MAP Sterilization Guide

| Operation | Operating mode | Warning | First use | Followin uses |
|---|--|---|-----------|------------------|
| Manual cleaning or assisted by an ultraso- nic device | Place the devices in a kit, box or container to avoid any contact between instruments. Immerse in the disinfecting solution with cleaning properties, assisted by an ultrasonic device if suitable. | No visible impurities should be observed on the instruments. Discard any instruments with large obvious defects (broken, bent, and twisted). Follow instructions and observe concentrations and time given by the manufacturer (see general recommendations under section 7). The disinfecting solution should be aldehyde free and without di- or triethanolamines as corrosion inhibitor. | | × |
| Rinsing | Abundant rinsing (at least 1 min) | Use quality water in accordance with local regulations. If a disinfecting solution contains a corrosion inhibitor, it is recommended to rinse the instruments just before the autoclaving. Dry on a single use non-woven cloth, or with a drying machine or filtered compressed air. | × | × |
| Inspection | Inspect devices and sort out those with defects. Assemble the devices (stops) | Ditry instruments must be cleaned and disinfected again. Discard instruments which show any deformations (bent, twisted), damages (broken, corroded) or defects (loss of color coding or marking) affecting the resistance, the safety or the performance of the instrument. | × | × |
| Packaging | Place the devices in a kit, box or container to avoid any contact between instruments and pack the devices in "Sterilization pouches". | Check the validity period of the pouch given by the manufacturer to determine the shelf life. Use packaging which are resistant up to a temperature of 141°C (286°F) and in accordance with EN ISO 11607. | × | × |
| Sterilization | Steam sterilization at: 134 °C / 273°F during 18 min. at 2, 1 bar. Check the success of the sterilisable cycle (use physico-chemical indicator for each performed cycle.) | The instruments, posts and the plastic supports must be sterilized according to the packaging labelling. *Use only autoclaves that are matching the requirements of EN 13060, EN 285. *The sterilization protocol has been validated by Produits Dentaires SA according to EN ISO 17665. *Respect the maintenance procedure of the autoclave device given by the manufacturer. *Use only this recommended sterilization procedure. *Control the efficiency (packaging integrity, no humidity, color change of sterilization indicators, physico-chemical integrators, digital records of cycles parameters). | × | × |
| Storage | Keep devices in sterilization packaging in a dry and clean environment | 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 validity period). | × | × |

