

Conical connection implants







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).

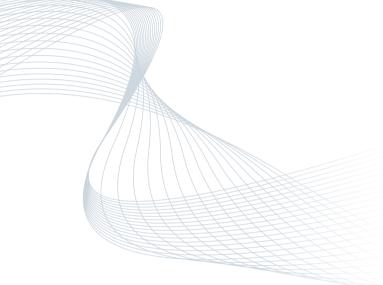
Warning. Not all Ziacom[®] products are available in all counties. Check availability in your country.

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Together for | Z



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The Company

Together for **health**

Ziacom[®] has been working for more than 15 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**.

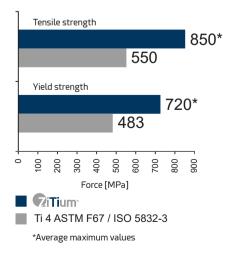
See the General Conditions for Accessing the Guarantee for Ziacom® products.

Zitium[®] titanium

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

Properties of Zitium® titanium



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.



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

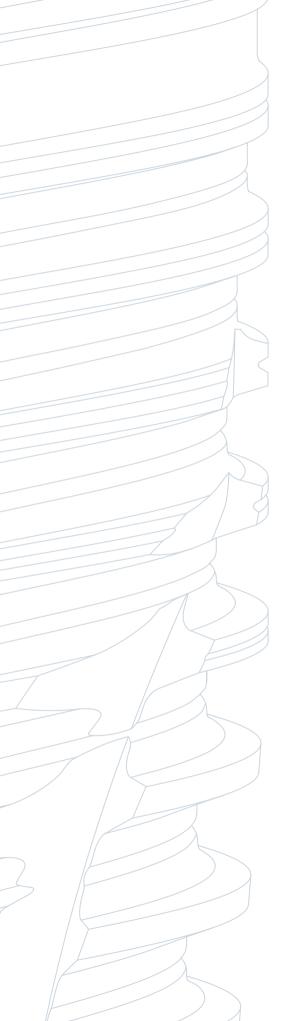
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Please see the up-to-date list of Ziacom® distributors at www.ziacom.com or email us at export@ziacom.com









GALAXY 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.

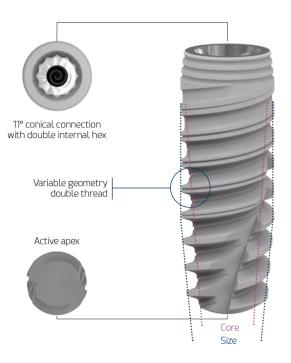
APEX

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

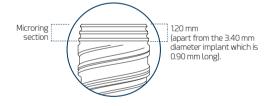
INDICATIONS

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



Dimensions of the implant's coronal section

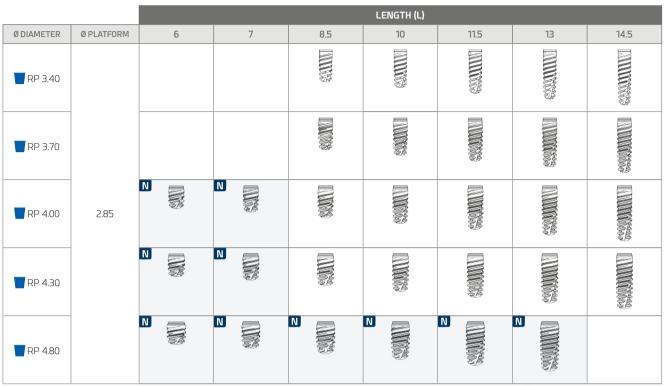


Advantages

- 1. The conical connection prevents micromovement and microfiltration at the implant-abutment interface.
- 2. The single platform provides a significant simplification of prosthetic procedures.
- 3. The reverse taper neck mitigates cortical stress during surgery.
- 4. The thread design confers a very high primary stability even in poor quality bone.
- 5. The active apex facilitates insertion axis correction in postextraction alveoli.



Diameters and lengths



Dimensions in mm.

New product. Check availability.

GALAXY implants

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.

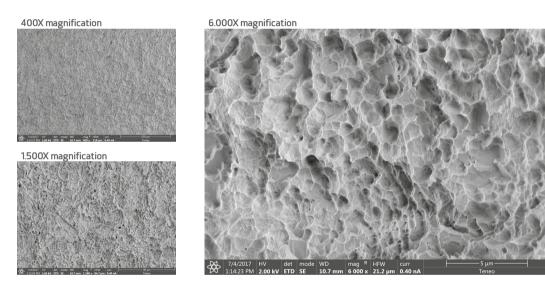
With our **Tibansure** surface treatment, at Ziacom Medical we have obtained a contaminant-free surface topography and optimal average macroand microporosity values, which are key specifications for achieving prompt and proper osseointegration and, in turn, extremely reliable and predictable implants.

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.

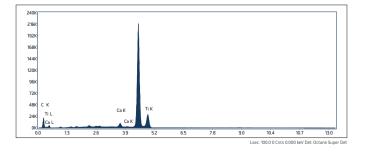
Surface morphology analysis

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 (%)
СК	9.32 (10.23)
AI 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 (%)).



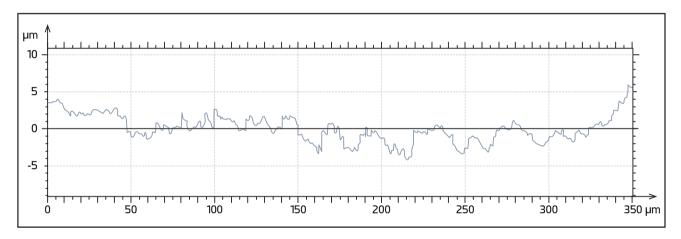
Surface roughness analysis

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].



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GALAXY implants

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 steam 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

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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

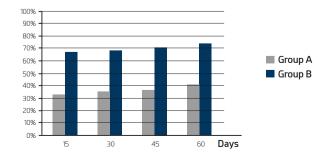
A study was conducted in the tibiae of 10 adult New Zealand rabbits after inserting four implants per rabbit (two in each tibia).

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 Tibansure 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%				



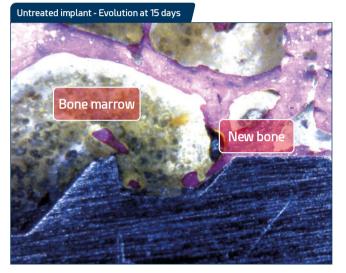


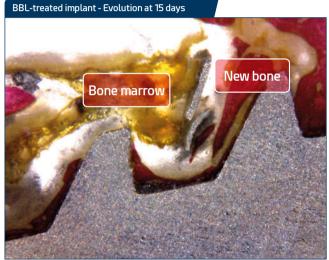
Conclusions

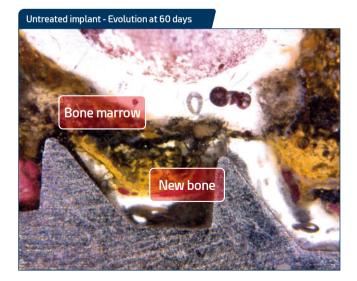
Within the scope of this study, the histomorphometric analysis demonstrated that the group B implants achieved quicker and more effective osseointegration than control group A. Nevertheless, an assessment of bone growth in the medullary portion of the subjects' tibiae revealed the new surface's potential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

GALAXY 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.

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.



IMPORTANT

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



New product. Check availability.

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.



Description of the symbology used

- CE MDD CE certification and notified body
- MD Name of the medical device
- LOT Number of product batch
- Patient information website
- UDI Unique device identification
- Temperature restriction
- Y Temperature restriction
- Caution, consult accompanying documents
- 🛞 Do not resterilise

Consult the instructions for use

Do not use if the packaging is damaged

Expiry date of the product

Non-reusable product

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 (\mathfrak{A})

- Date of manufacture
- Product manufacturer
- TT Titansure surface treatment
- TTA Titansure Active surface treatment
- **RxOnly** Caution: federal law prohibits dispensing without prescription

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.

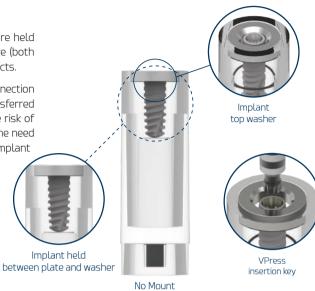




Ziacom[®] No Mount

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.

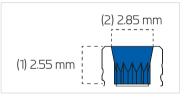
This packaging means that the pressure is applied directly to the connection so the implant can be safely and easily withdrawn from the vial and transferred to the surgical site. Therefore, Ziacom® No Mount implants eliminate the risk of reducing the primary stability caused by over instrumentation, 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	
			8.5	GLY3485	GLY3485A	
			10.0	GLY3410	GLY3410A	
	3.40	2.00/3.15	11.5	GLY3411	GLY3411A	-
			13.0	GLY3413	GLY3413A	
כערעאא			14.5	GLY3414	GLY3414A	-
			8.5	GLY3785	GLY3785A	
			10.0	GLY3710	GLY3710A	
	З.70	2.20/3.70	11.5	GLY3711	GLY3711A	-
			13.0	GLY3713	GLY3713A	3
			14.5	GLY3714	GLY3714A	-
			6.0	GLY4006	GLY4006A	
			7.0	GLY4007	GLY4007A	
	4.00	2.40/3.90	8.5	GLY4085	GLY4085A	
			10.0	GLY4010	GLY4010A	
			11.5	GLY4011	GLY4011A	
			13.0	GLY4013	GLY4013A	
			14.5	GLY4014	GLY4014A	
			6.0	GLY4306	GLY4306A	
			7.0	GLY4307	GLY4307A	-
			8.5	GLY4385	GLY4385A	
	4.30	2.60/4.05	10.0	GLY4310	GLY4310A	
			11.5	GLY4311	GLY4311A	-
			13.0	GLY4313	GLY4313A	
			14.5	GLY4314	GLY4314A	
			6.0	GLY4806	GLY4806A	
			7.0	GLY4807	GLY4807A	-
	4.80	2.90/4.40	8.5	GLY4885	GLY4885A	
	4.00	2.90/4.40	10.0	GLY4810	GLY4810A	
			11.5	GLY4811	GLY4811A	22
			13.0	GLY4813	GLY4813A	

Platform



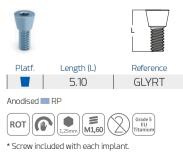
Single platform for all implants: (1) Height of inner cone (2) Diameter of the working platform

Metric



Unique metric of 1.60

Cover screw*



GALAXY implants

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.

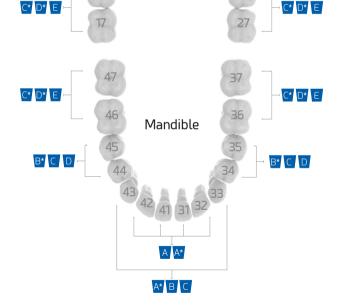
■ CLARIFICATIONS ON DRILLING MEASUREMENTS AND TECHNIQUES

- IMPLANT SIZE: identifies the diameter and length of the implant.
- IMPLANT BODY: diameter of the implant core.
- DRILL SIZE: drill bit diameter.

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• **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.

Periodontal chart GALAXY Implant diameter E RP A RP BRP C RP D RP Ø3.40 mm Ø3.70 mm Ø4.00 mm Ø4.30 mm Ø4.80 mm A* B C - Implants in positions AB marked with an "*" should be splinted or, in single restorations, alleviated of any occlusal loads. R* C D 11 21 B* C D B* C D 25 Maxilla 26



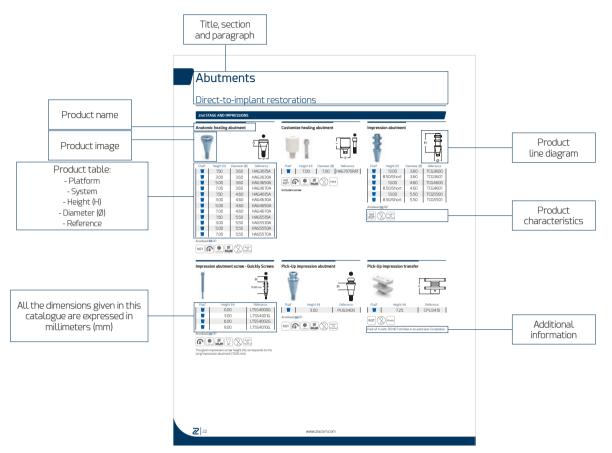
IMPORTANT

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



How to use this catalogue

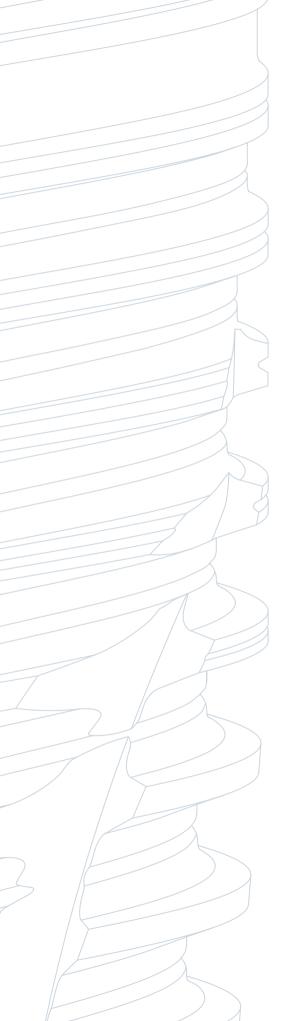
Product sheet



Symbology

Symbol N	Meaning	Symbol	Meaning	Symbol	Meaning
ROT	Rotatory element		Tx30 connection	Co-Cr +castable	Made from cobalt chromium + castable plastic
NO ROT	Non-rotatory element	MX,XX	Size in millimeters	Cobalt Chromium	Made from cobalt chromium
「 山」 ▼	Use with manual torque (see table on page 37)	45°	45° screw support	PEEK	Made from PEEK
Ncm N	Maximum operating torque	90°	90° screw support	Full	Made from castable plastic
Ncm 10 20 80 40 50 60 70	Ratchet torque range	\Diamond	Use in rotation with a CA	Plastic	Made from plastic
Galaxy (Galaxy connection	Rpm	Maximum rotation speed	XX° SSS	Recommended sterilisation temperature
1,25mm	Screw connection	XX USES	Maximum number of uses	Non sterile	Unsterilised product
Kirator k	Kirator connection	(2)	Single-use product		Use with abundant irrigation
Basic	Basic connection	Grade 5 ELI Titanium	Made from grade 5 ELI (extra-low interstitial) titanium	(XX°)	Maximum angle
XDrive >	XDrive connection	Stainless Steel	Made from stainless steel		

19 🖉









Direct-to-implant restorations

2nd STAGE AND IMPRESSIONS

Anatomic healing abutment

u			1	
l				
	(interest			
		una:	and the second se	



Platf.	Height (H)	Diameter (Ø)	Reference			
	1.50	3.60	HAG3615A			
	Э.00	3.60	HAG3630A			
	5.00	3.60	HAG3650A			
	7.00	3.60	HAG3670A			
	1.50	4.60	HAG4615A			
	Э.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						
ROT RT LIZERNAME RELATION RELATION						

Ъгн ø Platf. Diameter (Ø) Height (H) Reference 7.00 HAG7070RAT 7.00 NO ROT ۲

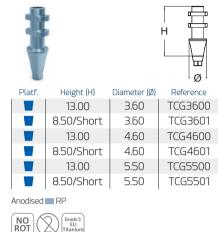
M1,60

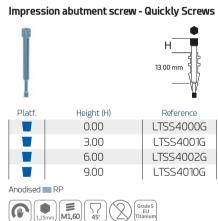
PEEK

Customize healing abutment

Includes screw

Impression abutment



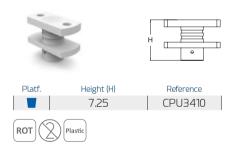


Н

Pick-Up impression abutment

Platf. Height (H) Reference 3.00 PUG3400 Anodised 🔜 RP () M1,60 ROT

Pick-Up impression transfer

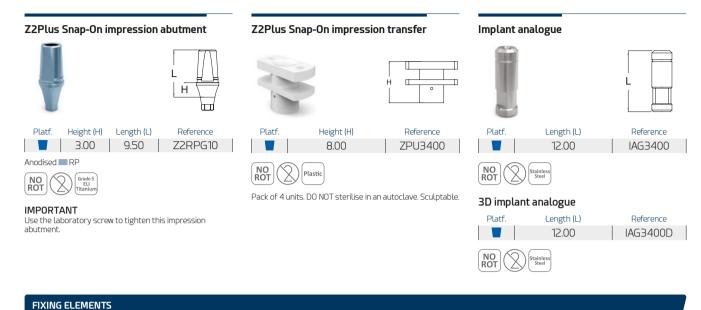


Pack of 4 units. DO NOT sterilise in an autoclave. Sculptable.

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



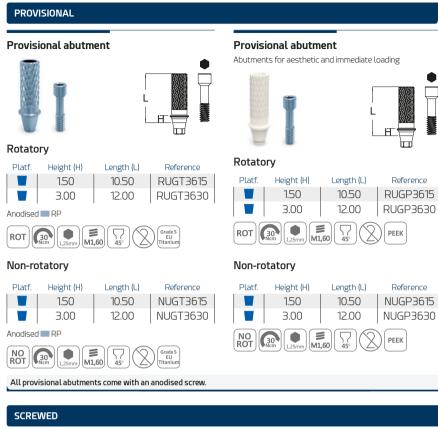






Special Kiran Tx30 screw with surface treatme Use only with Tx30 screwdrivers.

23 🖉





Mechanised base abutment

+ Castable abutment

 L		
	Ш	

Rotatory



Non-rotatory



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



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



Rotatory



Non-rotatory



All Tx30 variable rotation abutments come with a special Kiran Tx30 screw with surface treatment Ref. DSG4010TX.

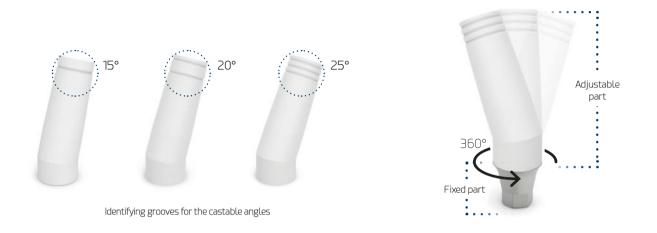


25 🖉

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.



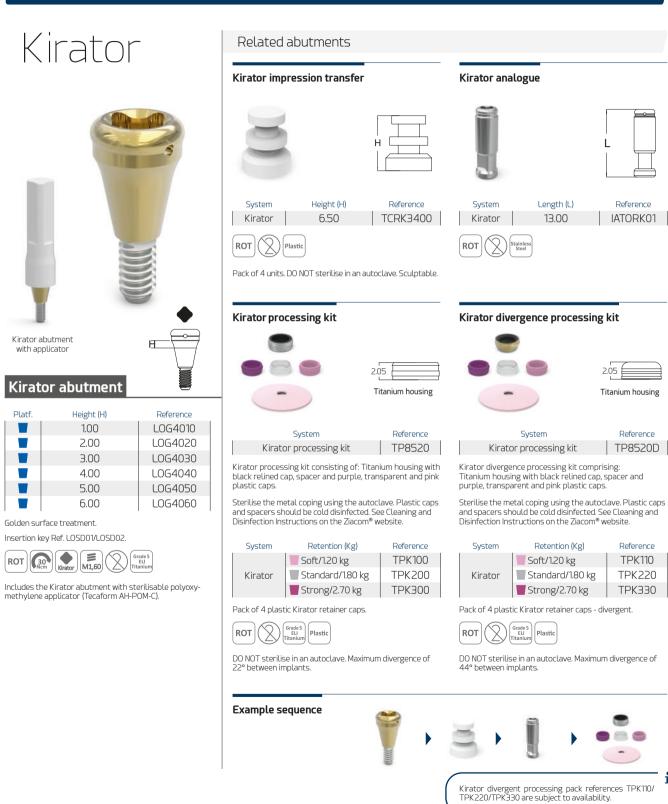
CEMENTED

Anatomic 15° angled abutment Anatomic 25° angled abutment Anatomic straight abutment Platf. Height (Hg/Ht) Length (L) Diameter (Ø) Reference Platf. Height (Hg/Ht) Length (L) Diameter (Ø) Reference Platf. Height (Hg/Ht) Length (L) Diameter (Ø) Reference 9.00 1.50/2.50 STG3615 1.50/2.50 9.00 A1G3615 3.60 A1G3625 3.60 3.60 1.50/2.50 9.00 3.00/4.00 10.50 STG3630 3.00/4.00 3.60 3.00/4.00 3.60 10.50 A2G3615 10.50 3.60 A2G3625 1.50/2.50 1.50/2.50 1.50/2.50 9.00 4.60 STG4615 9.00 4.60 A1G4615 9.00 A1G4625 4.60 3.00/4.00 10.50 4.60 STG4630 3.00/4.00 10.50 4.60 A2G4615 3.00/4.00 10.50 4.60 A2G4625 8.50 5.50 1.50/2.00 STG5515 Anodised 🔤 RP Anodised RP 3.00/3.50 10.00 5.50 STG5530 ROT NO ROT 1 1 M1,60 M1,60 Anodised RP M1,60 ELI ELI

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

Direct-to-implant restorations

OVERDENTURE





DIGITAL CAD-CAM

ZiaCam scanbody to implant



NO ROT Ŵ ELI PEEK M1.60

Height (Hg/Ht) Diameter (Ø)

M

M1,60

3.80

3.80

3.80

4.40

4.40

4.40

1.00/5.50

2.00/6.50

3.00/7.50

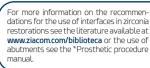
1.00/5.50

2.00/6.50

3.00/7.50

Indicated for the clinic.

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





ZiaCam Ti-Base



Rotatory

Platf.

ROT

NO ROT



Reference

FRUG305

FRUG315

FRUG330

FRUG405

FRUG415

FRUG430

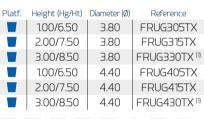
Grade 5 ELI itaniw

Frade 5 ELI

Tx30 ZiaCam Ti-Base



Rotatory



M1.60

Ht

Ha

Щ

Ø



Kirator abutment.Toolbar



Platf. Height (H) Reference Universal 1.80 LOTB100 Gold-coloured surface treatment.









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

45 M1,60

Non-rotatory

30

ROT

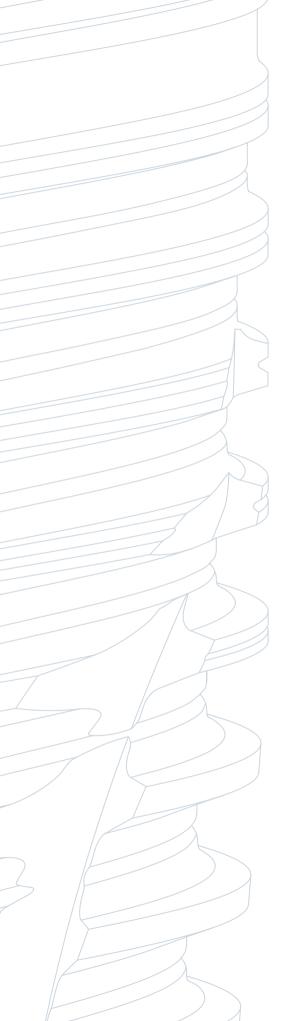
(1)

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)		
$ \begin{array}{c} NO \\ ROT \\ \textcircled{\texttt{NO}} \\ NCm \\ \end{array} \begin{array}{c} \textcircled{\texttt{I}} \\ \textcircled{\texttt{I}} \\ \textcircled{\texttt{I}} \\ I \\ I$					



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

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



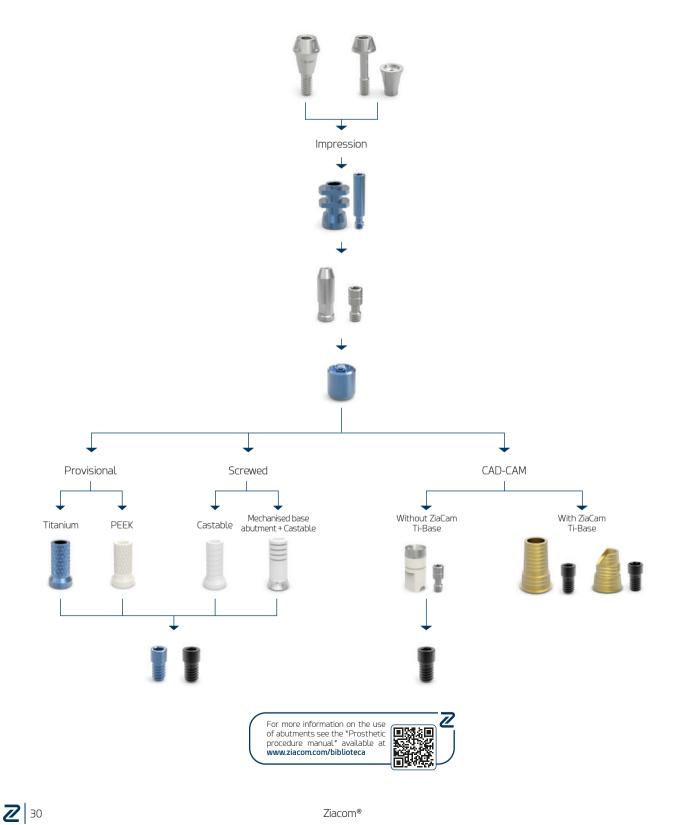


Abutments Restorations using transepithelials

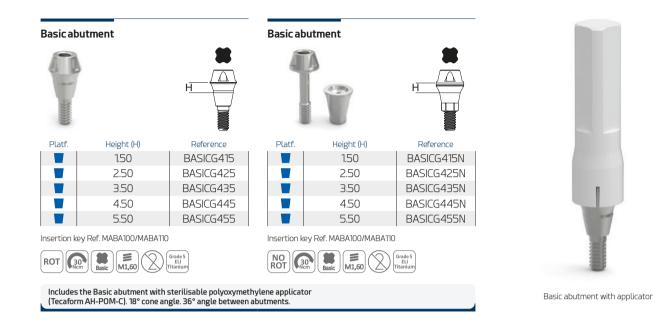


Restorations using transepithelials

Basic | Demonstrative sequence of use







Basic healing abutment



Basic impression abutment



Rotatory





Non-rotatory



All Basic impression abutments come with a screw.

Basic analogue



ROT

Reference

BATC134



Rotatory

System Length (L)
Basic 13.00

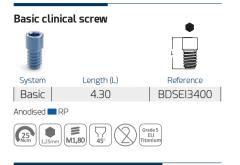
Reference BAIAEX34

Non-rotatory



Basic 3D analogue





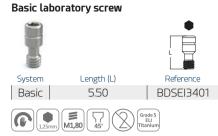
Kiran Tx30 Basic clinical screw



Kiran Basic clinical screw



Special Kiran screw with surface treatment



NOT apt for use as the final clinical screw.

Special Kiran Tx30 screw with surface treatment.

Basic provisional abutment

assessed a				
System	Length (L)	Reference		
Basic	8.50	BARUT10		
Anodised RP				

Basic provisional abutment





Rotatory

System

Basic

Т

Reference

Length (L)

Reference BARUP34



Non-rotatory

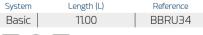


8.50

Abutment base mec. Basic + Abutment calcinable









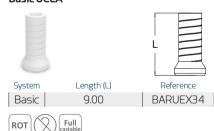
Non-rotatory







Basic UCLA





DIGITAL CAD-CAM

ZiaCam scanbody to Basic abutment



Rotatory





Indicated for clinical use.

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

ZiaCam to Basic Ti-Base





Rotatory



Non-rotatory



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

ZiaCam Tx30 to Basic Ti-Base





Rotatory





Non-rotatory

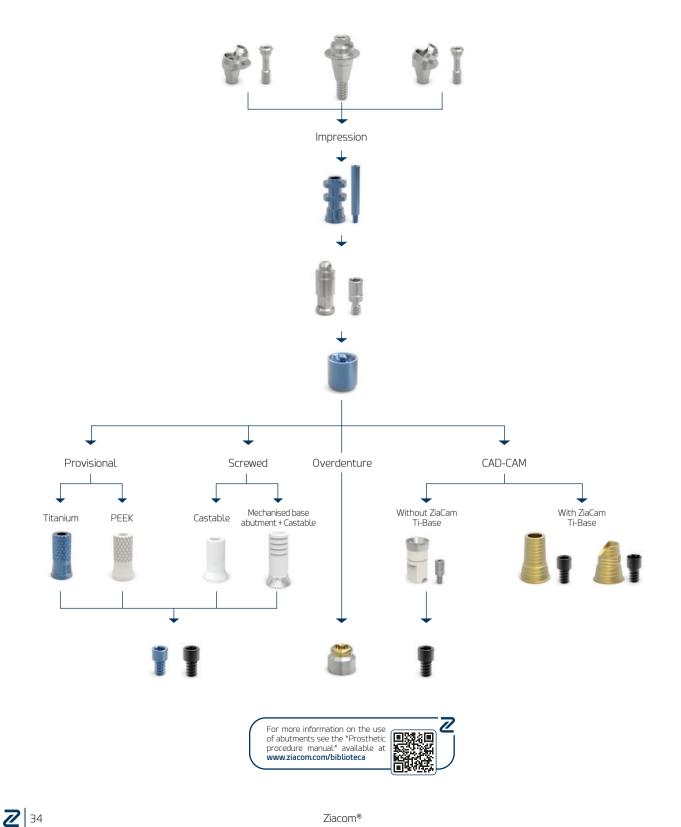
System	Height (Hg/Ht)	Reference		
Basic	0.30/5.70	BFNU341TX		

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

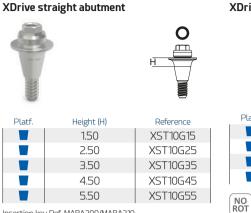


Restorations using transepithelials

• XDrive | Demonstrative sequence of use







Insertion key Ref. MABA200/MABA210.

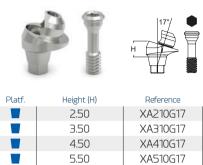
M rade ELI ROT M1,60

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



XDrive 30° angled abutment





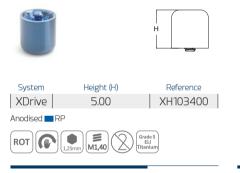
All angled XDrive abutments come with a stainless steel positioner and screw.

Grade 5 ELI Titaniur

XDrive healing abutment

Ô

(M1,60)



XDrive impression abutment



System

XDrive

Height (H) 10.50 XT103411

Reference



XDrive analogue





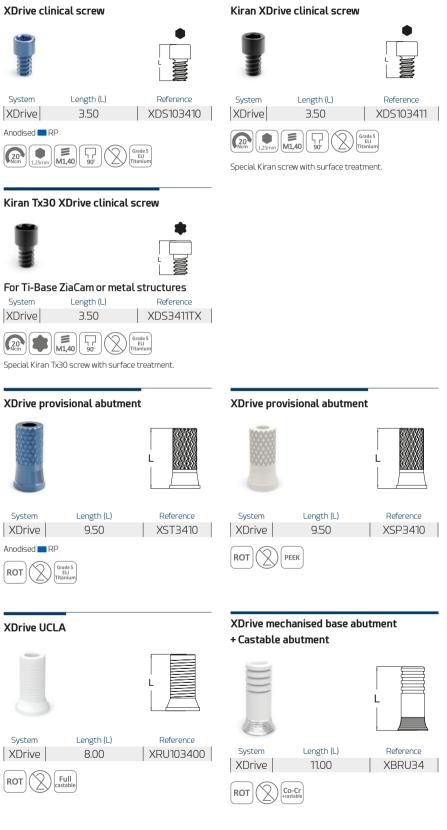
System	Length (L)	Reference
XDrive	13.00	XIA103400
ROT	Stainless Steel	

XDrive 3D analogue



Abutments

Z 36



XDrive laboratory screw

(Inclusion of the second secon

NOT apt for use as the final clinical screw.



Kirator XDrive abutment



Kirator abutment with gold surface treatment.





DIGITAL CAD-CAM

ZiaCam scanbody to XDrive abutment



Indicated for clinical use.

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

ZiaCam XDrive Ti-Base







ZiaCam Tx30 XDrive Ti-Base



System	Height (Hg/Ht)	Reference
XDrive	0.15/6.70	XFRU341
ROT 20	■ (● (■ (■ (■ (■ (■ (■ (■ (■ (■ (■ (■ (■ (■	Grade 5 ELI Titanium

Includes Kiran special screw with surface treatment Ref. XDS103411.

System	Height (Hg/Ht)	Reference
XDrive	0.15/5.70	XFRU341TX
ROT 20		Grade 5 EU Titanium

Includes Kiran Tx30 special screw with surface treatmen-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

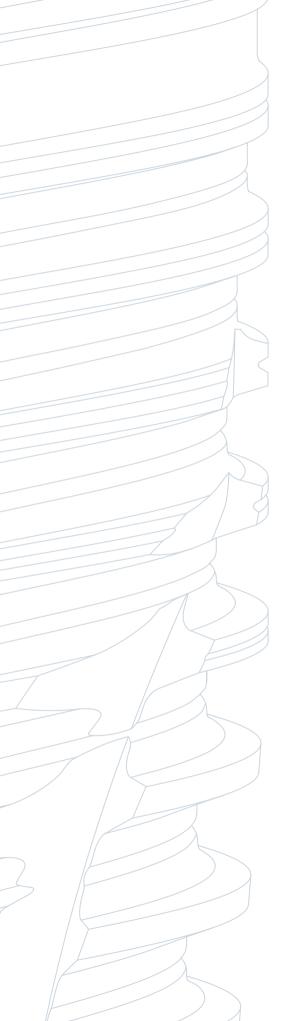
Exceeding the recommended tightening torque for screws and abutments compromises the prosthetic restoration and could damage the implant structure.

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.

37 🖉

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Surgical instruments

Surgical box

Contents of Galaxy boxes available

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

134° \$\$\$

Material: radel.

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



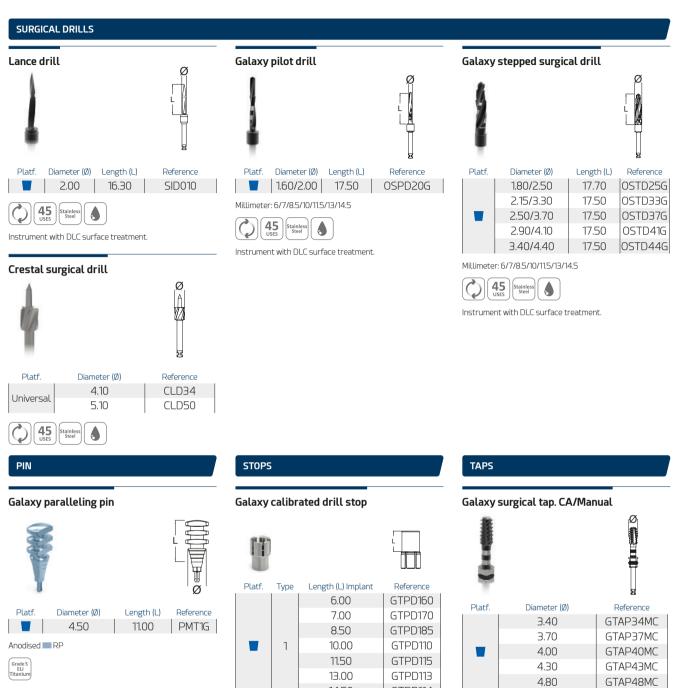


Contents	of surgica	al boxes
	0. 20.0.0.0	

Contents of surgical boxes		92006X08	BOX901GL
REF	Description	BO	BO
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. Ø3.70 mm. Millimeter. CA/Manual	•	
	Surgical tap. Ø4.00 mm. Millimeter. CA/Manual	•	
GTAP43MC	Surgical tap. Ø4.30 mm. Millimeter. CA/Manual	•	•
GTAP48MC		•	
MUR100G2	Probe/Paralleling pin. Millimeter. Grade 5 ELI titanium		•
	Probe/Paralleling pin. Millimeter. Grade 5 ELI titanium		
	Probe/Paralleling pin. Millimeter. Grade 5 ELI titanium		
	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. Ø 1.25 mm. Short. Manual	•	•
TORK50	Regulable torque wrench. 10/20/30/40/50/60/70 Ncm	•	•
UCANO	ריפעממטיב נטו קעב או פרוברו. וטו בטו סטו 40 סטוטטו ויט ואנדון		-

GLY GLY





Millimeter: 8.5/10/11.5/13/14.5



Instrument with DLC surface treatment.

41 2

* Complete pack of 14 calibrated stops.

2



Pack *

14.50

6.00

7.00

8.50

10.00

11.50

13.00

14.50

GTPD114

GTPD260

GTPD270

GTPD285

GTPD210

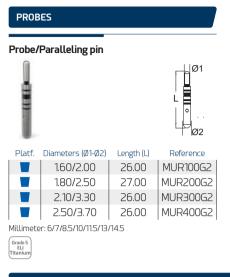
GTPD215

GTPD213

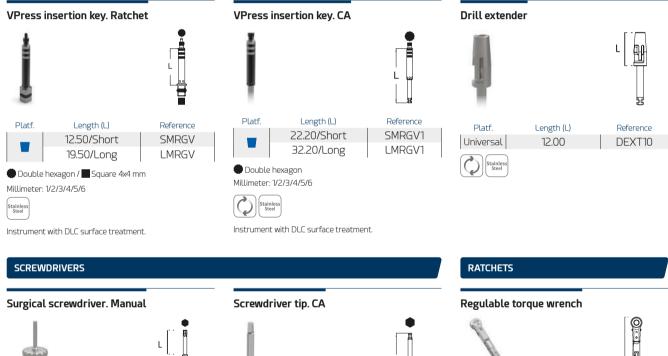
GTPD214

KSTPG120

Surgical instruments



KEYS



Platf.	Length (L)	Reference
	2.80/Mini	XSMSD *
Universal	9.50/Short	SMSD
Universal	14.50/Long	LMSD
	27.00/Extralong	XLMSD *
Hexagonal 1.25 mm		
Stainless 1,25mm		

 Platf.
 Length (L)
 Reference

 Universal
 20.00/Short
 MESD01*

 25.00/Long
 MESD

 Hexagonal 125 mm

 Image: State St



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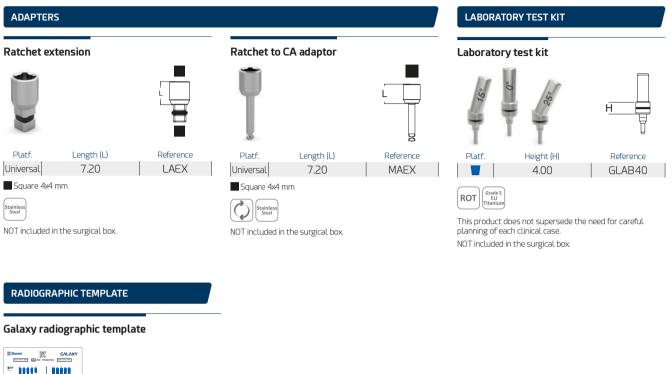
* Ref. MESD01 is NOT included in the surgical box.

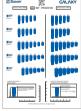
For more information on the use of surgical instruments see the section «Simplified surgical protocol» on page 50 of this catalogue.

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



Complementary instruments







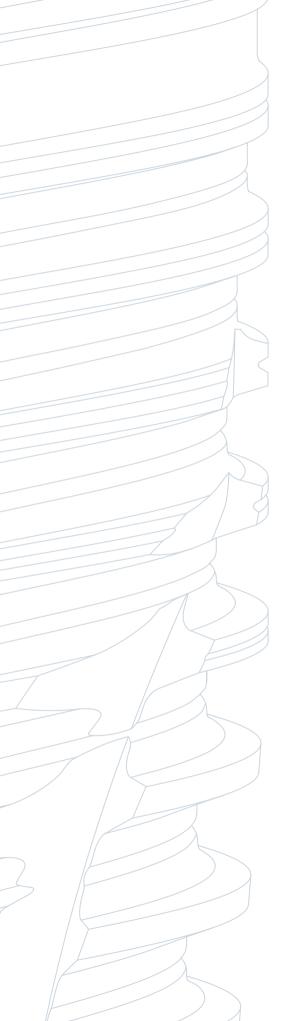
Scales 1:1 and 1:1.25

Material: transparent acetate. Non-sterilisable material.















Prosthetic instruments

Prosthetic box



Contents of prosthetic boxes available

Contents	Reference
Empty	BOXPN
Basic	BOXPSN
Complete	BOXPCN

134° \$\$\$

Material: Radel.

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



Contents of prosthetic boxes

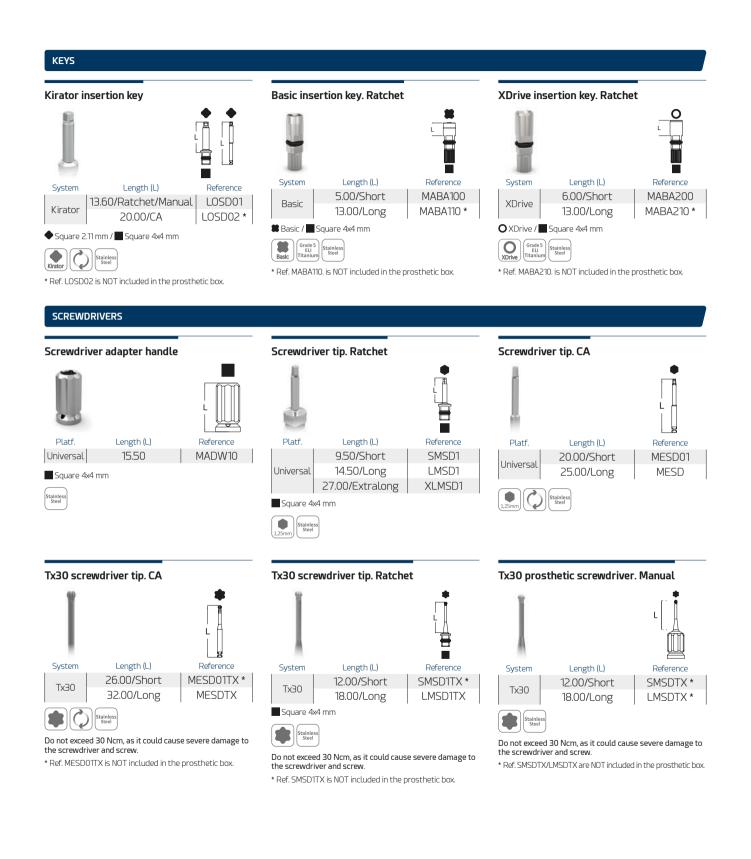
REF	Description	BOXPS	BOXPO
LOSD01	Kirator insert key. Ratchet		
MABA100	Basic insert key. Short. Ratchet. Grade 5 ELI titanium		
MABA200	XDrive insert key. Short. Ratchet. Grade 5 ELI titanium		
MADW10	Screwdriver adapter handle. 4x4. Manual		
SMSD1	Screwdriver tip. Ø1.25 mm. Short. Ratchet		
LMSD1	Screwdriver tip. Ø1.25 mm. Long. Ratchet		
XLMSD1	Screwdriver tip. Ø1.25 mm. Extralong. Ratchet		
MESD	Screwdriver tip. Ø1.25 mm. Long. CA.		
MESD01	Screwdriver tip. Ø1.25 mm. Short. CA.		
MESDTX	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		

S S

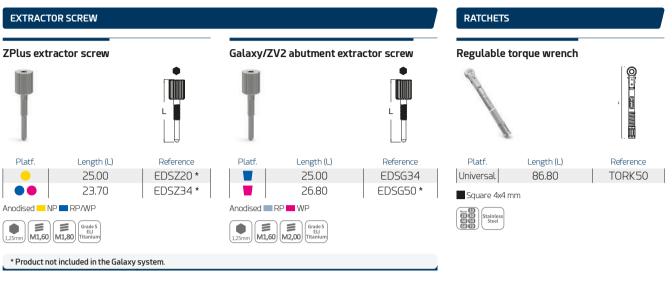
* Product not included in the Galaxy system.



47 🖉



Prosthetic instruments



Complementary instruments



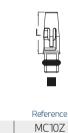


Platf.

Universal

Stainles: Steel

Square 4x4 mm



Extractor + Retainer inserter handle



Platf. A Length (L) B Length (L) Reference Kirator ZM-Equator 81.50 110.40 MBEI3610



NOT included in the prosthetic box.

Retainer inserter



Platf.	Length (L)	Reference
Kirator	32.00	MBEI3602
ZM-Equator	32.00	MBEI3603



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

Retentive joints instruments

NOT included in the prosthetic box.

Length (L)

12.00



Platf.	Measure	Reference
Universal	2x1	RREI0030

Pack of 10 units.

2 48

Simplified surgical protocol



Simplified 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 (milimeter drills). The drill tip is 0.5 mm long and is not included in the laser marked measurements.



14.5 mm 13 mm

11.5 mm 10 mm

8.5 mm

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.



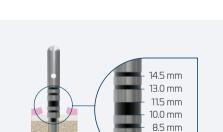
Check the depth of the surgical site, especially when not using drill stops. To check the surgical site axis, the paralleling pins are available in different diameters according to the drilling sequence.

VPress insertion keys

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

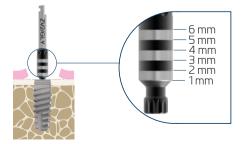




7.0 mm 6.5 mm

Ø3.70

Depth within the implant platform marked on the insertion keys





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DETAILS INSIDE THE GALAXY SURGICAL BOX



Recommendations on the maximum insertion torque



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.

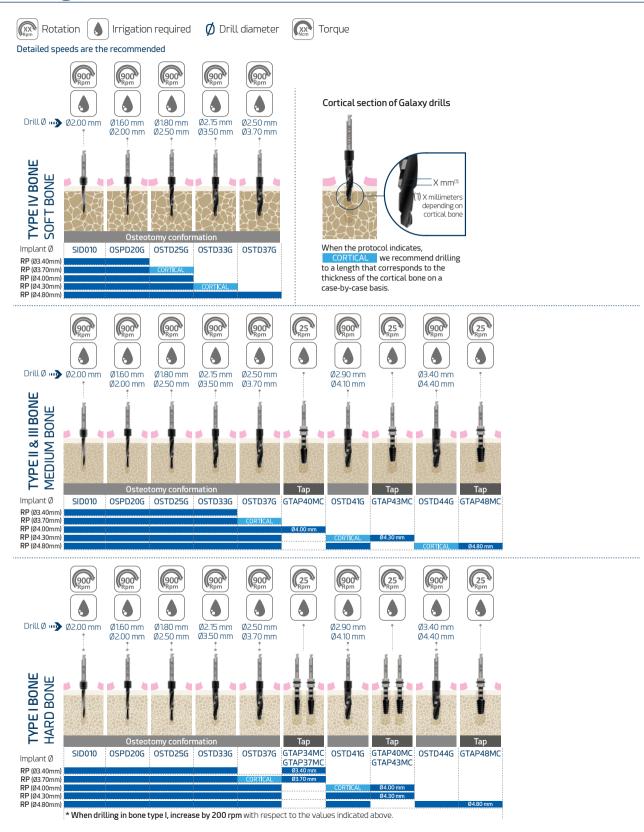
Control the final insertion torque with the adjustable dynamometric ratchet Ref. TORK50 or a contra-angle handpiece.

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.

Simplified surgical protocol

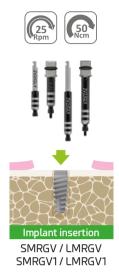
Drilling protocol - Ziacom® No Mount





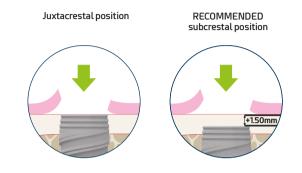
Galaxy implant insertion

Implant insertion



Crestal placement of implant

The drilling protocols are described so that the platform for the Galaxy implants is juxtacrestal. Nevertheless, recommendations are to leave the platform slightly subcrestal.



Bone types

Lekholm and Zarb classification (1985)



TYPE IV BONE - SOFT BONE

• Thin cortical layer surrounding a lowdensity trabecular bone.



TYPE II & III BONE - MEDIUM BONE

- Type II: thick layer of compact bone
- surrounding a dense trabecular bone.
- Type III: thin cortical layer surrounding a dense trabecular bone.



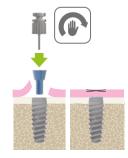
TYPE I BONE - HARD BONE

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 Composed almost entirely of homogeneous compact bone.

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.



Simplified surgical protocol

General recommendations

To consider during the intervention

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.

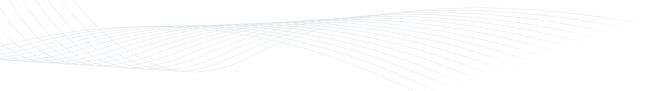
Damaged instruments must be disposed of according to local regulations.

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

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®.





Cleaning, disinfection and sterilisation



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:

- 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.

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IMPORTANT

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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

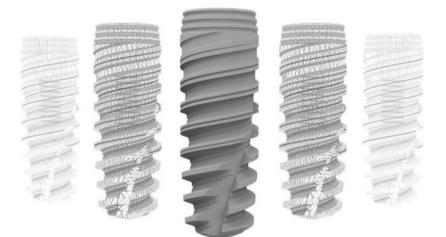
- Store the products in their original packaging and in a clean, dry location until they are used.
- After sterilisation, keep the products in the sealed sterilisation bags and in a clean, dry location.
- Never exceed the use by date indicated by the manufacturer of the sterilisation bags.
- Always follow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