

| Product Information —— | | | | | | |
|--|--|-------------------------|-----------------------------------|------------------------------|------------------------|-----------------------------------|
| Ref: | | Description: | | | | Lot: |
| Costumer Information — | SPACE FOR PROD | DUCT TAG | | SPACE FOR RAI | | |
| Name/Corporate name: | | | | | | |
| ID Number: | | CNES: | | | | |
| Addrres: | | | | | City: | UF: |
| Phone: | Cel Phone: | | E-mai | l: | | |
| Patient Information ——Name/ID: | | | | | Age: | Gender: ☐ Male ☐ Female |
| Clinical history: | ☐ Diabetes ☐ Hyperten ☐ Xerostom ☐ Bruxism ☐ Smoking | sion | | sensitivity ne Deficiency | | □Another disease: |
| Surgery Information —— | | | | | | |
| Reason of return: | | | sseointegration mary stability | | □Impossib □Fracture | ole to install e |
| ☐ Others (describe): | | | | | Date: | |
| Fill in all the fields below | in case of Fail | ure in osseoi | integration/Lacl | k of primary s | stability. | |
| Implant date: | _ | Bone type: □I □II | | | Immedia | ate loading implant: □Yes □ No |
| Removal date: | | ⊒ ⊒ ⊒ V | | | | |
| Fill in all the fields below Used instrumental S.I.N.? | | | uence of borers | were used? _ | | |
| Was a bone graft performed | | ☐ Yes If so | o, what material | was used? | | |
| Intraoral implantation region | on (tooth numb | per)? | | | | |



Dear Dentist,

Any occurrence related to our products is of the utmost importance to us. Therefore, we ask that the completion and submission of information for our analysis be done thoroughly. This information will be essential for a detailed analysis of the occurrence.

PREMISE

1. Warranty Scope

- S.I.N Implant System guarantees to all dental institutions that have acquired, provided that they are original products, that the instructions for use have been respected, as well as the following described situations:
- 1.1 The legitimate acquisition of products by the dental institution;
- 1.2 The careful selection of the patient with clinical indication for treatment and appropriate application of the technique;
- 1.3 Informed and signed consent by the patient, with proper guidance from the dentist;
- 1.4 That the patient does not present any contraindications described in the instructions for use;
- 1.5 That the use of the product has been carried out in strict compliance with the guidance and recommendations described in the instructions for use of each product.



The warranty **DOES NOT COVER** products that are sent without the following documents:

- 2.1 Fully completed (original) Product Evaluation Form;
- 2.3 Implant radiographs.

We remind you that S.I.N.'s quality is recognized by the most important certifications in the segment, such as ISO 13485/2016, ISO 9001/2015, RDC 665/2022, and MDD 93/42/EEC and MDR 745/2017, which enables marketing in Europe. In addition to these, other certificates received over the past more than 20 years attest to the continuous improvement of our work.

Still, for your greater satisfaction, we have a direct channel with a scientific dental consultant in the product research and development area, whom you can call to ask questions or even clarify this and other cases further.

| Declaration of Truthf | ulness — | | | | |
|--------------------------|--------------------------|-------------|--|--|--|
| I, document is true. | | ,declare th | ,declare that the information provided in this | | |
| Date: | | | | | |
| Signature and stamp of t | the dental professional: | | | | |
| Quality Control Form | Completion — | | | | |
| SAP Cliente | Ocorrência | Pedido | Laudo de Análise | | |
| | | | | | |