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CE

CE - DECLARATION OF CONFORMITY

(Directive 2007/47/EC amending Directive 93/42/EEC concerning medical de vices, Annex VII)

We

Manufacturer: CORICAMA SRL
Address: Via dei Fabbri 11 - 33085 MANIAGO (PN) – ITALY

Declare that the following medical devices:

MIRRORS, HANDLES (MIRRORS AND SCALPELS HANDLES), PROBES (PROBES AND EXPLORERS), CURETTES (CURETTES, HOES, SCALERS), PERIODONTAL KNIVES, FILES (PERIODONTAL AND BONE FILES), DENTAL DAM INSTRUMENTS (DAMS, FRAMES, PLIERS FOR DAM), DENTAL DAM CLAMPS, PLIERS FOR CLAMPS, MODELLING INSTRUMENTS, CONDENSERS (PLUGGERS, SPREADERS), SPATULAS FOR OBTURATIONS, AMALGAM CARRIERS, EXCAVATORS, PLACEMENT INSTRUMENTS, CROWNS AND BRIDGES INSTRUMENTS, MATRIX RETAINERS (AND MATRICES), IMPRESSION TRAYS, OSTEOTOMES, PERIOSTEALS ELEVATORS (AND SINUS LIFT INSTRUMENTS), CHISELS (PERIODONTAL AND BONE CHISELS), BONE SCRAPERS, WELLS (FOR AMALGAM AND BONE), NEEDLE HOLDERS, HAEMOSTATIC AND SURGICAL PLIERS, TWEEZERS (TOWEL/DRESSING, FOR SUTURES HAEMOSTATIC AND SURGICAL), SCISSORS (FOR DISSECTING, BANDAGE AND SUTURE), RETRACTORS, BONE RONGEURS, BONE CURETTES, SYRINGES, EXTRACTION FORCEPS, ROOT ELEVATORS, ORTHODONTIC INSTRUMENTS (PLIERS, CUTTERS, BAND AND BRACKETS INSTRUMENTS).

are classified as CLASS I, NON STERILE in accordance with Annex I to Directive 2007/47/EC amending

Directive 93/42/EEC concerning medical devices and are in compliance with the essential requirements of:

- Directive 2007/47/EC amending Directive 93/42/EEC concerning medical devices
- D. Lgs. 46/1997 amended by D. Lgs. 37/2010 applicable Italian legislation for Directive 2007/47/EC mending Directive 93/42/EEC concerning medical devices

Maniago, August 2020

Claudio Ghezzi
CEO & President

CORICAMA since 18