all sterilised using beta ray radiation at 25 kGy, apart from the DSQ orthodontic implants, which are supplied **unsterilised**.

IMPORTANT

All the products (except dental implants) listed in this Ziacom® catalogue are supplied unsterilised and must be sterilised before use.



Investment in innovation and training

In order to always offer the very best solutions for the **well-being of every patient**, and thanks to the experience and dedication of our **highly-qualified professionals** and **innovative Technological Centre**, our R&D&I team works incessantly in the field of **research and innovation** to **improve** our products and develop **new solutions** to meet the demands and needs of both patients and dentists.

We also invest in **research** and **ongoing training** as a way of providing **scientific support to the sector** and we firmly believe in training **young professionals** to ensure the best **advances in dentistry field**.

We therefore work closely with **training centres, universities and scientific bodies** to create a practical and specialised teaching environment to promote and strengthen their knowledge, abilities and professional growth.

In order to enhance our investment in the training and **development of dental professionals**, we have **specific areas at our facilities** for **hands-on training and practicals, state-of-the-art** training equipment and also a **physical and virtual showroom** where professionals can see all our dental solutions first hand.

Ziacom® around the world

We are committed to making oral health available to patients all over the world and have a solid **internal growth and expansion plan** to increase the company's **international presence** in those **areas where we our products are already available** and to add **new growth areas**.

In order to achieve this, we offer our **international associates** a **trusting and collaborative** partnership by adapting to their **local needs** and providing solutions that are specific to each market.

As part of our commitment to meet the specific **quality, regulatory and legal requirements of each country**, for both the registration and distribution of our products, we have **specific certifications** from each of the countries in which we trade.

Regional headquarter

Ziacom Medical SL

Madrid - ESPAÑA
Calle Búhos, 2 - 28320 Pinto
Tel: +34 91 723 33 06
info@ziacom.com

Subsidiaries

Ziacom Medical Portugal Lda

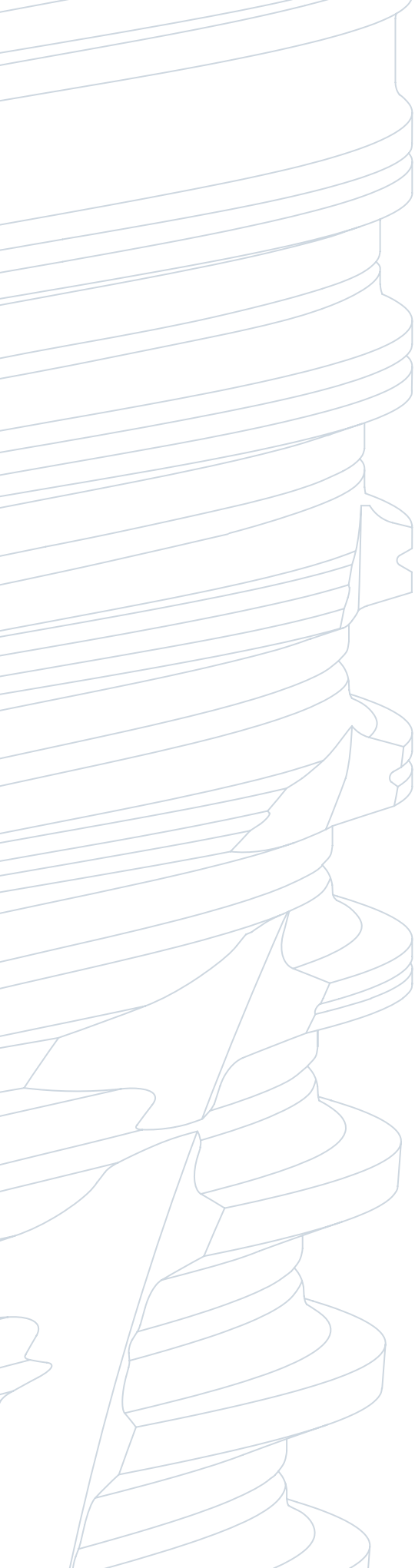
Av. Miguel Bombarda, 36 - 5º B
1050 -165 - Lisboa - PORTUGAL
Tel: +351 215 850 209
info.pt@ziacom.com

Ziacom Medical USA LLC

Miami - EEUU
333 S.E 2nd Avenue, Suite 2000
Miami, FL 33131 - USA
Phone: +1(786) 224 - 0089
info.usa@ziacom.com

Please see the up-to-date list of Ziacom® distributors at www.ziacom.com or email us at export@ziacom.com

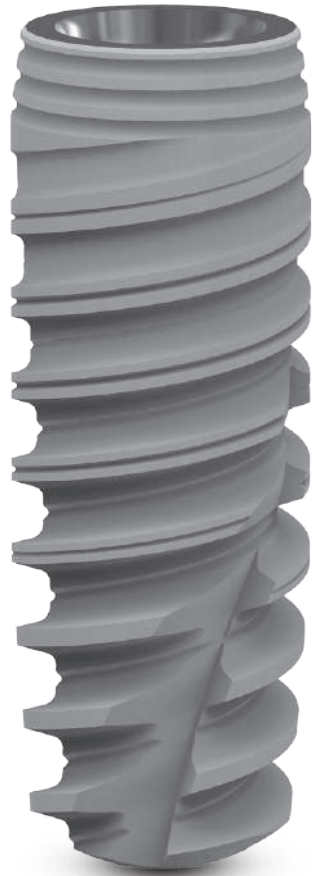




GALAXY

GALAXY

Conical connection implants



Characteristics

CONNECTION

- 11° conical connection with double internal hex.
- Single platform for all diameters.
- Platform switch.

CORTICAL ZONE

- Microrings.
- Inverted cone cortical macro-design.

CONICAL BODY

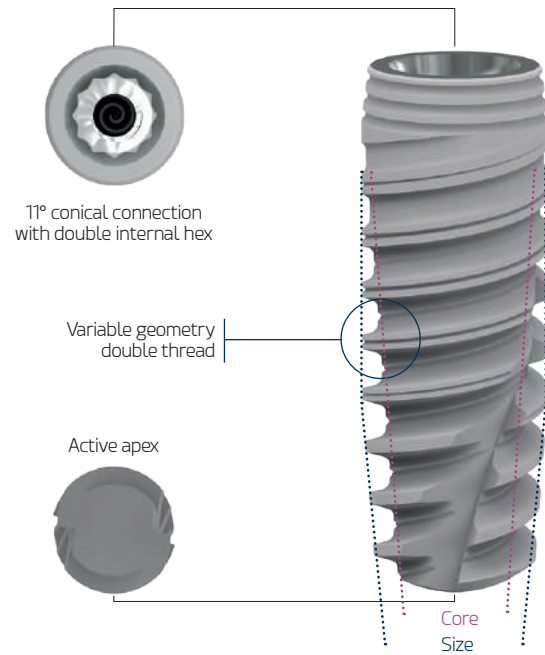
- Double threaded.
- Variable geometry:
 - » Coronal - thick trapezoidal thread.
 - » Middle - thinner trapezoidal thread.
 - » Apex - V-shaped thread.

APEX

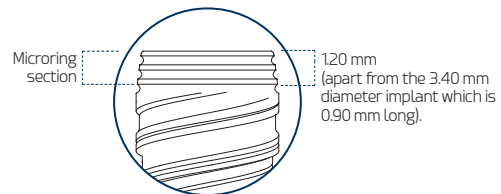
- Oblique apical windows.
- Self-tapping active apex.
- Atraumatic rounded apex.

INDICATIONS

- Bones of very poor quality.
- Immediate loading.
- Immediate postextraction implant placement.
- Aesthetic anterior segment.






































Dimensions of the implant's coronal section



Advantages

- Better sealing against leaks which means less bacterial load.
- Better distribution of forces directed towards the implant and not the connection.
- If the recommended torques are exceeded, the screw suffers the fracture, not the implant.
- Greater preservation of the crestal bone.
- Lower incidence of peri-implantitis.
- Better survival rate of conical connection implants.
- The conical connection prevents micromovement and microfiltration at the implant–abutment interface.
- The single platform provides a significant simplification of prosthetic procedures.
- The reverse taper neck mitigates cortical stress during surgery.
- The thread design confers a very high primary stability even in poor quality bone.
- The active apex facilitates insertion axis correction in postextraction alveoli.

Diameters and lengths

Ø DIAMETER	Ø PLATFORM	LENGTH (L)						
		6	7	8.5	10	11.5	13	14.5
 RP 3.40	2.85							
 RP 3.70								
 RP 4.00								
 RP 4.30								
 RP 4.80								

Dimensions in mm.

Surface treatments

■ Titansure surface

Implants inserted following surface treatment are known to benefit from improved osseointegration by increasing the bone-to-implant contact area. This is partly due to the implant's chemical composition and topographical characteristics.

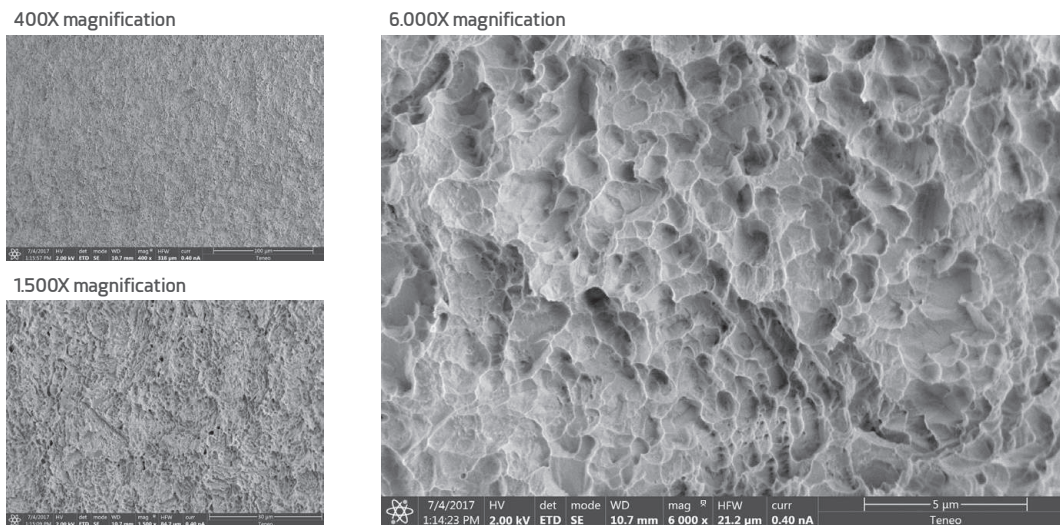
With our **Titansure** surface treatment, at Ziacom® we have obtained a contaminant-free surface topography and optimal average macro- and microporosity values, which are key specifications for achieving prompt and proper osseointegration and, in turn, extremely reliable and predictable implants.

■ TITANSURE SURFACE ANALYSIS

Titansure is an SLA surface treatment created through a subtraction process involving sandblasting with white aluminium oxide and double acid etching with hydrofluoric acid and a sulphuric/phosphoric acid mix.

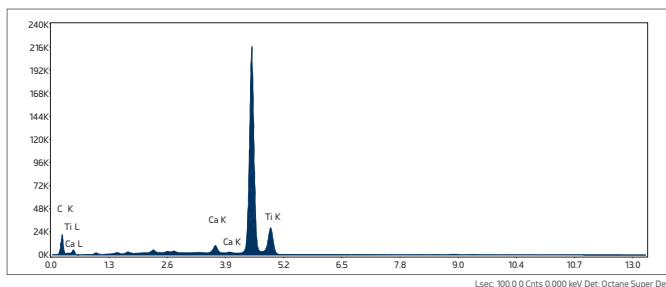
Surface morphology analysis

With the aid of a scanning electron microscope (FEI TENEO, Thermo Fisher Scientific Inc., Waltham, MA, USA), we can see the rough, porous surface creating numerous cavities with thin, sharp edges.



Surface elemental analysis

We used an energy-dispersive X-ray spectrometer (Octane Super, Edax-Ametek, Mahwah, NJ, USA) to analyse the chemical composition at the surface.



Compositional analysis of implant surface

ELEMENT	WEIGHT (%)
C K	9.32 (10.23)
Al K	-
Ti K	89.53 (11.77)

No aluminum was detected

Results are expressed as the mean and standard deviation of the mass percentage (WEIGHT %).

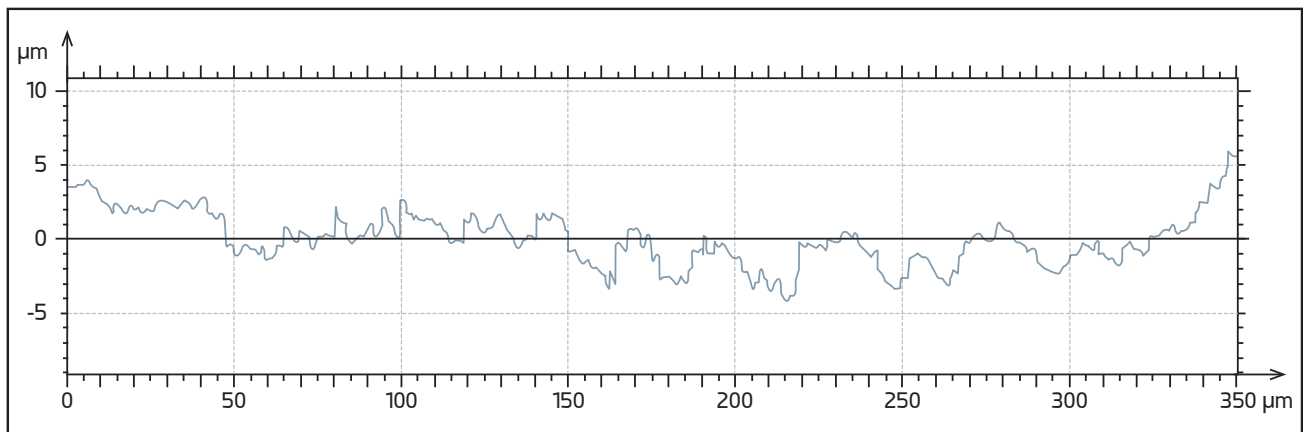
Surface roughness analysis

The roughness study was conducted with a Sensofar S NEOX interferometric-confocal microscope (Sensofar Medical, Terrasa, Spain) and SensoMAP Premium 7.4 software. The quantitative roughness profile parameters applied were: average roughness (Ra), root-mean-square roughness (Rq), maximum profile peak height roughness (Rp) and maximum profile valley depth roughness (Rv).

Ra (µm) (SD)	Rq (µm) (SD)	Rp (µm) (SD)	Rv (µm) (SD)
0.82 (0.10)	0.97 (0.08)	1.84 (0.04)	2.21 (0.01)

The 3D surface roughness (Sa), 3D root mean square height (Sq), maximum 3D peak height (Sp) and maximum 3D pit depth of the selected area (Sv) were also recorded.

Sa (µm) (SD)	Sq (µm) (SD)	Sp (µm) (SD)	Sv (µm) (SD)
0.76 (0.01)	0.97 (0.01)	4.20 (0.12)	4.62 (0.20)



The data were extracted from:

Rizo-Gorrita, M.; Fernandez-Asian, I.; Garcia-de-Frenza, A.; Vazquez-Pachon, C.; Serrera-Figallo, M.; Torres-Lagares, D.; Gutierrez-Perez, J. Influence of Three Dental Implant Surfaces on Cell Viability and Bone Behavior. An In Vitro and a Histometric Study in a Rabbit Model. Appl. Sci. 2020. 10(14), 4790

■ OPTIMAL OSSEOINTEGRATION

The **Titansure** surface has a three-dimensional surface structure with high peaks and broad troughs, which is known to be highly effective at promoting the coagulation cascade and the release of growth factors through platelet activation [Kim, H.; Choi, S.H.; Ryu, J.J.; Koh, S.Y.; Park, J.H.; Lee, I.S. The biocompatibility of SLA-treated titanium implants. Biomed. Mater. 2008. 3. 025011].

This type of surface may have an osteogenic effect thanks to its different topographical features at a micrometer and nanometer level, which has a very similar morphology to the osteoclastic bone resorption cavities [Le Guehennec, L.; Goyenvalle, E.; Lopez-Heredia, M.A.; Weiss, P.; Amouriq, Y.; Layrolle, P. Histomorphometric analysis of the osseointegration of four different implant surfaces in the femoral epiphyses of rabbits. Clin. Oral Implants Res. 2008. 19. 1103–1110].

For more information on the surface treatment see the literature available at www.ziacom.com/biblioteca





Surface treatments

■ Titansure Active surface treatment

Ziacom® presents the **Titansure Active** surface treatment with bone bioactive liquid (BBL) as the latest innovation for the presentation of our dental implants. The **Titansure Active** surface treatment is a combination of **Titansure** with BBL technology (Bone Bioactive Liquid), a patent acquired by Ziacom® and developed by the Biointelligence Systems research group led by Professor Maher Al-Atari Abou-Asi.

"BBL technology consists of a saline solution containing calcium chloride (CaCl₂) and magnesium chloride (MgCl₂·6H₂O) with a net negative charge and creates the ideal conditions for post-implant cell adhesion in the region with bone damage. What is more, surface treatment with BBL provides a significant increase in the density of hydroxyl groups on the surface of implants, thus improving their hydration considerably compared with other surfaces. This hydrophilic implant surface is precisely what enables active ion interaction with blood plasma and bone-forming cells long before the first stem cells can attach to the surface. Finally, this yields improved intercellular communication and a greater final bone-to-implant contact area in a significantly shorter time, thereby markedly reducing the postoperative inflammatory process."

Dr. Prof. Maher Al Atari

■ SURFACE STUDIES OF BBL-TREATED IMPLANTS

In vitro research

Dental pulp pluripotent-like stem cell (DPPSC) and dental pulp mesenchymal stem cell (DPMSC) cultures were prepared on titanium discs sandblasted with aluminium oxide and acid etched in an osteoblast differentiation medium.

The samples were divided into two treatment groups:

- **Group A.** Titanium discs - Traditional, untreated surface.
- **Group B.** Titanium discs - BBL-treated surface.

The surfaces were examined using energy-dispersive X-ray microanalysis (EDXMA) to determine the composition of surface elements.

Comparison of different elements in the two groups		
	Untreated surface	Treated surface Titansure Active
Carbon	32.22 ± 5.89	32.89 ± 1.76
Oxygen	14.34 ± 1.23	13.97 ± 1.45
Phosphorus	3.96 ± 2.8	3.89 ± 1.87
Calcium	5.86 ± 3.8	9.53 ± 4.04
Titanium	39.76 ± 1.65	41.34 ± 1.89
Ca/P	1.678	2.347

In vivo research

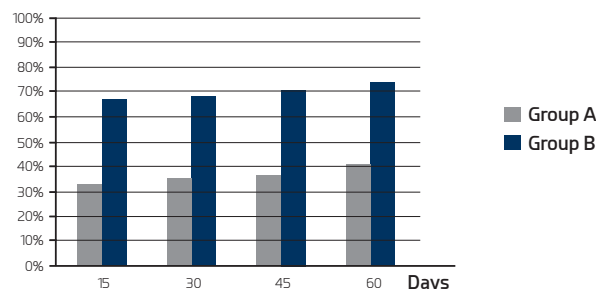
A study was conducted in the tibiae of 10 adult New Zealand rabbits after inserting four implants per rabbit (two in each tibia).

The subjects were assigned to two treatment groups with implants:

- **Group A.** Implants with a traditional, untreated surface.
- **Group B.** Implants with a traditional, BBL-treated surface.

In general, group B had higher BIC (bone-to-implant contact) values than group A.

Histomorphometric analysis - Bone-to-implant contact (BIC)		
Time of measurement	Group A Untreated surface (Control) mean + SD	Group B Treated surface Titansure Active mean + SD
15 days	33.7 ± 2.3%	68.92 ± 0.3%
30 days	35.8 ± 1.8%	69.35 ± 2.2%
45 days	37.9 ± 1.2%	70.34 ± 1.1%
60 days	41.2 ± 0.8%	73.89 ± 1.9%

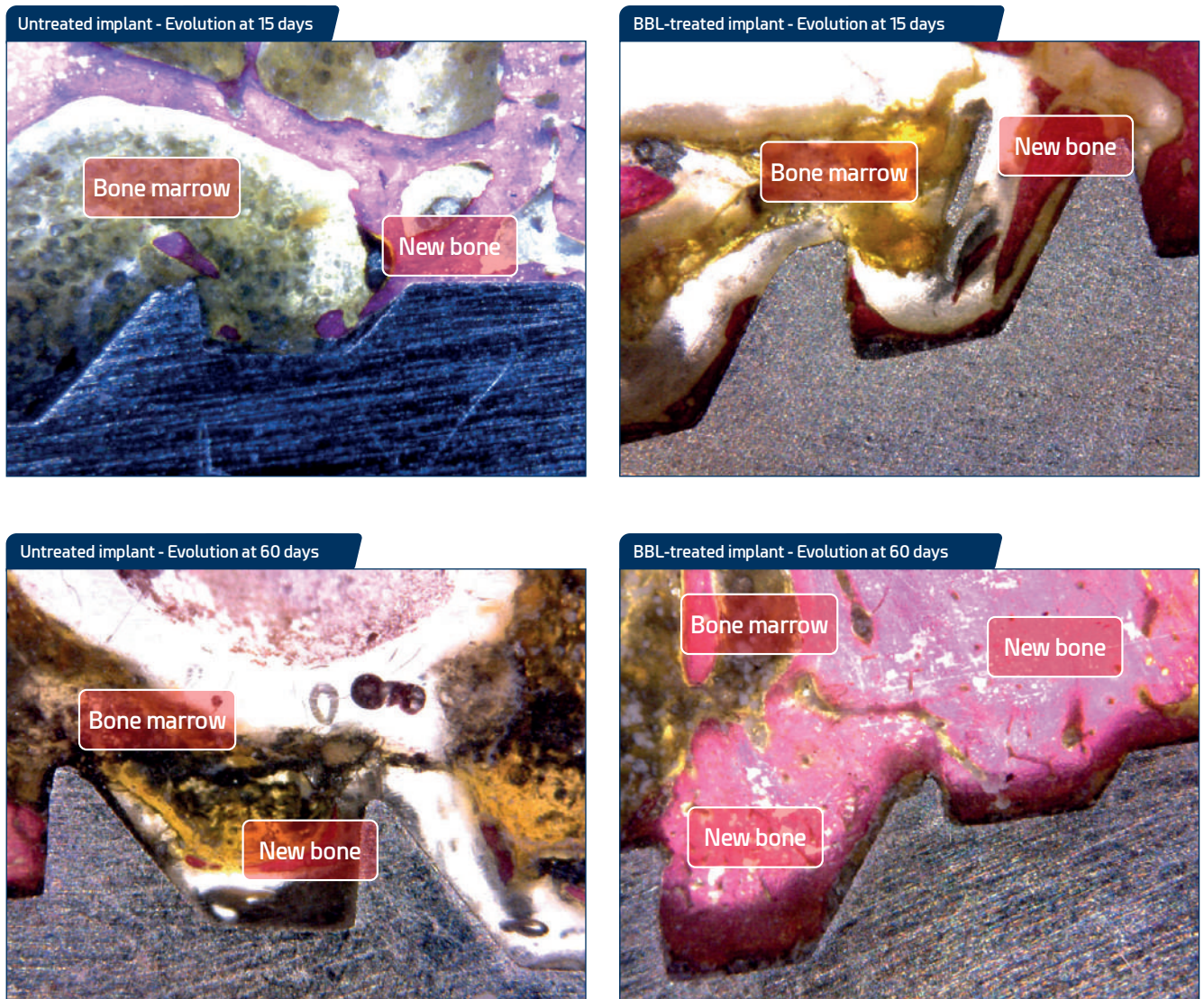


Conclusions

Within the scope of this study, the histomorphometric analysis demonstrated that the group B implants achieved quicker and more effective osseointegration than control group A. Nevertheless, an assessment of bone growth in the medullary portion of the subjects' tibiae revealed the new surface's potential for osteoinduction.

As explained by Dr. Sérgio Alexandre Gehrke, the histologist in charge of the study: "Within the study's limits, data from the histomorphometric analysis of the implants with a BBL-treated surface (78.92 + 0.3%) highlighted a much quicker and more effective osseointegration compared to the control group (53.8 + 2.3% of BIC). Assessment of bone growth in the medullary portion of the rabbits' tibiae showed the new test surface's potential for osteoinduction."

■ EVOLUTION OF OSSEOINTEGRATION



NOTE

The images are of Ziacom® implants manufactured specifically for use in the study of BBL-treated implants.

Product presentation

■ Packaging tailored to the type of surface

Ziacom® offers two different types of product packaging depending on the type of implant surface:

Blister packaging

Available for implants with **Titansure** surface treatment. The blisters are heat-sealed and include identification labels for product traceability and a flap for easy opening in the clinic but while preventing accidental opening.

Titansure



Bottle packaging

Available for implants with **Titansure Active** surface treatment. The sealed bottle contains bone bioactive liquid (BBL) to ensure the perfect preservation of the implant's properties. The bottles include identification labels for product traceability.

Titansure Active



IMPORTANT

Do not open the sterile container until just before inserting the implant.

■ Outer identification label

Ziacom® implants are supplied in a sealed cardboard box that includes a product identification label with a description of their main characteristics.

CE 1500	Ziacom®	MD	Implante Dental	ES	ZIACOM MEDICAL, S.L. Phis® - Madrid Soria 6/A - Tel: +34 91 723 33 00 www.ziacom.com
Rx Only	# XXXX - XXXXXXXXXXXXXXX	X Unid	Dental Implant	EN	
T	REF XXXXXXXXX	XX	Zahnimplantat	DE	
XXXXXXXXXX	LOT XXXXXXXX		Implant Dentaire	FR	
XXXXXXXXXX	Ø X,XXxXXmm		Implanto Dentale	IT	
XXXXXXXXXX	XXXXXXXX (01)XXXXXXXXXXXX(17)XXXXXX(11)XXXXXX(10)XXXXXXXXXX		Implante Dentário	PT	
XXXXXXXXXX	XXXXXXXX			UDI	
XXXXXXXXXX	XXXXXXXX				
XXXXXXXXXX	XXXXXXXX				
XXXXXXXXXX	XXXXXXXX				

Description of the symbology used

- CE marking and notified body number.
- Medical device indicator.
- Model code.
- Product name.
- Product batch number.
- Unique device identifier.
- One single sterile barrier system with protective outer packaging. Sterilised by radiation.
- One single sterile barrier system. Sterilised by radiation.
- Temperature limit.
- Caution, consult accompanying documents.
- Do not re-sterilize.
- Do not use if package is damaged.
- Single-use product.
- See instructions for use.
- Product expiration date.
- Date of manufacture.
- Product manufacturer.
- Titansure surface treatment.
- Titansure Active surface treatment.
- Prescription only.

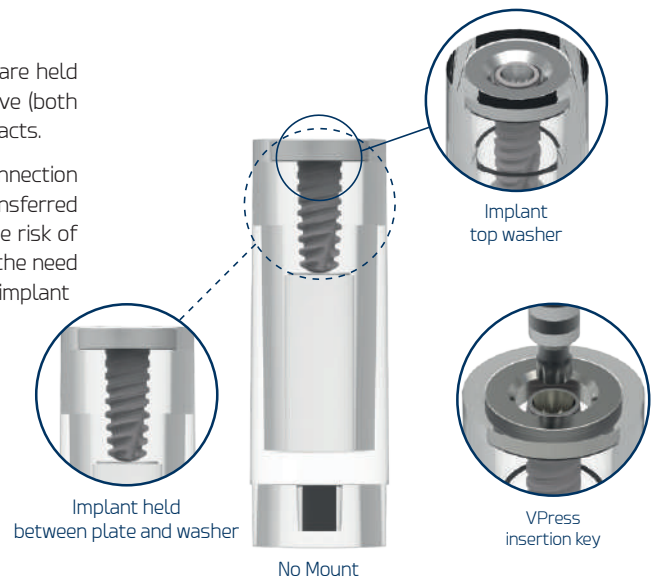
For full details on the product presentation and instructions for use (IFU) see www.ziacom.com/ifus or scan the QR code on the box.



■ Ziacom® No Mount

Galaxy implants are supplied in Ziacom® No Mount vials; the implants are held vertically inside a plastic vial between a plate below and a washer above (both made from titanium), thus preventing any movements or unwanted contacts.

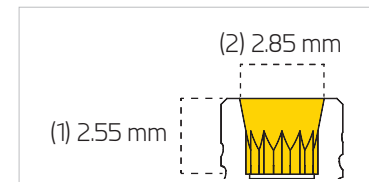
This packaging means that the pressure is applied directly to the connection so the implant can be safely and easily withdrawn from the vial and transferred to the surgical site. Therefore, Ziacom® No Mount implants eliminate the risk of reducing the primary stability caused by over instrumentation, squash the need to handle the implant when removing it from the mount, and simplify implant insertion in posterior areas with limited access.



Galaxy references

IMPLANT						
	Ø (mm)	Ø Core (mm)	Length (mm)	Ref. Titansure	Ref. Titansure Active	
GALAXY	3.40	2.00/3.15	8.5	GLY3485	GLY3485A	
			10.0	GLY3410	GLY3410A	
			11.5	GLY3411	GLY3411A	
			13.0	GLY3413	GLY3413A	
			14.5	GLY3414	GLY3414A	
3.70	2.20/3.70	8.5	GLY3785	GLY3785A		
		10.0	GLY3710	GLY3710A		
		11.5	GLY3711	GLY3711A		
		13.0	GLY3713	GLY3713A		
		14.5	GLY3714	GLY3714A		
4.00	2.40/3.90	6.0	GLY4006	GLY4006A		
		7.0	GLY4007	GLY4007A		
		8.5	GLY4085	GLY4085A		
		10.0	GLY4010	GLY4010A		
		11.5	GLY4011	GLY4011A		
		13.0	GLY4013	GLY4013A		
4.30	2.60/4.05	6.0	GLY4306	GLY4306A		
		7.0	GLY4307	GLY4307A		
		8.5	GLY4385	GLY4385A		
		10.0	GLY4310	GLY4310A		
		11.5	GLY4311	GLY4311A		
		13.0	GLY4313	GLY4313A		
4.80	2.90/4.40	6.0	GLY4806	GLY4806A		
		7.0	GLY4807	GLY4807A		
		8.5	GLY4885	GLY4885A		
		10.0	GLY4810	GLY4810A		
		11.5	GLY4811	GLY4811A		
		13.0	GLY4813	GLY4813A		

Platform



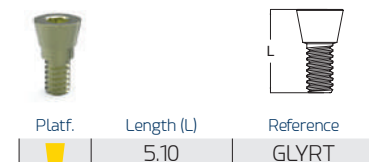
Single platform for all implants: (1) Height of inner cone (2) Diameter of the working platform

Metric



Unique metric of 1.60

Cover screw*



Anodised



* Screw included with each implant.

Recommendations for use

All implant treatments must respect the natural biomechanical stability of the oral cavity and allow the natural emergence of the dental crown through the soft tissue. The implantologist must assess the quantity and quality of bone currently in the implant area and consider the need for prior or simultaneous bone regeneration, as appropriate.

Ziacom® has a wide range of implants available to cover every reconstruction possibility. The inverted trapeziums on the periodontal chart represent the implant diameters and platforms recommended for each tooth position.

These recommendations are valid for the replacement of teeth with single restorations, bridges, hybrid work or overdentures.

Remember to maintain minimum distances between adjacent implants and between implants and teeth in order to preserve interdental papilla, bone vascularisation and natural emergence profiles.

Selection of the appropriate implant for each case is the sole responsibility of the implantologist. Ziacom® advises all clinicians to take into account the warnings based on scientific evidence which can be found in the product catalogues and our website.

■ CLARIFICATIONS ON DRILLING MEASUREMENTS AND TECHNIQUES

- **IMPLANT SIZE:** identifies the diameter and length of the implant.
- **IMPLANT BODY:** diameter of the implant core.
- **DRILL SIZE:** drill bit diameter.
- **DRILLING TECHNIQUE:** we have developed various drilling protocols to enable you to deal with different situations that arise in a schematic way when performing implant surgery.

For more information on implant size selection see the literature available at www.ziacom.com/biblioteca

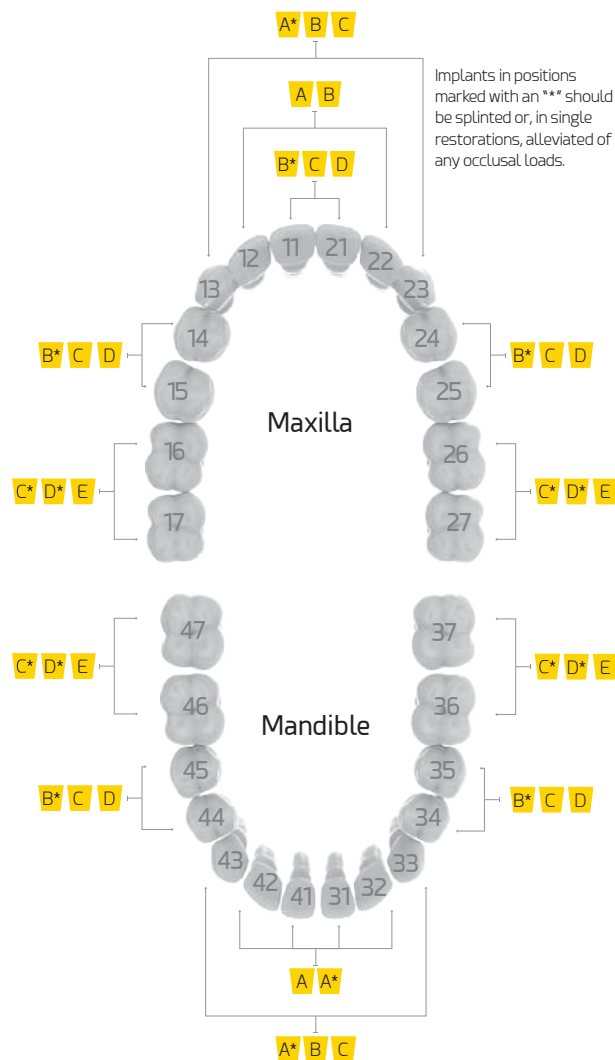


Periodontal chart

GALAXY

Implant diameter

A RP **B** RP **C** RP **D** RP **E** RP
 Ø3.40 mm Ø3.70 mm Ø4.00 mm Ø4.30 mm Ø4.80 mm



IMPORTANT

Short, 6.00 and 7.00 mm implants are ONLY recommended for splinted use in combination with normal length implants (≥ 10.00 mm).

How to use this catalogue

Product sheet

Title, section and paragraph

Product name

Product image

Product table:
 - Platform
 - System
 - Height (H)
 - Diameter (Ø)
 - Reference

Product line diagram

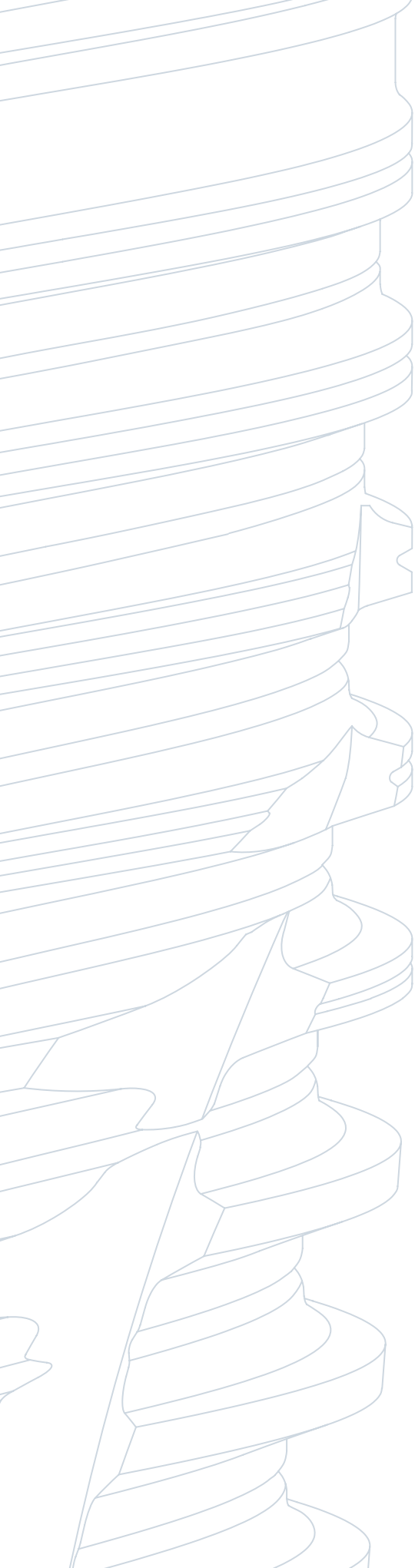
Product characteristics

Additional information

All the dimensions given in this catalogue are expressed in millimeters (mm)

Symbology

Symbol	Meaning	Symbol	Meaning	Symbol	Meaning
	Rotatory element		Size in millimeters		Made from cobalt chromium + castable plastic
	Non-rotatory element		45° screw support		Made from cobalt chromium
	Use with manual torque (see table on page 38)		90° screw support		Made from PEEK
	Maximum operating torque		Use in rotation with a CA		Made from castable plastic
	Ratchet torque range		Maximum rotation speed		Made from plastic
	Galaxy connection		Maximum number of uses		Recommended sterilisation temperature
	Screw connection		Single-use product		Unsterilised product
	Kirator connection		Made from grade 5 ELI (extra-low interstitial) titanium		Use with abundant irrigation
	Basic connection		Made from grade 2 titanium		Use with abundant irrigation
	XDrive connection		Made from stainless steel		
	Tx30 connection		Made from steel		



GALAXY

Abutments

Direct-to-implant
restorations

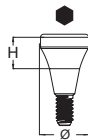


Abutments

Direct-to-implant restorations

2nd STAGE AND IMPRESSIONS

Anatomic healing abutment

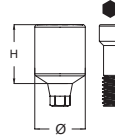


Platf.	Height (H)	Diameter (Ø)	Reference
■	1.50	3.60	HAG3615A
■	3.00	3.60	HAG3630A
■	5.00	3.60	HAG3650A
■	7.00	3.60	HAG3670A
■	1.50	4.60	HAG4615A
■	3.00	4.60	HAG4630A
■	5.00	4.60	HAG4650A
■	7.00	4.60	HAG4670A
■	1.50	5.50	HAG5515A
■	3.00	5.50	HAG5530A
■	5.00	5.50	HAG5550A
■	7.00	5.50	HAG5570A

Anodised ■ RP



Customize healing abutment

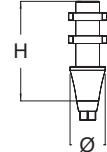


Platf.	Height (H)	Diameter (Ø)	Reference
■	7.00	7.00	HAG7070RAT



Includes screw.

Impression abutment

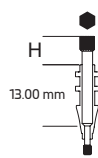


Platf.	Height (H)	Diameter (Ø)	Reference
■	13.00	3.60	TCG3600
■	8.50/Short	3.60	TCG3601
■	13.00	4.60	TCG4600
■	8.50/Short	4.60	TCG4601
■	13.00	5.50	TCG5500
■	8.50/Short	5.50	TCG5501

Anodised ■ RP



Impression abutment screw - Quickly Screws



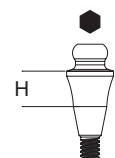
Platf.	Height (H)	Reference
■	0.00	LTSS4000G
■	3.00	LTSS4001G
■	6.00	LTSS4002G
■	9.00	LTSS4010G

Anodised ■ RP



The given impression screw height (H) corresponds to the long impression abutment (13.00 mm).

Pick-Up impression abutment

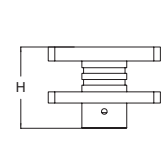


Platf.	Height (H)	Reference
■	3.00	PUG3400

Anodised ■ RP



Pick-Up impression transfer



Platf.	Height (H)	Reference
■	7.25	CPU3410



Pack of 4 units. DO NOT sterilise in an autoclave. Sculptable.

Z2Plus Snap-On impression abutment



Platf.	Height (H)	Length (L)	Reference
	3.00	9.50	Z2RPG10

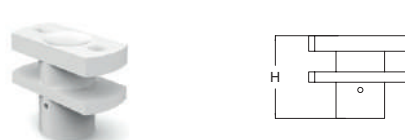
Anodised RP



IMPORTANT

Use the laboratory screw to tighten this impression abutment.

Z2Plus Snap-On impression transfer



Platf.	Height (H)	Reference
	8.00	ZPU3400



Pack of 4 units. DO NOT sterilise in an autoclave. Sculptable.

Implant analogue



Platf.	Length (L)	Reference
	12.00	IAG3400



3D implant analogue

Platf.	Length (L)	Reference
	12.00	IAG3400D



FIXING ELEMENTS

Kiran clinical screw



Platf.	Length (L)	Reference
	8.20	DSG4010



Special Kiran screw with surface treatment.

Laboratory screw



Platf.	Length (L)	Reference
	8.00	LBG4000



NOT apt for use as the final clinical screw.

Kiran Tx30 clinical screw



For abutments and Ti-Base ZiaCam Tx30

Platf.	Length (L)	Reference
	7.55	DSG4010TX



Special Kiran Tx30 screw with surface treatment.
Use only with Tx30 screwdrivers.

Abutments

PROVISIONAL

Provisional abutment



Rotatory

Platf.	Height (H)	Length (L)	Reference
	1.50	10.50	RUGT3615
	3.00	12.00	RUGT3630

Anodised RP



Non-rotatory

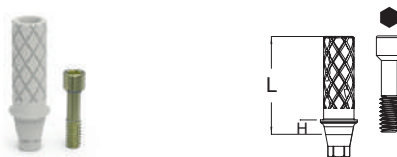
Platf.	Height (H)	Length (L)	Reference
	1.50	10.50	NUGT3615
	3.00	12.00	NUGT3630

Anodised RP



Provisional abutment

Abutments for aesthetic and immediate loading



Rotatory

Platf.	Height (H)	Length (L)	Reference
	1.50	10.50	RUGP3615
	3.00	12.00	RUGP3630



Non-rotatory

Platf.	Height (H)	Length (L)	Reference
	1.50	10.50	NUGP3615
	3.00	12.00	NUGP3630



All provisional abutments come with an anodised screw.

SCREWED

MECHANISED BASE UCLA

Mechanised base abutment + Castable abutment



Rotatory

Platf.	Length (L)	Reference
	10.60	BRUG36



Non-rotatory

Platf.	Length (L)	Reference
	10.60	BNUG36



All mechanised base UCLA abutments come with a special Kiran screw with surface treatment Ref. DSG4010.

Tx30 VARIABLE ROTATION ABUTMENT

Tx30 mechanised base abutment + 2 castable abutments (15° and 20°)



Rotatory

Platf.	15° Length (L)	20° Length (L)	Reference
	11.40	11.20	BRUG36TX



Non-rotatory

Platf.	15° Length (L)	20° Length (L)	Reference
	11.40	11.20	BNUG36TX



Tx30 mechanised base abutment + 2 castable abutments (20° and 25°)



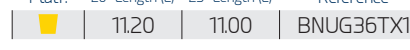
Rotatory

Platf.	20° Length (L)	25° Length (L)	Reference
	11.20	11.00	BRUG36TX1



Non-rotatory

Platf.	20° Length (L)	25° Length (L)	Reference
	11.20	11.00	BNUG36TX1

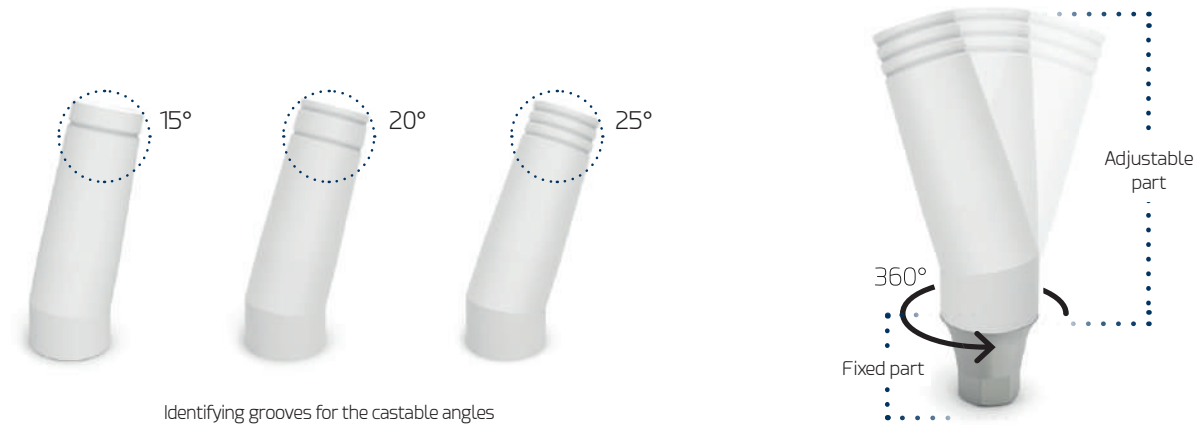


All Tx30 Variable Rotation abutments come with a special Kiran Tx30 screw with surface treatment Ref. DSG4010TX.

■ TX30 VARIABLE ROTATION ABUTMENT

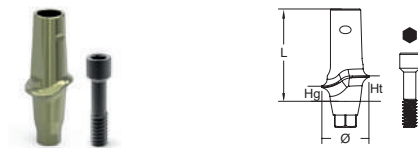
The Tx30 variable rotation abutment comprises a Cr-Co mechanised base that accepts 15°, 20° or 25° angled castable abutments and a Kiran clinical screw with a special Tx30 connection.

The Cr-Co base ensures a perfect fit and seal with the implant connection and the different angles of the castable abutments can be used to choose the best position for the correct emergence of the restoration screw access channel.



CEMENTED

Anatomic straight abutment



Platf.	Height (Hg/Ht)	Length (L)	Diameter (Ø)	Reference
■	1.50/2.50	9.00	3.60	STG3615
■	3.00/4.00	10.50	3.60	STG3630
■	1.50/2.50	9.00	4.60	STG4615
■	3.00/4.00	10.50	4.60	STG4630
■	1.50/2.00	8.50	5.50	STG5515
■	3.00/3.50	10.00	5.50	STG5530

Anodised ■ RP



Anatomic 15° angled abutment



Platf.	Height (Hg/Ht)	Length (L)	Diameter (Ø)	Reference
■	1.50/2.50	9.00	3.60	A1G3615
■	3.00/4.00	10.50	3.60	A2G3615
■	1.50/2.50	9.00	4.60	A1G4615
■	3.00/4.00	10.50	4.60	A2G4615

Anodised ■ RP



Anatomic 25° angled abutment



Platf.	Height (Hg/Ht)	Length (L)	Diameter (Ø)	Reference
■	1.50/2.50	9.00	3.60	A1G3625
■	3.00/4.00	10.50	3.60	A2G3625
■	1.50/2.50	9.00	4.60	A1G4625
■	3.00/4.00	10.50	4.60	A2G4625

Anodised ■ RP



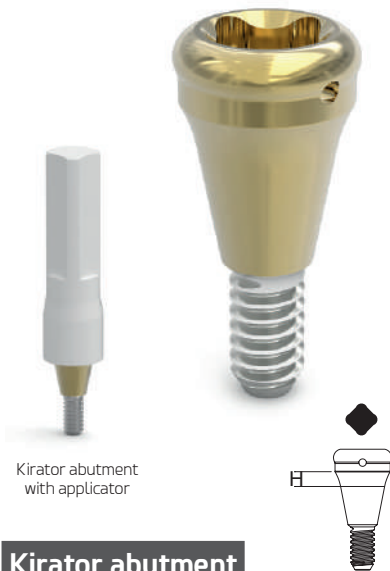
All cemented abutments come with a special Kiran screw with surface treatment Ref. DSG4010.

Abutments

Direct-to-implant restorations

OVERDENTURE

Kirator



Kirator abutment with applicator

Kirator abutment

Platf.	Height (H)	Reference
	1.00	LOG4010
	2.00	LOG4020
	3.00	LOG4030
	4.00	LOG4040
	5.00	LOG4050
	6.00	LOG4060

Golden surface treatment.

Insertion key Ref. LOSD01/LOSD02.



Includes the Kirator abutment with sterilisable polyoxymethylene applicator (Tecaform AH-POM-C).

Related abutments

Kirator impression transfer



System	Height (H)	Reference
Kirator	6.50	TCRK3400



Pack of 4 units. DO NOT sterilise in an autoclave. Sculptable.

Kirator analogue



System	Length (L)	Reference
Kirator	13.00	IATORK01



Kirator processing kit



System	Reference
Kirator processing kit	TPK8520

Kirator processing kit consisting of: Titanium housing with black lined cap, spacer and purple, transparent and pink plastic caps.

Sterilise the metal coping using the autoclave. Plastic caps and spacers should be cold disinfected. See Cleaning and Disinfection Instructions on the Ziacom® website.

System	Retention (Kg)	Reference
Kirator	Soft/1.20 kg	TPK100
	Standard/1.80 kg	TPK200
	Strong/2.70 kg	TPK300

Pack of 4 plastic Kirator retainer caps.



DO NOT sterilise in an autoclave. Maximum divergence of 22° between implants.

Kirator divergence processing kit



System	Reference
Kirator processing kit	TPK8520D

Kirator divergence processing kit comprising: Titanium housing with black lined cap, spacer and purple, transparent and pink plastic caps.

Sterilise the metal coping using the autoclave. Plastic caps and spacers should be cold disinfected. See Cleaning and Disinfection Instructions on the Ziacom® website.

System	Retention (Kg)	Reference
Kirator	Soft/1.20 kg	TPK110*
	Standard/1.80 kg	TPK220*
	Strong/2.70 kg	TPK330*

Pack of 4 plastic Kirator retainer caps - divergent.



DO NOT sterilise in an autoclave. Maximum divergence of 44° between implants.

Example sequence



The references with *(TPK110/TPK220/TPK330) of the Kirator divergent processing pack are subject to availability.

DIGITAL CAD-CAM
ZiaCam scanbody to implant


Platf.	Height (H)	Length (L)	Reference
	3.00	10.00	FNSYG41T

Anodised RP



Indicated for the clinic.

All ZiaCam scanbody to implant abutments include a screw Ref. LBG4000.

 For more information on the recommendations for the use of interfaces in zirconia restorations see the literature available at www.ziacom.com/biblioteca or the use of abutments see the "Prosthetic procedure" manual.

ZiaCam Ti-Base

Rotatory

Platf.	Height (Hg/Ht)	Diameter (Ø)	Reference
	1.00/5.50	3.80	FRUG305
	2.00/6.50	3.80	FRUG315
	3.00/7.50	3.80	FRUG330
	1.00/5.50	4.40	FRUG405
	2.00/6.50	4.40	FRUG415
	3.00/7.50	4.40	FRUG430


Non-rotatory

Platf.	Height (Hg/Ht)	Diameter (Ø)	Reference
	1.00/5.50	3.80	FNUG305
	2.00/6.50	3.80	FNUG315
	3.00/7.50	3.80	FNUG330
	1.00/5.50	4.40	FNUG405
	2.00/6.50	4.40	FNUG415
	3.00/7.50	4.40	FNUG430



All Ti-Base ZiaCam abutments come with a special Kiran screw with surface treatment Ref. DSG4010.

Tx30 ZiaCam Ti-Base

Rotatory

Platf.	Height (Hg/Ht)	Diameter (Ø)	Reference
	1.00/6.50	3.80	FRUG305TX
	2.00/7.50	3.80	FRUG315TX
	3.00/8.50	3.80	FRUG330TX ⁽¹⁾
	1.00/6.50	4.40	FRUG405TX
	2.00/7.50	4.40	FRUG415TX
	3.00/8.50	4.40	FRUG430TX ⁽¹⁾

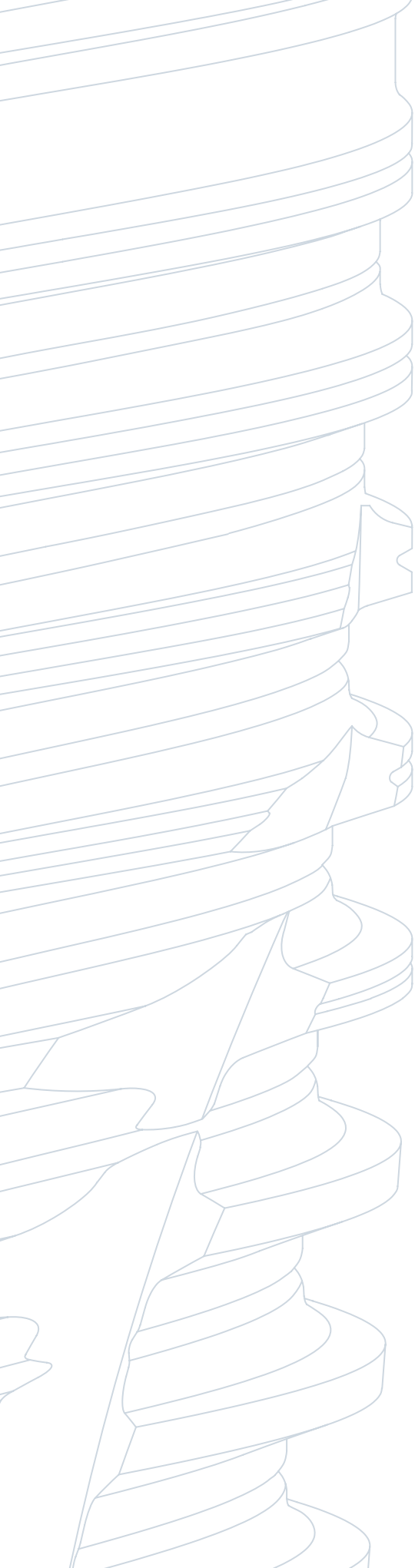

Non-rotatory

Platf.	Height (Hg/Ht)	Diameter (Ø)	Reference
	1.00/6.50	3.80	FNUG305TX
	2.00/7.50	3.80	FNUG315TX
	3.00/8.50	3.80	FNUG330TX ⁽¹⁾
	1.00/6.50	4.40	FNUG405TX
	2.00/7.50	4.40	FNUG415TX
	3.00/8.50	4.40	FNUG430TX ⁽¹⁾



All Ti-Base ZiaCam Tx30 abutments come with a special Kiran Tx30 screw with surface treatment Ref. DSG4010TX.

⁽¹⁾ Gingival heights of 3.00 mm have a maximum angle of 20° (all other heights have a maximum of 30°).



GALAXY

Abutments

Restorations
using transepithelials



Abutments

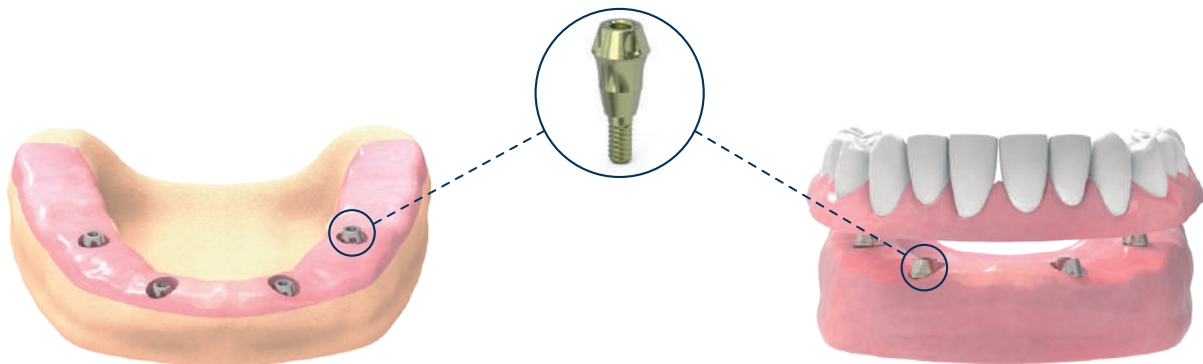
Restorations using transepithelials

■ Transepithelial abutments

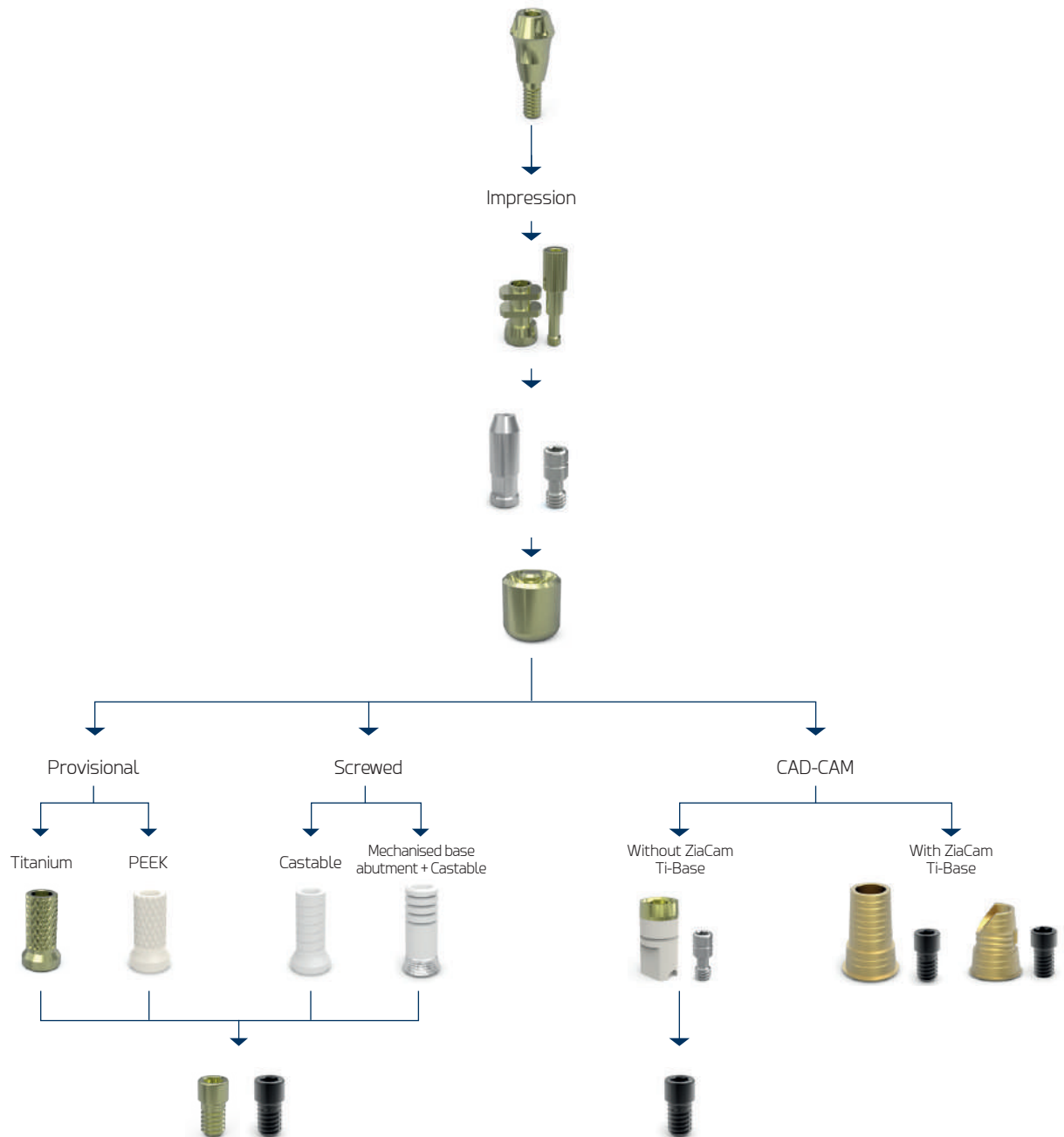
- Allows the formation and maturation of peri-implant tissue from the first 8 weeks.
- One abutment-one time, allows gingival adhesion to its surface as repeated disconnections are not necessary.
- It avoids the loss of bone and soft tissue as there is no mechanical rupture of the peri-implant interface.
- The prosthetic working area is above the gingival level, making the adhesion behaviour of the soft tissue more predictable and maintaining a good seal.
- Less formation of micro gaps at the implant/prosthetic component junction.
- Greater crestal bone preservation.
- Prosthesis try-in and anaesthesia-free definitive placement.
- If the recommended torques are exceeded, the screw fractures in the transepithelial and not inside the implant.

■ Attachment heights

- Higher abutment height equals greater marginal bone preservation in cemented prostheses.
- Taller abutments (≥ 2 mm) provide better soft tissue adaptation.
- Short abutments (< 2 mm) may compress the soft tissues resulting in greater crestal bone loss.
- Marginal bone loss will differ depending on the clinical decision on abutment height. Generally, for prosthetic abutments ≥ 2 mm there will be better crestal bone preservation.



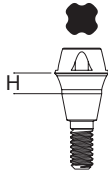
■ Basic | Demonstrative sequence of use



For more information on the use of abutments see the "Prosthetic procedure manual" available at www.ziacom.com/biblioteca

Abutments

Basic abutment



Platf.	Height (H)	Reference
	1.50	BASICG415
	2.50	BASICG425
	3.50	BASICG435
	4.50	BASICG445
	5.50	BASICG455

Insertion key Ref. MABA100/MABA110

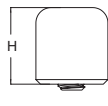


Includes the Basic abutment with sterilisable polyoxymethylene applicator (Tecaform AH-POM-C). 18° cone angle. 36° angle between abutments.



Basic abutment with applicator

Basic healing abutment

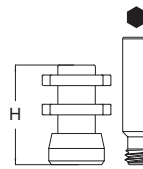


System	Height (H)	Reference
Basic	5.00	BAHAEX34

Anodised RP



Basic impression abutment



Rotatory

System	Height (H)	Reference
Basic	8.00	BATC134

Anodised RP



Non-rotatory

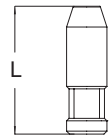
System	Height (H)	Reference
Basic	8.00	BATN134

Anodised RP



All Basic impression abutments come with a screw.

Basic analogue



Rotatory

System	Length (L)	Reference
Basic	13.00	BAIAEX34



Non-rotatory

System	Length (L)	Reference
Basic	13.00	BAIANEX34


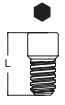


Basic 3D analogue

System	Length (L)	Reference
Basic	13.00	BAIAEX34D





Basic clinical screw

		
System	Length (L)	Reference
Basic	4.30	BDSEI3400

Anodised ■




Kiran Basic clinical screw

		
System	Length (L)	Reference
Basic	4.30	BDSEI3410



Special Kiran screw with surface treatment.



Basic laboratory screw

		
System	Length (L)	Reference
Basic	5.50	BDSEI3401



NOT apt for use as the final clinical screw.


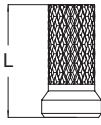
Kiran Tx30 Basic clinical screw

		
System	Length (L)	Reference
Basic	4.10	BDSEI34TX



Special Kiran Tx30 screw with surface treatment.


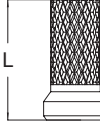
Basic provisional abutment

		
System	Length (L)	Reference
Basic	8.50	BARUT10

Anodised ■



Basic provisional abutment

		
System	Length (L)	Reference
Basic	8.50	BARUP34


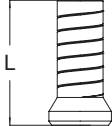


Non-rotatory

System	Length (L)	Reference
Basic	8.50	BANUP34


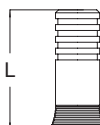


Basic UCLA

		
System	Length (L)	Reference
Basic	9.00	BARUEX34



Abutment base mec. Basic + Abutment calcinable

		
System	Length (L)	Reference
Basic	11.00	BBRU34



Non-rotatory

System	Length (L)	Reference
Basic	11.00	BBNU34



Basic

Abutments

DIGITAL CAD-CAM

ZiaCam scanbody to Basic abutment



Rotatory

System	Length (L)	Reference
Basic	8.70	FNSYB11T



Non-rotatory

System	Length (L)	Reference
Basic	8.70	FNSYB11NT



Indicated for clinical use.

All ZiaCam scanbody to Basic abutments include a screw
Ref. BDSEI3401.

ZiaCam to Basic Ti-Base



Rotatory

System	Height (Hg/Ht)	Reference
Basic	0.30/6.70	BFRU341



Non-rotatory

System	Height (Hg/Ht)	Reference
Basic	0.30/6.70	BFNU341



All ZiaCam to Basic Ti-Bases come with a Kiran special
screw with surface treatment Ref. BDSEI3410.

ZiaCam Tx30 to Basic Ti-Base



Rotatory

System	Height (Hg/Ht)	Reference
Basic	0.30/5.70	BFRU341TX



Non-rotatory

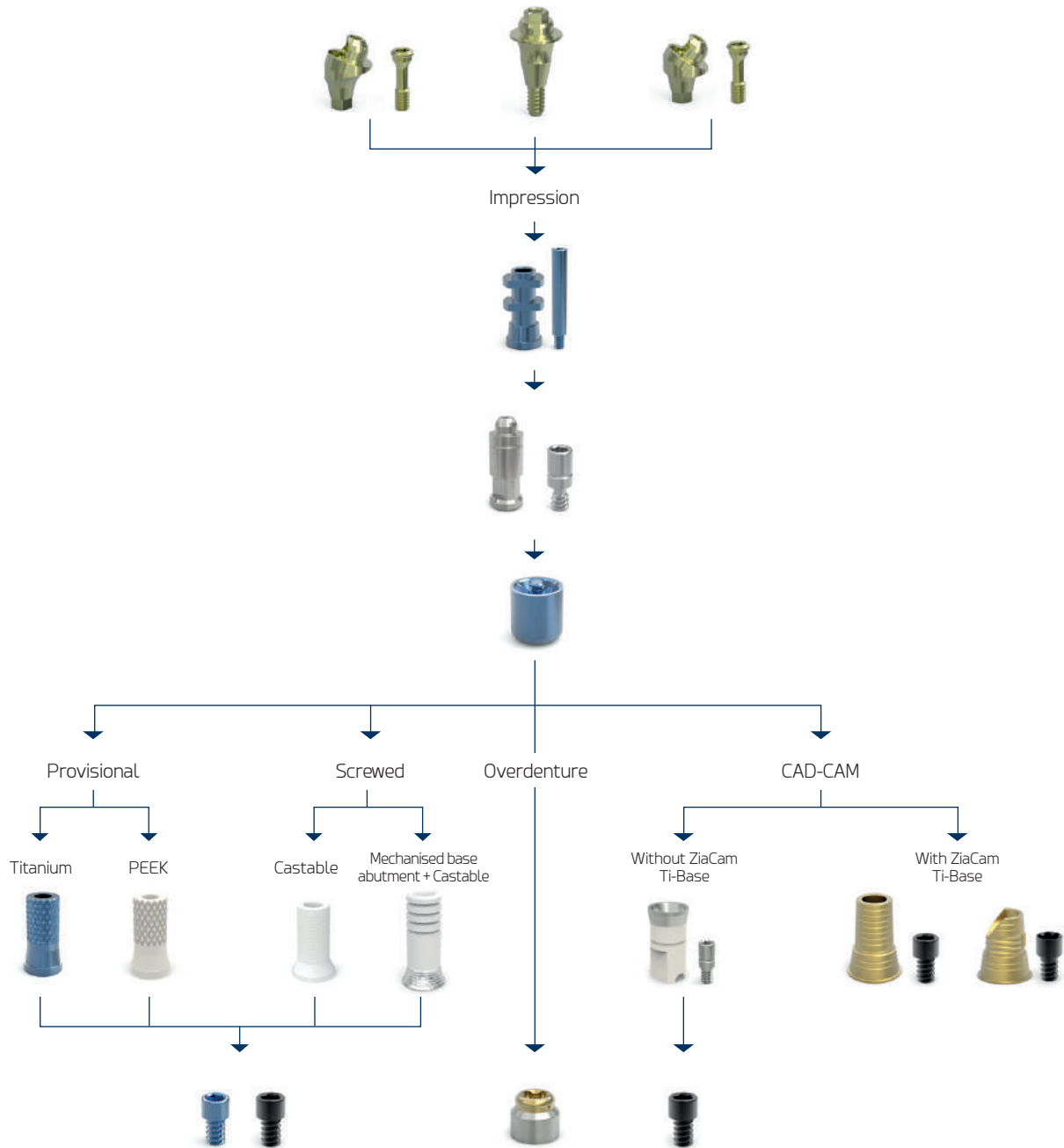
System	Height (Hg/Ht)	Reference
Basic	0.30/5.70	BFNU341TX





All ZiaCam Tx30 to Basic Ti-Bases come with a Kiran Tx30
special screw with surface treatment Ref. BDSEI34TX.

Restorations using transepithelials

■ XDrive | Demonstrative sequence of use



For more information on the use of abutments see the "Prosthetic procedure manual" available at www.ziacom.com/biblioteca

Abutments

XDrive straight abutment



Platf.	Height (H)	Reference
	1.50	XST10G15
	2.50	XST10G25
	3.50	XST10G35
	4.50	XST10G45
	5.50	XST10G55

Insertion key Ref. MABA200/MABA210.

Anodised



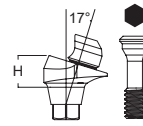
Includes the XDrive abutment with sterilisable polyoxymethylene applicator (Tecaform AH-POM-C).

21° cone angle. 42° angle between abutments.



XDrive abutment with applicator

XDrive 17° angled abutment



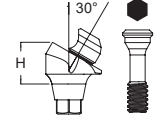
Platf.	Height (H)	Reference
	2.50	XA210G17
	3.50	XA310G17
	4.50	XA410G17
	5.50	XA510G17

Anodised



All angled XDrive abutments come with a stainless steel positioner and screw.

XDrive 30° angled abutment

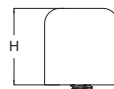


Platf.	Height (H)	Reference
	3.50	XA310G30
	4.50	XA410G30
	5.50	XA510G30

Anodised



XDrive healing abutment

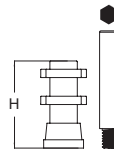


System	Height (H)	Reference
XDrive	5.00	XH103400

Anodised



XDrive impression abutment



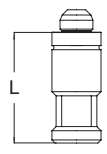
System	Height (H)	Reference
XDrive	10.50	XT103411

Anodised



Includes screw.

XDrive analogue



System	Length (L)	Reference
XDrive	13.00	XIA103400

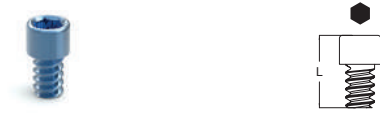


XDrive 3D analogue

System	Length (L)	Reference
XDrive	13.00	XIA103400D



XDrive clinical screw



System	Length (L)	Reference
XDrive	3.50	XDS103410

Anodised ■



Kiran XDrive clinical screw

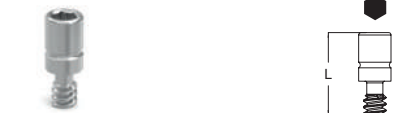


System	Length (L)	Reference
XDrive	3.50	XDS103411



Special Kiran screw with surface treatment.

XDrive laboratory screw



System	Length (L)	Reference
XDrive	5.10	XLB103410



NOT apt for use as the final clinical screw.

Kiran Tx30 XDrive clinical screw



For Ti-Base ZiaCam or metal structures

System	Length (L)	Reference
XDrive	3.50	XDS3411TX



Special Kiran Tx30 screw with surface treatment.

XDrive provisional abutment



System	Length (L)	Reference
XDrive	9.50	XST3410

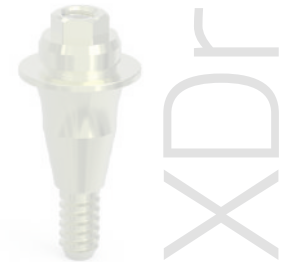
Anodised ■



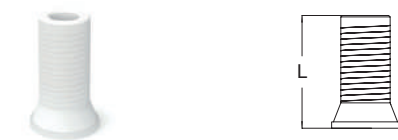
XDrive provisional abutment



System	Length (L)	Reference
XDrive	9.50	XSP3410



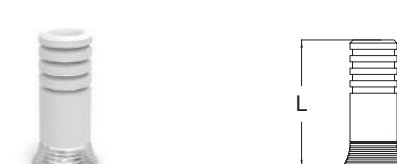
XDrive UCLA



System	Length (L)	Reference
XDrive	8.00	XRU103400



XDrive mechanised base abutment + Castable abutment



System	Length (L)	Reference
XDrive	11.00	XBRU34



Kirator XDrive abutment



System	Height (Hg/Ht)	Reference
XDrive	3.00/4.30	XLO3400

Kirator abutment with gold surface treatment.



Abutments

DIGITAL CAD-CAM

ZiaCam scanbody to XDrive abutment



System	Length (L)	Reference
XDrive	8.70	FNSYX11T



Indicated for clinical use.

All ZiaCam scanbody to XDrive abutments include a screw Ref. XLB103410.

ZiaCam XDrive Ti-Base



System	Height (Hg/Ht)	Reference
XDrive	0.15/6.70	XFRU341



Includes Kiran special screw with surface treatment Ref. XDS103411.

ZiaCam Tx30 XDrive Ti-Base



System	Height (Hg/Ht)	Reference
XDrive	0.15/5.70	XFRU341TX



Includes Kiran Tx30 special screw with surface treatment Ref. XDS3411TX.

Table of abutment torques

Element/Abutment	Instrument/Tool	Torque
Cover screws/Healing abutments	Hex screwdriver 1.25 mm	Manual
Impression abutment screws	Hex screwdriver 1.25 mm	Manual
Laboratory screws	Hex screwdriver 1.25 mm	Manual
Direct-to-implant clinical screws	Hex screwdriver 1.25 mm	30 Ncm
Direct-to-implant Kiran clinical screws	Hex screwdriver 1.25 mm	30 Ncm
Basic/XDrive abutments	Insertion keys: MABA100/MABA110/MABA200/MABA210	30 Ncm
Clinical screws on Basic	Hex screwdriver 1.25 mm	25 Ncm
Kiran clinical screws on Basic	Hex screwdriver 1.25 mm	25 Ncm
Clinical screws on XDrive	Hex screwdriver 1.25 mm	20 Ncm
Kiran clinical screws on XDrive	Hex screwdriver 1.25 mm	20 Ncm
ZiaCam scanbody + screw	Hex screwdriver 1.25 mm	Manual
Kirator abutments	Insertion keys: LOSD01/LOSD02	30 Ncm
Tx30 abutment/screw (Variable Rotation)	Tx30 Torx screwdriver	30 Ncm

ATTENTION

Exceeding the recommended tightening torque for screws and abutments compromises the prosthetic restoration and could damage the implant structure.



For immediate loading: DO NOT tighten manually, attach with the final torque. When using a screwdriver or adaptor for a contra-angle handpiece (CA), do not exceed a maximum speed of 25 rpm.

Surgical
instruments



Surgical instruments

Surgical box

■ Contents of Galaxy boxes available

Platf.	Contents	Reference
	Empty	BOX910
■	Basic, manual/CA	BOX900SGLY
	Complete, manual/CA	BOX901GLY

134°
SSS

Material: radel.

Ensure boxes do not touch the walls of the autoclave to avoid damage.



■ Contents of surgical boxes

REF	Description	BOX900SGLY	BOX901GLY
SID010	Lance drill. Ø2.00 mm	●	●
OSPD20G	Pilot drill. Ø1.6/2.00 mm. Millimeter	●	●
OSTD25G	Stepped surgical drill. Ø1.80/2.50 mm. Millimeter	●	●
OSTD33G	Stepped surgical drill. Ø2.15/2.60/3.30 mm. Millimeter	●	●
OSTD37G	Stepped surgical drill. Ø 2.50/3.10/3.70 mm. Millimeter	●	●
OSTD41G	Stepped surgical drill. Ø2.90/3.50/4.10 mm. Millimeter	●	●
OSTD44G	Stepped surgical drill. Ø 3.40/3.90/4.40 mm. Millimeter	●	●
CLD34	Crestal surgical drill. Ø4.10 mm		●
CLD50	Crestal surgical drill. Ø5.10 mm		●
PMT1G	Paralleling pin. Grade 5 ELI titanium		●
GTPD160	Calibrated drill stop. 1. H 6 mm. Grade 5 ELI titanium		●
GTPD170	Calibrated drill stop. 1. H 7 mm. Grade 5 ELI titanium		●
GTPD185	Calibrated drill stop. 1. H 8.50 mm. Grade 5 ELI titanium		●
GTPD110	Calibrated drill stop. 1. H 10 mm. Grade 5 ELI titanium		●
GTPD115	Calibrated drill stop. 1. H 11.50 mm. Grade 5 ELI titanium		●
GTPD113	Calibrated drill stop. 1. H 13 mm. Grade 5 ELI titanium		●
GTPD114	Calibrated drill stop. 1. H 14.5 mm. Grade 5 ELI titanium		●
GTPD260	Calibrated drill stop. 2. H 6 mm. Grade 5 ELI titanium		●
GTPD270	Calibrated drill stop. 2. H 7 mm. Grade 5 ELI titanium		●
GTPD285	Calibrated drill stop. 2. H 8.50 mm. Grade 5 ELI titanium		●
GTPD210	Calibrated drill stop. 2. H 10 mm. Grade 5 ELI titanium		●
GTPD215	Calibrated drill stop. 2. H 11.50 mm. Grade 5 ELI titanium		●
GTPD213	Calibrated drill stop. 2. H 13 mm. Grade 5 ELI titanium		●
GTPD214	Calibrated drill stop. 2. H 14.5 mm. Grade 5 ELI titanium		●
GTAP34MC	Surgical tap. Ø3.40 mm. Millimeter. CA/Manual	●	●
GTAP37MC	Surgical tap. Ø3.70 mm. Millimeter. CA/Manual	●	●
GTAP40MC	Surgical tap. Ø4.00 mm. Millimeter. CA/Manual	●	●
GTAP43MC	Surgical tap. Ø4.30 mm. Millimeter. CA/Manual	●	●
GTAP48MC	Surgical tap. Ø4.80 mm. Millimeter. CA/Manual	●	●
MUR100G2	Probe/Paralleling pin. Millimeter. Grade 5 ELI titanium		●
MUR200G2	Probe/Paralleling pin. Millimeter. Grade 5 ELI titanium		●
MUR300G2	Probe/Paralleling pin. Millimeter. Grade 5 ELI titanium		●
MUR400G2	Probe/Paralleling pin. Millimeter. Grade 5 ELI titanium		●
SMRGV1	VPress insertion key. Short. Millimeter. CA	●	●
LMRGV1	VPress insertion key. Long. Millimeter. CA	●	●
SMRGV	VPress insertion key. Short. Millimeter. Ratchet	●	●
LMRGV	VPress insertion key. Long. Millimeter. Ratchet	●	●
DEXT10	Drill extender	●	●
MESD	Screwdriver tip. Ø1.25 mm	●	●
LMSD	Surgical screwdriver. Ø1.25 mm. Long. Manual	●	●
SMSD	Surgical screwdriver. Ø1.25 mm. Short. Manual	●	●
TORK50	Regulable torque wrench. 10/20/30/40/50/60/70 Ncm	●	●

SURGICAL DRILLS

Lance drill



Platf.	Diameter (Ø)	Length (L)	Reference
	2.00	16.30	SID010



Instrument with DLC surface treatment.

Pilot drill



Platf.	Diameter (Ø)	Length (L)	Reference
	1.60/2.00	17.50	OSPD20G

Millimeter: 6/7/8.5/10/11.5/13/14.5



Instrument with DLC surface treatment.

Stepped surgical drill



Platf.	Diameter (Ø)	Length (L)	Reference
	1.80/2.50	17.50	OSTD25G
	2.15/3.30	17.50	OSTD33G
	2.50/3.70	17.50	OSTD37G
	2.90/4.10	17.50	OSTD41G
	3.40/4.40	17.50	OSTD44G

Millimeter: 6/7/8.5/10/11.5/13/14.5



Instrument with DLC surface treatment.

Crestal surgical drill



Platf.	Diameter (Ø)	Reference
Universal	4.10	CLD34
	5.10	CLD50



PIN

Paralleling pin



Platf.	Diameter (Ø)	Length (L)	Reference
	4.50	11.00	PMT1G

Anodised



STOPS

Calibrated drill stop



Platf.	Type	Length (L) Implant	Reference
	1	6.00	GTPD160
		7.00	GTPD170
		8.50	GTPD185
		10.00	GTPD110
		11.50	GTPD115
		13.00	GTPD113
		14.50	GTPD114
	2	6.00	GTPD260
		7.00	GTPD270
		8.50	GTPD285
		10.00	GTPD210
		11.50	GTPD215
		13.00	GTPD213
Pack *		--	KSTPG120

* Complete pack of 14 calibrated stops.



TAPS

Surgical tap. CA/Manual



Platf.	Diameter (Ø)	Reference
	3.40	GTAP34MC
	3.70	GTAP37MC
	4.00	GTAP40MC
	4.30	GTAP43MC
	4.80	GTAP48MC

Millimeter: 8.5/10/11.5/13/14.5



Instrument with DLC surface treatment.

Surgical instruments

PROBES

Probe/Paralleling pin



Platf.	Diameters (Ø1-Ø2)	Length (L)	Reference
	1.60/2.00	26.00	MUR100G2
	1.80/2.50	27.00	MUR200G2
	2.10/3.30	26.00	MUR300G2
	2.50/3.70	26.00	MUR400G2

Millimeter: 6/7/8.5/10/11.5/13/14.5



KEYS

VPress insertion key. Ratchet



Platf.	Length (L)	Reference
	12.50/Short	SMRGV
	19.50/Long	LMRGV

● Double hexagon / ■ Square 4x4 mm

Millimeter: 1/2/3/4/5/6



Instrument with DLC surface treatment.

VPress insertion key. CA



Platf.	Length (L)	Reference
	22.20/Short	SMRGV1
	32.20/Long	LMRGV1

● Double hexagon

Millimeter: 1/2/3/4/5/6



Instrument with DLC surface treatment.

Drill extender



Platf.	Length (L)	Reference
Universal	12.00	DEXT10



SCREWDRIVERS

Surgical screwdriver. Manual



Platf.	Length (L)	Reference
Universal	2.80/Mini	XSMSD *
	9.50/Short	SMSD
	14.50/Long	LMSD
	27.00/Extralong	XLMSD *

● Hexagonal 1.25 mm



* Ref. XSMSD/XLMSD are NOT included in the surgical box.

Screwdriver tip. CA



Platf.	Length (L)	Reference
Universal	20.00/Short	MESD01 *
	25.00/Long	MESD

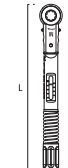
● Hexagonal 1.25 mm



* Ref. MESD01 is NOT included in the surgical box.

RATCHETS

Regulable torque wrench



Platf.	Length (L)	Reference
Universal	86.80	TORK50

■ Square 4x4 mm



Complementary instruments

ADAPTERS

Ratchet extension



Platf.	Length (L)	Reference
Universal	7.20	LAEX

■ Square 4x4 mm



NOT included in the surgical box.

Ratchet to CA adaptor



Platf.	Length (L)	Reference
Universal	7.20	MAEX

■ Square 4x4 mm



NOT included in the surgical box.

LABORATORY TEST KIT

Laboratory test kit



Platf.	Height (H)	Reference
Universal	4.00	GLAB40

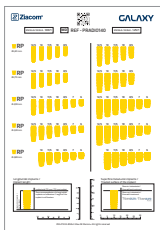


This product does not supersede the need for careful planning of each clinical case.

NOT included in the surgical box.

RADIOGRAPHIC TEMPLATE

Galaxy radiographic template

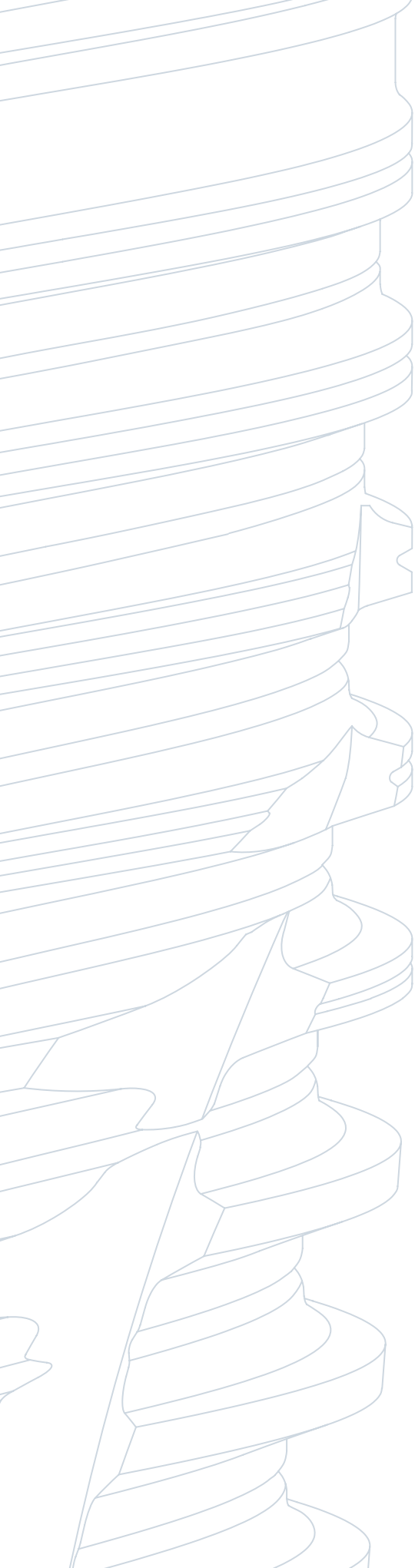


Platf.	Model	Reference
Universal	Galaxy	PRADIO140

Scales 1:1 and 1:1.25

Material: transparent acetate. Non-sterilisable material.

See the literature available at www.ziacom.com/biblioteca



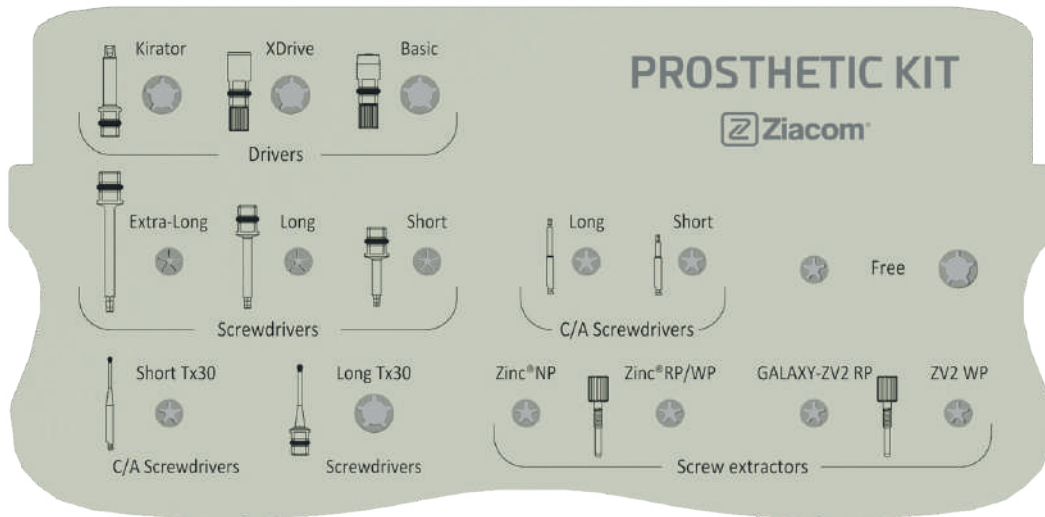
GALAXY

Prosthetic
instruments



Prosthetic instruments

Prosthetic box



■ Contents of prosthetic boxes available

Contents	Reference
Empty	BOXPN
Basic	BOXPSN
Complete	BOXPCN



Material: Radel

Ensure boxes do not touch the walls of the autoclave to avoid damage.



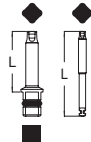
■ Contents of prosthetic boxes

REF	Description	BOXPSN	BOXPCN
LOSD01	Kirator insert key. Ratchet	●	●
MABA100	Basic insert key. Short. Ratchet. Grade 5 ELI titanium	●	●
MABA200	XDrive insert key. Short. Ratchet. Grade 5 ELI titanium	●	●
MADW10	Screwdriver adapter handle. 4x4. Manual	●	●
SMSD1	Screwdriver tip. Ø125 mm. Short. Ratchet	●	●
LMSD1	Screwdriver tip. Ø125 mm. Long. Ratchet	●	●
XLMSD1	Screwdriver tip. Ø125 mm. Extralong. Ratchet		●
MESD	Screwdriver tip. Ø125 mm. Long. CA.	●	●
MESD01	Screwdriver tip. Ø125 mm. Short. CA.	●	●
MESD1TX	Tx30 screwdriver tip. Long. CA.	●	●
LMSD1TX	Tx30 screwdriver tip. Long. Ratchet	●	●
EDSZ20 *	ZPlus extractor screw. Zinic®. NP. Grade 5 ELI titanium		●
EDSZ34 *	ZPlus extractor screw. Zinic®. RP/WP. Grade 5 ELI titanium		●
EDSG34	Abutment extractor screw. Galaxy/ZV2. RP. Grade 5 ELI titanium		●
EDSG50 *	Abutment extractor screw. ZV2. WP. Grade 5 ELI titanium		●
TORK50	Regulable torque wrench. 10/20/30/40/50/60/70 Ncm	●	●

* Product not included in the Galaxy system.

KEYS

Kirator insertion key



System	Length (L)	Reference
Kirator	13.60/Ratchet/Manual	LOSD01
	20.00/CA	LOSD02 *

◆ Square 2.11 mm / ■ Square 4x4 mm



* Ref. LOSD02 is NOT included in the prosthetic box.

Basic insertion key. Ratchet



System	Length (L)	Reference
Basic	5.00/Short	MABA100
	13.00/Long	MABA110 *

◆ Basic / ■ Square 4x4 mm



* Ref. MABA110. is NOT included in the prosthetic box.

XDrive insertion key. Ratchet



System	Length (L)	Reference
XDrive	6.00/Short	MABA200
	13.00/Long	MABA210 *

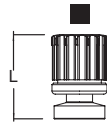
○ XDrive / ■ Square 4x4 mm



* Ref. MABA210. is NOT included in the prosthetic box.

SCREWDRIVERS

Screwdriver adapter handle

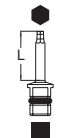


Platf.	Length (L)	Reference
Universal	12.90	MADW10

■ Square 4x4 mm



Screwdriver tip. Ratchet



Platf.	Length (L)	Reference
Universal	9.50/Short	SMSD1
	14.50/Long	LMSD1
	27.00/Extralong	XLMSD1

■ Square 4x4 mm



Screwdriver tip. CA



Platf.	Length (L)	Reference
Universal	20.00/Short	MESD01
	25.00/Long	MESD



Tx30 screwdriver tip. CA



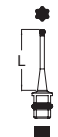
System	Length (L)	Reference
Tx30	26.00/Short	MESD01TX *
	32.00/Long	MESDTX



Do not exceed 30 Ncm, as it could cause severe damage to the screwdriver and screw.

* Ref. MESD01TX is NOT included in the prosthetic box.

Tx30 screwdriver tip. Ratchet



System	Length (L)	Reference
Tx30	12.00/Short	SMSD1TX *
	18.00/Long	LMSD1TX

■ Square 4x4 mm



Do not exceed 30 Ncm, as it could cause severe damage to the screwdriver and screw.

* Ref. SMSD1TX is NOT included in the prosthetic box.

Tx30 prosthetic screwdriver. Manual



System	Length (L)	Reference
Tx30	12.00/Short	SMSDTX *
	18.00/Long	LMSDTX *



Do not exceed 30 Ncm, as it could cause severe damage to the screwdriver and screw.

* Ref. SMSDTX/LMSDTX are NOT included in the prosthetic box.

Prosthetic instruments

EXTRACTOR SCREW

ZPlus extractor screw



Platf.	Length (L)	Reference
● Yellow	25.00	EDSZ20 *
● Blue ● Pink	23.70	EDSZ34 *

Anodised ■ NP ■ RP/WP



Galaxy/ZV2 abutment extractor screw



Platf.	Length (L)	Reference
■ Yellow	25.00	EDSG34
■ Pink	26.80	EDSG50 *

Anodised ■ RP ■ WP



* Product not included in the Galaxy system.

RATCHETS

Regulable torque wrench



Platf.	Length (L)	Reference
Universal	86.80	TORK50

■ Square 4x4 mm



Complementary instruments

CA to ratchet adapter



Platf.	Length (L)	Reference
Universal	12.00	MC10Z

■ Square 4x4 mm



NOT included in the prosthetic box.

Extractor + Retainer inserter handle



Platf.	A Length (L)	B Length (L)	Reference
Kirator	81.50	110.40	MBE13610
ZM-Equator			



NOT included in the prosthetic box.

Retainer inserter



Platf.	Length (L)	Reference
Kirator	32.00	MBE13602
ZM-Equator	32.00	MBE13603



Kirator / ZM-Equator plastic coping insertion tool.
NOT included in the prosthetic box.

Retentive joints instruments



Platf.	Measure	Reference
Universal	2x1	RRE10030

Pack of 10 units.

Surgical
protocol

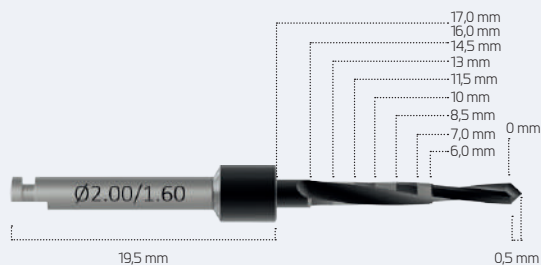


Surgical protocol

Characteristics of the Galaxy drilling system

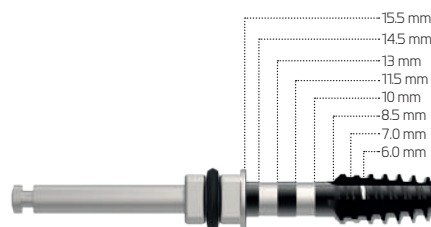
■ Ziacom® drill system - DLC surface

The drills for the Ziacom® implant systems are made from stainless steel coated with a diamond-like carbon (DLC) surface treatment which bestows them greater corrosion resistance during sterilisation, a low friction coefficient and increased wear resistance, thus increasing the service life of their cutting edge. Furthermore, they have a matte finish and therefore anti-reflective properties. A laser marking on the drill's shank identifies its inner and outer diameters and its length, while the horizontal laser marked bands on the active section corresponds to the different lengths of the implants (millimeter drills). The drill tip is 0.5 mm long and is not included in the laser marked measurements.



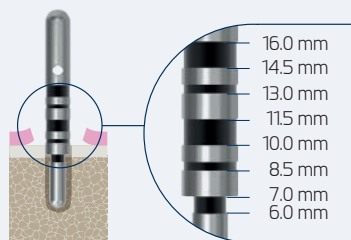
■ Ziacom® taps - DLC surface

Taps are available for contra-angle handpieces. The laser marking on the tap's shank identifies its diameter, while the horizontal laser marked bands on the active section corresponds to the different lengths.



■ Probe

Check the depth of the surgical site, especially when not using drill stops. To check the surgical site axis, the paralleling pins are available in different diameters according to the drilling sequence.

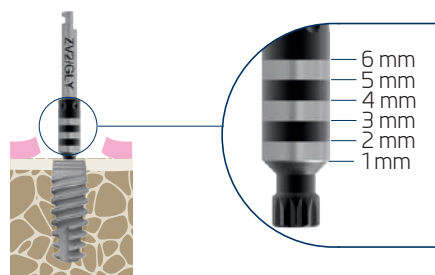


■ VPress insertion keys - DLC surface

The VPress insertion key for contra-angle handpieces or ratchets has been especially designed for transporting Galaxy implants from their No Mount vial to the surgical site ready for insertion.

Short and long insertion keys for ratchets and contra-angle handpieces

Depth within the implant platform marked on the insertion keys

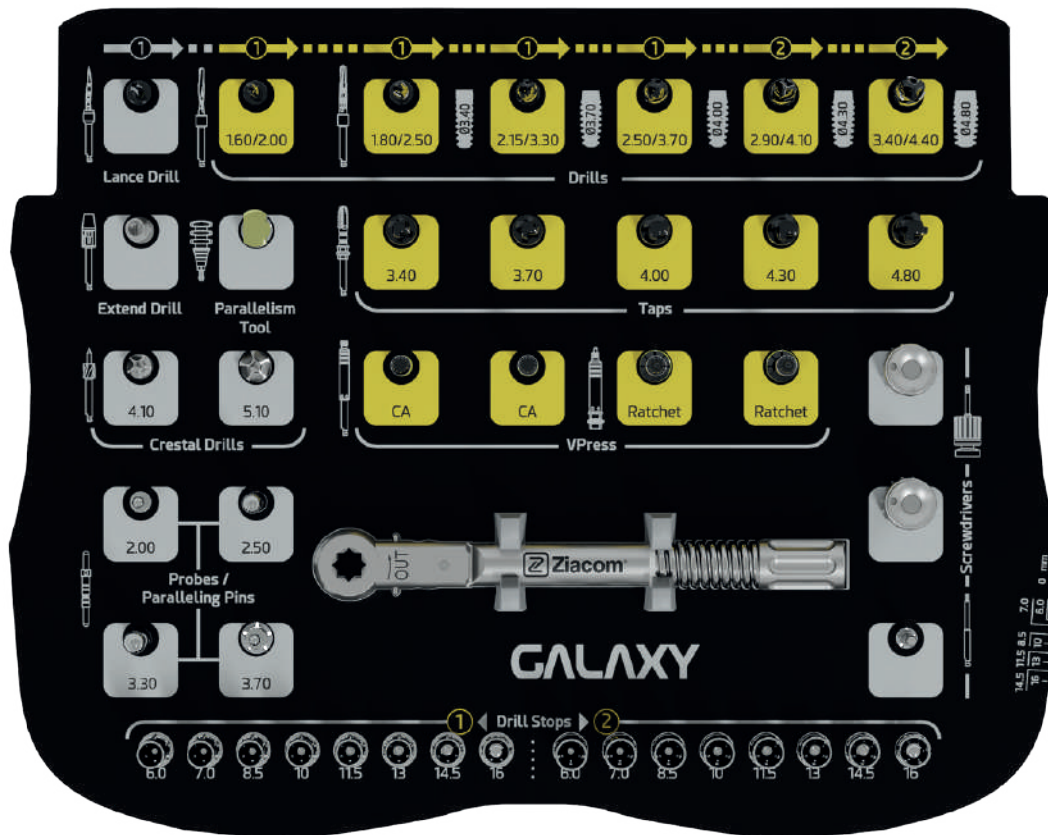


■ Drill stops

They are a surgical accessory that can be attached to the drills, thus facilitating the work by determining the depth of the osteotomy and providing additional safety in the preparation of the surgical site.



■ Details inside the Galaxy surgical box



Recommendation on the maximum insertion torque of the implant



The recommended insertion torque ranges between **35** and **50 Ncm** on a case-by-case basis.

To avoid deforming the key and/or implant connection, insertions performed with a contra-angle handpiece (CA) must respect the recommended maximum rpm (25 rpm) and maximum torque (50 Ncm).

If the implant cannot be fully inserted using the recommended maximum torque, withdraw the implant, repeat the drilling and then re-insert it.

Control the final insertion torque with the adjustable dynamometric ratchet Ref. TORK50 or a contra-angle handpiece.

Exceeding the maximum torque (50 Ncm) when inserting the implant can cause:

- Irreversible deformations in the implant's internal connection.
- Irreversible deformations in the implant insertion instruments.
- Difficulty or impossibility in dismounting the instrument/implant assembly.

Surgical protocol

Implant insertion using Ziacom® No Mount | Titansure

Ziacom® No Mount

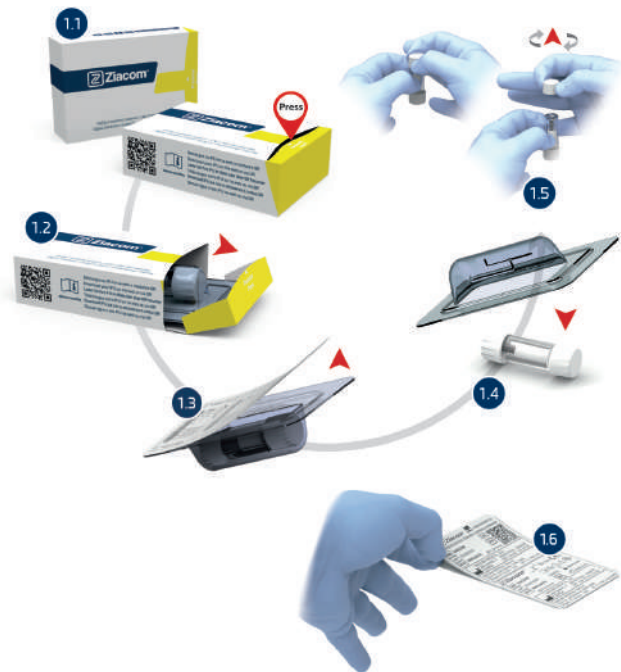
Surface treatment

Titansure



STEP 1 | Implant unpacking

- 1.1 Press the word "PRESS" and tear open the carton.
- 1.2 Remove the flap from the carton and pull out the blister.
- 1.3 Carefully remove the blister seal.
- 1.4 Drop the implant vial onto a sterile cloth in the surgical area.
- 1.5 Hold the vial with one hand in a vertical position. Remove the cap by turning it vertically.
- 1.6 Remember to remove the implant label in order to adhere it to the patient's implant card and medical record to allow the product to be traced.



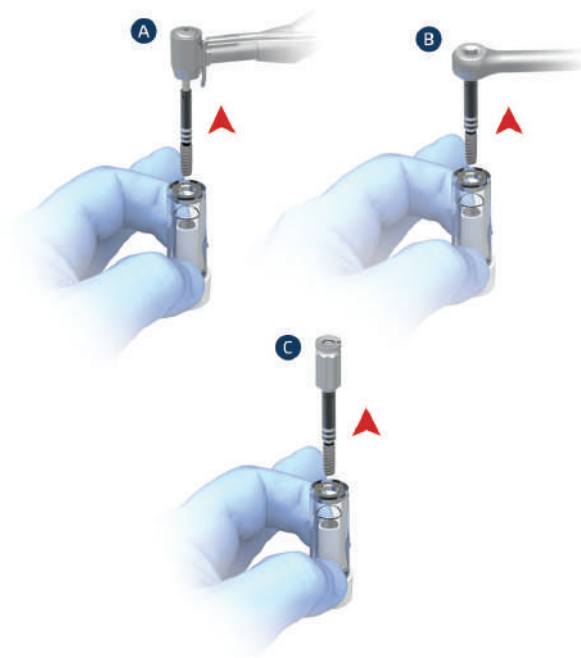
STEP 2 | Choice of insertion instrument

Depending on the clinical situation and access to the area, three different instruments can be chosen to insert the implant:

- Contra-angle.** Use VPress insertion key. CA of the length of your choice (Ref. SMRGV1 or LMRGV1) and insert it into the contra-angle.
- Ratchet Ref. TORK50.** Use VPress insertion key. Ratchet/Manual of the length of your choice (Ref. SMRGV or LMRGV) and insert it into the ratchet in function "IN".
- Screwdriver handle 4x4 Ref. MADW10.** Use VPress insertion key. Ratchet/Manual of the length of your choice (Ref. SMRGV or LMRGV) and insert it into the screwdriver handle.

STEP 3 | Remove the implant from the vial

Hold the implant carrier vial in one hand and insert the selected insertion instrument into the implant with the other hand. Remove the implant by pulling up the vial vertically.

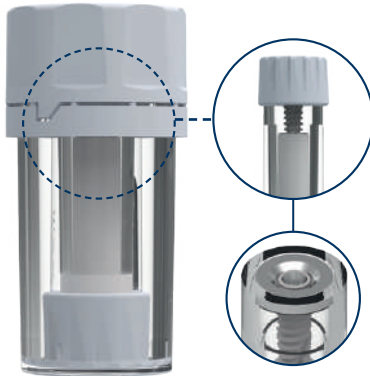


Implant insertion using Ziacom® No Mount | Titansure Active

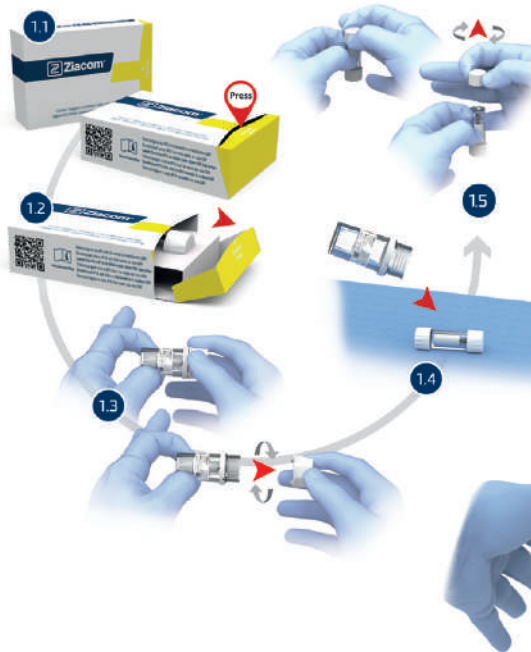
Ziacom® No Mount

Surface treatment

Titansure
Active



STEP 1 | Implant unpacking



- 1.1 Press the word "PRESS" and tear open the box.
- 1.2 Remove the container holding the vial containing the implant and the BBL.
- 1.3 Carefully remove the blister seal.
- 1.4 Drop the implant holder vial onto a sterile cloth in the surgical area.
- 1.5 Hold the vial with one hand in a vertical position. Remove the cap by turning and lifting it.

Note: Take care when opening the vial as the implant is submerged in a bioactive liquid.

- 1.6 Remember to remove the implant labels in order to adhere them to the patient's implant card and medical record to allow the implant reference number and lot number to be traced.

Note: Do not reuse any remaining liquid.

STEP 2 | Choice of insertion instrument

Depending on the clinical situation and access to the area, three different instruments can be chosen to insert the implant:

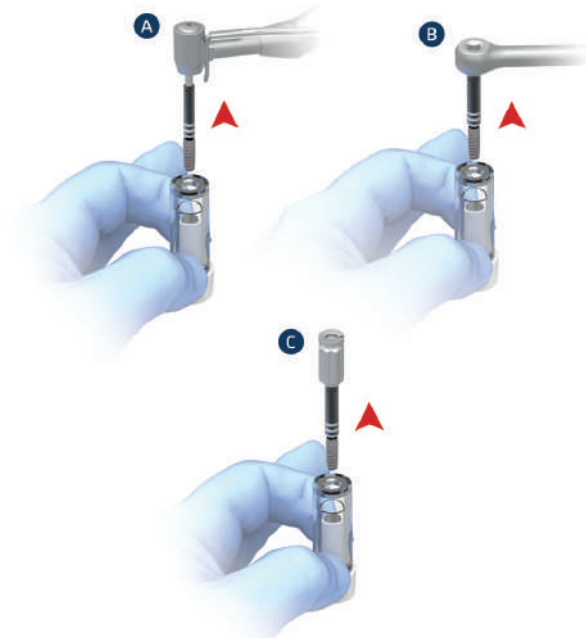
- A Contra-angle.** Use VPress insertion key. CA of the length of your choice (Ref. SMRGV1 or LMRGV1) and insert it into the contra-angle.
- B Ratchet Ref. TORK50.** Use VPress insertion key. Ratchet/Manual of the length of your choice (Ref. SMRGV or LMRGV) and insert it into the ratchet in function "IN".
- C Screwdriver handle 4x4 Ref. MADW10.** Use VPress insertion key. Ratchet/Manual of the length of your choice (Ref. SMRGV or LMRGV) and insert it into the screwdriver handle.

STEP 3 | Remove the implant from the vial

Hold the implant carrier vial in one hand and insert the selected insertion instrument into the implant with the other hand. Remove the implant by pulling up the vial vertically.

Note:

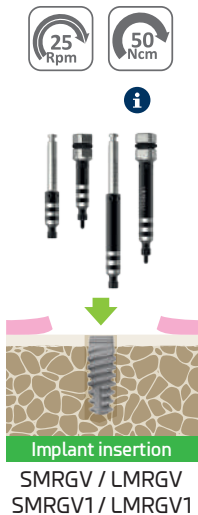
Take care when opening the vial. The Bioactive Liquid may spill. Any remaining Bioactive Liquid cannot be reused.



Surgical protocol

Galaxy implant insertion

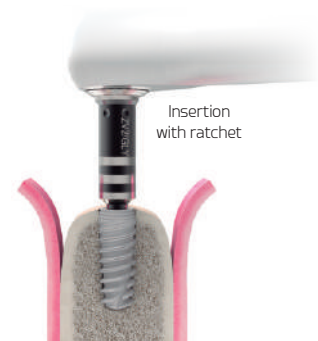
STEP 4 | Implant insertion



When inserting with contra-angle, use a maximum speed of 25 Rpm.

The recommended insertion torque is between 35 and 50 Ncm.

If there is resistance during insertion, it is recommended that the implant be rotated in the opposite direction to the insertion direction and after seconds of pause continue with insertion. Repeat this process as many times as necessary.



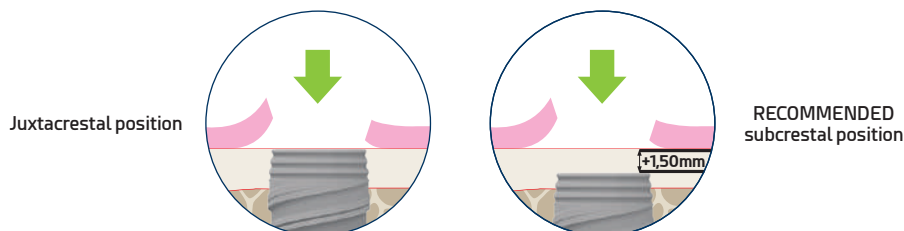
IMPORTANT



The maximum torque for insertion of the dental implants is 50 Ncm. Exceeding the maximum insertion torque indicated for the implants can cause serious damage to the dental implant, its connection, the Mount and the clinical screw included. Refer to the surgical protocol for specific Mount removal considerations, according to implant connection type and bone type.

STEP 5 | Crestal placement of implant

The drilling protocols are described so that the platform of the Galaxy implants is in a juxtacrestal position. However, it is recommended to leave the platform at a subcrestal level of +1.5mm.

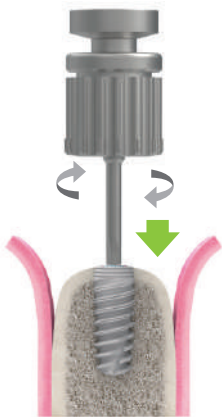


■ Subcrestal position

- Improves mucosal preservation.
- Improves the thickness of keratinised tissue.
- Suggests improved bone preservation when combined with conical connection.
- Helps to obtain an ideal emergence profile in aesthetic areas.
- Prevents the implant surface from being exposed, which can facilitate bacterial growth.
- Prevents fibrous connective tissue formation at the implant interface.
- Better preservation of the crestal bone.
- Allows the use of taller abutments. Recommended for preserving bone tissue in thin biotype gingiva (≤ 1.0 mm).
- Reduces the risk of peri-implant pathologies.

■ Soft tissue conditioning

STEP 1 | Cover screw placement



Insert manual surgical screwdriver Ref. SMSD or LMSD into the cover screw. Approach it to the implant avoiding the fall and accidental screw swallowing. Insert it into the implant until it locks, with manual torque and clockwise.

A second surgery is required to place a cover screw in order to uncover the implant and fit the required abutment.

Depending on the individual case, the professional may decide not to fit a cover screw but to directly fit a healing abutment.



STEP 2 | Soft tissue closure



Close and suture the soft tissue, fitting the flaps carefully.

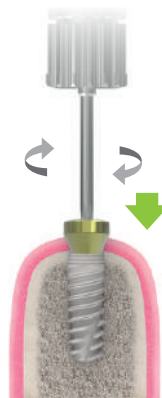
STEP 3 | Uncovering and removing the cover screw



Locate the implant and make an incision until the cover screw is exposed or use the tissue punch Ref. MPU34 on the soft tissue. Remove the screw with the manual surgical screwdriver Ref. SMSD or LMSD.



STEP 4 | Healing abutment placement



Insert selected healing abutment with manual surgical screwdriver Ref. SMSD or LMSD.

The choice of the healing abutment will depend on each case. It must match the implant platform and be in accordance with the gingival tissue height to avoid abutment occlusion. Excessive height could subject the implant to premature loading, compromising the osseointegration process.



Surgical protocol

■ Bone types

Misch classification (1988)



TYPE D1 BONE

- Dense cortical and dense trabecular bone.
- > 1250 HU



TYPE D2 BONE

- Porous cortex and dense trabecular bone.
- 850 - 1250 HU



TYPE D3 BONE

- Porous cortex and thin trabecular bone.
- 350 - 850 HU



TYPE D4 BONE

- Sparse crestal cortex and thin trabecular bone.
- 150 - 350 HU

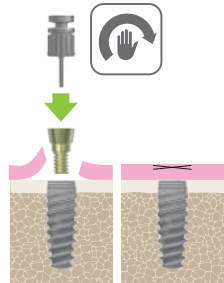
HU = Hounsfield Units

IMPORTANT NOTE

In order to simplify the surgical drilling protocols we have created quick drilling guides, in which the criteria for D1-D2 bones as "High Density" bones and D3-D4 bones as "Low Density" bones are unified.

■ Handling cover screw

Place the cover screw in the screwdriver. Move it towards the implant while taking care that it does not fall and become accidentally ingested. Place the screw in the implant applying manual torque in a clockwise direction.



■ Considerations on provisionalisation and immediate loading

Immediate provisionalisation and immediate loading are procedures that involve the placement of the prosthesis within 72 hours after implant surgery. The fundamental difference between these procedures is whether or not the prosthesis will be functionally loaded.

Adequate primary stability of the implant at the time of insertion is crucial to consider placement of a provisional prosthesis or immediate loading. This stability can be measured objectively by insertion torque, which should be equal to or greater than 40-45 Ncm or by analysis of the resonance frequency (ISQ value), which should be equal to or greater than 70.

■ IMMEDIATE PROVISIONALISATION

Immediate provisionalisation implies an exhaustive control of the occlusion, both in the centric (closing) position and during lateral or dynamic movements that occur during mastication. By freeing the provisional from any kind of contact in these situations, the transmission of forces to the implant is prevented.

The main objectives of immediate provisionalisation are:

- Immediate closure of edentulous spaces in aesthetic areas.
- Guided regeneration of the gingival emergence profile thanks to the presence of the temporary crown or bridge.

■ IMMEDIATE LOADING

The principle of immediate loading involves, in a controlled manner, the transmission of contacts from the moment of placement of the restoration while the restoration is in occlusion, therefore we distinguish between:

- Immediate progressive loading, using a temporary acrylic restoration as the first restoration (released in dynamic occlusion).
- Definitive immediate loading, with rigid material and active occlusion from day one.

Both processes involve risks in the success of the osseointegration of the implant, so it is up to the professional, based on their clinical experience and the case in question, to decide whether or not to place immediate provisionalisation and/or immediate loading.

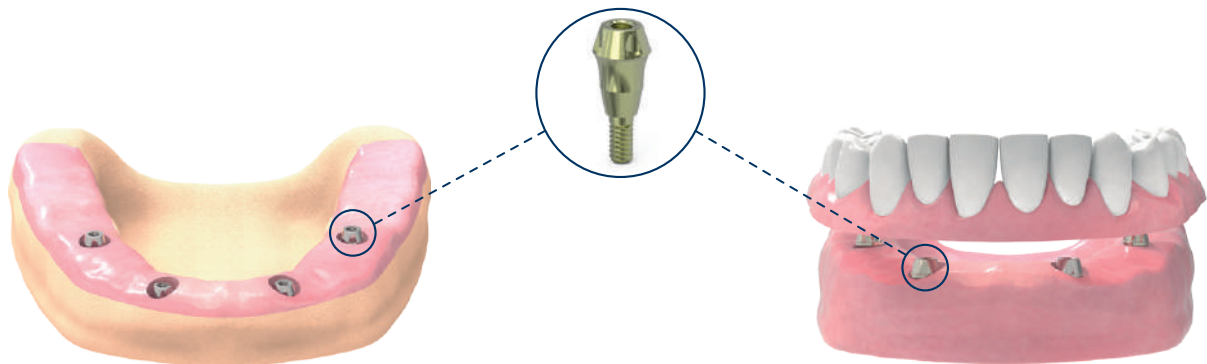
Restorations using transepithelials

■ Transepithelial abutments

- Allows the formation and maturation of peri-implant tissue from the first 8 weeks.
- One abutment-one time, allows gingival adhesion to its surface as repeated disconnections are not necessary.
- It avoids the loss of bone and soft tissue as there is no mechanical rupture of the peri-implant interface.
- The prosthetic working area is above the gingival level, making the adhesion behaviour of the soft tissue more predictable and maintaining a good seal.
- Less formation of micro gaps at the implant/prosthetic component junction.
- Greater crestal bone preservation.
- Prosthesis try-in and anaesthesia-free definitive placement.
- If the recommended torques are exceeded, the screw fractures in the transepithelial and not inside the implant.

■ Attachment heights

- Higher abutment height equals greater marginal bone preservation in cemented prostheses.
- Taller abutments (≥ 2 mm) provide better soft tissue adaptation.
- Short abutments (< 2 mm) may compress the soft tissues resulting in greater crestal bone loss.
- Marginal bone loss will differ depending on the clinical decision on abutment height. Generally, for prosthetic abutments ≥ 2 mm there will be better crestal bone preservation.

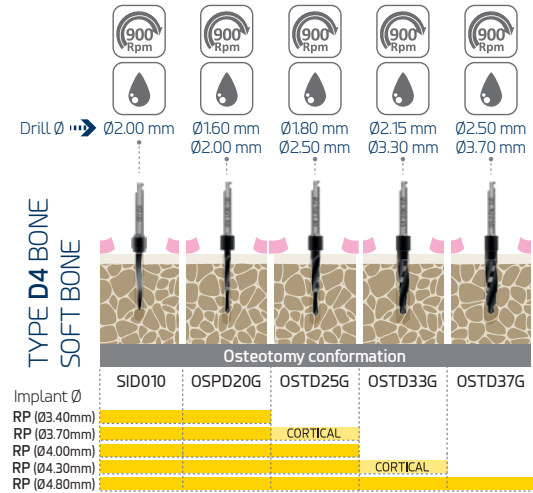


Simplified surgical protocol

Drilling protocol - Ziacom® No Mount

Rotation
 Irrigation required
 Drill diameter
 Torque

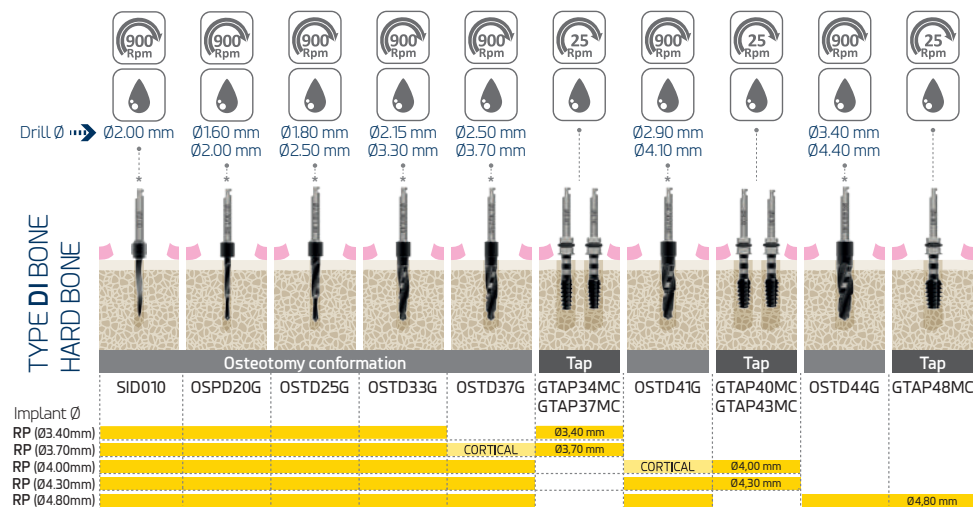
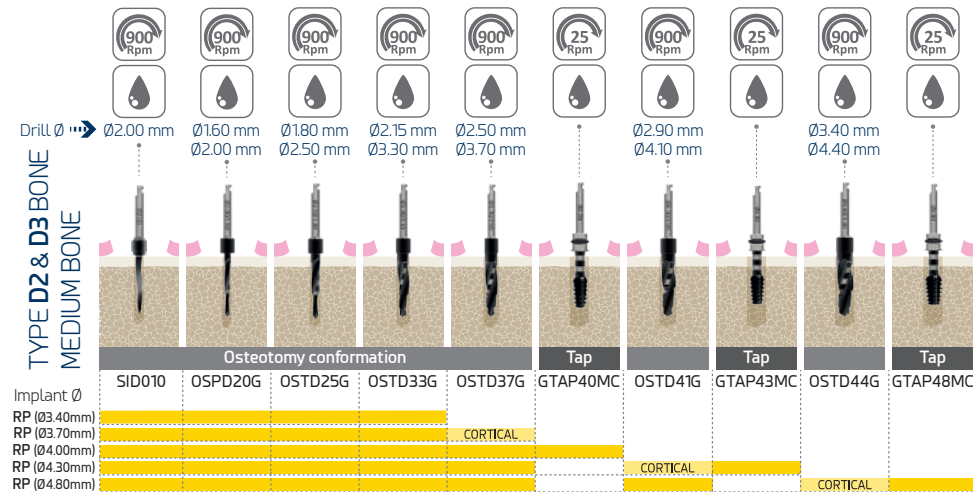
Detailed speeds are the recommended



Cortical section of Galaxy drills



When the protocol indicates, **CORTICAL** we recommend drilling to a length that corresponds to the thickness of the cortical bone on a case-by-case basis.



* When drilling in bone type I, increase by 200 rpm with respect to the values indicated above.

General recommendations

■ To consider during the intervention

1

The surgical drills must be inserted in the contra-angle handpieces when the motor is stopped and ensure they are attached and rotate correctly before starting to drill. Treat the drills with the utmost care; the slightest damage to the tips could compromise their effective operation.

2

Damaged instruments must be disposed of according to local regulations.

3

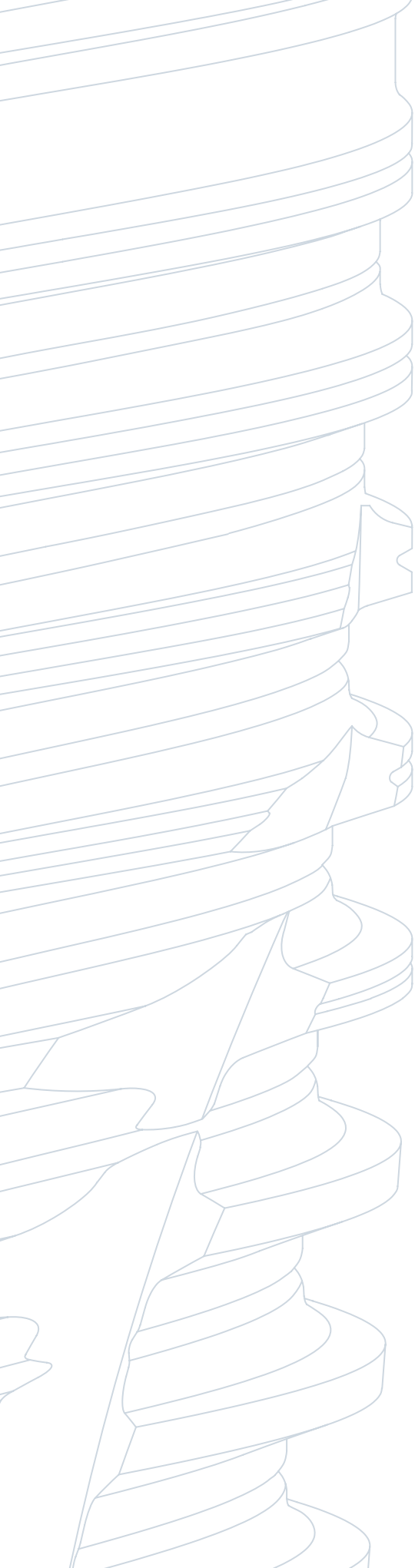
Implantologists should keep one of the identification labels supplied with the product in the patient's file so that it may be traced correctly.

4

Each instrument must only be used for the specific use recommended by the manufacturer.

Always consult the surgical and prosthetic protocols published in this catalogue, as well as the other documents available in the "Reference literature" section of our website www.ziacom.com/biblioteca which explained the procedures, protocols and instructions for use before using the Galaxy system by Ziacom®.





GALAXY

Cleaning, disinfection and sterilisation

The protocols described in this section must only be carried out by personnel qualified to clean, disinfect and sterilise the dental materials specified here in.

Cleaning and disinfection instructions

Applicable for instruments, surgical and prosthetic boxes and plastic retainer caps.

■ Disassembly

1. Dismount* the appropriate instruments, for example manual ratchets, drills or drill stops.
2. Remove the various components from the surgical or prosthetic box for correct cleaning.

■ Cleaning and disinfection

For disinfecting instruments and surgical boxes:

1. Submerge the instruments in a detergent/disinfectant solution** suitable for dental instruments to help eliminate any adhered biological residues. If an ultrasound bath is available***, confirm that the detergent/disinfectant solution is indicated for use with this type of equipment.
2. Manually remove any biological residues with a non-metallic brush and pH-neutral detergent.
3. Rinse with copious water.
4. When cleaning the surgical and prosthetic boxes, always use a pH-neutral detergent and non-abrasive utensils to avoid damaging the surface of the boxes.
5. Dry the materials with disposable cellulose, lint-free clothes or compressed air.

For disinfecting plastic caps and spacers:

1. Submerge in a neat benzalkonium chloride solution for 10 minutes.
2. Rinse with distilled water.
3. Dry the caps and spacer before use.

■ Inspection

1. Check that the instruments are perfectly clean; if not, repeat the cleaning and disinfection steps.
2. Discard any instruments with imperfections and replace them before the next procedure.
3. Check that the instruments and the surgical and prosthetic boxes are perfectly dry before reassembling the parts and proceeding to their sterilisation.

* See the assembly disassembly manuals at www.ziacom.com/biblioteca

** Follow the instructions from the disinfectant's manufacturer to determine the correct concentrations and times.

*** Follow the instructions from the ultrasound bath's manufacturer to determine the correct temperature, concentration and times.

Sterilisation instructions for steam autoclave

Applicable to orthodontic implants, abutments, and surgical and prosthetic instruments and boxes.

1. Introduce each material separately in individual sterilisation bags, then seal the bags. For joint sterilisation, place the instruments in their surgical box, introduce the box into a sterilisation bag and seal the bag.
2. Place the bags to be sterilised in the autoclave.
3. Sterilise in a steam autoclave at 134°C/273°F (max. 137°C/276°F) for 4 min (minimum) and at 2 atm. Torque wrenches must be sterilised in 3 vacuum cycles at 132°C/270°F for a minimum of 1.5 minutes and vacuum-dried for a minimum of 20 minutes.

For the United States only: The validated and recommended sterilisation cycle for the US must be performed in a steam autoclave at 132°C/270°F for at least 15 min and with the drying time of at least 15 - 30 min.

IMPORTANT

Make sure the drying stage is allowed to run to completion, otherwise the products may be damp.

Check the sterilisation equipment if the materials or sterilisation bags are damp at the end of the sterilisation cycle.

Perform the necessary maintenance actions on the autoclave according to the established periodicity and following the manufacturer's instructions.



Storage of Ziacom® products

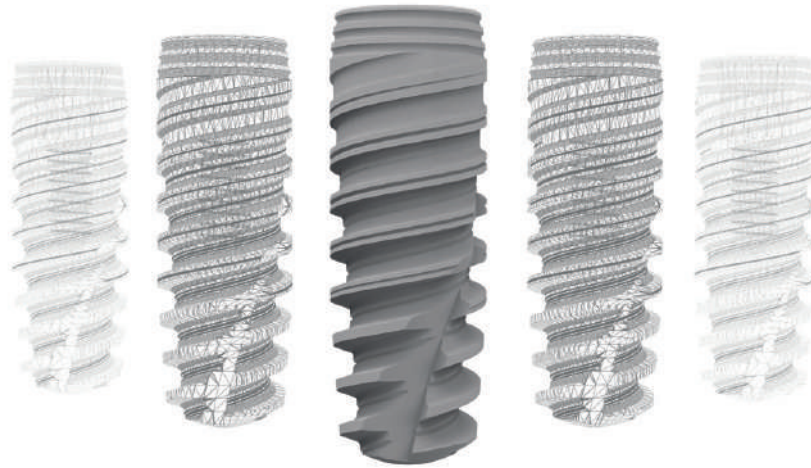
- Store the products in their original packaging and in a clean, dry location until they are used.
- After sterilisation, keep the products in the sealed sterilisation bags and in a clean, dry location.
- Never exceed the use by date indicated by the manufacturer of the sterilisation bags.
- Always follow the indications of the manufacturer of the sterilisation bags.

General recommendations

- Never use damaged or dirty material; never reuse single-use products. The user is responsible for following the instructions described in this document correctly.
- The attention to piercing or sharp elements. Gloves should be worn when cleaning the materials to avoid accidents during handling.
- Follow the safety instructions indicated by the manufacturer of the disinfectant agent.
- The product's sterility cannot be guaranteed if the sterilisation bag is open, damaged or damp.
- Respect all stages of the sterilisation process. If the materials or sterilisation bags contain traces of water or moisture, check the autoclave and repeat the sterilisation.
- Orthodontic abutments and implants are supplied UNSTERILISED and must always be sterilised before use.
- Instruments and surgical and prosthetic boxes are supplied UNSTERILISED and must always be sterilised before use and cleaned and disinfected after use.
- The sterilisation, cleaning and disinfection processes gradually deteriorate the instruments. Inspect the instruments thoroughly to detect any signs of deterioration.
- Avoid contact between products made from different materials (steel, titanium, etc.) during the cleaning, disinfection and sterilisation processes.
- Ziacom Medical SL recommends these instructions are implemented for the correct maintenance and safety of their products; accordingly, the company refuses any liability for any damage to the products that could arise if the user applies alternative cleaning, disinfection and sterilisation procedures.

See www.ziacom.com/biblioteca for the latest version of the cleaning, disinfection and sterilisation instructions.





See the latest version of the general conditions of sale on our website www.ziacom.com.

Check the availability of each product in your country.

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www.ziacom.com

Ziacom Medical SL

Calle Búhos, 2
28320 Pinto - Madrid - ESPAÑA
Tfno.: +34 91 723 33 06
info@ziacom.com

Ziacom Medical Portugal Lda

Av. Miguel Bombarda, 36 - 5° B
1050 -165 - Lisboa - PORTUGAL
Tel: +351 215 850 209
info.pt@ziacom.com

Ziacom Medical USA LLC

333 S.E 2nd Avenue, Suite 2000
Miami, FL 33131 - USA
Phone: +1 (786) 224 - 0089
info.usa@ziacom.com