

DAS ■

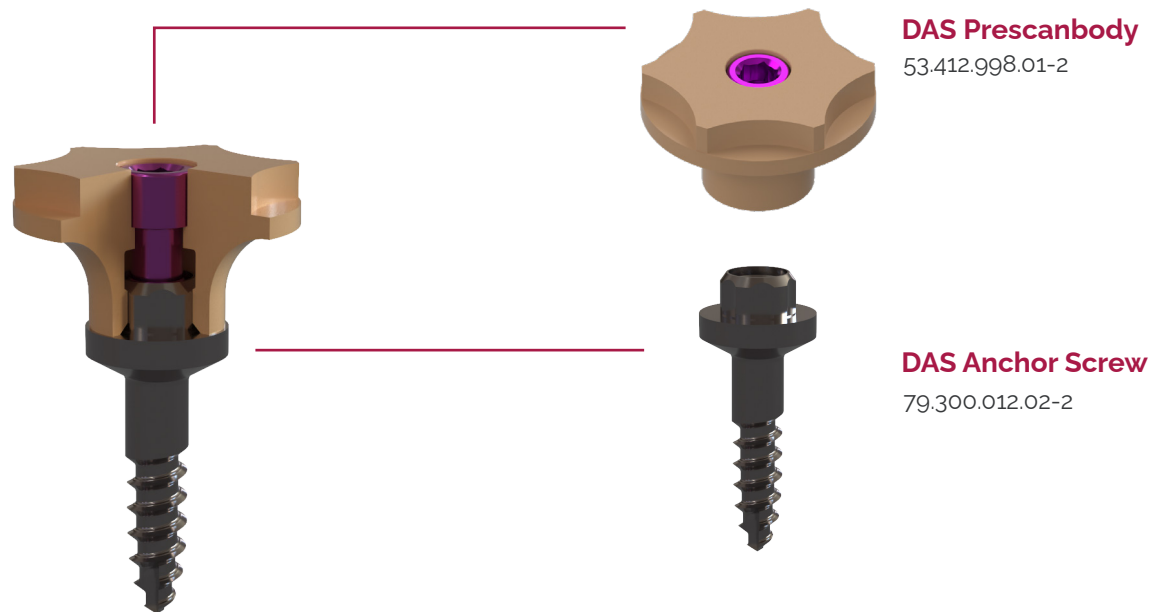
PRESCANBODY



DAS PRESCANBODY

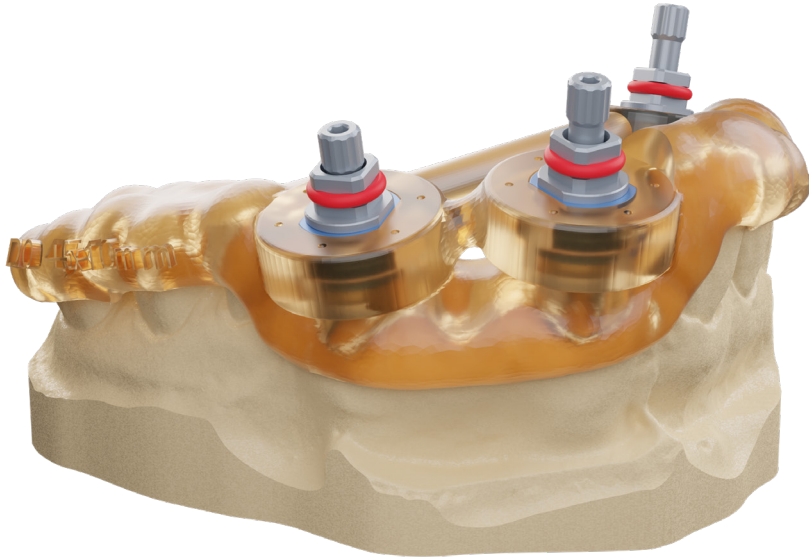
A method that provides an initial reference by means of a scan conducted prior to or following implant placement.

COMPONENTS OF THE PREOPERATIVE PHASE



PHASE ○ PLANNING

The clinician develops the treatment plan and designs the surgical guide.



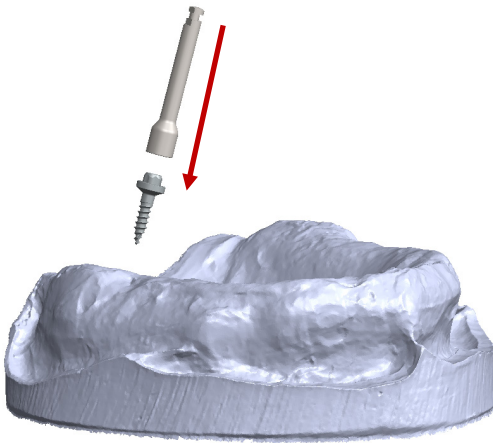
SURGICAL GUIDE DESIGN

Create the **surgical guide** according to the preoperative planning using the appropriate guided surgery software and the correct library corresponding to the selected implant brand, model, diameter, and length.

PHASE 1 **PREOPERATIVE PREPARATION**

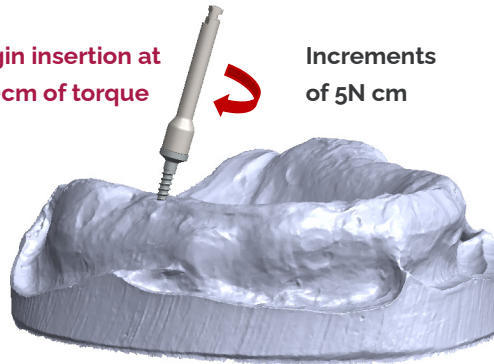
Once the surgical planning and guide design have been completed, the placement process is initiated prior to the surgery.

PLACEMENT OF THE DAS ANCHOR SCREW IN THE DESIGNATED AREA



Using the compatible screwdriver DAS MU (43.321.316.01-2), the **DAS Anchor Screw** must be placed vertically on the cusp of the arch and progressively screwed in, following the torque protocol outlined.

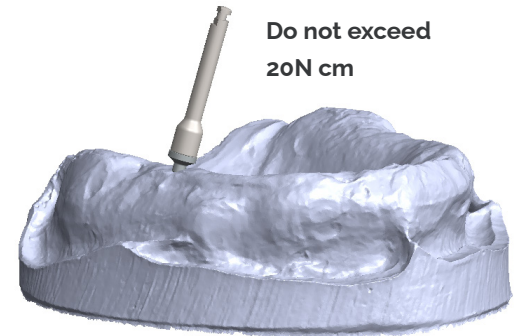
**Begin insertion at
5 N·cm of torque**



**Increments
of 5N cm**

Begin with a torque of 5 N cm until the screw no longer turns freely. Continue increasing the torque in 5 N cm increments until the **DAS Anchor Screw** is fully threaded.

**Do not exceed
20N cm**



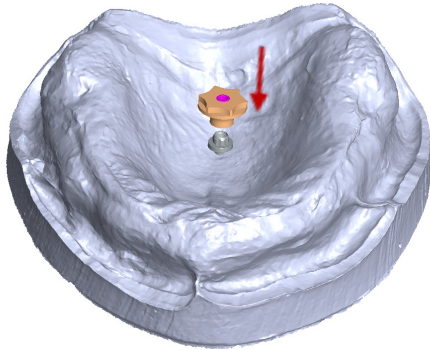
Throughout the entire process, the maximum torque of 20 N cm must not be exceeded.

NOTES

- Do not exceed the maximum torque of 20 N·cm to avoid bone damage or loss of system stability.
- If during the insertion of the component the required position is not reached without exceeding 20 N·cm torque, stop, carefully remove the screw, and reposition it again at 20 N·cm.
- It is recommended to have the surgical guide ready at this stage (in accordance with phase 3).

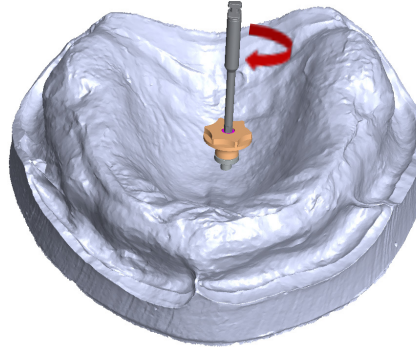
PHASE 2 PREOPERATIVE SCAN

Step 1



Place the **DAS Prescanbody** onto the DAS Anchor Screw, ensuring it is properly seated.

Step 2

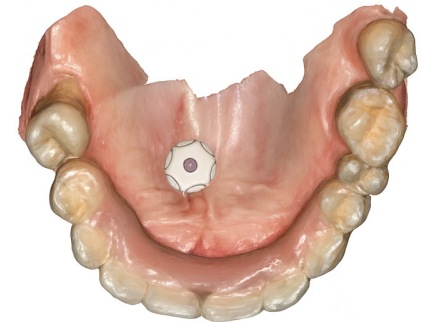


Secure the **DAS Prescanbody** to the DAS Anchor Screw using the compatible screw provided, applying a maximum torque of 5 N·cm.

Use a clinical mirror to verify that there are no visible gaps between the base of the **DAS Prescanbody** and the surface of the DAS Anchor Screw.

If any separation is detected, check the correct placement of the **DAS Prescanbody**.

Step 3



Proceed with the preoperative scan to record the exact location of the devices.

Perform a complete scan of the arch or arches.

Record the bite and preexisting anatomical conditions.

Place the designed surgical guide and proceed with the surgery.



Warning:

Excessive torque beyond the recommended limits may fracture or damage the screw.

NOTES

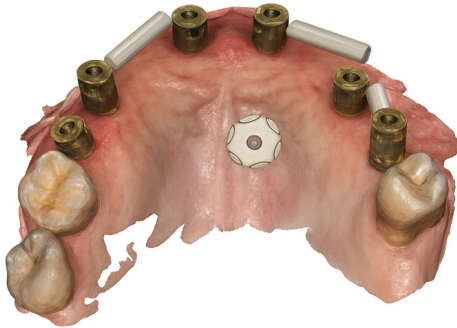
- If the DAS Prescanbody interferes with the placement of the surgical guide, it can be removed and placed after the surgery. Due to its geometry, it will always retain its original position.

PHASE 3 POSTOPERATIVE SCAN

Once the surgery is completed, **the DAS Prescanbody is scanned again.**

Step 1

Postoperative scan (POST)

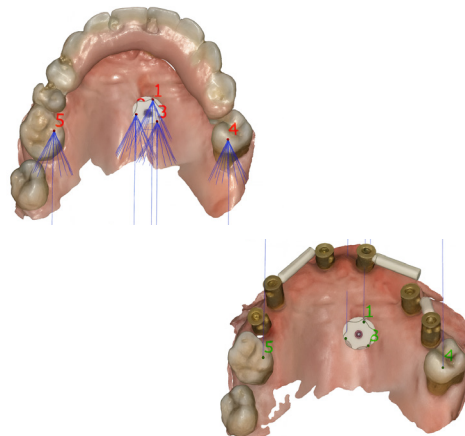


The results of this scan can be aligned with those of the preoperative scan using mesh alignment.

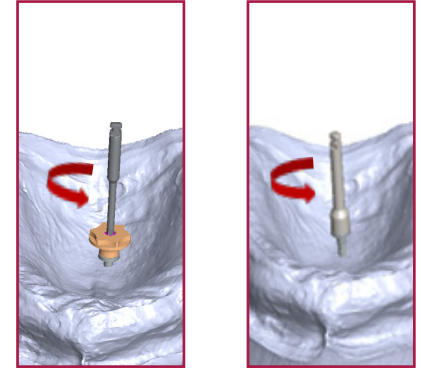
These results can be used to finalize the prosthetic design.

Step 2

Selection of matching points



Warning: Excessive torque beyond the recommended limits may fracture or damage the screw.



Step 3

Remove the Prescanbody and the DAS Anchor Screw.

For the DAS Anchor Screw, use the compatible DAS MU screwdriver, applying a maximum torque of 30 N·cm.

PRE-SCANBODY DAS COMPONENTS



Prescanbody

53.412.998.01-2



DAS Anchor Screw

79.300.012.02-2



Compatible Screwdriver
DAS MU

43.321.316.01-2



Screwdriver Hex. 1.20

43.601.103.02-2

TALLADIUM GUARANTEE

TERMS AND CONDITIONS

These guarantee terms and conditions ("T&C") cover the entire range of Talladium products ("Products"), manufactured by TALLADIUM ESPAÑA S.L. and distributed by Geoda Medical S.L. or official dealers. The guarantee described in these T&C is exclusively in benefit of the clinician ("Clinician") and of the dental technician ("Technician") and not for the benefit of third parties or institutions, including patients.

GUARANTEE PERIOD

TALLADIUM ESPAÑA S.L. offers a lifelong guarantee for its entire range of products starting from the date of issue of the invoice.

GUARANTEE SCOPE

Subject to the limitations and exceptions described in these T&C, TALLADIUM ESPAÑA S.L. will offer the following benefits:

QUALITY: If there are defects in the materials or in the manufacturing of the Product, TALLADIUM ESPAÑA S.L. will replace the Product with no additional cost.

SAFETY: If, having complied with all the product indications, the prosthesis should have to be made again, due to a fault in the Dynamic Abutment or Dynamic Titanium Base system, TALLADIUM ESPAÑA S.L. will replace the abutments and screws necessary to remake the prosthesis, as well as the costs derived from its manufacturing.

In case of having used our products and having complied with all the product indications, the implants suffer any damage, TALLADIUM ESPAÑA S.L. will pay the cost of the implants. This coverage will only be valid during the first 6 months after the collocation of the prosthesis which includes our products.

CLAIM REQUIREMENTS AND PROCEDURE

To receive the benefits indicated in these T&C, the treating Clinician must satisfy the following requirements:

- a) The claim must be notified to TALLADIUM ESPAÑA S.L. within (30) days since the date the claimed defect was detected.
 - b) This requires that the Clinician or Technician must contact the customer service department by telephone or by e-mail to make the claim.
 - c) A claim form will be completed, which, together with a document or report which justifies the faulty Product and the faulty Product itself, will be sent by the customer to TALLADIUM ESPAÑA S.L. offices, within the previously indicated period.
 - d) Clinicians or Technicians presenting a claim in agreement with these T&C must be up to date in any payments owing to TALLADIUM ESPAÑA S.L. or to any of its subsidiaries, at the time when the claim form is presented.
 - e) All the use procedures of our Products must be carried out in agreement with the instructions of TALLADIUM ESPAÑA S.L. as well as in accordance with commonly accepted dentistry practices.
 - f) The expenses derived from this procedure will be assumed by the customer. The return shipping costs will be assumed by TALLADIUM ESPAÑA S.L. in all those cases covered by these T&C.
- Regardless of the guarantee rights, claims should be notified as soon as possible in order to comply with regulatory requirements.

GENERAL LIMITATIONS OF THIS GUARANTEE

With the exception of the guarantee described in these T&C, neither TALLADIUM ESPAÑA S.L. nor its representatives, nor third parties manufacturing or distributing the Products, represent or offer a guarantee, agreement or any other express or implicit, oral or written, commitment, with respect to the Products (without limitation), including guarantees involved in the marketing, durability or suitability for individual uses or purposes. In addition and within the maximum extent permitted by the relative law, TALLADIUM ESPAÑA S.L. rejects (on its own behalf, and on behalf of its representatives and third parties that manufacture or distribute Products) any responsibility with respect to any direct or indirect damage caused, which may result from or be a consequence of the design, composition of the dental prosthesis into which the Products are integrated.

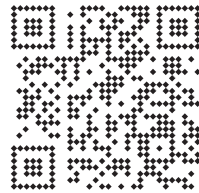
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TALLADIUM ESPAÑA S.L. limits this guarantee to:

- Transformed abutments that form part of the dental prosthesis. But not the screws used to anchor them.
- Those products that are not used with the accessories and parts marketed by Talladium España

AMENDMENT OR SUSPENSION OF THE GUARANTEE

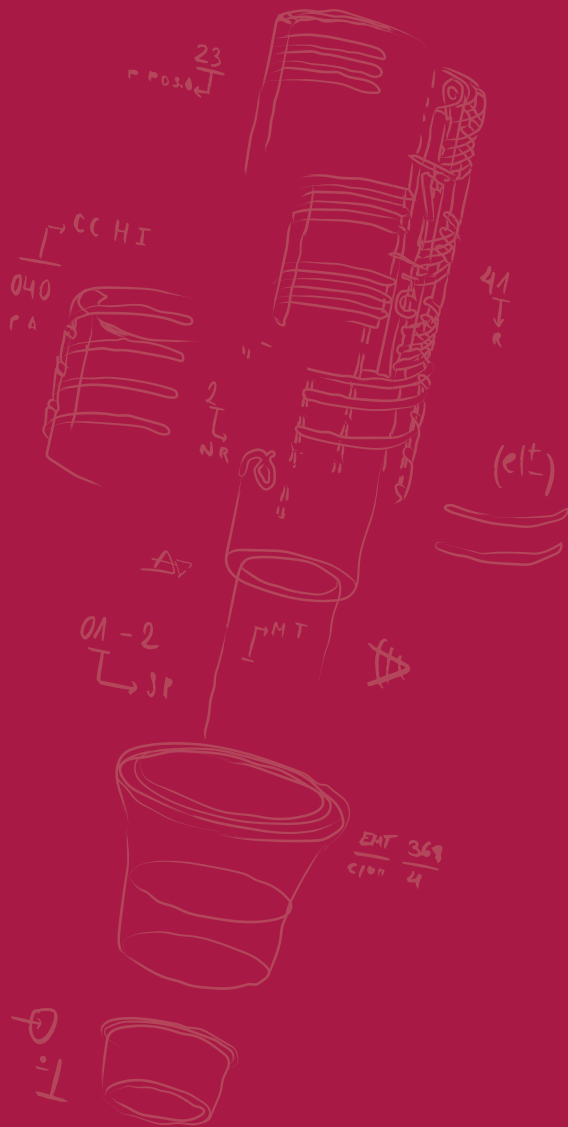
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