







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

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

ZIACOM[®], Zinic[®], Zinic[®]MT, ZMK[®], ZMR[®], ZM1[®], ZM1[®]MT, ZM4[®], ZM4[®]MT, ZM8[®], ZM8[®]N, ZM8[®]S, Galaxy[®], ZV2[®], Ziacom[®]3D, Kiran[®], Kirator[®], ZM-Equator[®], Basic[®], XDrive[®], ZiaCam[®], ZIACOR[®], Tx30[®], Zellplex[®], DSQ[®], Titansure[®] and Ziasure[®] are some of the trademarks registered of Ziacom Medical SLU. Consult the website for the full list and their corresponding logos.

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▲The Company

Making future together

ZIACOM[®] is a **Spanish multinational company** specialising in the design and manufacture of dental implants, abutments and surgical instruments and providing **top-quality**, **integral solutions to dental professionals**.

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

In 2015. ZIACOM® introduced its **diversification strategy** with the development of **new lines of business and product families** and the launch of two of our most emblematic implant models – **ZM4**® and **Zinic**®. This helped us reach a **15% share of the Spanish market** by 2016 with the sale of over 230.000 implants.

The following year, the company embarked on an **ambitious programme of international expansion**, which has been updated and broadened in 2021.

ZIACOM[®] excellence of quality

A **constant search for quality** has been one of the cornerstones guiding our development since the company's creation and is one of our **main hallmarks**.

That's why ZIACOM® has state-of-the-art technology which we use 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 exclusively using high-quality raw materials and after applying strict controls to the selection of our main suppliers.

Ziacom Medical SLU is a **licensed manufacturer of medical devices** and an AEMPS (Spanish Agency for Medicines and Medical Devices)

6425-PS **commercial authorisation holder**. Our **quality management system is certified** in accordance with the requirements of ISO standards 9001:2015 and 13485:2016. and is also GMP 21 CFR 820 compliant.



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

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

Zitium® titanium

We ensure maximum quality by manufacturing all of our implants in extra-high-strength, grade 4 Zitium[®] titanium which bestows them with a substantially improved elastic limit and mechanical properties.

Properties of Zitium® titanium



Thanks to **Zitium**[®] titanium, our implants meet the requirements of standards ASTM F67 and ISO 5832-2 and are certified in accordance with Directive 2007/47/EC on medical devices amending 93/42/EEC 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

Loyal to our pioneering spirit, we **invest heavily in continual research and training** as a vehicle for providing the **sector with scientific support** and developing **new solutions and products** to satisfy our clients' demands.

Accordingly, in recent years we have **increased our R&D budget by 50%** in order to optimise production process efficiency. We have also incorporated a team of **dentists as product specialists** and conducted studies with implantology experts working in the field.

This dedication to innovation spurred us to **retrofit our central facilities** in 2018. So our headquarters are now housed in a **smart building** that was specifically designed and constructed as a **biomedical equipment manufacturing plant**. The facilities, covering **4.000** m² of usable floor area, accommodate the group's central services and all of the **infrastructure** needed to perform the company's **research**, **production** and **training** functions.

Our commitment to research also manifests in our **support for scientific societies**, participation in **national and international conferences** and engagement in **research collaborations**.

ZIACOM® around the world

In line with our philosophy and **pioneering spirit**, ZIACOM[®] is immersed in a considerable **expansion programme** that we initiated in 2017. The aim is to develop our **international presence** in **previously consolidated areas** and open up other areas of **new growth**. By adapting to and obtaining the **appropriate country-specific certificates**, ZIACOM® has managed to export its **successful model** across the globe via various **distribution networks** and with products tailored to each market. ZIACOM® therefore strives to meet each country's **specific quality, regulatory and legal requirements** with respect to the registration and distribution of our products.

Headquarters



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Please consult the updated list of ZIACOM distributors at www.ziacom.es or email export@ziacom.es





▲Implants Zv2[®]

Characteristics

CONNECTION

- Conical connection: 11º morse taper with double internal hexagon.
- Conical sealing: no infiltration.
- Friction fit: no microfiltrations.
- RP and WP platforms.
- Platform switch: soft tissue formation and emergence profile shaping.

CORTICAL AREA

- Shoulder implant design for crestal bone placement.
- Slightly tapered core in coronal area: high cortical compression.
- Thread in reduction up to platform.
- 0.2 mm bevel (except for 3.40mm diameter implant whose bevel is 0.15 mm).

BODY

- Lead threads: provide stability during insertion with 0.8 mm thread pitch.
- Optimised cylindrical morphology: high primary stability.
- · Atraumatic apex: protects anatomical structures.

CYLINDRICAL DESIGN

- Versatile, suitable for all positions.
- Available in narrow 3.40mm diameter.
- Morphology allows surgical compatibility.



Dimensions of the implant's coronal section





Diameters and lengths available

				LENGTH (L)		
ØDIAMETER	Ø PLATFORM	6	8	10	12	14
RP 3.40	2.05					
RP 4.10	2.02					
WP 4.80	3.85					

When choosing the correct implant length, consider the overdrill due to the length of the drill tip:

Length of drill tip



Total implant dimensions



Length (L)

▲Implants Zv2[®]

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 Medical 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.

CHARACTERISATION OF THE TITANSURE® SURFACE TREATMENT

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 (%)
CK	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 (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 roughness profile featured peaks and valleys in the range of $3-4 \,\mu\text{m}$.



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].



▲Implants Zv2[®]

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

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 anal	vsis - Bone-to-im	plant contact (BIC)
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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





▲Implants Zv2[®]

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.

Tibansure

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.

ZIACOM



ZIACOM® No Mount

IMPORTANT

ZV2[®] 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.

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.

	Implante Dental ES Dental Implant EN	ZIACOT 333 S.I Miamij
MD ZV24110 Lor 2000000 Und RP Ø4.10X10mm ● VPress® (01)0843548124367/11 ■ VPress® www.ziacom.es ■ TEREER / ^{max} S S > TEREER / ^{max} S S S	Zahninghanta te Implant Dentaire m Implanto Dentaire m Implanto Dentairo m 7/0000011/00000010/2000000 000 0 0 0 0 0 0 0 0 0 0 0 0 0 0	ZIACOM MEDICAL, SIU Cale Búhos, 2 28320 Pinto - Madrid ESPAÑA Telf: +34 91 723 33 06 M MEDICAL USALIC E. 2nd Avenue, SiaLic E. 2nd Avenue, SiaLic J, FL 33131 - USA Tel: +1 (786) 224-0089

Description of the symbology used

- Image: Construction and notified body
 I
 - 💫 Do not resterilise
 - Do not use if the packaging is damaged
 - (2) Non-reusable product
 - Consult the instructions for use
 - Expiry date of the product
 - Date of manufacture
 - Product manufacturer
 - **RxOnly** Caution: federal law prohibits dispensing without prescription

 $rac{1}{2}$ For full details on the product presentation and instructions for use (IFU) see www.ziacom.es/ifus or scan the QR code on the box.

References: ZV2® with ZIACOM® No Mount - Titansure®/Titansure® Active

	IMPLANT					
	Ø (mm)	Ø Core (mm)	Length	Titansure®ref.	Titansure® Active ref.	
e			8.0	ZV23408	ZV23408A	-
S	2 40	700	10.0	ZV23410	ZV23410A	
Ń	3.40	2.00	12.0	ZV23412	ZV23412A	
			14.0	ZV23414	ZV23414A	
			6.0	ZV24106	ZV24106A	
4.10	4.10 3.40	8.0	ZV24108	ZV24108A	-	
		.10 3.40	10.0	ZV24110	ZV24110A	
			12.0	ZV24112	ZV24112A	
			14.0	ZV24114	ZV24114A	
			10.0	ZV24810	ZV24810A	
	4.80	4.10	12.0	ZV24812	ZV24812A	
			14.0	ZV24814	ZV24814A	



Metrics of 1.60 (RP) and 2.00 (WP)

Cover screw*



* Screw included with each implant.

Platform



(1) Height of inner cone (2) Diameter of the working platform

▲Implants Zv2[®]

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.
- UNDERDRILLING TECHNIQUE: comprises the preparation of the implant bed using a final drill bit with a smaller diameter than the implant core. Technique associated with a high insertion torque and an increase in primary stability.

IMPORTANT

Possible increased risk of bone necrosis due to pressure.

 SIMPLIFIED DRILLING TECHNIQUE: technique proposed by Coelho and Cols in 2013 (1). It consists of the use of pilot drill and final drill corresponding to the size of the implant. It reduces drilling sequence but with risk of bone necrosis due to thermal increase.

Periodontal chart

Implant diameter (1)

A RP	BRP	WP
Ø3.40 mm	Ø4.10 mm	Ø4,80 mm

(1) Diameters are available for analog platforms.

Coronal implant diameter







📍 For more information on implant size selection see the literature available at www.ziacomes/en/bibliography

▲How to use this catalogue



Product sheet Title, section and paragraph ABUTMENTS | Direct-to-implant restorations Product name 2nd stage and impr <u>Å</u> . Product Ť Product image line diagram Enter (4) 3.60 3.60 3.60 4.60 4.60 4.60 5.50 5.50 5.50 5.00 5 Placf. Height (H) Diameter (2) Reference 1 7.00 7.00 HAG7070RAT 1 7.00 7.00 HAG7070WAT 150 3.00 7.00 150 3.00 7.00 150 3.00 7.00 150 3.00 7.00 150 3.00 7.00 150 3.00 7.00 150 3.00 7.00 150 3.00 7.00 HAG3615A HAG3630A Plat. Height (H) 13.00 850/Short 13.00 850/Short 13.00 850/Short 11.00 800/Short 11.00 3.60 3.60 4.60 5.50 5.50 5.00 5.00 5.00 6.50 6.50 HAG3630A HAG3670A HAG4675A HAG4670A HAG4670A HAG4670A HAG5575A HAG5530A TCG3601 TCG4600 TCG4601 TCG5500 TCG5501 TCG5500 Product table: - Platform - System - Height (H) HAGSSSO - Diameter (Ø) Product HAGSOBOA HAGSOSOA - Reference characteristics Pick-Up im on abutment screw - Quickly Screws Pick-Up impres H_____ 8 -0.00 3.00 6.00 9.00 0.00 3.00 6.00 9.00 Platf. Height (H) Paderence 7.25 CPU3410 PUG3400 3.00 All the dimensions given in this catalogue are expressed in Additional millimetres (mm) information

Key to symbols

Symbol	Meaning	Symbol	Meaning	Symbol	Meaning
ROT	Rotatory element	MX.XX	Metric in millimetres	Cobalt Chromium	Made from cobalt chromium
NOROT	Non-rotatory element	45°	45° screw support	PEEK	Made from PEEK
	Use with manual torque (consult table on page 39)	90°	90° screw support	Full castable	Made from castable plastic
XX Ncm	Maximum operating torque	\Diamond	Use in rotation with a CA	Plastic	Made from plastic
Ncm 10 20 30 40 50 60 70	Ratchet torque range	Rpm	Maximum rotation speed	XX° SSS	Recommended sterilisation temperature
1.25mm	Screw connection	XX USES	Maximum number of uses	Non sterile	Non-sterile product
Kirator*	Kirator [®] connection	(Single-use product		Use with abundant irrigation
Basic [®]	Basic [®] connection	Grade 5 ELI Titanium	Made from grade 5 ELI (extra-low interstitial) titanium	∑xx₀	Maximum angle
O XDrive*	XDrive [®] connection	Stainless Steel	Made from stainless steel		
	Tx30® connection	Co-Cr +castable	Made from cobalt chromium + castable plastic		



Abutments Direct-to-implant restorations



. _ _ _ _ _ _ _ _

atomic healing abut	tment	Customize healing abutment	Improcession abutment
*		5	inipression abutment
Height (H) Diar 1.50 1.50 3.00 1.50 1.50 1.50 3.00 1.50 3.00 1.50 3.00 1.50 3.00 1.50 3.00 1.50 1.50 1.50 3.00 1.50 1.50 1.50 3.00 1.50 1.50 1.50 3.00 1.50 3.00 1.50 1.50 1.50 3.00 1.50 3.00 1.50 3.00 1.50 3.00 1.50 3.00 1.50 3.00 1.50 3.00 1.50 3.00 1.50 3.00 1.50 3.00 1.50 3.00 1.50	meter (v) Reference 3.60 HAG3615A 3.60 HAG3650A 3.60 HAG3650A 3.60 HAG3650A 3.60 HAG3650A 3.60 HAG3650A 3.60 HAG3650A 4.60 HAG4615A 4.60 HAG4650A 4.60 HAG4650A 4.60 HAG4650A 5.50 HAG5515A 5.50 HAG5510A 5.50 HAG5510A 5.50 HAG5510A 5.50 HAG5510A 5.50 HAG5510A 5.50 HAG550A 5.50 HAG5015A 5.00 HAG501A 5.00 HAG5070A 5.00 HAG5070A 5.00 HAG5070A 6.50 HAG5515A 6.50 HAG550A 6.50 HAG550A	Platf. Height (H) Diameter (Ø) Reference 7.00 7.00 HAG7070R 7.00 7.00 HAG7070W NO (I) (I) (I) Includes screw. (I) (I) (I)	AT Platf. Height (H) Diameter (Ø) Referen /AT 13.00 3.60 TCG36 Image: State St

Impression abutment screw - Quickly Screws

		H
Platf.	Height (H)	Reference
	0.00	LTSS4000G
	3.00	LTSS4001G
	6.00	LTSS4002G
	9.00	LTSS4010G
	0.00	LTSS5000G
	3.00	LTSS5001G
	6.00	LTSS5002G
	9.00	LTSS5010G
Anodised		



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 Э.00 PUG5000 Anodised 🔤 RP 💻 WP



Pick-Up impression abutment







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

▲ ABUTMENTS | Direct-to-implant restorations

2 nd stage and impressions	Fixing elements	Provisional				
Z2Plus® Snap-On impression abut	tment Z2Plus	® Snap-On impres	sion transfer	Implant a	nalogue	
			H O			
Platf. Height (H) Length (L) Re Image: Constraint of the state o	eference Platf. RPG10 WPG10	Height (H)	Reference ZPU3400	Platf.	Length (L) 12.00 12.00	Reference IAG3400 IAG5000
Anodised RP WP	Pack of 4	units. DO NOT sterilise in	an autoclave. Sculptable.	NO ROT () 3D implar	Stainless Steel	
IMPORTANT Use the laboratory screw to tighten this impro-	ession			Platf.	Length (L)	Reference

abutment.

Fixing elements

Kiran® clinical screw

Platf.	Length (L)	Reference			
	8.20	DSG4010			
	10.40	DSG5010			
G30 1,25mm M1,60 M2,00 45° C Grade 5 EU Titanium					

Special Kiran® screw with surface treatment.

Laboratory screw

10



45°

Grade 5 ELI Titaniur

S M1,60 **M**2,00 NOT apt for use as the final clinical screw.

Kiran® Tx30® clinical screw

12.00

12.00



NO



IAG3400D

IAG5000D

For abutments and Ti-Base ZiaCam® Tx30®

Platf.	Length (L)	Reference			
	7.55	DSG4010TX			
	8.65	DSG5010TX			
Image: State					

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

▲ ABUTMENTS | Direct-to-implant restorations

Provisional abutment Abutments for aesthetic and immediate loading Image: Constraint of the sector
Provisional abutment Abutments for aesthetic and immediate loading Image: Constraint of the second
Abutments for aesthetic and immediate loading Image: Constraint of the second
Platf. Height (H) Length (L) Reference 1.50 10.50 RUGP3615 3.00 12.00 RUGP5015 3.00 12.00 RUGP5030
Rotatory Platf. Height (H) Length (L) Reference 1.50 10.50 RUGP3615 3.00 12.00 RUGP3630 1.50 10.50 RUGP5015 3.00 12.00 RUGP5030 801 3.00 12.00 RUGP5030
Platf. Height (H) Length (L) Reference 1.50 10.50 RUGP3615 3.00 12.00 RUGP3630 1.50 10.50 RUGP5015 3.00 12.00 RUGP5030 3.00 12.00 RUGP5030
Grade S ELI TRanium
Non-rotatory
Platf. Height (H) Length (L) Reference Image:

All provisional abutments come with an anodised screw.

ABUTMENTS | Direct-to-implant restorations



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.



▲ ABUTMENTS | Direct-to-implant restorations

	mpressions	Fixing ele				wed Cer	mented	Overdenture		
Cemented										
Anatomic straight abutment			Anatomic 15° angled abutment				Anatomic 25° angled abutment			
			ł		↓ ↓ ↓ Hg			1	L Hg	
Platf. Height (Hg/Ht) L 1 1,50/2.50 1 1 3,00/4.00 1 1 1,50/2.50 1 1 1,50/2.50 1 1 3,00/4.00 1 1 3,00/4.00 1 1 1,50/2.00 1 1 1,50/2.50 1 1 1,50/2.50 1	ength (L) Diamet 9.00 3.6 10.50 3.6 9.00 4.6 10.50 5.5 10.000 5.5 8.500 5.5 8.500 5.5 8.500 5.5	er (Ø) Reference 50 STG3615 50 STG4615 50 STG4615 50 STG5515 50 STG5530 50 STG5015	Platf. Height (Hg/F 1.50/2.50 3.00/4.00 1.50/2.50 3.00/4.00 3.00/4.00 3.00/4.00 3.00/4.00 3.00/4.00	t) Length (L)) 9.00) 10.50) 9.00) 10.50) 10.50 WP	Diameter (Ø) 3.60 3.60 4.60 4.60 5.00	Reference A1G3615 A2G3615 A1G4615 A2G4615 A2G5015	Platf. F	Height (Hg/Ht) Length (1.50/2.50 9.00 3.00/4.00 10.50 1.50/2.50 9.00 3.00/4.00 10.50 3.00/4.00 10.50 1.50 8.00 1.50 9.00 1.50 9.00 3.00/4.00 10.50 1.50 9.00 1.50 9.00 1.50 9.00 1.50 9.00 1.50 9.00 1.50 9.00 1.50 9.00 1.50 9.00 1.50 9.00 1.50 10.50) Diameter (3.60 3.60 4.60 4.60	 Reference A1G3625 A2G3625 A1G4625 A2G4625 A2G4625
Image: 3.00/4.00 Image: 3.00/4.00 Image: 3.00/3.50 Anodised RP Image: NOT Image: 3.00/3.50	10.50 5.0 8.50 6.5 10.00 6.5 1,60 M2,00 45	00 STG5030 00 STG6515 00 STG6530 00 STG6530		n M1,60 M2,0		Titanium				

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

ABUTMENTS | Direct-to-implant restorations



▲ ABUTMENTS | Direct-to-implant restorations

						Digital CAD-CAM
Digital CAD-CA	Μ					
ZiaCam® scanbody to i	mplant	ZiaCam®	° scanbody to imp	olant		
1			Î			
Platf. Length (L) 9.00 9.00 9.00 9.00	Reference FNSYG40 FNSYG50	Platf.	Length (L) 8.00 8.00	Reference FNSYG40T FNSYG50T		
NO ROT (1.25mm) (1.25mm) (1.400 (N Indicated for the laboratory.	E2,00 PEEK	Anodised	RP WP	0 PEEK Grade ELI Titaniuu	ŋ	

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



▲ ABUTMENTS | Direct-to-implant restorations

2nd stage and impression

Digital CAD-CAM

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
	1.00/5.50	4.80	FRUG505
	2.00/6.50	4.80	FRUG515
	3.00/7.50	4.80	FRUG530
	1.00/5.50	6.30	FRUG605
	2.00/6.50	6.30	FRUG615
	3.00/7.50	6.30	FRUG630
ROT	30 125mm S M1.60	M2.00	Grade 5 ELI Titanium

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		
	1.00/5.50	4.80	FNUG505		
	2.00/6.50	4.80	FNUG515		
	3.00/7.50	4.80	FNUG530		
	1.00/5.50	6.30	FNUG605		
	2.00/6.50	6.30	FNUG615		
	3.00/7.50	6.30	FNUG630		

All Ti-Base ZiaCam $\textcircled{\sc or}$ abutments come with a special Kiran $\textcircled{\sc or}$ screw with surface treatment Ref. DSG4010.



Ш - а

Rotatory

Non-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			
	1.00/6.50	4.40	FRUG405TX			
	2.00/7.50	4.40	FRUG415TX			
	3.00/8.50	4.40	FRUG430TX			
	1.00/6.50	4.80	FRUG505TX			
	2.00/7.50	4.80	FRUG515TX			
	3.00/8.50	4.80	FRUG530TX			
	1.00/6.50	6.30	FRUG605TX			
	2.00/7.50	6.30	FRUG615TX			
	3.00/8.50	6.30	FRUG630TX			
ROT (30) (1) (1) (1) (1) (1) (1) (1) (1) (1) (1						
$\begin{bmatrix} 30^{\circ} \\ 20^{\circ} \end{bmatrix} \begin{bmatrix} 20^{\circ} \\ 1 \end{bmatrix} $						

ZiaCam[®] scanbody to Ti-Base ZiaCam[®]

Digital CAD-CAM



 \star (2) Model 1 (Mod.1) must be used with the 3.80 mm diameter Ti-Base ZiaCam®..

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
	1.00/6.50	4.40	FNUG405TX
	2.00/7.50	4.40	FNUG415TX
	3.00/8.50	4.40	FNUG430TX
	1.00/6.50	4.80	FNUG505TX
	2.00/7.50	4.80	FNUG515TX
	3.00/8.50	4.80	FNUG530TX
	1.00/6.50	6.30	FNUG605TX
	2.00/7.50	6.30	FNUG615TX
	3.00/8.50	6.30	FNUG630TX
D			

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°).

Ti-Base ZiaCam® products marked with * can only be used with 3.80 mm diameter ZiaCam® scanbody Mod. 1 Ref. FNSFEX201.

指 For more information on the recommendations for the use of interfaces in zirconia restorations see the literature available at www.ziacom.es/en/bibliography





Abutments Restorations using transepithelials



Basic[®] XDrive[®]

Basic®

Demonstrative sequence of use







👖 For more information on the use of abutments see the "Prosthetic procedure manual" available at www.ziacom.es/en/download-eng



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

Basic[®] healing abutment



System	Height (H)	Reference
Basic®	5.00	BAHAEX34
Anodised R	P	



Basic® impression abutment



Rotatory





Non-rotatory



All Basic[®] impression abutments come with a screw.

Rotatory

System Length (L) Basic® 13.00 ROT

Basic[®] analogue



Reference BAIAEX34

Non-rotatory



Basic® 3D analogue







M M1,80

4		
System	Length (L)	Reference
Basic®	4.30	BDSEI3400
Anodised R	D	

Kiran® Tx30® Basic® clinical screw



Special Kiran® Tx30® screw with surface treatment.

Basic[®] provisional abutment

System	Length (L)	Refer
Basic®	8.50	BAR
Anodised R F)	

Length (L)

9.00

Full astable



Basic[®] UCLA

System

Basic®

ROT

ence UT10

Reference

BARUEX34



Basic[®]

25

Kiran® Basic® clinical screw



M Frade 5 ELI 45° M1,80

Special Kiran® screw with surface treatment.

Basic[®] laboratory screw



NOT apt for use as the final clinical screw.

Basic[®] provisional abutment





Reference

Reference

BANUP34

BARUP34

Rotatory

System Length (L) Basic® 8.50



Non-rotatory



Basic® mechanised base abutment + Castable abutment



Non-rotatory

NO ROT



Co-Cr

Reference BBNU34





👖 For more information on the use of abutments see the "Prosthetic procedure manual" available at www.ziacom.es/en/download-eng

Reference

XA310G30

XA410G30

XA510G30

Grade 5 ELI Titaniun



Includes the XDrive® abutment with sterilisable polyoxymethylene applicator (Tecaform AH-POM-C). 21° cone angle. 42° angle between abutments.

XDrive® healing abutment



System	Height (H)	Reference	
XDrive®	5.00	XH103400	
Anodised 🗖 RP			

ROT RELI

XDrive® impression abutme	l	t
---------------------------	---	---



Height (H) System XDrive[®] 10.50 Anodised 🔜 RP



Includes screw

XDrive® analogue



System

XDrive®

ROT

Reference

XT103411

L	
	Ľ.

Length (L) 13.00

Reference

XDrive® 3D analogue

inle

	0	
System	Length (L)	Reference
XDrive®	13.00	XIA103400D
ROT	Stainless Steel	









ZiaCam[®] scanbody to XDrive[®] abutment







ZiaCam[®] scanbody to XDrive[®] abutment

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



Includes special Kiran[®] screw with surface treatment Ref. XDS103411.

Ti-Base ZiaCam® Tx30® XDrive®



Height (Hg/Ht) Reference XFRU341TX ר 90

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

ZiaCam[®] scanbody to Ti-Base XDrive[®] abutment





System Length (L) XDrive[®] 7.00

PEEK

NO ROT

Reference FNSFX11

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

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.

ATTENTION

System

XDrive[®]

ROT

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





▲ SURGICAL INSTRUMENTS

Surgical box

Surgical box

Contents of ZV2[®] boxes available

Contents of surgical boxes

Platf.	Contents	Reference
-	Empty	B0X920
-	Basic, manual/CA	BOX900SZV2
	Complete, manual/CA	BOX901ZV2

134° \$\$\$

Material: radel.

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





	-	2V2	2
		00S2	01Z/
DEE	Description	9X08	9X08
SID010	Lance drill. Ø2.00 mm		
0SPD22Z	Pilot drill. Ø16/2.00 mm. Millimeter		
OSTD28Z	Surgical drill. Ø1.80/2.50 mm. Millimeter		
OSTD35Z	Surgical drill. Ø3.50 mm. Millimeter	•	
OTD42Z	Surgical drill. Ø4.20 mm. Millimeter	•	
OTDZ1CA	Cortical drill. Ø3.50 mm	•	
OTDZ2CA	Cortical drill. Ø4.10 mm		
OTDZ3CA	Cortical drill. Ø4.80 mm	•	
CLD34	Crestal surgical drill. Ø4.10 mm		
CLD50	Crestal surgical drill. Ø5.10 mm		
PMT1G	Paralleling pin. ZV2®. RP. Manual. Grade 5 ELI titanium		
PMT2G	Paralleling pin. ZV2®. WP. Manual. Grade 5 ELI titanium		
VTPD106	Calibrated drill stop. ZV2®. 3. H6 mm. Grade 5 ELI titanium		
VTPD108	Calibrated drill stop. ZV2®. 3. H8 mm. Grade 5 ELI titanium		
VTPD110	Calibrated drill stop. ZV2®. 3. H10 mm. Grade 5 ELI titanium		
VTPD112	Calibrated drill stop. ZV2®. 3. H12 mm. Grade 5 ELI titanium		
VTPD114	Calibrated drill stop. ZV2®. 3. H14 mm. Grade 5 ELI titanium		
VTPD210	Calibrated drill stop. ZV2®. 4. H10 mm. Grade 5 ELI titanium		
VTPD212	Calibrated drill stop. ZV2®. 4. H12 mm. Grade 5 ELI titanium		
VTPD214	Calibrated drill stop. ZV2®. 4. H14 mm. Grade 5 ELI titanium		
VTAP34M	Surgical tap. ZV2®. RP. Ø3.40 mm. Millimeter. CA/Manual		
VTAP41M	Surgical tap. ZV2®. RP. Ø4.10 mm. Millimeter. CA/Manual		
VTAP48M	Surgical tap. ZV2®. WP. Ø4.80 mm. Millimeter. CA/Manual		
MUR100V2	Depth gauge/Paralleling pin. Ø2.20 mm. Millimeter. Grade 5 ELI titanium		•
MUR200V2	Depth gauge/Paralleling pin. Ø2.20/2.80 mm. Millimeter. Grade 5 ELI titanium		
MUR300V2	MDepth gauge/Paralleling pin. Ø3.50 mm. Millimeter. Grade 5 ELI titanium		
MUR400V2	Depth gauge/Paralleling pin. Ø4.20 mm. Millimeter. Grade 5 ELI titanium		
SMRGV	VPress® insertion key. Millimeter. Ratchet		
LMRGV	VPress® insertion key. Long. Millimeter. Ratchet		
SMWGV	VPress® insertion key. Short. Millimeter. Ratchet		
SMRGV1	VPress® insertion key. Short. Millimeter. CA		
LMRGV1	VPress® insertion key. Long. Millimeter. CA		
SMWGV1	VPress® insertion key. Short. Millimeter. CA		
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		
TORK70	Regulable torque wrench. 10/20/30/40/50/60/70 Ncm		

▲ SURGICAL INSTRUMENTS



Grade 5 ELI Fitaniur

▲ SURGICAL INSTRUMENTS

						Depth gauges	Keys	Screwdrivers	Ratchets
Dept	Depth gauges								
ZV2® depth gauge/Paralleling pins									
Platf.	Diameter (Ø)	Length (L)	Reference						
	2.20	26.00	MUR100V2						
	2.80	27.00	MUR200V2						
	3.50	26.00	MUR300V2						
	4.20	26.00	MUR400V2						
Millimeter	: 6/8/10/12/14								

VPress[®] insertion key. CA

Grade 5 ELI Titanium

Keys

VPress® insertion key. Ratchet



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

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

🖠 For more information on the use of surgical instruments see the section "Surgical protocols" on page 54 of this catalogue.

Drill extender

▲ SURGICAL INSTRUMENTS | Complementary instruments



Laboratory test kit

Laboratory test kit



ROT Grade 5 ELI Titanium

This product does not supersede the need for careful planning of each clinical case. NOT included in the surgical box.

Radiographic templates

ZV2® radiographic template



Scales 1:1 and 1:1.25

Material: transparent acetate. Non-sterilisable material

See the literature available at www.ziacom.es/en/bibliography





Prosthetic instruments



▲ PROSTHETIC INSTRUMENTS

Prosthetic box

atchots Kovs

Extractor screw

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

		XPSN	XPCN
REF	Description	B	BO
TORK50	Regulable torque wrench. 10/20/30/40/50/60/70 Ncm		
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	٠	
EDSZ20 *	ZPlus®/Z2Plus® extractor screw. Zinic®. NP. Grade 5 ELI titanium		
EDSZ34 *	ZPlus®/Z2Plus® 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		

* Product not included in the ZV2® system.



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

▲ PROSTHETIC INSTRUMENTS										
					ewdrivers	Extractor screw				
Extractor screw										
ZPlus®/Z2P	lus® extract	or screw	(Galaxy®/ZV2® abutment extractor screw						
Ì										
Platf.	Length (L)	Reference	e	Platf.	Length (L)	Reference				
	13.50	EDSZ20)*		15.00	EDSG34				
	13.50	EDSZ34	*		15.00	EDSG50				
Anodised NP RP/WP And					Anodised 🔤 RP 🚾 WP					
(1.25mm) (1.60) (M	Grade 5 ELI Titanium		125mm M1.60 M2.00 Grade S ELI Titanium							

* Product not included in the ZV2 $^{\odot}$ system.

▲ PROSTHETIC INSTRUMENTS | Complementary instruments



Retentive joints instruments









▲Surgical protocol





traceability

The clinician must keep

with the product, for proper

in the patient's file the identification label supplied.

General recommendations

Suplementary instrument



Depth gauge/Paralleling pin

Check the surgical site depth, especially if stoppers were not used.

To check the surgical site axis, the paralleling pins have different diameters according to the drilling sequence.

Consider during intervention



Surgical drills should be inserted in the

contra-angle with the surgical motor stopped, ensuring correct anchoring and rotation before starting drilling. Treat the drills with great care: the slightest damage to the tips can compromise their effectiveness.

Cover screw handling



Position the cover screw on the screwdriver. Approach the implant by avoiding accidental dropping and ingestion of the screw. Insert it into the implant with manual torque and clockwise.

Maximum insertion torque and speed

The recommended insertion torque is between **35 and 50 Ncm** according to each case without being limited to a single torque



The Implant placement should be performed with controlled torque and according to the density and bone.

Insertion instruments or contra-angle (CA) screwdrivers use maximum speed of:



Healing abutment placement

Damaged

instruments

must be disposed

local regulations.

Second phase surgical procedure

of according to

Each instrument

must be used only

for the specific use

recommended by

the manufacturer



The healing abutment should correspond to the implant platform, considering the option of applying the platform switch technique with anatomical abutments and be in accordance with the height of the gingival tissue to avoid abutment occlusion. Excessive height could expose the implant to premature loading, compromising the osseointegration process.

IMPORTANT WARNINGS

About implant placement

Excessive compression to the bone can lead a non-osseointegration of the implant.

Failure to follow the steps described in the surgical sequence may result in:

- Lack of primary stability due to loss of support bone.
- Difficulties during the implant placement.

Exceeding the torque (50 Ncm) at the implant insertion can produce:

- Irreversible distortions in the
- internal/external connection.
- Irreversible deformations in the instruments indicated for insertion of the implact
- the implant.
- Difficulty of disassembling the instrument/implant assembly

ZV2® implants

ZIACOM® surgical protocol establishes a crestal position of the implant platform.

To avoid cortical stress and deformation of the key and/or connection of the implant, insertion with contra-angle (CA) must respect the maximum recommended rpm (25 Rpm) and the maximum indicated torque (50 Ncm).

If resistance is encountered during insertion, it is recommended to turn the insertion anticlockwise and after seconds of pause continue with the insertion. Repeat this process as many times as necessary.

Check the final insertion torque with the regulable torque wrench Ref. TORK 70 or with CA

Make sure that the entire implant with **Titansure® / Titansure® Active** surface treatment is completely covered with bone.

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.es/en/download-eng which explained the procedures, protocols and instructions for use before using the ZV2® 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 herein.

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.es/en/download-eng
 - ** 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 autoclaves

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 SLU 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.







Consulte las condiciones generales de venta actualizadas en nuestra página web www.ziacom.es

Consulte la disponibilidad de cada producto por país.

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