**ProRoot® MTA Carrier**

FOR DENTAL USE ONLY

**DIRECTIONS FOR USE - PROROOT® MTA CARRIER**

REF A0407 - A0408

1) **INDICATIONS FOR USE**
   Instruments are only to be used in a clinical or hospital setting by qualified users.

   **Application field**
   **General remarks:**
   • Carriers have been designed to manually dispense ProRoot® MTA.
   • The transparent dispensing tubes have been designed for loading and dispensing ProRoot® MTA on the site of surgery.
   • The carriers and tubes are to be chosen in accordance with the defect to obturate.
   • The grey carrier is for coronal endodontic applications. The brown carrier is for surgical obturations (repair of root resorption).
   • The tubes for coronal endodontics are longer (8 mm) and of a greater volume than the tubes designed for surgical endodontics (4 mm long).
   • All tools have been designed for manual use.
   • The round end is to be used to pack down and condense ProRoot® MTA.

2) **CONTRAINDICATIONS**
   No known.

3) **WARNINGS**
   No known.

4) **PRECAUTIONS**
   Sterilize instruments before use.
5) ADVERSE REACTIONS
In the present technical state, no adverse reaction has been reported so far.

6) STEP BY STEP INSTRUCTIONS
6.1) Usage in Coronal Endodontics (repair of perforations, creation of apical plug, pulp capping)
1) Select the grey endodontic carrier.
2) Select the dispensing tube for endodontics (8 mm).
3) Pull one end of the tube over around 1.5 mm of the tip of the carrier.
4) Dispense the ProRoot® MTA powder onto a mixing plate.
5) Open the sterile water ampoule.
6) Gradually incorporate the liquid into the ProRoot® MTA powder using a mixing stick. The volume of water provided is the exact quantity needed for 1 g of powder.
7) Fill the other end of the tube with the ProRoot® MTA mixture, by pressing down vertically.
8) Position the filled dispensing tube over the defect to repair.
9) Press down on the carrier to eject the ProRoot® MTA from the tube by using it like a piston.
10) Repeat steps 7 to 9 until the defect is sealed.
11) Lightly condense the ProRoot® MTA using the round end of the carrier, and remove any excess material with an endodontic excavator. For pulp capping or for repairing perforations, ProRoot® MTA can also be tamped down with a moist cotton pellet.
12) Place a moist cotton pellet in the canal, and seal the access preparation with a temporary restoration for a minimum of 4 (four) hours to allow ProRoot® MTA to set.

Note: If the ProRoot® MTA mixture is too moist to be tamped down after coming out of the dispensing tube, quickly dab it with a paper point or a cotton pellet to remove any excess water. This will give it a better consistency for setting.

6.2) Usage in surgical endodontics (root-end filling)
1) Use the brown surgical carrier designed for this purpose.
2) Select the dispensing tube for surgical application (4 mm long).
3) Pull one end of the tube over around 1.5 mm of the tip of the carrier.
4) Dispense the ProRoot® MTA powder onto a sterile mixing plate.
5) Open the sterile water ampoule.
6) Gradually incorporate the liquid into the ProRoot® MTA powder using a mixing stick. The volume of water provided is the exact quantity needed for 1 g of powder.
7) Fill the other end of the tube with the ProRoot® MTA mixture, by pressing down vertically.
8) Position the filled dispensing tube over the cavity to obturate.
9) Press down on the carrier to eject the ProRoot® MTA from the tube by using it like a piston.
10) Repeat steps 7 to 9 until the cavity is completely filled.
11) Condense the ProRoot® MTA using the round end of the carrier, and remove any excess material with an endodontic excavator.
12) Smooth the ProRoot® MTA using a moist cotton pellet.

Note: If the ProRoot® MTA mixture is too moist to condense after application, quickly dab it with a paper point or a cotton pellet to remove any excess water. This will give it a better consistency for setting.
7) DISINFECTION, CLEANING AND STERILIZATION
Reprocessing procedure for dental instruments

I - FOREWORD
Devices that are marked as “sterile” do not require any specific treatment before the first use. For all other devices not labelled “Sterile”, cleaning and sterilization prior first use is required according to section III - STEP-BY-STEP INSTRUCTIONS part 4 to 8 of this DFU.

For those devices that are not labelled “single use”, re-processing of the devices should be carried out as per this DFU. For hygiene and sanitary safety purposes, these instruments must be cleaned and sterilized before each re-use to prevent any contamination.

Excluded devices:
Uniclip and Moser Calcineable plastic posts cannot be sterilized and must be disinfected by immersion NaOCl (2,5 % at least) during 5 min. at ambient temperature.

II - GENERAL RECOMMENDATION
1) Use only a detergent solution, with disinfecting effect, which is approved for its efficacy (VAH/DGHM-listing, CE marking, FDA approval) and in accordance with the DFU of the detergent solution manufacturer. For all metal devices, it is recommended to use anticorrosion disinfecting and cleaning agents.

2) For your own safety, please wear personal protective equipment (gloves, glasses, mask).

3) The user is responsible for the cleaning and sterilization of the product for the first cycle and each further usage as well as for the usage of damaged or dirty devices where applicable after sterilization.

4) It is safest for the practitioner to use our devices only once. Should our devices be reused, we recommend that they should not be used more than 5 times. After each processing they should be carefully inspected before use: the appearance of defects such as deformations (bent, unwound), breakage, corrosion, loss of colour coding or marking, indicate that the devices are not able to fulfil the intended use with the required safety level and must therefore be discarded.

For our root canal shaping instruments we recommend not to exceed the following maximum number of uses;

<table>
<thead>
<tr>
<th>Type of canal</th>
<th>Stainless Steel instruments with a diameter ≤ISO 015</th>
<th>Stainless Steel instruments with a diameter &gt;ISO 015</th>
<th>NiTi instruments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Extremely curved (&gt;30°) or S-shaped canals</td>
<td>1 canal max.</td>
<td>2 canals max.</td>
<td>2 canals max.</td>
</tr>
<tr>
<td>Moderately curved canals (10° to 30°)</td>
<td>1 canal max.</td>
<td>4 canals max.</td>
<td>4 canals max.</td>
</tr>
<tr>
<td>Slightly curved (&lt;10°) or straight canals</td>
<td>1 canal max.</td>
<td>8 canals max.</td>
<td>8 canals max.</td>
</tr>
</tbody>
</table>

5) Single use marked devices are not approved for re-use.

6) For the final rinsing step deionised water use is mandatory, whether using an automated washer-disinfector or a manual cleaning method. Tap water is permissible for the other rinsing steps.
7) Instruments with plastic handles, and NiTi instruments should not be used with Hydrogen Peroxide ($H_2O_2$) solution which is known to degrade them.
8) Only the active part of the NiTi instrument, which is in contact with the patient should be immersed in a NaOCl solution concentrate at NOT more than 5%.
9) Avoid device to dry out, prior to, or during pre-disinfection, or cleaning. Dried biological material can be difficult to remove.
10) Use only device appropriated support for reprocessing.
11) Do not use label systems or identification markers directly on the device.

III - STEP-BY-STEP INSTRUCTIONS

<table>
<thead>
<tr>
<th>Operation</th>
<th>Activities</th>
<th>Warning and remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Disassembling</td>
<td>- Disassemble the device, if applicable.</td>
<td>- Remove and discard silicone stops.</td>
</tr>
</tbody>
</table>
| 2. Pre-Disinfection  | - Soak all devices immediately after use in a disinfection solution (We recommend the use of Prolystica® 2X Concentrate Enzymatic Presoak and Cleaner at 0.4% for a minimum of 15 minutes). Use a tray made from high density polyethylene or stainless steel. | - Follow instructions and respect concentrations and immersion times given by the manufacturer (an excessive concentration may cause corrosion or others defects on devices).  
- The pre-disinfection solution should be a specific solution targeted by the supplier for pre-disinfection. It should be used at the dilution specified by the supplier. It should contain, or be combined with a proteolytic enzyme.  
- The pre-disinfection solution should be aldehyde free (to avoid blood impurities fixation) and without di- or triethanolamines as corrosion inhibitor. Change the pre-disinfection solution regularly i.e. When it becomes soiled, or when efficacy is diminished due to exposure to microbial loads.  
- Do not use pre-disinfecting solutions containing Phenol or any products, which are not compatible with the devices.  
- For visible impurities observed on instruments a pre-cleaning is recommended with a soft brush (made from either nylon, polypropylene, acrylic). Manually brush the device until visible impurities are removed. |
| 3. Rinsing           | - Abundant rinsing (at least 1 min) under running water (ambient temperature). | - Use tap water for rinsing.  
- If a pre-disinfectant solution contains a corrosion inhibitor, it is recommended to do the rinsing step just before starting the cleaning step. |
| 4a. Automated Cleaning with washer-disinfector | - Place the devices in a kit, support, or container (made from stainless steel or titanium) to avoid any contact between devices or posts.  
- Place the devices in the washer-disinfector and execute the defined cycle (Ao value > 3000 or, at least 5 min at 90°C (194°F)).  
- Use a detergent solution with cleaning properties (we recommend Neodisher Mediclean Forte at 0.4%). | - Discard any devices with defects (broken, bent,…).  
- Avoid any contact between instruments or posts when placing in the washer-disinfector use kits, supports or containers.  
- Follow instructions and concentrations given by the manufacturer of the detergent solution.  
- Follow the instructions of the washer-disinfector and verify the success criteria after each cycle have been met as stated by the manufacturer.  
- The final rinse step should be with deionised water. For other steps follow the water quality defined by the manufacturer.  
- Use only approved washer-disinfector according to EN ISO 15883, maintained and validated regularly.  
- It is recommended to use an alkaline detergent with tensides, which has grease removal, disinfection (against bacteria/ fungi) and corrosion inhibition properties. The detergent should be approved for its efficacy (VAH/DGHM-listing, CE marking, FDA approval) and used in accordance with its DFU The detergent should be aldehyde free and without di- or triethanolamines as corrosion inhibitor. |
### 4b.i Manual Cleaning assisted by an ultrasonic device

- Place the devices in a kit, support or container (made from stainless steel, polypropylene or titanium) to avoid any contact between devices.
- Immerse in the detergent solution with cleaning properties (we recommend Neodisher Mediclean Forte at 2%), assisted by an ultrasonic device if suitable for at least 15 min.
- No visible impurities should be observed on the devices.
- If visible impurities are observed on the devices, the device must be manually brushed with a soft brush (made from either nylon, polypropylene, acrylic) until visible impurities are removed.
- Discard any devices with defects (broken, bent, and unwound).
- Follow instructions, observe water quality, concentrations and cleaning time stated by the manufacturer of the cleaning solution.
- It is recommended to use an alkaline detergent with tensides, which has grease removal, disinfection (against bacteria/fungi) and corrosion inhibition properties. The detergent should be approved for its efficacy (VAH/DGHM-listing, CE marking, FDA approval) and used in accordance with the DFU of the detergent solution manufacturer.
- The detergent should be aldehyde free and without di- or triethanolamines as corrosion inhibitor.

### 4b.ii Rinsing

- Abundant rinsing (at least 1 min) under running water (ambient temperature).
- Use deionised water for rinsing.
- If the previously used cleaning solution contains a corrosion inhibitor, it is recommended to do the rinsing step just before starting the autoclaving.

### 4b.iii Drying

- Devices should be thoroughly dried before inspection and packaging.
- Dry with a single use non-woven cloth.
- Devices should be dried until visual traces of moisture are eliminated.
- Particular attention has to be paid to effectively dry joints or cavities within a device.

### 5. Inspection

- If applicable assemble the devices (including the placement of new silicon stops).
- Inspect the devices functionality.
- Visually inspect devices with naked eye under appropriate lighting (min 500 lux) and sort out those with defects.
- Dirty devices must be cleaned again.
- Do not re-use silicon stops.
- Discard devices, which show any defect as described in the General Recommendation above (point 4).

### 6. Packaging

- Place the devices in a kit, support or container to avoid any contact between instruments or posts and pack the devices in "Sterilization pouches".
- Device must be double-packaged using paper-plastic pouches for steam sterilization prior sterilization. Ensure that the pouches are suitable for steam sterilization and were validated and manufactured as per ISO 11607 and EN 868-5.
- Use an appropriate packaging, moist-heat resistant (141°C, 286°F) and compliant with ISO 11607.
- Avoid any contact between instruments or posts during sterilization. Use kits, supports or containers.
- For sharp devices that are not contained within a box, silicon tubes should be placed around the devices to prevent packaging piercing.
- Seal the pouches according to the recommendation of the pouch manufacturer.
- If a thermo-sealer is used, the process must be validated and the thermosealer must be calibrated and qualified.
- Check the validity period of the pouch given by the pouch manufacturer to determine the shelf life.
### Sterilization

- The following sterilization cycles can be used:
  - 132°C (269.6°F), 4 minutes;
  - 134°C (273.2°F), 3 minutes;
  - 134°C (273.2°F), 18 minutes.
  
  We recommend a steam sterilization at 134°C / 273.2°F during 18 minutes for the purpose of de-activating potential prions.

- The instruments and posts must be sterilized according to the packaging labelling.
- When sterilizing multiple instruments in one autoclave cycle ensure that the sterilizer's maximum load is not exceeded.
- Place the pouches in the steam sterilizer according to the recommendation given by the sterilizer manufacturer.
- Use only Pre-Vaccum air Removal steam sterilizer that are matching the requirements of EN 13060 (class B, small sterilizer) and EN 285 (full size sterilizer), with saturated steam.
- Use a validated sterilization procedure according to ISO 17665 with a minimum drying time of 20 min.
- Respecting the maintenance procedure of the sterilizer is under the responsibility of the owner and should be performed following the requirements for medical devices sterilization (examples: planning of maintenance, qualification, acceptance criteria of condensate and water as per EN 285, annex 2).
- Control the efficiency and acceptance criteria of the sterilization procedure (packaging integrity, no humidity, no colour change of packaging, positive physico-chemical indicators, conformity of actual cycle parameters, to reference cycle parameters). A special attention should be paid to the packaging integrity if the sterilization cycle 134°C (273.2°F), 18 minutes was used.
- Store traceability records and define shelf-life according to packaging manufacturer guidelines.
- Shorter sterilization cycles according to local regulations are possible but are not guaranteed to de-activate prions.

### Storage

- Keep devices in sterilization packaging in a clean environment, away from sources of moisture and direct sunlight.
- Store at ambient temperature (typically 15 - 25°C (59 - 77°F)).

- After sterilization, the product should be manipulated with care in order to keep the integrity of the packaging (sterile barrier).
- Sterility cannot be guaranteed if packaging is open, damaged or wet.
- Check the packaging and the medical devices before using them (packaging integrity, no humidity and use by date). In case of damage, a complete rework should be performed.

### Symbols

<table>
<thead>
<tr>
<th>EN</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>SSt</td>
<td>Stainless steel</td>
</tr>
<tr>
<td>Sil</td>
<td>Silicone</td>
</tr>
<tr>
<td><img src="134%C2%B0C" alt="134°C" /></td>
<td>Autoclavable at the specified temperature</td>
</tr>
<tr>
<td><img src="Caution" alt="Caution" /></td>
<td>Caution: See directions for use</td>
</tr>
<tr>
<td>![Opened packages are not replaced](Opened packages are not replaced)</td>
<td>Opened packages are not replaced</td>
</tr>
</tbody>
</table>