PFL

Technical specifications

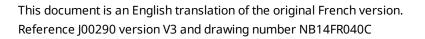
| Intended use | Periodontics | | | | | | |
|----------------------|---|--------------------|-------------------------|---|--|-------|--|
| Indications for use | Biofilm removal | | | | | | |
| Description | Left-angled tip for the pre-molar and molar sectors, complementing the PFR tip, for the interproximal scaling of thin periodontiums | | | | | | |
| Tip type | Smooth | | | | | | |
| | Tip made of su 12b | rgical stainless s | | | | | |
| Allergenic factor | Surgical stainless steel contains: | | | | | | |
| | nickelchromium | | | | | | |
| Commercial reference | PFL: F02171 | | | | | achia | |
| Medical class | Class of medical device: IIa according to 93/42/EEC directive | | | | | 8 | |
| Adjustments | Newtron range | P5 Range | Suprasson P5 Booster | | Prophy Max - P-Max - Implant Center 2 - Piezotome 2 | | |
| | 12 | 12 | 9 | 2 | | Ń | |

Service life

As it is not possible to establish a maximum number of uses (that may be determined by many parameters such as duration of use, hardness of tissue, the force applied, wear), we recommend that medical devices being routinely used are replaced at least once a year. Replace the medical device if the tip does not vibrate at the expected frequency, the treatment is not progressing as normal and is taking longer or is at a standstill.

Associated documentation

| Document title | References |
|--|------------|
| Consulting electronic user instructions | J00007 |
| Cleaning, disinfection and sterilisation instructions for tips | J02001 |
| Periofine clinical tips | J02171 |
| Ultrasonic generator power settings table | J58000 |
| General instructions relating to the complete range of standard ultrasonic dental tips | J02101 |

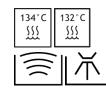




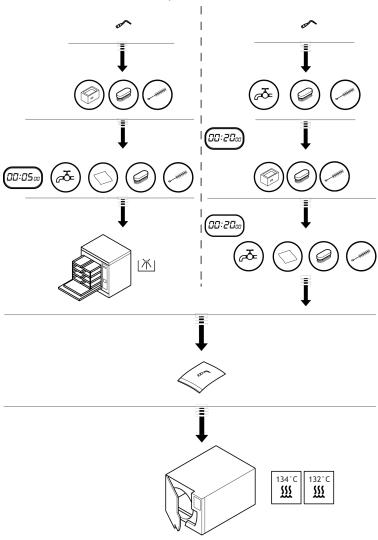


Sterilisation

Sterilise prior to each use



The tips must be cleaned, disinfected and sterilised prior to each use. Prior to each use, please refer to the cleaning, disinfection and sterilisation instructions for tips J02001.



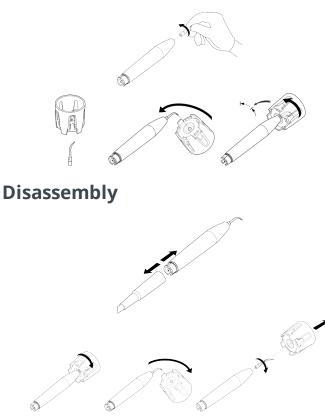


The practitioner has to continually monitor the clinical site to immediately identify any heating risk.

Tip | PFL | J00291 | (98) | V3 | 05/2021 | NB14EN040C

SATELEC A Company of ACTEON Group 17 av. Gustave Eiffel ZI du Phare 33700 MERIGNAC FRANCE Tel. +33 (0) 556 34 06 07 Fax. +33 (0) 556 34 92 92 satelec@acteongroup.com

Installation



Manufacturer identification



SATELEC

A Company of ACTEON Group 17, avenue Gustave Eiffel ZI du Phare 33700 MERIGNAC France Tel. +33 (0) 556.34.06.07 Fax. +33 (0) 556.34.92.92 E.mail : satelec@acteongroup.com www.acteongroup.com



Date of first CE marking

1998



