

1. PRACTITIONER/PATIENT'S DETAILS

Customer Code (See PL and/or invoice):
 Name (who encountered the malfunction) :
 Patient ID : (optional)

THE FOLLOWING MUST BE ENCLOSED

- **Panorama or sectoral X-RAY** for each step of the treatment plan (at least post-malfunction)
- **The X-RAY of the last visit** before the occurrence of the malfunction.

2. TRACEABILITY

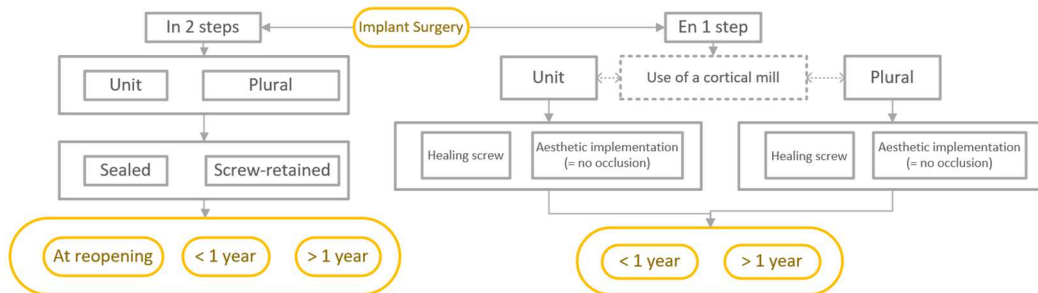
WARNING : the implant and associated prosthetic component shall be returned decontaminated, cleaned and sterilized

Site N°:.....
Malfunctioning implant: Reference: unknown Placement date
 Batch n°: unknown Removal date:
Prosthetic component associated*: Yes (fill in below) No (n/a)
 *to implant at the time of malfunction Reference: unknown Placement date:
 Batch n°: unknown Removal date:

3. TYPE OF MALFUNCTION

Blocking/Unsuitability Fracture Activation (3.0) Other:

4. SURGERY INFORMATION (tick below or n/a if not applicable) n/a (prothèse)



5. LOCALIZATION OF THE MALFUNCTION

Cover screw Implant collar Internal thread Implant connection Other:

6. PRE/PER/POST SURGERY SITUATION

Bruxism/Biomechanical excess Non-compliance to the instructions Periodontal disease Trauma / Accident
 Restricted mouth opening Overall good health Local infection
 Complications during the site preparation Other:

Bone quality : D1 D2 D3 D4 Unkown
 If non adequate bone volume: ROG Graft: type..... Brand..... & date

7. PROSTHESIS INFORMATION

Antagonist situation

Natural tooth Fixed implant supported prosthesis Removable braces
 Fixed tooth-supported prosthesis Missing tooth Non-restored curve of Spee

Material used on the restored tooth

Resin Laminate zirconium Hybrid material
 Ceramo-metal (CCM) Full zirconium Other:

Follow-up and monitoring of the patient

Nr of visits: Date of last visit: Occlusion monitoring and treatment during each visit: Yes No Unknown

Additional comments: