Ministero della Salute

DIREZIONE GENERALE DEI DISPOSITIVI MEDICI E DEL SERVIZIO FARMACEUTICO

DGDMF/III/P/I.5.1.e.1/2019/992
HAVING REGARD to Directive 93/42/EEC concerning medical devices;
HAVING REGARD to the Legislative Decree n. 46/97 and its following amendments implementing Directive 93/42/EEC;
HAVING REGARD to the request ref. 31635– A - 30/05/2019, submitted by the Company **VANNINI DENTAL INDUSTRY S.r.l.**, located in Via di Campigliano 55/A, 50012 Grassina (FI), Italy; VAT N°. 03651220489;
WHEREAS this Company paid the fees required by Ministerial Decree (D.M.) January 16, 2019;
HAVING REGARD to the official deeds:

IT IS ATTESTED

that, according to the Directive 93/42/EEC, the Company **VANNINI DENTAL INDUSTRY S.r.l.**, located in Via di Campigliano 55/A, 50012 Grassina (FI), Italy; is the manufacturer and has marked CE as medical devices, the following products:

Product	**Code**
PRESTIGE PUTTY	055001
PRESTIGE PUTTY SOFT	055002
PRESTIGE A PLUS PUTTY	055101
PRESTIGE REGULAR	055105 – 055205 - 055405
PRESTIGE REGULAR FAST	055505 – 055605 - 055420
PRESTIGE LIGHT	055106 – 055206 - 055425
PRESTIGE A PLUS LIGHT	055108 – 055208 - 055408
PRESTIGE HYDROLIGHT	055107 - 055207 - 055407
PRESTIGE MINI KIT	055028
PRESTIGE MINI KIT SOFT	055029
PRESTIGE A PLUS MINI KIT	055030
PRESTIGE MONOPHASE	055010 – 055210 - 055410
PRESTIGE BITE CAD CAM	055015 – 055215 - 055415
PRESTIGE VDX 5:1 IMPLANT	055112-055113
PROTESIL PUTTY	056004 -056002
PROTESIL LIGHT	056011 - 056116 - 056006A
PROTESIL CATALYST GEL	056021
PROTESIL MINI KIT	056217 - 056017A
SUPERFORM TRAY SYSTEM	101024 - 101025
KROMALTROPIC	002030 - 002031
CLIP ALGIN	002020 - 002025
CROMATIC	002010 - 002015

KROMALGIN PIU'	002000 - 002003
PROTESIL CHROMATIC	005011
PROTESIL NORMAL RIGID	005006
PROTESIL ELASTIC RAPID	005002
PRESTIGE UNIVERSAL ADHESIVE	058005

The above mentioned products, according to the art. 4 of Directive 93/42/EEC, can freely circulate and can be placed on the market in Italy and all over the European Union.
This document has been issued in a unique original version upon request of the manufacturer in order to export medical devices to **Countries outside European Union.**
It is not allowed any reproduction or publication of this document by paper, press, electronic base or websites.
It is only allowed to show or to delivery it, upon request of the customs or Health Competent Authorities of the importing country.

The Executive Manager
Dott. Marco Musella

DP