



Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

NOV - 7 2005

Mr. Michael Wierz
Export Sales Manager
Schutz-Dental GMBH
Dieselstrasse 5-6
Rosbach, Hessen
GERMANY D-61191

Re: K052073

Trade/Device Name: FUTURAPRESS LT, N AND HP, FUTURAJET, FUTURASELF,
FUTRAGEN

Regulation Number: 21 CFR 872.3760

Regulation Name: Denture relining, repairing, or rebasing resin

Regulatory Class: II

Product Code: EBI

Dated: July 28, 2005

Received: August 1, 2005

Dear Mr. Wierz:

This letter corrects our substantially equivalent letter of October 26, 2005

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

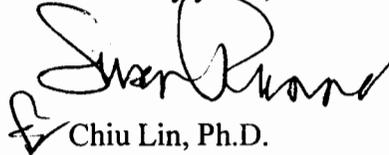
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (240) 276-0115. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/industry/support/index.html>.

Sincerely yours,



Chiu Lin, Ph.D.

Director

Division of Anesthesiology, General Hospital,

Infection Control and Dental Devices

Office of Device Evaluation

Center for Devices and

Radiological Health

Enclosure

Indications for Use

510(k) Number (if known): _____

Device Name: _____

Indications for Use:

FuturaPress LT	Cold-curing, methyl methacrylate resin for adding the saddles to cobalt chrome denture bases, repairs, additions and relines.
FuturaPress N	Cold-curing, methyl methacrylate resin for adding the saddles to cobalt chrome dentures bases, repairs, additions and relines.
FuturaJet	Cold-curing, methyl methacrylate resin intended specifically for injection moulding techniques (e. g. Unipress).
FuturaSelf	Cold-curing repair resin for additions, replacing teeth and other repairs.
FuturAcryl 2000	Heat-curing, methyl methacrylate resin for fabricating partial and full dentures using flasks and clamps or injection moulding systems.
FuturaPress HP	Heat-curing, methyl methacrylate resin, with liquid phase, for adding the saddles to cobalt chrome denture bases and relining dentures.
FuturaGen	Cold-curing, methyl methacrylate resin for the completion of full and partial dentures.

Prescription Use (Part 21 CFR 801 Subpart D)

AND/OR

Over-The-Counter Use _____ (21 CFR 801 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE OF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)



Page ___ of ___

(Signature Sign-Off)
Department of Anesthesiology, General Hospital,
Infection Control, Dental Devices

510(k) Number: 1052073